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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2005-0001	FDA-2005-0001-0001	Cold, Cough, Allergy, Bronchodilator, and Antiasthmatic Drug Products for Over-the-Counter Human Use; Proposed Amendment of Monograph for Over-the-Counter Bronchodilator Drug Products	7/13/2005	null date	7/13/2005	0910-AF32	05-13709
FDA	FDA-2005-0002	FDA-2005-0002-0001	High Chemical Co. et al.; Withdrawal of Approval of 13 New Drug Applications	9/21/2005	null date	9/21/2005		05-18873
FDA	FDA-2005-0003	FDA-2005-0003-0001	Drug Approvals: Circumstances Under Which an Active Ingredient May Be Simultaneously Marketed in Both a Prescription Drug Product and an Over-the-Counter Drug Product	9/1/2005	null date	9/1/2005	0910-AF72	05-17390
FDA	FDA-2005-0004	FDA-2005-0004-0001	Use of Materials Derived From Cattle in Human Food and Cosmetics		null date	9/7/2005	0910-AF47	05-17693
FDA	FDA-2005-0005	FDA-2005-0005-0001	Psychopharmacologic Drugs Advisory Committee; Notice of Meeting	9/21/2005	null date	9/21/2005		05-18872
FDA	FDA-2005-0006	FDA-2005-0006-0001	Industry Exchange Workshop on Food and Drug Administration Clinical Trial Requirements; Public Workshop	9/21/2005	null date	9/21/2005		05-18871
FDA	FDA-2005-0007	FDA-2005-0007-0002	Cold, Cough, Allergy, Bronchodilator, and Antiasthmatic Drug Products for Over-the-Counter Human Use; Technical Amendment	3/19/2007	null date	3/19/2007		E7-04957
FDA	FDA-2005-0007	FDA-2005-0007-0001	Cold, Cough, Allergy, Bronchodilator, and Antiasthmatic Drug Products for Over-the-Counter Human Use; Proposed Amendment of the Tentative Final Monograph for Combination Drug Products	7/13/2005	null date	7/13/2005	0910-AF33	05-13708
FDA	FDA-2005-0008	FDA-2005-0008-0001	Current Good Manufacturing Practice for Positron Emission Tomography Drugs		null date	9/20/2005		05-18510
FDA	FDA-2005-0009	FDA-2005-0009-0001	Stakeholder Meeting on the Implementation of A New Direction for the Food and Drug Administrations Radiological Health Program; Public Meeting	9/23/2005	null date	9/23/2005		05-19077
FDA	FDA-2005-0010	FDA-2005-0010-0001	Preparation for International Conference on Harmonization Meetings in Chicago, Illinois; Public Meeting	9/23/2005	null date	9/23/2005		05-19017
FDA	FDA-2005-0011	FDA-2005-0011-0001	Memorandum of Understanding Between the Food and Drug Administration and the Food and Drug Administration Alumni Association	9/23/2005	null date	9/23/2005		05-19016

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Food and Drug Administration (FDA) -- 1547 Documents

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Food and Drug Administration (FDA)								
FDA	FDA-2005-0012	FDA-2005-0012-0001	Frozen Desserts; Petition to Revoke Standards for Goats Milk Ice Cream and Mellorine and to Amend Standards for Ice Cream and Frozen Custard, Sherbet, and Water Ices; Petition to Amend Standards for Parmesan and Reggiano Cheese	9/27/2005	null date	9/27/2005		05-19194
FDA	FDA-2005-0013	FDA-2005-0013-0003	Designation of New Animal Drugs for Minor Uses or Minor Species		null date	7/26/2007	0910-AF60	E7-14444
FDA	FDA-2005-0013	FDA-2005-0013-0001	Designation of New Animal Drugs for Minor Uses or Minor Species		null date	9/27/2005	0910-AF60	05-19196
FDA	FDA-2005-0013	FDA-2005-0013-0002	Designation of New Animal Drugs for Minor Uses or Minor Species; Reopening of the Comment Period	12/28/2005	1/27/2002	12/28/2005	0910-AF60	05-24512
FDA	FDA-2005-0014	FDA-2005-0014-0001	International Conference on Harmonisation Workshop on Oncolytic Viruses; Public Workshop	9/27/2005	null date	9/27/2005		05-19195
FDA	FDA-2005-0015	FDA-2005-0015-0001	Advisory Committee for Pharmaceutical Science; Notice of Meeting	9/27/2005	null date	9/27/2005		05-19193
FDA	FDA-2005-0016	FDA-2005-0016-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Veterinary Feed Directive	9/28/2005	null date	9/28/2005		05-19393
FDA	FDA-2005-0017	FDA-2005-0017-0001	Memorandum of Understanding between the Food and Drug Administration, Forensic Chemistry Center and the Federal Bureau of Investigation	9/28/2005	null date	9/28/2005		05-19339
FDA	FDA-2005-0018	FDA-2005-0018-0001	Memorandum of Understanding Between the Food and Drug Administration and the National Library of Medicine	9/28/2005	null date	9/28/2005		05-19340
FDA	FDA-2005-0019	FDA-2005-0019-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Dissemination of Information on Unapproved/New Uses for Marketed Drugs, Biologics, and Devices	9/28/2005	null date	9/28/2005		05-19394
FDA	FDA-2005-0020	FDA-2005-0020-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Extralabel Drug Use in Animals	9/28/2005	null date	9/28/2005		05-19392
FDA	FDA-2005-0021	FDA-2005-0021-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Reprocessed Single-Use Device Labeling	9/29/2005	null date	9/29/2005		05-19509

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Food and Drug Administration (FDA) -- 1547 Documents

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Food and Drug Administration (FDA)								
FDA	FDA-2005-0022	FDA-2005-0022-0001	Medical Devices; Reprocessed Single-Use Devices; Termination of Exemptions From Premarket Notification; Requirement for Submission of Validation Data	9/29/2005	null date	9/29/2005		05-19510
FDA	FDA-2005-0023	FDA-2005-0023-0001	Food Labeling; Nutrient Content Claims, Definition of Sodium Levels for the Term ``Healthy	9/29/2005	null date	9/29/2005	0910-AC49	05-19511
FDA	FDA-2005-0024	FDA-2005-0024-0001	Draft Guidance for Industry on Using Electronic Means to Distribute Certain Product Information; Availability	9/30/2005	null date	9/30/2005		05-19731
FDA	FDA-2005-0025	FDA-2005-0025-0001	Draft Guidance for Industry and FDA Review Staff on Collection of Platelets by Automated Methods; Availability	10/3/2005	null date	10/3/2005		05-19727
FDA	FDA-2005-0026	FDA-2005-0026-0001	Registration of Food Facilities Under the Public Health Security and Bioterrorism Preparedness and Response Act of 2002	10/3/2005	null date	10/3/2005	0910-AC40	05-19730
FDA	FDA-2005-0027	FDA-2005-0027-0001	Iceberg Water Deviating From Identity Standard; Temporary Permit for Market Testing	10/3/2005	null date	10/3/2005		05-19728
FDA	FDA-2005-0028	FDA-2005-0028-0001	International Conference on Harmonisation; Draft Guidance on E2B(R) Clinical Safety Data Management: Data Elements for Transmission of Individual Case Safety Reports; Availability	10/3/2005	null date	10/3/2005		05-19655
FDA	FDA-2005-0029	FDA-2005-0029-0001	Food and Drug Administrations Communication of Drug Safety Information; Public Hearing	10/3/2005	null date	10/3/2005		05-19759
FDA	FDA-2005-0030	FDA-2005-0030-0001	Guidance for Industry and Food and Drug Administration Staff; Class II Special Controls Guidance Document: AFP-L3% Immunological Test Systems; Availability	10/4/2005	null date	10/4/2005		05-19853
FDA	FDA-2005-0031	FDA-2005-0031-0001	Third Annual Stakeholder Meeting on the Medical Device User Fee and Modernization Act of 2002; Public Meeting	10/4/2005	null date	10/4/2005		05-19864
FDA	FDA-2005-0032	FDA-2005-0032-0001	Medical Devices; Immunology and Microbiology Devices; Classification of AFP-L3% Immunological Test Systems	10/4/2005	null date	10/4/2005		05-19863
FDA	FDA-2005-0033	FDA-2005-0033-0001	New Animal Drugs; Change of Sponsor	10/4/2005	null date	10/4/2005		C5-17472
FDA	FDA-2005-0034	FDA-2005-0034-0001	Establishing a Docket for the Biological Products for Treatment of Rare Plasma Protein Disorders Public Workshop; Availability	10/4/2005	null date	10/4/2005		05-19852

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Food and Drug Administration (FDA)								
FDA	FDA-2005-0035	FDA-2005-0035-0001	Vision 2006--A Conversation With the American Public; Notice of Public Meetings on Specific Food and Drug Administration Issues; Notice of Cancellation of Meetings	10/5/2005	null date	10/5/2005		05-19956
FDA	FDA-2005-0036	FDA-2005-0036-0001	Draft Guidance for Industry and Food and Drug Administration Staff; Functional Indications for Implantable Cardioverter Defibrillators; Availability	10/6/2005	null date	10/6/2005		05-20092
FDA	FDA-2005-0037	FDA-2005-0037-0001	Substances Prohibited From Use in Animal Food or Feed		null date	10/6/2007	0910-AF46	05-20196
FDA	FDA-2005-0038	FDA-2005-0038-0001	The Essentials of Food and Drug Administration Device Regulations: A Primer for Manufacturers and Suppliers; Public Workshop	10/6/2005	null date	10/6/2005		05-20093
FDA	FDA-2005-0039	FDA-2005-0039-0001	Pediatric Advisory Committee; Notice of Meeting	10/7/2005	null date	10/7/2005		05-20303
FDA	FDA-2005-0040	FDA-2005-0040-0001	Solicitation of Public Review and Comment on Research Protocol: Gonadotropin-releasing Hormone Agonist Test in Disorders of Puberty	10/7/2005	null date	10/7/2005		05-20301
FDA	FDA-2005-0040	FDA-2005-0040-0002	Pediatric Ethics Subcommittee of the Pediatric Advisory Committee; Notice of Meeting	10/7/2005	null date	10/7/2005		05-20302
FDA	FDA-2005-0041	FDA-2005-0041-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Regulations Under the Federal Import Milk Act	10/7/2005	null date	10/7/2005		05-20148
FDA	FDA-2005-0041	FDA-2005-0041-0002	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Regulations Under the Federal Import Milk Act	12/30/2005	null date	12/30/2005		E5-08114
FDA	FDA-2005-0042	FDA-2005-0042-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Guidance for Industry: Notification of a Health Claim or Nutrient Content Claim Based on an Authoritative Statement of a Scientific Body	10/11/2005	null date	10/11/2005		05-20308
FDA	FDA-2005-0043	FDA-2005-0043-0001	Draft Guidance for Industry and FDA Staff: Compliance With the Medical Device User Fee and Modernization Act of 2002, as amended--Prominent and Conspicuous Mark of Manufacturers on Single-Use Devices; Availability	10/11/2005	null date	10/11/2005		05-20329

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Food and Drug Administration (FDA)								
FDA	FDA-2005-0043	FDA-2005-0043-0002	Guidance for Industry and Food and Drug Administration Staff: Compliance With the Medical Device User Fee and Modernization Act of 2002, as amended--Prominent and Conspicuous Mark of Manufacturers on Single-Use Devices; Availability	5/1/2006	null date	5/1/2006		E6-06458
FDA	FDA-2005-0044	FDA-2005-0044-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Cosmetic Product Voluntary Reporting Program	10/11/2005	null date	10/11/2005		05-20307
FDA	FDA-2005-0044	FDA-2005-0044-0002	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Cosmetic Product Voluntary Reporting Program	12/30/2005	null date	12/30/2005		E5-08112
FDA	FDA-2005-0045	FDA-2005-0045-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; General Licensing Provisions: Biologics License Application, Changes to an Approved Application, Labeling, Revocation and Suspension, and Forms FDA 356h and 2567	10/11/2005	null date	10/11/2005		05-20306
FDA	FDA-2005-0046	FDA-2005-0046-0001	Cold, Cough, Allergy, Bronchodilator, and Antiasthmatic Drug Products for Over-the-Counter Human Use; Amendment of Final Monograph for Over-the-Counter Nasal Decongestant Drug Products	10/11/2005	null date	10/11/2005	0910-AF34	05-20304
FDA	FDA-2005-0047	FDA-2005-0047-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Guidance for Industry: Fast Track Drug Development Programs--Designation, Development, and Application Review	10/11/2005	null date	10/11/2005		05-20305
FDA	FDA-2005-0048	FDA-2005-0048-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Investigational New Drug Regulations	10/12/2005	null date	10/12/2005		05-20362
FDA	FDA-2005-0049	FDA-2005-0049-0001	Blood Products Advisory Committee; Notice of Meeting	10/14/2005	null date	10/14/2005		05-20560
FDA	FDA-2005-0050	FDA-2005-0050-0001	Oncologic Drugs Advisory Committee; Notice of Meeting	10/14/2005	null date	10/114/2005		05-20559
FDA	FDA-2005-0051	FDA-2005-0051-0001	Transmissible Spongiform Encephalopathies Advisory Committee; Notice of Meeting	10/14/2005	null date	10/14/2005		05-20558

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Food and Drug Administration (FDA)								
FDA	FDA-2005-0052	FDA-2005-0052-0001	Prescription Drug User Fee Act; Public Meeting	10/18/2005	null date	10/18/2005		05-20875
FDA	FDA-2005-0052	FDA-2005-0052-0002	Prescription Drug User Fee Act; Public Meeting; Correction	10/28/2005	null date	10/28/2005		05-21525
FDA	FDA-2005-0053	FDA-2005-0053-0001	Guidance for Industry on Providing Regulatory Submissions in Electronic Format--Human Pharmaceutical Product Applications and Related Submissions Using the Electronic Common Technical Document Specifications; Availability	10/19/2005	null date	10/19/2005		05-20921
FDA	FDA-2005-0054	FDA-2005-0054-0001	Assessing Consumer Perceptions of Health Claims; Public Meeting; Request for Comments	10/19/2005	null date	10/19/2005		05-20969
FDA	FDA-2005-0055	FDA-2005-0055-0001	Cheeses and Related Cheese Products; Proposal to Permit the Use of Ultrafiltered Milk	10/19/2005	null date	10/19/2005		05-20874
FDA	FDA-2005-0056	FDA-2005-0056-0001	International Conference on Harmonisation; Guidance on S7B Nonclinical Evaluation of the Potential for Delayed Ventricular Repolarization (QT Interval Prolongation) by Human Pharmaceuticals; Availability	10/20/2005	null date	10/20/2005		05-20959
FDA	FDA-2005-0057	FDA-2005-0057-0001	International Conference on Harmonisation; Guidance on E14 Clinical Evaluation of QT/QTc Interval Prolongation and Proarrhythmic Potential for Non-Antiarrhythmic Drugs; Availability	10/20/2005	null date	10/20/2005		05-20971
FDA	FDA-2005-0058	FDA-2005-0058-0001	Anesthetic and Life Support Drugs Advisory Committee; Notice of Meeting	10/20/2005	null date	10/20/2005		05-20970
FDA	FDA-2005-0059	FDA-2005-0059-0001	Draft Guidance for Industry on Recommendations for Implementing a Collection Program for Source Plasma Containing Disease-Associated and Other Immunoglobulin Antibodies; Availability	10/20/2005	null date	10/20/2005		05-20958
FDA	FDA-2005-0060	FDA-2005-0060-0001	Science Board to the Food and Drug Administration; Notice of Meeting	10/21/2005	null date	10/21/2005		05-21036
FDA	FDA-2005-0061	FDA-2005-0061-0001	Agency Information Collection Activities: Proposed Collection; Comment Request; Guidance for Industry on Formal Meetings With Sponsors and Applicants for Prescription Drug User Fee Act Products	10/24/2005	null date	10/24/2005		05-21151
FDA	FDA-2005-0062	FDA-2005-0062-0001	MicroArray Quality Control Project Meeting on MicroArray Quality Control; Public Meeting	10/24/2005	null date	10/24/2005		05-21152

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Food and Drug Administration (FDA)								
FDA	FDA-2005-0063	FDA-2005-0063-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Medical Devices: Humanitarian Use Devices	10/24/2005	null date	10/24/2005		05-21158
FDA	FDA-2005-0064	FDA-2005-0064-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Food Contact Substances Notification	10/24/2005	null date	10/24/2005		05-21155
FDA	FDA-2005-0065	FDA-2005-0065-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Guidance for Industry on Formal Dispute Resolution; Appeals Above the Division Level	10/24/2005	null date	10/24/2005		05-21156
FDA	FDA-2005-0066	FDA-2005-0066-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Current Good Manufacturing Practices and Related Regulations for Blood and Blood Components; and Requirements for Donor Testing, Donor Notification, and ``Lookback	10/24/2005	null date	10/24/2005		05-21153
FDA	FDA-2005-0067	FDA-2005-0067-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Waivers of In Vivo Demonstration of Bioequivalence of Animal Drugs in Soluble Powder Oral Dosage Form Products and Type A Medicated Articles	10/24/2005	null date	10/24/2005		05-21154
FDA	FDA-2005-0068	FDA-2005-0068-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; 2005 Food Safety Survey	10/24/2005	null date	10/24/2005		05-21157
FDA	FDA-2005-0068	FDA-2005-0068-0002	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; 2005 Food Safety Survey; Correction	11/4/2005	null date	11/4/2005		05-21974
FDA	FDA-2005-0069	FDA-2005-0069-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Generic Food and Drug Administration Rapid Response Surveys	10/25/2005	null date	10/25/2005		05-21240
FDA	FDA-2005-0070	FDA-2005-0070-0001	Clinical Pharmacology Subcommittee of the Advisory Committee for Pharmaceutical Science; Notice of Meeting	10/25/2005	null date	10/25/2005		05-21241

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Food and Drug Administration (FDA)								
FDA	FDA-2005-0071	FDA-2005-0071-0001	Medical Devices; Immunology and Microbiology Devices; Classification of Cystic Fibrosis Transmembrane Conductance Regulator Gene Mutation Detection System	10/26/2005	null date	10/26/2005		05-21348
FDA	FDA-2005-0072	FDA-2005-0072-0001	Vaccines and Related Biological Products Advisory Committee; Notice of Meeting	10/26/2005	null date	10/26/2005		05-21350
FDA	FDA-2005-0073	FDA-2005-0073-0001	Guidance for Industry on Potassium Chloride Modified-Release Tablets and Capsules: In Vivo Bioequivalence and In Vitro Dissolution Testing; Availability	10/26/2005	null date	10/26/2005		05-21347
FDA	FDA-2005-0074	FDA-2005-0074-0001	Guidance for Industry and Food and Drug Administration Staff; Class II Special Controls Guidance Document: Cystic Fibrosis Transmembrane Conductance Regulator Gene Mutation Detection Systems; Availability	10/26/2005	null date	10/26/2005		05-21349
FDA	FDA-2005-0075	FDA-2005-0075-0001	Psychopharmacologic Drugs Advisory Committee; Notice of Meeting	10/28/2005	null date	10/28/2005		05-21524
FDA	FDA-2005-0076	FDA-2005-0076-0001	Oncologic Drugs Advisory Committee; Amendment of Notice	10/28/2005	null date	10/28/2005		05-21493
FDA	FDA-2005-0077	FDA-2005-0077-0001	Oral Dosage Form New Animal Drugs; Ivermectin and Praziquantel Paste	11/1/2005	null date	11/1/2005		05-21641
FDA	FDA-2005-0078	FDA-2005-0078-0001	Guidance for Industry: A Notice from the Food and Drug Administration to Growers, Food Manufacturers, Food Warehouse Managers, and Transporters of Food Products on Decontamination of Transport Vehicles; Availability	11/1/2005	null date	11/1/2005		05-21642
FDA	FDA-2005-0079	FDA-2005-0079-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Notice of Participation	11/1/2005	null date	11/1/2005		05-21774
FDA	FDA-2005-0080	FDA-2005-0080-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Study to Measure the Compliance of Prescribers With the Contraindication of the Use of Triptans in Migraine Headache Patients With Vascular Disease	11/2/2005	null date	11/2/2005		05-21807
FDA	FDA-2005-0081	FDA-2005-0081-0001	New Animal Drugs for Use in Animal Feeds; Melengestrol	11/2/2005	null date	12/10/4971		05-21808

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2005-0082	FDA-2005-0082-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Emergency Shortages Data Collection System (Formerly the Emergency Medical Device Shortage Program Survey)	11/4/2005	null date	11/4/2005		05-21973
FDA	FDA-2005-0083	FDA-2005-0083-0001	Advisory Committee on Special Studies Relating to the Possible Long-Term Health Effects of Phenoxy Herbicides and Contaminants (Ranch Hand Advisory Committee); Notice of Meeting	11/4/2005	null date	11/4/2005		05-22013
FDA	FDA-2005-0084	FDA-2005-0084-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; ZOMETA; Correction	11/4/2005	null date	11/4/2005		05-22012
FDA	FDA-2005-0085	FDA-2005-0085-0001	Pediatric Advisory Committee; Notice of Meeting	11/4/2005	null date	1/4/2005		05-22014
FDA	FDA-2005-0086	FDA-2005-0086-0001	Medical Devices; General and Plastic Surgery Devices; Classification of the Low Energy Ultrasound Wound Cleaner	11/7/2005	null date	11/7/2005		05-22068
FDA	FDA-2005-0087	FDA-2005-0087-0001	Oral Dosage Form New Animal Drugs; Tetracycline Hydrochloride Soluble Powder	11/7/2005	null date	11/7/2005		05-21889
FDA	FDA-2005-0088	FDA-2005-0088-0001	Meeting To Discuss Possible Changes to the Regulatory Jurisdiction of Certain Food Products Containing Meat and Poultry	11/7/2005	null date	11/7/2005		05-22123
FDA	FDA-2005-0089	FDA-2005-0089-0001	Guidance for Industry and Food and Drug Administration Staff; Class II Special Controls Guidance Document: Low Energy Ultrasound Wound Cleaner; Availability	11/7/2005	null date	11/7/2005		05-22069
FDA	FDA-2005-0090	FDA-2005-0090-0001	Change of Name; Technical Amendment	11/8/2005	null date	11/8/2005		05-22167
FDA	FDA-2005-0091	FDA-2005-0091-0001	Draft Guidance for Industry and Food and Drug Administration Staff; Class II Special Controls Guidance Document: Tinnitus Masker Devices; Availability	11/8/2005	null date	11/8/2005		05-22268
FDA	FDA-2005-0092	FDA-2005-0092-0001	Guidance for Industry: Validation of Analytical Procedures for Type C Medicated Feeds; Availability	8/1/2005	null date	8/1/2005		05-22222
FDA	FDA-2005-0093	FDA-2005-0093-0001	Food and Drug Administration Modernization Act of 1997: Modifications to the List of Recognized Standards, Recognition List Number: 013	11/8/2005	null date	11/8/2005		05-22267
FDA	FDA-2005-0094	FDA-2005-0094-0001	Ear, Nose, and Throat Devices; Tinnitus Masker; Designation of Special Controls	1/8/005	2/6/2006	1/8/2005		05-22269

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2005-0095	FDA-2005-0095-0001	Medical Devices: A Pilot Program to Evaluate a Proposed Globally Harmonized Alternative for Premarket Procedures; Guidance for Industry and FDA Staff; Availability	11/10/2005	null date	11/10/2005		05-22387
FDA	FDA-2005-0096	FDA-2005-0096-0001	Obstetrical and Gynecological Devices; Designation of Special Control for Condom and Condom With Spermicidal Lubricant		11/14/2005	11/14/2005	0910-AF21	05-22611
FDA	FDA-2005-0097	FDA-2005-0097-0001	Draft Guidance for Industry and Food and Drug Administration Staff; Class II Special Controls Guidance Document: Labeling for Male Condoms Made of Natural Rubber Latex; Availability	11/14/2005	null date	11/14/2005		05-22610
FDA	FDA-2005-0098	FDA-2005-0098-0001	Revised Compliance Policy Guide Regarding Prior Notice of Imported Food Under the Public Health Security and Bioterrorism Preparedness and Response Act of 2002; Availability	11/14/2005	null date	11/14/2005		05-22500
FDA	FDA-2005-0099	FDA-2005-0099-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Record Retention Requirements for the Soy Protein and Risk of Coronary Heart Disease Health Claim	11/15/2005	null date	11/15/2005		05-22636
FDA	FDA-2005-0100	FDA-2005-0100-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Survey on Program Funding	11/15/2005	null date	11/15/2005		05-22637
FDA	FDA-2005-0101	FDA-2005-0101-0001	Environmental Assessment; Categorical Exclusions	11/15/2005	null date	11/15/2005		05-22563
FDA	FDA-2005-0102	FDA-2005-0102-0001	Request for Nominations for Nonvoting Member Representing Industry Interests on a Public Advisory Committee; Nonprescription Drugs Advisory Committee	11/15/2005	null date	11/15/2005		05-22562
FDA	FDA-2005-0103	FDA-2005-0103-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Importers Entry Notice	11/16/2005	null date	11/16/2005		05-22671
FDA	FDA-2005-0104	FDA-2005-0104-0001	Oral Dosage Form New Animal Drugs; Tylosin	11/16/2005	null date	11/16/2005		05-22752
FDA	FDA-2005-0105	FDA-2005-0105-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; General Administrative Procedures: Citizen Petitions; Petition for Reconsideration or Stay of Action; Advisory Opinions	11/16/2005	null date	11/16/2005		05-22668

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2005-0106	FDA-2005-0106-0001	Draft Guidance for Industry on Noncontraceptive Estrogen Drug Products for the Treatment of Vasomotor Symptoms and Vulvar and Vaginal Atrophy Symptoms--Recommended Prescribing Information for Health Care Providers and Patient Labeling; Availability	11/16/2005	null date	11/16/2005		05-22754
FDA	FDA-2005-0107	FDA-2005-0107-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Filing Objections and Requests for a Hearing on a Regulation or Order	11/16/2005	null date	11/16/2005		05-22753
FDA	FDA-2005-0108	FDA-2005-0108-0001	Food Additives Permitted for Direct Addition to Food for Human Consumption; Vitamin D3		null date	11/16/2005		05-22670
FDA	FDA-2005-0109	FDA-2005-0109-0001	Drug and Biological Product Consolidation; Investigational New Drug Application Number Conversion	11/17/2005	null date	11/17/2005		05-22802
FDA	FDA-2005-0110	FDA-2005-0110-0001	New Animal Drugs; Florfenicol		null date	11/21/2005		05-22935
FDA	FDA-2005-0111	FDA-2005-0111-0001	Notice of Approval of Supplemental New Animal Drug Application; Ivermectin and Praziquantel Paste	11/21/2005	null date	11/21/2005		05-22941
FDA	FDA-2005-0112	FDA-2005-0112-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Regulations for In Vivo Radiopharmaceuticals Used for Diagnosis and Monitoring	11/22/2005	null date	11/22/2005		05-23039
FDA	FDA-2005-0113	FDA-2005-0113-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Guidance for Requesting an Extension to Use Existing Label Stock After the Trans Fat Labeling Effective Date of January 1, 2006	11/22/2005	null date	11/22/2005		05-23040
FDA	FDA-2005-0114	FDA-2005-0114-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Guidance for Requesting an Extension to Use Existing Label Stock After the Trans Fat Labeling Effective Date of January 1, 2006	11/22/2005	null date	11/22/2005		05-23041
FDA	FDA-2005-0115	FDA-2005-0115-0001	Guidance for Industry: Questions and Answers Regarding the Final Rule on Establishment and Maintenance of Records (Edition 2); Availability	11/22/2005	null date	11/22/2005		05-23062

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2005-0115	FDA-2005-0115-0002	Guidance for Industry: Questions and Answers Regarding the Final Rule on Establishment and Maintenance of Records (Edition 3); Availability	6/29/2006	null date	6/29/2006		E6-10239
FDA	FDA-2005-0115	FDA-2005-0115-0003	Guidance for Industry: Questions and Answers Regarding the Final Rule on Establishment and Maintenance of Records (Edition 4); Availability	9/26/2006	null date	9/26/2006		06-08241
FDA	FDA-2005-0116	FDA-2005-0116-0001	Investigational New Drugs: Export Requirements for Unapproved New Drug Products	11/23/2005	null date	11/23/2005	0910-AA61	05-23120
FDA	FDA-2005-0117	FDA-2005-0117-0001	New Animal Drugs; Flunixin	11/25/2005	null date	11/25/2005		
FDA	FDA-2005-0118	FDA-2005-0118-0002	Food Labeling: Nutrient Content Claims, Expansion of the Nutrient Content Claim ``Lean	1/12/2007	null date	1/12/2007	0910-ZA27	E7-00330
FDA	FDA-2005-0118	FDA-2005-0118-0001	Food Labeling: Nutrient Content Claims, Expansion of the Nutrient Content Claim ``Lean	11/25/2005	2/8/2006	11/25/2005		05-23293
FDA	FDA-2005-0119	FDA-2005-0119-0001	New Animal Drugs; Change of Sponsors Address	11/25/2005	null date	11/25/2005		05-23296
FDA	FDA-2005-0120	FDA-2005-0120-0001	New Animal Drugs; Change of Sponsors Name	11/25/2005	null date	11/25/2005		05-23297
FDA	FDA-2005-0121	FDA-2005-0121-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Focus Groups as Used by the Food and Drug Administration	11/25/2005	null date	11/25/2005		05-23248
FDA	FDA-2005-0122	FDA-2005-0122-0001	Implantation or Injectable Dosage Form New Animal Drugs; Boldenone	11/25/2005	null date	11/25/2005		05-23295
FDA	FDA-2005-0123	FDA-2005-0123-0001	Vaccines and Related Biological Products Advisory Committee; Notice of Meeting	11/29/2005	null date	11/29/2005		05-23373
FDA	FDA-2005-0124	FDA-2005-0124-0001	Nonprescription Drugs Advisory Committee and Pulmonary-Allergy Drugs Advisory Committee; Notice of Meeting and Request for Comments	11/29/2005	null date	11/29/2005		05-23372
FDA	FDA-2005-0125	FDA-2005-0125-0001	Guidance for Industry and Food and Drug Administration Staff, Guidance for Records Access Authority Provided in Title III, Subtitle A, of the Public Health Security and Bioterrorism Preparedness and Response Act of 2002; Availability	11/30/2005	null date	11/30/2005		05-23504

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2005-0126	FDA-2005-0126-0001	Draft Guidance for Industry on Safety, Efficacy, and Pharmacokinetic Studies to Support Marketing of Immune Globulin Intravenous (Human) as Replacement Therapy for Primary Humoral Immunodeficiency; Availability	12/1/2005	null date	12/1/2005		05-23520
FDA	FDA-2005-0127	FDA-2005-0127-0001	Change of Address; Technical Amendment	12/1/2005	null date	12/1/2005		05-23521
FDA	FDA-2005-0128	FDA-2005-0128-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Medical Device Recall Authority	12/1/2005	null date	12/1/2005		05-23519
FDA	FDA-2005-0129	FDA-2005-0129-0003	Revocation of Status of Specific Products; Group A Streptococcus; Confirmation of Effective Date	4/21/2006	null date	4/21/2006	0910-AF20	06-03790
FDA	FDA-2005-0129	FDA-2005-0129-0001	Revocation of Status of Specific Products; Group A Streptococcus; Companion Document to Direct Final Rule		12/2/2005	12/2/2005	0910-AF20	05-23545
FDA	FDA-2005-0129	FDA-2005-0129-0002	Revocation of Status of Specific Products; Group A Streptococcus	12/2/2005	1/15/2006	12/2/2005	0910-AF20	05-23546
FDA	FDA-2005-0130	FDA-2005-0130-0001	Over-the-Counter Drug Products; Safety and Efficacy Review; Additional Sunscreen Ingredients	12/5/2005	null date	12/5/2005		05-23576
FDA	FDA-2005-0131	FDA-2005-0131-0001	Over-the-Counter Drug Products; Safety and Efficacy Review; Additional Dandruff Control Ingredient	12/5/2005	null date	12/5/2005		05-23569
FDA	FDA-2005-0132	FDA-2005-0132-0001	Over-the-Counter Drug Products; Safety and Efficacy Review; Additional Acne Ingredient	12/5/2005	null date	12/5/2005		05-23570
FDA	FDA-2005-0133	FDA-2005-0133-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; MedWatch: Food and Drug Administration Medical Products Reporting Program	12/7/2005	null date	12/7/2005		05-23676
FDA	FDA-2005-0134	FDA-2005-0134-0001	Industry Exchange Workshop on Food and Drug Administration Clinical Trial Requirements; Public Workshop; Amendment of Notice	12/7/2005	null date	12/7/2005		05-23675
FDA	FDA-2005-0135	FDA-2005-0135-0001	Risk Management, Corrective and Preventive Actions, and Training: An Educational Forum; Public Workshop	12/7/2005	null date	12/7/2005		05-23677

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2005-0136	FDA-2005-0136-0001	Draft Guidance for Industry and Food and Drug Administration; Nucleic Acid Based In Vitro Diagnostic Devices for Detection of Microbial Pathogens; Availability	12/8/2005	null date	12/8/2005		05-23746
FDA	FDA-2005-0137	FDA-2005-0137-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Notice of a Claim for Generally Recognized as Safe Exemption Based on a Generally Recognized as Safe Determination	12/8/2005	null date	12/8/2005		05-23747
FDA	FDA-2005-0138	FDA-2005-0138-0001	Food Additives Permitted for Direct Addition to Food for Human Consumption; Synthetic Fatty Alcohols	12/8/2005	null date			05-23745
FDA	FDA-2005-0139	FDA-2005-0139-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; State Enforcement Notifications	12/8/2005	null date	12/8/2005		05-23744
FDA	FDA-2005-0140	FDA-2005-0140-0001	Ophthalmic and Topical Dosage Form New Animal Drugs; Miconazole Nitrate Cream; Miconazole Nitrate Lotion; Miconazole Nitrate Spray	12/9/2005	null date	12/9/2005		05-23811
FDA	FDA-2005-0141	FDA-2005-0141-0002	Dandruff, Seborrheic Dermatitis, and Psoriasis Drug Products Containing Coal Tar and Menthol for Over-the-Counter Human Use; Amendment to the Monograph	3/6/2007	null date	3/6/2007	0910-AF49	E7-03808
FDA	FDA-2005-0141	FDA-2005-0141-0001	Dandruff, Seborrheic Dermatitis, and Psoriasis Drug Products Containing Coal Tar and Menthol for Over-the-Counter Human Use; Proposed Amendment to the Monograph	12/9/2005	null date	12/9/2005	0910-AF49	05-23839
FDA	FDA-2005-0142	FDA-2005-0142-0001	Oral Dosage Form New Animal Drugs; Sulfadimethoxine Soluble Powder	12/9/2005	null date	12/9/2005		05-23813
FDA	FDA-2005-0143	FDA-2005-0143-0001	Food Ingredient Solutions, LLC; Filing of Color Additive Petition	12/9/2005	null date	12/9/2005		05-23812
FDA	FDA-2005-0144	FDA-2005-0144-0001	International Drug Scheduling; Convention on Psychotropic Substances; Single Convention on Narcotic Drugs; Butorphanol; Delta-9-tetrahydrocannabinol (Dronabinol); Gamma-Hydroxybutyric Acid; Ketamine; Khat; Tramadol; Zopiclone; Buprenorphine; Oripavine	12/13/2005	null date	12/13/2005		05-23958

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2005-0145	FDA-2005-0145-0001	Guidance for Industry and Food and Drug Administration; Requesting an Extension to Use Existing Label Stock After the Trans Fat Labeling Effective Date of January 1, 2006; Availability	12/14/2005	null date	12/14/2005		05-23987
FDA	FDA-2005-0146	FDA-2005-0146-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Reprocessed Single- Use Device Labeling	12/15/2005	null date	12/15/2005		05-24041
FDA	FDA-2005-0147	FDA-2005-0147-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Evaluation of Consumer-Friendly Formats for Brief Summary in Direct-to-Consumer Print Advertisements for Prescription Drugs: Study 1	12/15/2005	null date	12/15/2005		05-24040
FDA	FDA-2005-0148	FDA-2005-0148-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Food and Drug Administration Recall Regulations (Guidelines)	12/15/2005	null date	12/15/2005		05-24042
FDA	FDA-2005-0149	FDA-2005-0149-0001	Advisory Committees; Filing of Annual Reports	12/15/2005	null date	12/15/2005		05-24039
FDA	FDA-2005-0150	FDA-2005-0150-0001	Animal Drugs, Feeds, and Related Products; Withdrawal of Approval of New Animal Drug Applications	12/16/2005	null date	12/16/2005		05-24103
FDA	FDA-2005-0151	FDA-2005-0151-0001	Joint Meeting of the Nonprescription Drugs Advisory Committee and the Endocrinologic and Metabolic Drugs Advisory Committee; Notice of Meeting	12/16/2005	null date	12/16/2005		05-24101
FDA	FDA-2005-0152	FDA-2005-0152-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Research Study Complaint Form	12/16/2005	null date	12/16/2005		05-24102
FDA	FDA-2005-0153	FDA-2005-0153-0001	Memorandum of Understanding Between the United States Food and Drug Administration and the C-Path Institute	12/16/2005	null date	12/16/2005		05-24100
FDA	FDA-2005-0154	FDA-2005-0154-0001	Animal Drugs, Feeds, and Related Products; Withdrawal of Approval of New Animal Drug Applications	12/16/2005	null date	12/16/2005		05-24104
FDA	FDA-2005-0155	FDA-2005-0155-0001	Oral Dosage Form New Animal Drugs; Moxidectin Gel; Moxidectin and Praziquantel Gel	12/19/2005	null date	12/19/2005		05-24166

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Food and Drug Administration (FDA) -- 1547 Documents

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Food and Drug Administration (FDA)								
FDA	FDA-2005-0156	FDA-2005-0156-0001	Biological Products; Bacterial Vaccines and Toxoids; Implementation of Efficacy Review; Anthrax Vaccine Adsorbed; Final Order	12/19/2005	null date	12/19/2005		05-24223
FDA	FDA-2005-0157	FDA-2005-0157-0001	Biological Products; Bacterial Vaccines and Toxoids; Implementation of Efficacy Review	12/19/2005	null date	12/19/2005		05-24224
FDA	FDA-2005-0158	FDA-2005-0158-0001	Clinical Studies of Safety and Effectiveness of Orphan Products; Availability of Grants; Request for Applications	12/19/2005	null date	12/19/2005		05-24164
FDA	FDA-2005-0159	FDA-2005-0159-0001	New Animal Drugs; Change of Sponsor; Tiamulin	12/19/2005	null date	12/19/2005		05-24165
FDA	FDA-2005-0160	FDA-2005-0160-0001	New Animal Drugs; Change of Sponsor; Chloramphenicol Capsules	12/20/2005	null date	12/20/2005		05-24270
FDA	FDA-2005-0161	FDA-2005-0161-0001	Regulatory Process for Pediatric Mechanical	12/20/2005	null date	12/20/2005		05-24271
FDA	FDA-2005-0162	FDA-2005-0162-0001	Notice of Approval of Supplemental New Animal Drug Application; Tilmicosin	12/20/2005	null date	12/20/2005		05-24269
FDA	FDA-2005-0163	FDA-2005-0163-0001	Phenylpropanolamine-Containing Drug Products for Over-the-Counter Human Use; Tentative Final Monographs	12/22/2005	null date	12/22/2005	0910-AF34	E5-07646
FDA	FDA-2005-0164	FDA-2005-0164-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Food and Drug Administration Public Health Notification (formerly known as Safety Alert/Public Health Advisory) Readership Survey	12/22/2005	null date	12/22/2005		E5-07642
FDA	FDA-2005-0165	FDA-2005-0165-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Voluntary Hazard Analysis and Critical Control Point Manuals for Operators and Regulators of Retail and Food Service Establishments	12/22/2005	null date	12/22/2005		E5-07644
FDA	FDA-2005-0166	FDA-2005-0166-0001	Advisory Committees; Tentative Schedule of Meetings for 2006	12/22/2005	null date	12/22/2005		E5-07645
FDA	FDA-2005-0167	FDA-2005-0167-0001	Hand-Held, Doppler Ultrasound Prenatal Listening Devices	12/22/2005	null date	12/22/2005		E5-07643
FDA	FDA-2005-0168	FDA-2005-0168-0001	New Animal Drugs; Moxidectin	12/23/2005	null date	12/23/2005		05-24386
FDA	FDA-2005-0169	FDA-2005-0169-0002	Food Labeling: Health Claims; Soluble Dietary Fiber From Certain Foods and Coronary Heart Disease	5/22/2006	null date	5/22/2006		06-04703
FDA	FDA-2005-0169	FDA-2005-0169-0001	Food Labeling: Health Claims; Soluble Dietary Fiber From Certain Foods and Coronary Heart Disease	12/23/2005	null date	12/23/2005		05-24387

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2005-0170	FDA-2005-0170-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Medical Device Reporting: Manufacturer Reporting, Importer Reporting, User Facility Reporting, and Distributor Reporting	12/23/2005	null date	12/23/2005		E5-07726
FDA	FDA-2005-0171	FDA-2005-0171-0001	Oral Dosage Form New Animal Drugs; Furosemide	12/27/2005	null date	12/27/2005		05-24440
FDA	FDA-2005-0172	FDA-2005-0172-0001	Guidance for Industry: Questions and Answers Regarding Food Allergens, Including the Food Allergen Labeling and Consumer Protection Act of 2004 (Edition 2); Availability	12/27/2005	null date	12/27/2005		E5-07803
FDA	FDA-2005-0173	FDA-2005-0173-0001	Draft Guidance for Industry: Lead in Candy Likely To Be Consumed Frequently by Small Children: Recommended Maximum Level and Enforcement Policy; Draft Supporting Document: Supporting Document for Recommended Maximum Level for Lead in Candy Likely To Be Consumed Frequently by Small Children; Availability	12/27/2005	null date	12/27/2005		05-24494
FDA	FDA-2005-0174	FDA-2005-0174-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Reclassification Petitions for Medical Devices	12/27/2005	null date	12/27/2005		E5-07804
FDA	FDA-2005-0175	FDA-2005-0175-0001	Food Labeling: Ingredient Labeling of Dietary Supplements That Contain Botanicals; Withdrawal	12/28/2005	null date	12/28/2005		05-24511
FDA	FDA-2005-0176	FDA-2005-0176-0001	Determination That DECADRON (Dexamethasone) Tablets, 1.5 Milligrams, Were Not Withdrawn From Sale for Reasons of Safety or Effectiveness	12/28/2005	null date	12/28/2005		E5-07875
FDA	FDA-2005-0177	FDA-2005-0177-0001	Animal Drug User Fee Act; Public Meeting	12/28/2005	null date	12/28/2005		E5-07876
FDA	FDA-2005-0178	FDA-2005-0178-0001	Training Program for Regulatory Project Managers; Information Available to Industry	11/29/2005	null date	12/29/2005		E5-08017
FDA	FDA-2005-0179	FDA-2005-0179-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Medical Devices; Humanitarian Use Devices	12/30/2005	null date	12/30/2005		E5-08110

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2005-0180	FDA-2005-0180-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Mammography Quality Standards Act Final Regulations; Modifications and Additions to Policy Guidance Help System<greek-i>9	12/30/2005	null date	12/30/2005		E5-08111
FDA	FDA-2005-0181	FDA-2005-0181-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Draft Guidance for Clinical Trial Sponsors: Establishment and Operation of Clinical Trial Data Monitoring Committees	12/30/2005	null date	12/30/2005		E5-08115
FDA	FDA-2006-0001	FDA-2006-0001-0001	New Animal Drugs; Monensin	1/3/2006	null date	1/3/2006		05-24671
FDA	FDA-2006-0002	FDA-2006-0002-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Requirements for Collection of Data Relating to the Prevention of Medical Gas Mixups at Health Care Facilities--Survey	1/3/2006	null date	1/3/2006		E5-08113
FDA	FDA-2006-0003	FDA-2006-0003-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Current Good Manufacturing Practices and Related Regulations for Blood and Blood Components; and Requirements for Donor Testing, Donor Notification, and ``Lookback	1/3/2006	null date	1/3/2006		E5-08134
FDA	FDA-2006-0004	FDA-2006-0004-0001	Guidance for Industry on Development of Target Animal Safety and Effectiveness Data to Support Approval of Non-Steroidal Anti- Inflammatory Drugs for Use in Animals; Availability	1/4/2006	null date	1/4/2006		E5-08223
FDA	FDA-2006-0005	FDA-2006-0005-0001	Guidance for Industry and Review Staff on Recommended Approaches to Integration of Genetic Toxicology Study Results; Availability	1/4/2006	null date	1/4/2006		E5-08224
FDA	FDA-2006-0006	FDA-2006-0006-0001	University of Arkansas/Food and Drug Administration Food Labeling; Public Workshop	1/4/2006	null date	1/4/2006		E5-08225
FDA	FDA-2006-0007	FDA-2006-0007-0001	International Cooperation on Harmonisation of Technical Requirements for Registration of Veterinary Medicinal Products; Draft Revised Guidance for Industry on Impurities in New Veterinary Drug Substances (Revision); Request for Comments; Availability	1/4/2006	null date	1/4/2006		E5-08222

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2006-0008	FDA-2006-0008-0001	Implementation Plan for the Memorandum of Understanding Regarding the Sharing and Exchange of Information About Therapeutic Products Between the Food and Drug Administration Department of Health and Human Services of the United States of America and Health Products and Food Branch, Health Canada of Canada	1/6/2006	null date	1/6/2006		06-00113
FDA	FDA-2006-0009	FDA-2006-0009-0001	Oral Dosage Form New Animal Drugs; Phenylbutazone Powder	1/6/2006	null date	1/6/2006		06-00090
FDA	FDA-2006-0010	FDA-2006-0010-0001	Drug Safety and Risk Management Advisory Committee; Notice of Meeting	1/6/2006	null date	1/6/2006		E6-00006
FDA	FDA-2006-0011	FDA-2006-0011-0001	Oncologic Drugs Advisory Committee; Notice of Meeting	1/6/2006	null date	1/6/2006		E5-08333
FDA	FDA-2006-0012	FDA-2006-0012-0001	Pediatric Oncology Subcommittee of the Oncologic Drugs Advisory Committee; Notice of Meeting	1/6/2006	null date	1/6/2006		E5-08332
FDA	FDA-2006-0013	FDA-2006-0013-0001	Guidance for Industry and Food and Drug Administration: Requesting an Extension to Use Existing Label Stock After the Trans Fat Labeling Effective Date of January 1, 2006; Addendum December 30, 2005; Availability	1/6/2006	null date	1/6/2006		06-00116
FDA	FDA-2006-0014	FDA-2006-0014-0001	Medical Devices; Availability of Safety and Effectiveness Summaries for Premarket Approval Applications	1/9/2006	null date	1/9/2006		E6-00059
FDA	FDA-2006-0015	FDA-2006-0015-0001	International Cooperation on Harmonization of Technical Requirements for Registration of Veterinary Medicinal Products; Final Guidance for Industry on Environmental Impact Assessments for Veterinary Medicinal Products--Phase II; Availability	1/9/2006	null date	1/9/2006		E6-00039
FDA	FDA-2006-0016	FDA-2006-0016-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Survey of Healthcare Practitioners Regarding Their Preferences for Public Health Notifications	1/9/2006	null date	1/9/2006		E6-00072
FDA	FDA-2006-0017	FDA-2006-0017-0001	Cellular, Tissue and Gene Therapies Advisory Committee; Notice of Meeting	1/9/2006	null date	1/9/2006		E6-00071
FDA	FDA-2006-0018	FDA-2006-0018-0001	Draft Guidance for Industry and Food and Drug Administration Staff; Class II Special Controls Guidance Document: Herpes Simplex Virus Types 1 and 2 Serological Assays; Availability	1/9/2006	null date	1/9/2006		06-00174

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AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2006-0019	FDA-2006-0019-0001	Agency Emergency Processing Under Office of Management and Budget Review; Guidance on Informed Consent for In Vitro Diagnostic Device Studies Using Leftover Human Specimens That are Not Individually Identifiable	1/9/2006	null date	1/9/2006		E6-00073
FDA	FDA-2006-0020	FDA-2006-0020-0002	Immunology and Microbiology Devices; Reclassification of Herpes Simplex Virus (Types 1 and/or 2) Serological Assays; Correction	3/13/2006	null date	3/13/2006		E6-03522
FDA	FDA-2006-0020	FDA-2006-0020-0001	Immunology and Microbiology Devices; Reclassification of Herpes Simplex Virus (Types 1 and/or 2) Serological Assays	1/9/2006	null date	1/9/2006		06-00173
FDA	FDA-2006-0021	FDA-2006-0021-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Experimental Study of Carbohydrate Content Claims on Food Labels	1/10/2006	null date	1/10/2006		E6-00094
FDA	FDA-2006-0022	FDA-2006-0022-0001	International Cooperation on Harmonisation of Technical Requirements for Registration of Veterinary Medicinal Products; Draft Revised Guidance for Industry on Impurities in New Veterinary Medicinal Products (Revised); Request for Comments; Availability	1/10/2006	null date	1/10/2006		E6-00090
FDA	FDA-2006-0023	FDA-2006-0023-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Food Contact Substances Notification	1/10/2006	null date	1/10/2006		E6-00091
FDA	FDA-2006-0024	FDA-2006-0024-0001	Medical Device Reporting	1/10/2006	null date	1/10/2006		06-00172
FDA	FDA-2006-0025	FDA-2006-0025-0001	New Animal Drugs For Use in Animal Feeds; Monensin	1/11/2006	null date	1/11/2006		06-00228
FDA	FDA-2006-0026	FDA-2006-0026-0001	Implantation or Injectable Dosage Form New Animal Drugs; Hyaluronate Sodium Injection	1/11/2006	null date	1/11/2006		06-00229
FDA	FDA-2006-0027	FDA-2006-0027-0001	Anti-Counterfeit Drug Initiative Workshop and Vendor Display	1/11/2006	null date	1/11/2006		06-00249
FDA	FDA-2006-0028	FDA-2006-0028-0001	Memorandum of Understanding Between the United States Food and Drug Administration Department of Health and Human Services and the Australian Pesticides and Veterinary Medicines Authority, Australia	1/12/2006	null date	1/12/2006		06-00251

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Food and Drug Administration (FDA) -- 1547 Documents

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Food and Drug Administration (FDA)								
FDA	FDA-2006-0029	FDA-2006-0029-0001	Determination That Celestone Soluspan (Betamethasone Sodium Phosphate and Betamethasone Acetate) Injection and Celestone (Betamethasone Sodium Phosphate) Injection Were Not Withdrawn From Sale for Reasons of Safety or Effectiveness	1/12/2006	null date	1/12/2006		E6-00178
FDA	FDA-2006-0030	FDA-2006-0030-0001	Guidance for Industry on Formal Dispute Resolution: Scientific and Technical Issues Related to Pharmaceutical Current Good Manufacturing Practice; Availability	1/12/2006	null date	1/12/2006		E6-00233
FDA	FDA-2006-0031	FDA-2006-0031-0001	Protocol Regarding the Sharing of the Phonetic and Orthographic Computer Analysis Tool to Support Review and Evaluate Proprietary Names of Therapeutic Products Between the Food and Drug Administration Department of Health and Human Services of the United States of America and Health Products and Food Branch, Health Canada of Canada	1/12/2006	null date	1/12/2006		06-00252
FDA	FDA-2006-0032	FDA-2006-0032-0001	Human drugs: Phenylpropanolamine-containing products (OTC); tentative final monographs; correction	1/13/2006	null date	1/13/2006		Z5-07646
FDA	FDA-2006-0033	FDA-2006-0033-0001	New Animal Drugs	11/13/20006	null date	11/13/2006		06-55502
FDA	FDA-2006-0033	FDA-2006-0033-0002	New Animal Drugs For Use in Animal Feeds	6/22/2006	null date	6/22/2006		06-55520
FDA	FDA-2006-0033	FDA-2006-0033-0003	General Hospital and Personal Use Devices	9/12/2006	null date	9/12/2006		06-55527
FDA	FDA-2006-0033	FDA-2006-0033-0004	Food Labeling		null date	3/14/2007		07-55502
FDA	FDA-2006-0033	FDA-2006-0033-0005	Recordkeeping Requirements for Human Food and Cosmetics Manufactured From, Processed With, or Otherwise Containing, Material From Cattle	8/15/2007	null date	8/15/2007		07-55510
FDA	FDA-2006-0034	FDA-2006-0034-0001	University of Arkansas/Food and Drug Administration Food Labeling; Public Workshop; Correction	1/13/2006	null date	1/13/200		E6-00268
FDA	FDA-2006-0035	FDA-2006-0035-0003	Current Good Manufacturing Practice Regulation and Investigational New Drugs; Withdrawal	: 5/2/2006	null date	5/2/2006		06-04091
FDA	FDA-2006-0035	FDA-2006-0035-0002	Current Good Manufacturing Practice Regulation and Investigational New Drugs	1/17/2006	4/3/2020	1/17/2006		06-00353
FDA	FDA-2006-0035	FDA-2006-0035-0001	Current Good Manufacturing Practice Regulation and Investigational New Drugs; Companion Document to Direct Final Rule	1/17/2006	null date	1/17/2006		06-00350

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Food and Drug Administration (FDA)								
FDA	FDA-2006-0036	FDA-2006-0036-0001	Guidance for Industry on Exploratory Investigational New Drug Studies; Availability	17/2006	null date	1/17/2006		06-00354
FDA	FDA-2006-0037	FDA-2006-0037-0001	Draft Guidance for Industry on Investigational New Drugs; Approaches to Complying with Current Good Manufacturing Practice During Phase 1; Availability	1/17/2006	null date	1/17/2006		06-00352
FDA	FDA-2006-0038	FDA-2006-0038-0001	Institutional Review Boards: Requiring Sponsors and Investigators to Inform Institutional Review Boards of Any Prior Institutional Review Board Reviews; Withdrawal	1/17/2006	null date	1/17/2006		E6-00357
FDA	FDA-2006-0039	FDA-2006-0039-0001	Agency Information Collection Activities; Proposed Collection;Comment Request; Cosmetic Labeling Regulations	1/18/2006	null date	1/18/2006		E6-00443
FDA	FDA-2006-0040	FDA-2006-0040-0001	Able Laboratories, Inc.; Withdrawal of Approval of 43 Abbreviated New Drug Applications	1/19/2006	null date	1/19/2006		E6-00506
FDA	FDA-2006-0041	FDA-2006-0041-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Guidance for Industry on Formal Dispute Resolution; Appeals Above the Division Level	1/24/2006	null date	1/24/2006		E6-00763
FDA	FDA-2006-0042	FDA-2006-0042-0001	Requirements on Content and Format of Labeling for Human Prescription Drug and Biological Products	1/24/2006	null date	1/24/2006	0910-AA94	06-00545
FDA	FDA-2006-0043	FDA-2006-0043-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Guidance for Industry on Formal Meetings With Sponsors and Applicants for Prescription Drug User Fee Act Products	1/24/2006	null date	1/24/2006		E6-00765
FDA	FDA-2006-0044	FDA-2006-0044-0001	Two Guidances for Industry on the Content and Format of Labeling for Human Prescription Drug and Biological Products; Availability	1/24/2006	null date	1/24/2006		06-00544
FDA	FDA-2006-0045	FDA-2006-0045-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Good Laboratory Practice Regulations for Nonclinical Studies	1/24/2006	null date	1/24/2006		E6-00768
FDA	FDA-2006-0046	FDA-2006-0046-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Request for Samples and Protocols	1/24/2006	null date	1/24/2006		E6-00764

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Food and Drug Administration (FDA)								
FDA	FDA-2006-0047	FDA-2006-0047-0001	Draft Guidances for Industry on the Content and Format of Labeling for Human Prescription Drug and Biological Products; Availability	1/24/2006	null date	1/24/2006		06-00543
FDA	FDA-2006-0048	FDA-2006-0048-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Blood Establishment Registration and Product Listing, Form FDA 2830	1/25/2006	null date	1/25/2006		E6-00844
FDA	FDA-2006-0049	FDA-2006-0049-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Export Certificates for FDA Regulated Products	1/25/2006	null date	1/25/2006		E6-00845
FDA	FDA-2006-0050	FDA-2006-0050-0001	Global Harmonization Task Force, Study Groups 1, 2, 3, and 4; New Proposed and Final Documents; Availability	1/25/2006	null date	1/25/2006		E6-00846
FDA	FDA-2006-0051	FDA-2006-0051-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; SPIRIVA HANDIHALER	1/27/2006	null date	1/27/2006		E6-01050
FDA	FDA-2006-0052	FDA-2006-0052-0001	Peripheral and Central Nervous System Drugs Advisory Committee; Notice of Meeting	1/27/2006	null date	1/27/2006		E6-01006
FDA	FDA-2006-0053	FDA-2006-0053-0001	Oncologic Drugs Advisory Committee; Amendment of Notice	1/17/2006	null date	1/17/2006		E6-01003
FDA	FDA-2006-0054	FDA-2006-0054-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; ENABLEX	1/30/2006	null date	1/30/2006		E6-01072
FDA	FDA-2006-0055	FDA-2006-0055-0001	Blood Products Advisory Committee; Notice of Meeting	1/30/2006	null date	1/30/2006		E6-01075
FDA	FDA-2006-0056	FDA-2006-0056-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; XOLAIR	1/30/2006	null date	1/30/2006		E6-01078
FDA	FDA-2006-0057	FDA-2006-0057-0001	Listing of Color Additives Exempt From Certification; Food, Drug, and Cosmetic Labeling: Cochineal Extract and Carmine Declaration	1/30/2006	null date	1/30/2006	0910-AF12	E6-01104
FDA	FDA-2006-0058	FDA-2006-0058-0001	Anti-Infective Drugs Advisory Committee; Notice of Meeting	1/30/2006	null date	1/30/2006		E6-01069
FDA	FDA-2006-0059	FDA-2006-0059-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; HUMIRA	2/1/2006	null date	2/1/2006		E6-01313
FDA	FDA-2006-0060	FDA-2006-0060-0001	Pediatric Advisory Committee; Notice of Meeting	2/1/2006	null date	2/1/2006		E6-01223

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2006-0061	FDA-2006-0061-0001	Distribution of Blood Derivatives by Registered Blood Establishments that Qualify as Health Care Entities; Prescription Drug Marketing Act of 1987; Prescription Drug Amendments of 1992; Policies, Requirements and Administrative Procedures	2/1/2006	null date	2/1/2006		E6-01225
FDA	FDA-2006-0062	FDA-2006-0062-0001	Vaccines and Related Biological Products Advisory Committee; Notice of Meeting	1/1/2006	null date	1/1/2006		E6-01224
FDA	FDA-2006-0063	FDA-2006-0063-0001	Termination, By Expiration, of Declaration of Emergency Justifying Emergency Use Authorization of Anthrax Vaccine Adsorbed	2/1/2006	null date	2/1/2006		E6-01311
FDA	FDA-2006-0064	FDA-2006-0064-0001	Psychopharmacologic Drugs Advisory Committee; Notice of Meeting	2/1/2006	null date	2/1/2006		E6-01222
FDA	FDA-2006-0065	FDA-2006-0065-0001	Determination That CLARITIN (Loratadine) Hives Relief Syrup, 5 Milligrams per 5 Milliliters, Was Not Withdrawn From Sale for Reasons of Safety or Effectiveness	2/2/2006	null date	2/2/2006		E6-01364
FDA	FDA-2006-0066	FDA-2006-0066-0001	Summaries of Medical and Clinical Pharmacology Reviews of Pediatric Studies; Availability	2/2/2006	null date	2/2/2006		E6-01366
FDA	FDA-2006-0067	FDA-2006-0067-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; OMACOR	2/2/2006	null date	2/2/2006		E6-01365
FDA	FDA-2006-0068	FDA-2006-0068-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; ALOXI	2/2/2006	null date	2/2/2006		06-00903
FDA	FDA-2006-0069	FDA-2006-0069-0001	Summaries of Medical and Clinical Pharmacology Reviews of Pediatric Studies; Availability	2/3/2006	null date	2/3/2006		E6-01435
FDA	FDA-2006-0070	FDA-2006-0070-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; SURPASS	3/1/2006	null date	2/3/2006		E6-01434
FDA	FDA-2006-0071	FDA-2006-0071-0001	Oral Dosage Form New Animal Drugs; Firocoxib Paste	2/3/2006	null date	2/3/2006		06-00993
FDA	FDA-2006-0072	FDA-2006-0072-0001	Human Subject Protection--Information for Institutional Review Boards, Clinical Investigators, and Sponsors; Rescission, Reissuance, and Development of Food and Drug Administration Guidance Documents; Availability	2/3/2006	null date	2/3/2006		E6-01476
FDA	FDA-2006-0073	FDA-2006-0073-0001	Draft Guidance for Industry on Patient-Reported Outcome Measures: Use in Medical Product Development to Support Labeling Claims; Availability	2/3/2006	null date	2/3/2006		E6-01433

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2006-0074	FDA-2006-0074-0001	Determination That TEQUIN (Gatifloxacin) Injection, 10 Milligrams per Milliliter (200 Milligrams), Was Not Withdrawn From Sale for Reasons of Safety or Effectiveness	2/3/2006	null date	2/3/2006		E6-01475
FDA	FDA-2006-0075	FDA-2006-0075-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; CYPHER	2/3/2006	null date	2/3/2006		E6-01436
FDA	FDA-2006-0076	FDA-2006-0076-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Irradiation in the Production, Processing, and Handling of Food	2/6/2006	null date	2/6/2006		E6-01516
FDA	FDA-2006-0077	FDA-2006-0077-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Record Retention Requirements for the Soy Protein and Risk of Coronary Heart Disease Health Claim	2/6/2006	null date	2/6/2006		E6-01518
FDA	FDA-2006-0078	FDA-2006-0078-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Impact of Coupons on Consumer Perceptions of Products in Prescription Drugs in Direct-to-Consumer Prescription Drug Print Advertisements	2/6/2006	null date	2/6/2006		E6-01521
FDA	FDA-2006-0079	FDA-2006-0079-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Proposed Experimental Study of Trans Fat Claims on Foods	2/6/2006	null date	2/6/2006		E6-01517
FDA	FDA-2006-0080	FDA-2006-0080-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Requirements on Content and Format of Labeling for Human Prescription Drugs and Biologics; Requirements for Prescription Drug Product Labels	2/6/2006	null date	2/6/2006		E6-01519
FDA	FDA-2006-0081	FDA-2006-0081-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Experimental Study of Possible Footnotes and Cueing Schemes to Help Consumers Interpret Quantitative Trans Fat Disclosure on the Nutrition Facts Panel	2/6/2006	null date	2/6/2006		E6-01522

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2006-0081	FDA-2006-0081-0002	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Experimental Study of Possible Footnotes and Cueing Schemes to Help Consumers Interpret Quantitative Trans Fat Disclosure on the Nutrition Facts Panel	3/7/2007	null date	3/7/2007		E7-03904
FDA	FDA-2006-0082	FDA-2006-0082-0001	Referral of ZINECARD (dexrazoxane) and RELPAX (eletriptan) Written Requests for the Conduct of Pediatric Studies	2/6/2006	null date	2/6/2006		E6-01520
FDA	FDA-2006-0083	FDA-2006-0083-0001	Change of Address; Technical Amendment	2/6/2006	null date	2/6/2006		06-01040
FDA	FDA-2006-0084	FDA-2006-0084-0001	Behavior-Based Blood Donor Deferrals in the Era of Nucleic Acid Testing; Public Workshop; Request for Comments	2/7/2006	null date	2/7/2006		E6-01588
FDA	FDA-2006-0085	FDA-2006-0085-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Postmarketing Adverse Drug Experience Reporting	2/7/2006	null date	2/7/2006		E6-01587
FDA	FDA-2006-0086	FDA-2006-0086-0001	Independent Evaluation of the Food and Drug Administrations First Cycle Review Performance--Retrospective Analysis Final Report; Availability	2/7/2006	null date	2/7/2006		E6-01605
FDA	FDA-2006-0087	FDA-2006-0087-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; ALIMTA	2/8/2006	null date	2/8/2006		E6-01642
FDA	FDA-2006-0088	FDA-2006-0088-0001	Microbiology Devices; Reclassification of Hepatitis A Virus Serological Assays	9/1/2006	null date	9/1/2006		06-01206
FDA	FDA-2006-0088	FDA-2006-0088-0002	Microbiology Devices; Reclassification of Hepatitis A Virus Serological Assays; Correction	3/1/2006	null date	3/1/2006		06-01871
FDA	FDA-2006-0089	FDA-2006-0089-0001	Guidance for Industry and Food and Drug Administration Staff; Class II Special Controls Guidance Document; Hepatitis A Virus Serological Assays; Availability	2/9/2006	null date	2/9/2006		06-01207
FDA	FDA-2006-0090	FDA-2006-0090-0001	Advisory Committee on Special Studies Relating to the Possible Long-Term Health Effects of Phenoxy Herbicides and Contaminants (Ranch Hand Advisory Committee); Notice of Meeting	2/9/2006	null date	2/9/2006		E6-01737
FDA	FDA-2006-0091	FDA-2006-0091-0002	Orthopedic Devices; Reclassification of the Intervertebral Body Fusion Device	null date	null date	2/1/2007		E7-11240

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2006-0091	FDA-2006-0091-0001	Orthopedic Devices; Reclassification of the Intervertebral Body Fusion Device	2/9/2006	2/9/2006	2/9/2006		E6-01736
FDA	FDA-2006-0092	FDA-2006-0092-0001	Draft Guidance for Industry and Food and Drug Administration Staff; Pharmacogenetic Tests and Genetic Tests for Heritable Markers; Availability	2/9/2006	null date	2/9/2006		E6-01787
FDA	FDA-2006-0093	FDA-2006-0093-0001	Draft Guidance for Industry and Food and Drug Administration Staff; Draft Class II Special Controls Guidance Document: Intervertebral Body Fusion Device; Availability	2/9/2006	null date	2/9/2006		E6-01735
FDA	FDA-2006-0094	FDA-2006-0094-0001	Oral Dosage Form New Animal Drugs; Praziquantel, Pyrantel Pamoate, and Febantel Tablets	2/9/2006	null date	2/9/2006		06-01205
FDA	FDA-2006-0095	FDA-2006-0095-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Recommendations for the Early Food Safety Evaluation of New Non-Pesticidal Proteins Produced by New Plant Varieties Intended for Food Use	2/10/2006	null date	2/10/2006		E6-01806
FDA	FDA-2006-0096	FDA-2006-0096-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Financial Disclosure by Clinical Investigators	2/10/2006	null date	2/10/2006		E6-01807
FDA	FDA-2006-0097	FDA-2006-0097-0001	Determination That PEPTAVLON (Pentagastrin) for Subcutaneous Injection, 0.25 Milligrams per Milliliter, Was Not Withdrawn From Sale for Reasons of Safety or Effectiveness	2/10/2006	null date	2/10/2006		E6-01847
FDA	FDA-2006-0098	FDA-2006-0098-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; General Administrative Procedures: Citizen Petitions; Petition for Reconsideration or Stay of Action; Advisory Opinions	2/10/2006	null date	2/10/2006		E6-01846
FDA	FDA-2006-0099	FDA-2006-0099-0001	Implantation or Injectable Dosage Form New Animal Drugs; Moxidectin Solution	2/13/2006	null date	2/13/2006		06-01264
FDA	FDA-2006-0100	FDA-2006-0100-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Pharmaceutical Development Study	2/13/2006	null date	2/13/2006		E6-01918

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2006-0101	FDA-2006-0101-0001	Emerging Clostridial Disease; Public Workshop	2/14/2006	null date	2/14/2006		06-01371
FDA	FDA-2006-0102	FDA-2006-0102-0001	Guidance for Industry: Considerations for Developmental Toxicity Studies for Preventive and Therapeutic Vaccines for Infectious Disease Indications; Availability	2/14/2006	null date	2/14/2006		E6-01998
FDA	FDA-2006-0103	FDA-2006-0103-0001	Danisco USA, Inc.; Filing of Food Additive Petition	2/15/2006	null date	2/15/2006		E6-02130
FDA	FDA-2006-0103	FDA-2006-0103-0002	Danisco USA, Inc.; Filing of Food Additive Petition; Amendment	4/27/2006	null date	4/27/2006		E6-06370
FDA	FDA-2006-0103	FDA-2006-0103-0003	Food Additives Permitted for Direct Addition to Food for Human Consumption; Polydextrose	8/21/2007	9/20/2007	8/21/2007		E7-16322
FDA	FDA-2006-0104	FDA-2006-0104-0001	Medical Devices; Cardiovascular Devices; Classification of Implantable Intra-Aneurysm Pressure Measurement System	2/15/2006	null date	2/15/2006		06-01417
FDA	FDA-2006-0105	FDA-2006-0105-0001	ARCH Chemicals, Inc.; Filing of Food Additive Petition	2/15/2006	null date	2/15/2006		E6-02137
FDA	FDA-2006-0106	FDA-2006-0106-0001	Guidance for Industry on Nonclinical Safety Evaluation of Pediatric Drug Products; Availability	2/15/2006	null date	2/15/2006		E6-02139
FDA	FDA-2006-0107	FDA-2006-0107-0001	Guidance forIndustry and Food and Drug Administration; Class II Special Controls Guidance Document: Implantable Intra-Aneurysm Pressure Measurement System; Availability	2/15/2006	null date	2/15/2006		E6-02142
FDA	FDA-2006-0108	FDA-2006-0108-0001	Medical Devices; Radiology Devices; Reclassification of Bone Sonometers	5/16/2006	1/0/1900	2/15/2006		E6-02076
FDA	FDA-2006-0109	FDA-2006-0109-0001	Draft Guidance for Industry and Food and Drug Administration Staff; Medical Devices: Radiology Devices; Class II Special Controls Guidance Document: Bone Sonometers; Availability	2/15/2006	null date	2/15/2006		E6-02078
FDA	FDA-2006-0110	FDA-2006-0110-0001	Request for Nominations for Voting Members on	2/15/2006	null date	2/15/2006		E6-02071
FDA	FDA-2006-0111	FDA-2006-0111-0001	Manufacturing Subcommittee of the Advisory Committee for Pharmaceutical Science; Notice of Meeting	2/16/2006	null date	2/16/2006		E6-02237
FDA	FDA-2006-0112	FDA-2006-0112-0001	Guidance for Industry on Reports on the Status of Postmarketing Study Commitments--Implementation of Section 130 of the Food and Drug Administration Modernization Act of 1997; Availability	2/16/2006	null date	2/16/2006		E6-02184

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AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2006-0113	FDA-2006-0113-0001	Implantation or Injectable Dosage Form New Animal Drugs; Estradiol Benzoate	2/17/2006	null date	2/17/2006		06-01488
FDA	FDA-2006-0114	FDA-2006-0114-0001	Guidance for Industry on Waivers of In Vivo Demonstration of Bioequivalence of Animal Drugs in Soluble Powder Oral Dosage Form Products and Type A Medicated Articles; Availability	2/17/2006	null date	2/17/2006		E6-02291
FDA	FDA-2006-0115	FDA-2006-0115-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Investigational New Drug Regulations	2/17/2006	null date	2/17/2006		E6-02289
FDA	FDA-2006-0116	FDA-2006-0116-0001	Draft Guidance for Industry and FDA Staff: Whole Grains Label Statements; Availability	2/17/2006	null date	2/17/2006		06-01509
FDA	FDA-2006-0117	FDA-2006-0117-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; ERBITUX	2/17/2006	null date	2/17/2006		E6-02354
FDA	FDA-2006-0118	FDA-2006-0118-0001	Animal Drug User Fee Act; Public Meeting; Cancellation	2/21/2006	null date	2/21/2006		06-01571
FDA	FDA-2006-0119	FDA-2006-0119-0001	Cardiovascular and Renal Drugs Advisory Committee; Notice of Meeting	2/23/2006	null date	2/23/2006		E6-02542
FDA	FDA-2006-0120	FDA-2006-0120-0001	Peripheral and Central Nervous System Drugs Advisory Committee; Amendment of Notice	2/23/2006	null date	2/23/2006		E6-02541
FDA	FDA-2006-0121	FDA-2006-0121-0001	Listing of Color Additives Exempt From Certification; Tomato Lycopene Extract and Tomato Lycopene Concentrate	2/24/2006	null date	2/24/2006		06-01710
FDA	FDA-2006-0122	FDA-2006-0122-0001	Memorandum of Understanding Between the United States Food and Drug Administration and the United States General Services Administration	2/24/2006	null date	2/24/2006		06-01746
FDA	FDA-2006-0123	FDA-2006-0123-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Focus Groups as Used by the Food and Drug Administration	2/27/2006	null date	2/27/2006		E6-02726
FDA	FDA-2006-0124	FDA-2006-0124-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Aluminum in Large and Small Volume Parenterals Used in Total Parenteral Nutrition	2/27/2006	null date	2/27/2006		E6-02727
FDA	FDA-2006-0125	FDA-2006-0125-0001	Guidance for Industry on Internal Radioactive Contamination-- Development of Decorporation Agents; Availability	3/2/2006	null date	3/2/2006		E6-02942

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2006-0126	FDA-2006-0126-0001	Report on the Performance of Drug and Biologics Firms in Conducting Postmarketing Commitment Studies; Availability	3/3/2006	null date	3/3/2006		E6-03019
FDA	FDA-2006-0127	FDA-2006-0127-0001	Peripheral and Central Nervous System Drugs Advisory Committee; Notice of Meeting	3/3/2006	null date	3/3/2006		E6-03021
FDA	FDA-2006-0128	FDA-2006-0128-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Filing Objections and Requests for a Hearing on a Regulation or Order	3/3/2006	null date	3/3/2006		E6-03020
FDA	FDA-2006-0129	FDA-2006-0129-0001	Draft Guidance for Industry: Guide to Minimize Food Safety Hazards of Fresh-Cut Fruits and Vegetables; Availability	3/6/2006	null date	3/6/2006		E6-03084
FDA	FDA-2006-0130	FDA-2006-0130-0001	Industry Exchange Workshop on Food and Drug Administration Clinical Trial Requirements; Public Workshop	3/7/2006	null date	3/7/2006		E6-03229
FDA	FDA-2006-0131	FDA-2006-0131-0001	Draft Guidance for Industry on Clinical Data Needed to Support the Licensure of Trivalent Inactivated Influenza Vaccines; Availability	3/10/2006	null date	3/10/2006		E6-03370
FDA	FDA-2006-0132	FDA-2006-0132-0001	Draft Guidance for Industry on Clinical Data Needed to Support the Licensure of Pandemic Influenza Vaccines; Availability	3/10/2006	null date	3/10/2006		E6-03371
FDA	FDA-2006-0133	FDA-2006-0133-0001	Guidance for Industry and Food and Drug Administration; Hospital Bed System Dimensional and Assessment Guidance to Reduce Entrapment; Availability	3/10/2006	null date	3/10/2006		E6-03369
FDA	FDA-2006-0134	FDA-2006-0134-0001	Pediatric Advisory Committee; Amendment of Notice	3/10/2006	null date	3/10/2006		E6-03435
FDA	FDA-2006-0135	FDA-2006-0135-0001	Food Additives Permitted For Direct Addition to Food for Human Consumption; Glycerides and Polyglycides	3/13/2006	null date	3/13/2006		06-02354
FDA	FDA-2006-0136	FDA-2006-0136-0001	Guidance for Industry on Prescription Drug Marketing Act-- Donation of Prescription Drug Samples to Free Clinics; Availability	3/14/2006	null date	3/14/2006		E6-03532
FDA	FDA-2006-0137	FDA-2006-0137-0001	Oral Dosage Form New Animal Drugs; Sulfamerazine, Sulfamethazine, and Sulfaquinoxaline Powder	3/14/2006	null date	3/14/2006		06-02396
FDA	FDA-2006-0138	FDA-2006-0138-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; CRESTOR	3/15/2006	null date	3/15/2006		E6-03641
FDA	FDA-2006-0139	FDA-2006-0139-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; OVIDREL	3/15/2006	null date	3/15/2006		E6-03640

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2006-0140	FDA-2006-0140-0001	Guidance for Industry on Nonclinical Safety Evaluation of Drug or Biologic Combinations; Availability	2/15/2006	null date	3/15/2006		E6-03713
FDA	FDA-2006-0141	FDA-2006-0141-0001	Science Board to the Food and Drug Administration; Notice of Meeting	3/15/2006	null date	3/15/2006		E6-03639
FDA	FDA-2006-0142	FDA-2006-0142-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; PRIALT	5/1/2006	null date	5/1/2006		E6-03712
FDA	FDA-2006-0143	FDA-2006-0143-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; RELPAX	5/1/2006	null date	3/15/2006		E6-03711
FDA	FDA-2006-0144	FDA-2006-0144-0001	New Animal Drugs; Change of Sponsors Drug Labeler Code		null date			06-02554
FDA	FDA-2006-0145	FDA-2006-0145-0001	Obstetrics and Gynecology Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting	3/16/2006	null date	3/16/2006		E6-03786
FDA	FDA-2006-0146	FDA-2006-0146-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Emergency Shortages Data Collection System (Formerly the Emergency Medical Device Shortage Program Survey)	3/16/2006	null date	3/16/2006		E6-03820
FDA	FDA-2006-0147	FDA-2006-0147-0001	Guidance for Industry on Using a Centralized IRB Process in Multicenter Clinical Trials; Availability	3/16/2006	null date	3/16/2006		E6-03785
FDA	FDA-2006-0148	FDA-2006-0148-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; TYSABRI	3/16/2006	null date	3/16/2006		E6-03781
FDA	FDA-2006-0149	FDA-2006-0149-0001	Agency Information Collection Activities: Proposed Collection; Comment Request; Prescription Drug Marketing Act of 1987	3/16/2006	null date	3/16/2006		E6-03818
FDA	FDA-2006-0149	FDA-2006-0149-0002	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Prescription Drug Marketing Act of 1987	6/2/2006	null date	6/2/2006		E6-08569
FDA	FDA-2006-0150	FDA-2006-0150-0001	Confidentiality Arrangement Between the United States Food and Drug Administration and the French Health Products Safety Agency	3/16/2006	null date	3/16/2006		06-02539
FDA	FDA-2006-0151	FDA-2006-0151-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; CommentRequest; Notice of Participation	3/16/2006	null date	3/16/2006		E6-03819

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Food and Drug Administration (FDA) -- 1547 Documents

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Food and Drug Administration (FDA)								
FDA	FDA-2006-0151	FDA-2006-0151-0002	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Notice of Participation	6/2/2006	null date	6/2/2006		E6-08567
FDA	FDA-2006-0152	FDA-2006-0152-0001	Medical Devices; Availability of Safety and Effectiveness Summaries for Premarket Approval Applications	3/17/2006	null date	3/17/2006		E6-03850
FDA	FDA-2006-0153	FDA-2006-0153-0001	Manufacturing Subcommittee of the Advisory Committee for Pharmaceutical Science; Cancellation	3/17/2006	null date	3/17/2006		E6-03851
FDA	FDA-2006-0154	FDA-2006-0154-0001	Memorandum of Understanding Between the United States Food and Drug Administration, the National Cancer Institute, and the Centers for Medicare and Medicaid Services	3/20/2006	null date	3/20/2006		06-02656
FDA	FDA-2006-0155	FDA-2006-0155-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; MYCAMINE	3/20/2006	null date	3/20/2006		E6-03956
FDA	FDA-2006-0156	FDA-2006-0156-0001	Guidance for Industry and Food and Drug Administration Staff; Class II Special Controls Guidance Document: Reagents for Detection of Specific Novel Influenza A Viruses: Availability	3/22/2006	null date	3/22/2006		06-02743
FDA	FDA-2006-0157	FDA-2006-0157-0001	Medical Devices; Immunology and Microbiology Devices; Classification of Reagents for Detection of Specific Novel Influenza A Viruses	3/22/2006	null date	3/22/2006		06-02742
FDA	FDA-2006-0158	FDA-2006-0158-0001	New Animal Drugs; Adamantane and Neuraminidase Inhibitor Anti- influenza Drugs; Extralabel Animal Drug Use; Order of Prohibition	3/22/2006	5/22/2006	3/22/2006		06-02689
FDA	FDA-2006-0159	FDA-2006-0159-0001	The Ninth Annual Food and Drug Administration-Orange County Regulatory Affairs Educational Conference	3/22/2006	null date	3/22/2006		E6-04092
FDA	FDA-2006-0160	FDA-2006-0160-0001	Oral Dosage Form New Animal Drugs; Orbifloxacin	3/23/2006	null date	3/23/2006		06-02791
FDA	FDA-2006-0161	FDA-2006-0161-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; MYCAMINE	3/23/2006	null date	3/23/2006		E6-04165
FDA	FDA-2006-0162	FDA-2006-0162-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; ALAMAST	3/23/2006	null date	3/23/2006		E6-04163
FDA	FDA-2006-0163	FDA-2006-0163-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; VESICARE	3/23/2006	null date	3/23/2006		E6-04164

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Food and Drug Administration (FDA) -- 1547 Documents

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Food and Drug Administration (FDA)								
FDA	FDA-2006-0164	FDA-2006-0164-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Generic Food and Drug Administration Rapid Response Surveys	3/24/2006	null date	3/24/2006		E6-04262
FDA	FDA-2006-0165	FDA-2006-0165-0001	Eligibility Determination for Donors of Human Cells, Tissues, and Cellular and Tissue-Based Products; Correction	3/24/2006	null date	3/24/2006	0910-AB27	06-02841
FDA	FDA-2006-0166	FDA-2006-0166-0001	Memorandum of Understanding Between the Food and Drug Administration, Department of Health and Human Services, of the United States of America and the Certification and Accreditation Administration of the Peoples Republic of China Covering Ceramicware Intended for Use in the Preparation, Serving or Storage of Food or Drink and Offered for Export to the United States of America	3/27/2006	null date	3/27/2006		06-02894
FDA	FDA-2006-0167	FDA-2006-0167-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Medical Device Reporting: Manufacturer Reporting, Importer Reporting, User Facility Reporting, and Distributor Reporting	3/28/2006	null date	3/28/2006		E6-04426
FDA	FDA-2006-0168	FDA-2006-0168-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Guidance for Industry on Formal Meetings With Sponsors and Applicants for Prescription Drug User Fee Act Product	3/28/2006	null date	3/28/2006		E6-04424
FDA	FDA-2006-0169	FDA-2006-0169-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Guidance for Reagents for Detection of Specific Novel Influenza A Viruses	3/28/2006	null date	3/28/2006		E6-04427
FDA	FDA-2006-0170	FDA-2006-0170-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Guidance on Informed Consent for In Vitro Diagnostic Device Studies Using Leftover Human Specimens That are Not Individually Identifiable	3/28/2006	null date	3/28/2006		E6-04425

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Food and Drug Administration (FDA)								
FDA	FDA-2006-0171	FDA-2006-0171-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Survey of Health Care Practitioners Regarding Their Preferences for Public Health Notifications	3/28/2006	null date	3/28/2006		E6-04440
FDA	FDA-2006-0172	FDA-2006-0172-0001	Annual Comprehensive List of Guidance Documents at the Food and Drug Administration	28/2006	null date	3/28/2006		06-02941
FDA	FDA-2006-0173	FDA-2006-0173-0001	Guidance for Clinical Trial Sponsors: Establishment and Operation of Clinical Trial Data Monitoring Committees; Availability	3/28/2006	null date	3/28/2006		E6-04428
FDA	FDA-2006-0174	FDA-2006-0174-0001	Food Labeling: Health Claims; Dietary Noncariogenic Carbohydrate Sweeteners and Dental Caries	3/29/2006	null date	3/29/2006		06-03007
FDA	FDA-2006-0175	FDA-2006-0175-0001	Agency Information Collection Activities: Proposed Collection; Comment Request; Environmental Impact Considerations	3/29/2006	null date	3/29/2006		E6-04507
FDA	FDA-2006-0176	FDA-2006-0176-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Requirements for Submission of Labeling for Human Prescription Drugs and Biologics in Electronic Format	3/29/2006	null date	3/29/2006		E6-04506
FDA	FDA-2006-0177	FDA-2006-0177-0001	Implantation or Injectable Dosage Form New Animal Drugs; Flunixin	3/29/2006	null date	3/29/2006		06-03006
FDA	FDA-2006-0178	FDA-2006-0178-0001	Change of Telephone Number; Technical Amendment	3/30/2006	null date	3/30/2006		06-03046
FDA	FDA-2006-0179	FDA-2006-0179-0001	Draft Guidance for Industry on Providing Regulatory Submissions in Electronic Format--Orphan-Drug and Humanitarian Use Device Designation Requests and Related Submissions; Availability	3/31/2006	null date	3/31/2006		E6-04709
FDA	FDA-2006-0180	FDA-2006-0180-0001	Implantation or Injectable Dosage Form New Animal Drugs; Flunixin	3/31/2006	null date	3/31/2006		06-03118
FDA	FDA-2006-0181	FDA-2006-0181-0001	New Animal Drugs; Removal of Obsolete and Redundant Regulations	3/31/2006	null date	3/31/2006		06-03121
FDA	FDA-2006-0181	FDA-2006-0181-0002	New Animal Drugs for Use in Animal Feeds; Bacitracin; Nicarbazin; Oxytetracycline and Neomycin; Penicillin	3/31/2006	null date	3/31/2006		06-03120
FDA	FDA-2006-0181	FDA-2006-0181-0003	New Animal Drugs for Use in Animal Feeds; Bacitracin; Nitarsone; Zoalene	3/31/2006	null date	3/31/2006		06-03122

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Food and Drug Administration (FDA)								
FDA	FDA-2006-0182	FDA-2006-0182-0001	Medical Device Reporting; Premarket Approval of Medical Devices; Quality System Regulation; Technical Amendment	3/31/2006	null date	3/31/2006		06-03089
FDA	FDA-2006-0183	FDA-2006-0183-0001	Food and Drug Administration Modernization Act of 1997: Modifications to the List of Recognized Standards, Recognition List Number: 014	3/31/2006	null date	3/31/2006		E6-04695
FDA	FDA-2006-0184	FDA-2006-0184-0001	Ophthalmic and Topical Dosage Form New Animal Drugs; Gentamicin Sulfate, Betamethasone Valerate, Clotrimazole Ointment	4/3/2006	null date	4/3/2006		06-03149
FDA	FDA-2006-0185	FDA-2006-0185-0001	Joint Meeting of the Dental Products Panel of the Medical Devices Advisory Committee of the Center for Devices and Radiological Health and the Peripheral and Central Nervous System Drugs Advisory Committee of the Center for Drug Evaluation and Research; Notice of Meeting	4/3/2006	null date	4/3/2006		E6-04760
FDA	FDA-2006-0186	FDA-2006-0186-0001	Draft Guidance for Industry and Food and Drug Administration Staff; Class II Special Controls Guidance Document: Topical Oxygen Chamber for Extremities; Availability	4/6/2006	null date	4/6/2006		E6-04961
FDA	FDA-2006-0187	FDA-2006-0187-0001	General and Plastic Surgery Devices; Reclassification of the Topical Oxygen Chamber for Extremities	4/6/2006	4/6/2006	4/6/2006		E6-04962
FDA	FDA-2006-0188	FDA-2006-0188-0001	New Animal Drugs; Change of Sponsor; Soluble Bacitracin Methylene Disalicylate and Streptomycin Sulfate Oral Powder	4/7/2006	null date	4/7/2006		06-03353
FDA	FDA-2006-0189	FDA-2006-0189-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Substances Generally Recognized as Safe: Notification Procedure	4/7/2006	null date	4/7/2006		E6-05088
FDA	FDA-2006-0190	FDA-2006-0190-0001	Cooperative Agreement to Support a Single-Source Application--The Critical Path Institute: Collaborative Cardiovascular Drug Safety and Biomarker Research Program--ACTION; Availability of Sole Source Cooperative Agreement; Request for Application	4/7/2006	null date	4/7/2006		06-03408
FDA	FDA-2006-0191	FDA-2006-0191-0001	New Animal Drugs for Use in Animal Feeds; Chlortetracycline	4/7/2006	null date	4/7/2006		06-03352

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Food and Drug Administration (FDA)								
FDA	FDA-2006-0192	FDA-2006-0192-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Guidance for Requesting an Extension to Use Existing Label Stock After the Trans Fat Labeling Effective Date of January 1, 2006	4/10/2006	null date	4/10/2006		E6-05199
FDA	FDA-2006-0193	FDA-2006-0193-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Blood Establishment Registration and Product Listing, Form FDA 2830	4/10/2006	null date	4/10/2006		E6-05146
FDA	FDA-2006-0194	FDA-2006-0194-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Export Certificates for Food and Drug Administration-Regulated Products	4/10/2006	null date	4/10/2006		E6-05148
FDA	FDA-2006-0195	FDA-2006-0195-0001	Medical Gas Containers and Closures; Current Good Manufacturing Practice Requirements	4/10/2006	4/10/2006	4/10/2006		06-03370
FDA	FDA-2006-0196	FDA-2006-0196-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Voluntary Hazard Analysis and Critical Control Point Manuals for Operators and Regulators of Retail and Food Service Establishments	4/10/2006	null date	4/10/2006		E6-05142
FDA	FDA-2006-0197	FDA-2006-0197-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Reprocessed Single-Use Device Labeling	4/10/2006	null date	4/10/2006		E6-05150
FDA	FDA-2006-0198	FDA-2006-0198-0001	Guidance for Industry: Gamma Irradiation of Blood and Blood Components: A Pilot Program for Licensing; Withdrawal of Guidance	4/10/2006	null date	4/10/2006		E6-05204
FDA	FDA-2006-0199	FDA-2006-0199-0001	Guidance for Industry and Food and Drug Administration Staff; In Vitro Diagnostic Devices to Detect Influenza A Viruses: Labeling and Regulatory Path; Availability	4/10/2006	null date	4/10/2006		E6-05203
FDA	FDA-2006-0200	FDA-2006-0200-0001	Draft Guidance for Industry: Center for Biologics and Evaluation Pilot Licensing Program for Immunization of Source Plasma Donors Using Immunogen Red Blood Cells Obtained from an Outside Supplier; Withdrawal of Guidance	4/11/2006	null date	4/11/2006		E6-05220

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Food and Drug Administration (FDA)								
FDA	FDA-2006-0201	FDA-2006-0201-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Interstate Shellfish Dealers Certificate	4/11/2006	null date	4/11/2006		E6-05222
FDA	FDA-2006-0202	FDA-2006-0202-0001	Regulatory Site Visit Training Program	4/11/2006	null date	4/11/2006		E6-05221
FDA	FDA-2006-0203	FDA-2006-0203-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Food Labeling; Trans Fatty Acids in Nutrition Labeling	4/11/2006	null date	4/11/2006		E6-05219
FDA	FDA-2006-0204	FDA-2006-0204-0001	Radiological Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting	4/12/2006	null date	4/12/2006		E6-05411
FDA	FDA-2006-0205	FDA-2006-0205-0001	Oncologic Drugs Advisory Committee; Notice of Meeting	4/12/2006	null date	4/12/2006		E6-05413
FDA	FDA-2006-0206	FDA-2006-0206-0001	Stakeholder Meeting to Discuss the Possible Implementation of Two Review Performance Goals Referenced in the Medical Device User Fee and Modernization Act of 2002; Public Meeting	4/13/2006	null date	4/13/2006		E6-05494
FDA	FDA-2006-0207	FDA-2006-0207-0001	International Conference on Harmonisation; Guidance on S8 Immunotoxicity Studies for Human Pharmaceuticals; Availability	4/13/2006	null date	4/13/2006		E6-05495
FDA	FDA-2006-0208	FDA-2006-0208-0001	Food and Drug Administration-Regulated Products Containing Nanotechnology Materials; Planning of Public Meeting	4/14/2006	null date	4/14/2006		E6-05526
FDA	FDA-2006-0209	FDA-2006-0209-0001	Guidance for Industry on Exocrine Pancreatic Insufficiency Drug Products--Submitting New Drug Applications; Availability	4/14/2006	null date	4/14/2006		E6-05528
FDA	FDA-2006-0210	FDA-2006-0210-0001	Oral Dosage Form New Animal Drugs; Fenbendazole Granules	4/14/2006	null date	4/14/2006		06-03586
FDA	FDA-2006-0211	FDA-2006-0211-0001	International Cooperation on Harmonisation of Technical Requirements for Registration of Veterinary Medicinal Products; Draft Revised Guidance for Industry on Stability Testing of New Veterinary Drug Substances and Medicinal Products (Revision); Request for Comments; Availability	4/14/2006	null date	4/14/2006		E6-05525
FDA	FDA-2006-0212	FDA-2006-0212-0001	Draft Guidance for Industry: Recommended Study Design and Evaluation of Effectiveness Studies for Swine Respiratory Disease Claims; Availability	4/14/2006	null date	4/14/2006		E6-05527

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Food and Drug Administration (FDA)								
FDA	FDA-2006-0213	FDA-2006-0213-0001	Guidance for Industry and FDA Staff; The Mammography Quality Standards Act Final Regulations: Modifications and Additions to Policy Guidance Help System<greek-i>9; Availability	4/19/2006	null date	4/19/2006		E6-05785
FDA	FDA-2006-0214	FDA-2006-0214-0001	Orthopaedic and Rehabilitation Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting	4/19/2006	null date	4/19/2006		E6-05783
FDA	FDA-2006-0215	FDA-2006-0215-0001	Preparation for International Conference on Harmonization Meetings in Yokohama, Japan; Public Meeting	4/20/2006	null date	4/20/2006		E6-05905
FDA	FDA-2006-0216	FDA-2006-0216-0001	New Animal Drugs for Use in Animal Feeds; Melengestrol and Monensin	4/21/2006	null date	4/21/2006		06-03820
FDA	FDA-2006-0217	FDA-2006-0217-0001	MicroArray Quality Control Project on the Evaluation of Analysis Protocols for Deoxyribonucleic Acid Microarray Data	4/21/2006	null date	4/21/2006		E6-05995
FDA	FDA-2006-0218	FDA-2006-0218-0001	Vaccine Adverse Event Reporting; Revised Form VAERS-2; Withdrawal of Proposed Revised Form	4/21/2006	null date	4/21/2006		E6-05970
FDA	FDA-2006-0219	FDA-2006-0219-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; FASLODEX; Correction	4/24/2006	null date	4/24/2006		E6-06083
FDA	FDA-2006-0220	FDA-2006-0220-0001	Guidance for Sponsors, Institutional Review Boards, Clinical Investigators, and Food and Drug Administration Staff; Guidance on Informed Consent for In Vitro Diagnostic Device Studies Using Leftover Human Specimens That Are Not Individually Identifiable; Availability	4/25/2006	null date	4/25/2006		E6-06145
FDA	FDA-2006-0221	FDA-2006-0221-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Experimental Evaluation of Variations in Content and Format of the Brief Summary in Direct-to-Consumer Print Advertisements for Prescription Drugs	4/25/2006	null date	4/25/2006		E6-06142
FDA	FDA-2006-0222	FDA-2006-0222-0001	Industry Exchange Workshop on Food and Drug Administration Clinical Trial Requirements; Public Workshop; Correction	4/25/2006	null date	4/25/2006		E6-06119
FDA	FDA-2006-0223	FDA-2006-0223-0001	Regulatory Site Visit Training Program; Correction	4/25/2006	null date	4/25/2006		E6-06120

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2006-0224	FDA-2006-0224-0001	Guidance for Industry on Bar Code Label Requirements--Questions and Answers; Availability	4/27/2006	null date	4/27/2006		E6-06312
FDA	FDA-2006-0225	FDA-2006-0225-0001	New Animal Drugs for Use in Animal Feeds; Lasalocid and Chlortetracycline	4/27/2006	null date	4/27/2006		06-03953
FDA	FDA-2006-0226	FDA-2006-0226-0001	Draft ``Guidance for Industry: Informed Consent Recommendations for Source Plasma Donors Participating in Plasmapheresis and Immunization Programs; Availability	4/27/2006	null date	4/27/2006		E6-06314
FDA	FDA-2006-0227	FDA-2006-0227-0001	Research Review Subcommittee of the Vaccines and Related Biological Products Advisory Committee; Notice of Meeting	5/1/2006	null date	5/1/2006		E6-06508
FDA	FDA-2006-0228	FDA-2006-0228-0001	Vaccines and Related Biological Products Advisory Committee; Notice of Meeting	1/1/2006	null date	1/1/2006		E6-06509
FDA	FDA-2006-0229	FDA-2006-0229-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Research Study Complaint Form	5/1/2006	null date	5/1/2006		E6-06457
FDA	FDA-2006-0230	FDA-2006-0230-0001	Agency Emergency Processing Under the Office of Management and Budget Review; MedWatch--The Food and Drug Administration Safety Information and Adverse Event Reporting Program; Proposal to Survey MedWatch Partners Organizations	5/1/2006	null date	5/1/2006		E6-06461
FDA	FDA-2006-0231	FDA-2006-0231-0001	International Cooperation on Harmonisation of Technical Requirements for Approval of Veterinary Medicinal Products; Draft Revised Guidance for Industry on Pharmacovigilance of Veterinary Medicinal Products: Management of Adverse Event Reports; Request for Comments; Availability	5/2/2006	null date	5/2/2006		E6-06602
FDA	FDA-2006-0232	FDA-2006-0232-0001	International Cooperation on Harmonisation of Technical Requirements for Registration of Veterinary Medicinal Products (VICH); Draft Guidance for Industry on Pharmacovigilance of Veterinary Medicinal Products; Data Elements for Submission of Adverse Event Reports (VICH GL42); Request for Comments; Availability	5/2/2006	null date	5/2/2006		E6-06601

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2006-0233	FDA-2006-0233-0001	Guidance for Industry: Guidance on the Labeling of Certain Uses of Lecithin Derived From Soy Under the Federal Food, Drug, and Cosmetic Act; Availability	5/2/2006	null date	5/2/2006		E6-06551
FDA	FDA-2006-0234	FDA-2006-0234-0001	Industry Exchange Workshop to Celebrate Food and Drug Administration Centennial: Past, Present, and Future of Regulated Food, Drugs, Nutritional Supplements, and Medical Devices; Public Workshop	5/3/2006	null date	5/3/2006		06-04185
FDA	FDA-2006-0235	FDA-2006-0235-0001	Memorandum of Understanding Between the Food and Drug Administration, United States Department of Health and Human Services, the Animal and Plant Health Inspection Service, the United States Department of Agriculture, and The National Institutes of Health, United States Department of Health and Human Services Concerning Laboratory Animal Welfare	5/3/2006	null date	5/3/2006		06-04184
FDA	FDA-2006-0236	FDA-2006-0236-0001	Summaries of Medical and Clinical Pharmacology Reviews of Pediatric Studies; Availability	5/3/2006	null date	5/3/2006		E6-06706
FDA	FDA-2006-0237	FDA-2006-0237-0001	Guidance for Industry on Using Electronic Means to Distribute Certain Product Information; Availability	5/3/2006	null date	5/3/2006		E6-06705
FDA	FDA-2006-0238	FDA-2006-0238-0001	Food Safety and Defense Workshop; Public Workshop	5/10/2006	null date	5/10/2006		06-04366
FDA	FDA-2006-0239	FDA-2006-0239-0001	Draft Guidance for Clinical Investigators, Institutional Review Boards, and Sponsors; Process for Handling Pediatric Referrals to the Food and Drug Administration: Additional Safeguards for Children in Clinical Investigations	5/10/2006	null date	5/10/2006		E6-07058
FDA	FDA-2006-0240	FDA-2006-0240-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; General Administrative Procedures: Citizen Petitions; Petition for Reconsideration or Stay of Action; Advisory Opinions	5/11/2006	null date	5/11/2006		E6-07157

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2006-0241	FDA-2006-0241-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Irradiation in the Production, Processing, and Handling of Food	5/11/2006	null date	5/11/2006		E6-07178
FDA	FDA-2006-0241	FDA-2006-0241-0002	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Irradiation in the Production, Processing, and Handling of Food	7/25/2006	null date	7/25/2006		E6-11776
FDA	FDA-2006-0242	FDA-2006-0242-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Adverse Drug Experience Reporting	5/11/2006	null date	5/11/2006		E6-07159
FDA	FDA-2006-0243	FDA-2006-0243-0003	Blood Vessels Recovered With Organs and Intended for Use in Organ Transplantation; Withdrawal	9/14/2006	null date	9/14/2006		06-07644
FDA	FDA-2006-0243	FDA-2006-0243-0004	Blood Vessels Recovered With Organs and Intended for Use in Organ Transplantation		null date	3/12/2007	0910-AF65	07-01131
FDA	FDA-2006-0243	FDA-2006-0243-0001	Blood Vessels Recovered With Organs and Intended for Use in Organ Transplantation	5/12/2006	7/26/2006	5/12/2006		06-04369
FDA	FDA-2006-0243	FDA-2006-0243-0002	Blood Vessels Recovered With Organs and Intended for Use in Organ Transplantation; Companion Document to Direct Final Rule	5/12/2006	7/26/2006	5/12/2006		06-04370
FDA	FDA-2006-0244	FDA-2006-0244-0001	New Animal Drugs for Use in Animal Feeds; Melengestrol and Tylosin	5/12/2006	null date	5/12/2006		06-04426
FDA	FDA-2006-0245	FDA-2006-0245-0001	New Animal Drugs; Change of Sponsor	5/15/2006	null date	5/15/2006		06-04505
FDA	FDA-2006-0246	FDA-2006-0246-0001	New Animal Drugs; Change of Sponsor; Fomepizole	5/16/2006	null date	5/16/2006		06-04534
FDA	FDA-2006-0247	FDA-2006-0247-0001	Product Stability Data; Notice of Pilot Project	5/16/2006	null date	5/16/2006		E6-07391
FDA	FDA-2006-0248	FDA-2006-0248-0001	Guidance for Industry: Preparing a Claim of Categorical Exclusion or an Environmental Assessment for Submission to the Center for Food Safety and Applied Nutrition; Availability	5/18/2006	null date	5/18/2006		E6-07528
FDA	FDA-2006-0249	FDA-2006-0249-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Records and Reports Concerning Experience With Approved New Animal Drugs	5/19/2006	null date	5/19/2006		E6-07616

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2006-0250	FDA-2006-0250-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Guidance on Informed Consent for In Vitro Diagnostic Device Studies Using Leftover Human Specimens That are Not Individually Identifiable	5/19/2006	null date	5/19/2006		E6-07617
FDA	FDA-2006-0251	FDA-2006-0251-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Experimental Study of Qualified Health Claims: Consumer Inferences About Omega-3 Fatty Acids, Monounsaturated Fatty Acids From Olive Oil, and Green Tea	5/22/2006	null date	5/22/2006		E6-07692
FDA	FDA-2006-0252	FDA-2006-0252-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Focus Groups as Used by the Food and Drug Administration	5/22/2006	null date	5/22/2006		E6-07698
FDA	FDA-2006-0253	FDA-2006-0253-0001	International Conference on Harmonisation; Guidance on Q8 Pharmaceutical Development; Availability	5/22/2006	null date	5/22/2006		E6-07727
FDA	FDA-2006-0254	FDA-2006-0254-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Guidance on Reagents for Detection of Specific Novel Influenza A Viruses	5/22/2006	null date	5/22/2006		E6-07708
FDA	FDA-2006-0255	FDA-2006-0255-0001	Draft Compliance Policy Guide; Guidance Levels for 3-MCPD (3- chloro-1,2-propanediol) in Acid-Hydrolyzed Protein and Asian-Style Sauces; Availability	5/23/2006	null date	5/23/2006		E6-07796
FDA	FDA-2006-0256	FDA-2006-0256-0001	Draft Guidance for Industry and Food and Drug Administration Staff; Guidance for the Use of Bayesian Statistics in Medical Device Clinical Trials; Availability	5/23/2006	null date	5/23/2006		E6-07855
FDA	FDA-2006-0257	FDA-2006-0257-0001	Testing for Malarial Infections in Blood Donors; Public Workshop	5/23/2006	null date	5/23/2006		E6-07854
FDA	FDA-2006-0258	FDA-2006-0258-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Aluminum in Large and Small Volume Parenterals Used in Total Parenteral Nutrition	5/25/2006	null date	5/25/2006		E6-07984

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2006-0259	FDA-2006-0259-0001	Agency Information Collection Activities; Announcement of Officeof Management and Budget Approval; Filing Objections and Requests for a Hearing on a Regulation or Order	5/25/2006	null date	5/25/2006		E6-07991
FDA	FDA-2006-0260	FDA-2006-0260-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Guidance for Industry on Submitting and Reviewing Complete Responses to Clinical Holds	5/25/2006	null date	5/25/2006		E6-07983
FDA	FDA-2006-0260	FDA-2006-0260-0002	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Submitting and Reviewing Complete Responses to Clinical Holds	8/13/2007	null date	8/13/2007		E7-15740
FDA	FDA-2006-0261	FDA-2006-0261-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Requirements for Collection of Data Relating to the Prevention of Medical Gas Mixups at Health Care Facilities--Survey	5/25/2006	null date	5/25/2006		E6-07988
FDA	FDA-2006-0262	FDA-2006-0262-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Financial Disclosure by Clinical Investigators	5/25/2006	null date	5/25/2006		E6-07987
FDA	FDA-2006-0263	FDA-2006-0263-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; User Fee Cover Sheet; Form FDA 3397	5/25/2006	null date	5/25/2006		E6-07985
FDA	FDA-2006-0264	FDA-2006-0264-0001	Orthopaedic and Rehabilitation Devices Panel of the Medical Devices Advisory Committee; Amendment of Notice	5/26/2006	null date	5/26/2006		E6-08088
FDA	FDA-2006-0265	FDA-2006-0265-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Investigational Device Exemptions Reports and Records	5/26/2006	null date	5/26/2006		E6-08125
FDA	FDA-2006-0266	FDA-2006-0266-0001	Oral Dosage Form New Animal Drugs; Trimethoprim and Sulfadiazine Oral Paste	5/31/2006	null date	5/31/2006		E6-08303
FDA	FDA-2006-0267	FDA-2006-0267-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Prior Notice of Imported Food Under the Public Health Security and Bioterrorism Preparedness and Response Act of 2002	5/31/2006	null date	5/31/2006		E6-08311

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2006-0268	FDA-2006-0268-0001	Implantation or Injectable Dosage Form New Animal Drugs; Trimethoprim and Sulfadiazine	5/31/2006	null date	5/31/2006		E6-08309
FDA	FDA-2006-0269	FDA-2006-0269-0001	New Animal Drugs for Use in Animal Feeds; Melengestrol, Ractopamine, Monensin, and Tylosin	6/1/2006	null date	6/1/2006		E6-08420
FDA	FDA-2006-0270	FDA-2006-0270-0001	Guidance for Industry on Chemistry, Manufacturing, and Controls Information; Withdrawal and Revision of Seven Guidances	6/1/2006	null date	6/1/2006		E6-08417
FDA	FDA-2006-0271	FDA-2006-0271-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Applications for Food and Drug Administration Approval to Market a New Drug: Patent Submission and Listing Requirements and Application of 30-Month Stays on Approval of Abbreviated New Drug Applications Certifying That a Patent Claiming a Drug Is Valid or Will Not Be Infringed	6/2/2006	null date	6/2/2006		E6-08570
FDA	FDA-2006-0272	FDA-2006-0272-0001	Agency Information Collection Activities: Proposed Collection; Comment Request; Survey of Health Care Professionals on the Food Safety and Nutrition Information That They Provide to Pregnant Women	6/2/2006	null date	6/2/2006		E6-08566
FDA	FDA-2006-0273	FDA-2006-0273-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Draft Guidance for Industry and Food and Drug Administration Staff on Class II Special Controls Guidance Document: Automated Blood Cell Separator Device Operating by Centrifugal or Filtration Separation Principle	6/2/2006	null date	6/2/2006		E6-08571
FDA	FDA-2006-0274	FDA-2006-0274-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Investigational New Drug Regulations	6/2/2006	null date	6/2/2006		E6-08568
FDA	FDA-2006-0275	FDA-2006-0275-0001	International Conference on Harmonisation; Guidance on Q9 Quality Risk Management; Availability	6/2/2006	null date	6/2/2006		E6-08573
FDA	FDA-2006-0276	FDA-2006-0276-0002	Listing of Color Additives Exempt From Certification; Mica-Based Pearlescent Pigments; Confirmation of Effective Date	9/15/2006	null date	9/15/2006		E6-15275

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Food and Drug Administration (FDA) -- 1547 Documents

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Food and Drug Administration (FDA)								
FDA	FDA-2006-0276	FDA-2006-0276-0001	Listing of Color Additives Exempt From Certification; Mica-Based Pearlescent Pigments	6/2/2006	7/3/2006	6/2/2006		E6-08575
FDA	FDA-2006-0277	FDA-2006-0277-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Registration of Food Facilities Under the Public Health Security and Bioterrorism Preparedness and Response Act of 2002	6/2/2006	null date	6/2/2006		E6-08574
FDA	FDA-2006-0278	FDA-2006-0278-0001	Guidance for Industry on Chronic Cutaneous Ulcer and Burn Wounds--Developing Products for Treatment; Availability	6/2/2006	null date	6/2/2006		E6-08572
FDA	FDA-2006-0279	FDA-2006-0279-0001	Guidance for Industry on Antiviral Product Development-- Conducting and Submitting Virology Studies to the Agency; Availability	6/5/2006	null date	6/5/2006		E6-08635
FDA	FDA-2006-0280	FDA-2006-0280-0001	Implantation or Injectable Dosage Form New Animal Drugs; Oxytetracycline Injection, 200 Milligram/Milliliter	6/6/2006	null date	6/6/2006		E6-08694
FDA	FDA-2006-0281	FDA-2006-0281-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Administrative Detention and Banned Medical Devices	6/7/2006	null date	6/7/2006		E6-08838
FDA	FDA-2006-0282	FDA-2006-0282-0001	Medical Devices; Exception From General Requirements for Informed Consent	8/7/2006	1/0/1900	6/7/2006	0910-AC25	E6-08790
FDA	FDA-2006-0283	FDA-2006-0283-0001	Medical Devices; Ear, Nose, and Throat Devices; Classification of Olfactory Test Device	6/7/2006	null date	6/7/2006		E6-08791
FDA	FDA-2006-0284	FDA-2006-0284-0001	Guidance for Industry and Food and Drug Administration Staff; Class II Special Controls Guidance Document: Olfactory Test Device; Availability	6/7/2006	null date	6/7/2006		E6-08792
FDA	FDA-2006-0285	FDA-2006-0285-0001	Industry Exchange Workshop on Food and Drug Administration Clinical Trial Requirements; Public Workshop	6/8/2006	null date	6/8/2006		E6-08896
FDA	FDA-2006-0286	FDA-2006-0286-0001	Oral Dosage Form New Animal Drugs; Oxibendazole Suspension	6/8/2006	null date	6/8/2006		E6-08953
FDA	FDA-2006-0287	FDA-2006-0287-0001	Oral Dosage Form New Animal Drugs; Oxibendazole Paste	6/8/2006	null date	6/8/2006		E6-08894
FDA	FDA-2006-0288	FDA-2006-0288-0001	Guidance on Marketed Unapproved Drugs; Compliance Policy Guide; Availability	6/9/2006	null date	6/9/2006		E6-09032
FDA	FDA-2006-0289	FDA-2006-0289-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; CETROTIDE	6/9/2006	null date	6/9/2006		E6-09031

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Food and Drug Administration (FDA) -- 1547 Documents

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Food and Drug Administration (FDA)								
FDA	FDA-2006-0290	FDA-2006-0290-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Evaluation of Consumer-Friendly Formats for Brief Summary in Direct-to-Consumer Print Advertisements for Prescription Drugs: Study 1	6/9/2006	null date	6/9/2006		E6-08981
FDA	FDA-2006-0291	FDA-2006-0291-0001	Carbinoxamine Products; Enforcement Action Dates	6/9/2006	null date	6/9/2006		E6-09033
FDA	FDA-2006-0292	FDA-2006-0292-0001	Alltech, Inc.; Withdrawal of Food Additive Petition	6/9/2006	null date	6/9/2006		E6-08982
FDA	FDA-2006-0293	FDA-2006-0293-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; UROXATRAL	6/13/2006	null date	6/13/2006		E6-09201
FDA	FDA-2006-0294	FDA-2006-0294-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; INCRELEX	6/13/2006	null date	6/13/2006		E6-09138
FDA	FDA-2006-0295	FDA-2006-0295-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; RESTYLANE	6/13/2006	null date	6/13/2006		E6-09213
FDA	FDA-2006-0296	FDA-2006-0296-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; LUVERIS	6/13/2006	null date	6/13/2006		E6-09139
FDA	FDA-2006-0297	FDA-2006-0297-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; TYGACIL	6/13/2006	null date	6/13/2006		E6-09214
FDA	FDA-2006-0298	FDA-2006-0298-0002	Prescription Drug Marketing Act Pedigree Requirements under 21 CFR Part 203 Compliance Policy Guide and Guidance for Industry: Prescription Drug Marketing Act Pedigree Requirements Questions and Answers; Notice of Availability	11/15/2006	null date	11/15/2006		06-09211
FDA	FDA-2006-0298	FDA-2006-0298-0001	Prescription Drug Marketing Act Pedigree Requirements; Effective Date and Compliance Policy Guide; Request for Comment	6/14/2006	7/14/2006	6/14/2006		06-05362
FDA	FDA-2006-0299	FDA-2006-0299-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; CUBICIN	6/14/2006	null date	6/14/2006		E6-09225
FDA	FDA-2006-0300	FDA-2006-0300-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; DUTASTERIDE	6/14/2006	null date	6/14/2006		E6-09224
FDA	FDA-2006-0301	FDA-2006-0301-0001	International Cooperation on Harmonisation of Technical Requirements for Registration of Veterinary Medicinal Products; Final Guidance for Industry on ``Specifications: Test Procedures and Acceptance Criteria for New Biotechnological/Biological Veterinary Medicinal Products; Availability	6/15/2006	null date	6/15/2006		E6-09324
FDA	FDA-2006-0302	FDA-2006-0302-0001	New Animal Drugs for Use in Animal Feeds; Lasalocid; Correction	6/15/2006	null date	6/15/2006		E6-09321

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2006-0303	FDA-2006-0303-0001	International Cooperation on Harmonisation of Technical Requirements for Registration of Veterinary Medicinal Products; Final Guidance for Industry on ``Specifications: Test Procedures and Acceptance Criteria for New Veterinary Drug Substances and New Medicinal Products: Chemical Substances; Availability	6/15/2006	null date	6/15/2006		E6-09327
FDA	FDA-2006-0304	FDA-2006-0304-0001	Georgia-Pacific Resins, Inc.; Filing of Food Additive Petition	6/15/2006	null date	6/15/2006		E6-09319
FDA	FDA-2006-0305	FDA-2006-0305-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; INSPRA	6/1/2006	null date	6/16/2006		E6-09412
FDA	FDA-2006-0306	FDA-2006-0306-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; SYMLIN	6/1/2006	null date	6/16/2006		E6-09414
FDA	FDA-2006-0307	FDA-2006-0307-0001	Merck and Co., Inc., et al.; Withdrawal of Approval of 65 New Drug Applications and 52 Abbreviated New Drug Applications	6/16/2006	null date	6/16/2006		E6-09440
FDA	FDA-2006-0308	FDA-2006-0308-0001	Guidance for Industry and Food and Drug Administration Staff; the Review and Inspection of Premarket Approval Application Manufacturing Information and Operations; Availability	6/19/2006	null date	6/19/2006		E6-09505
FDA	FDA-2006-0309	FDA-2006-0309-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; ROZEREM	6/19/2006	null date	6/19/2006		E6-09509
FDA	FDA-2006-0310	FDA-2006-0310-0001	Draft Guidance for Industry and Food and Drug Administration Staff; the Review and Inspection of Premarket Approval Applications Under the Bioresearch Monitoring Program; Availability	6/20/2006	null date	6/20/2006		E6-09653
FDA	FDA-2006-0311	FDA-2006-0311-0001	Ophthalmic Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting	6/20/2006	null date	6/20/2006		E6-09601
FDA	FDA-2006-0312	FDA-2006-0312-0001	The Essentials of Food and Drug Administration Device Regulations: A Primer for Manufacturers and Suppliers; Public Workshop	6/21/2006	null date	6/21/2006		06-05570
FDA	FDA-2006-0313	FDA-2006-0313-0001	Guidance for Industry; Recommendations for the Early Food Safety Evaluation of New Non-Pesticidal Proteins Produced by New Plant Varieties Intended for Food Use; Availability	6/21/2006	null date	6/21/2006		E6-09688

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2006-0314	FDA-2006-0314-0001	Medical Devices; Availability of Safety and Effectiveness Summaries for Premarket Approval Applications	6/22/2006	null date	6/22/2006		E6-09898
FDA	FDA-2006-0315	FDA-2006-0315-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Guidance for Requesting an Extension to Use Existing Label Stock After the Trans Fat Labeling Effective Date of January 1, 2006	6/22/2006	null date	6/22/2006		E6-09824
FDA	FDA-2006-0316	FDA-2006-0316-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; CIALIS	2/1/2006	null date	6/22/2006		E6-09899
FDA	FDA-2006-0317	FDA-2006-0317-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Substances Generally Recognized as Safe: Notification Procedure	6/22/2006	null date	6/22/2006		E6-09827
FDA	FDA-2006-0318	FDA-2006-0318-0001	Over-the-Counter Drug Products; Safety and Efficacy Review; Additional Laxative Ingredient	6/22/2006	null date	6/22/2006		E6-09896
FDA	FDA-2006-0319	FDA-2006-0319-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; BONIVA	2/1/2006	null date	6/22/2006		E6-09817
FDA	FDA-2006-0320	FDA-2006-0320-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Recommendations for Early Food Safety Evaluation of New Non-Pesticidal Proteins Produced by New Plant Varieties Intended for Food Use	6/22/2006	null date	6/22/2006		E6-09826
FDA	FDA-2006-0321	FDA-2006-0321-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Product Jurisdiction: Assignment of Agency Component for Review of Premarket Applications	6/22/2006	null date	6/22/2006		E6-09900
FDA	FDA-2006-0322	FDA-2006-0322-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Infectious Disease Issues in Xenotransplantation	6/22/2006	null date	6/22/2006		E6-09816
FDA	FDA-2006-0323	FDA-2006-0323-0001	Food and Drug Administration Modernization Act of 1997: Modifications to the List of Recognized Standards, Recognition List Number: 015	6/23/2006	null date	6/23/2006		E6-09959
FDA	FDA-2006-0324	FDA-2006-0324-0001	Oral Dosage Form New Animal Drugs; Oxytetracycline	6/27/2006	null date	6/27/2006		E6-10053
FDA	FDA-2006-0325	FDA-2006-0325-0001	Blood Products Advisory Committee; Notice of Meeting	6/28/2006	null date	6/28/2006		06-05870

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2006-0326	FDA-2006-0326-0001	Draft Guidance for Industry: Analytical Methods Description for Type C Medicated Feeds; Availability	6/28/2006	null date	6/28/2006		06-05860
FDA	FDA-2006-0327	FDA-2006-0327-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Request for Samples and Protocols	6/29/2006	null date	6/29/2006		06-05805
FDA	FDA-2006-0328	FDA-2006-0328-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Medical Device User Fee Cover Sheet	6/29/2006	null date	6/29/2006		06-05806
FDA	FDA-2006-0329	FDA-2006-0329-0001	Program Priorities in the Center for Food Safety and Applied Nutrition; Request for Comments	6/29/2006	null date	6/29/2006		E6-10241
FDA	FDA-2006-0330	FDA-2006-0330-0001	The Use of Bayesian Statistics in Medical Device Clinical Trials; Public Meeting	6/29/2006	null date	6/29/2006		06-05804
FDA	FDA-2006-0331	FDA-2006-0331-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Importers Entry Notice	6/30/2006	null date	6/30/2006		E6-10271
FDA	FDA-2006-0332	FDA-2006-0332-0001	Emerging Clostridial Disease; Public Workshop; Reopening of the Administrative Record	7/5/2006	null date	7/5/2006		E6-10409
FDA	FDA-2006-0333	FDA-2006-0333-0001	Oral Dosage Form New Animal Drugs; Oxytetracycline Hydrochloride Soluble Powder	7/5/2006	null date	7/5/2006		E6-10445
FDA	FDA-2006-0334	FDA-2006-0334-0001	Oral Dosage Form New Animal Drugs; Griseofulvin	7/5/2006	null date	7/5/2006		E6-10406
FDA	FDA-2006-0335	FDA-2006-0335-0001	Oral Dosage Form New Animal Drugs; Ivermectin Liquid	7/5/2006	null date	7/5/2006		E6-10444
FDA	FDA-2006-0336	FDA-2006-0336-0001	Determination ofRegulatory Review Period for Purposes of Patent Extension; TAXUS EXPRESS Paclitaxel-Eluting Coronary Stent System	7/5/2006	null date	7/5/2006		E6-10408
FDA	FDA-2006-0337	FDA-2006-0337-0001	Ophthalmic and Topical Dosage Form New Animal Drugs; Copper Naphthenate Solution	7/5/2006	null date	7/5/2006		E6-10407
FDA	FDA-2006-0338	FDA-2006-0338-0001	Ophthalmic and Topical Dosage Form New Animal Drugs; Gentamicin Sulfate, Betamethasone Valerate, Clotrimazole Ointment	7/6/2006	null date	7/6/2006		E6-10496
FDA	FDA-2006-0339	FDA-2006-0339-0001	Streptomycin Residues in Cattle Tissues; Withdrawal of Compliance Policy Guide	7/1/2006	null date	7/1/2006		E6-10671
FDA	FDA-2006-0340	FDA-2006-0340-0001	Human-Labeled Drugs Distributed and Used in Animal Medicine; Withdrawal of Compliance Policy Guide	7/7/2006	null date	7/7/2006		E6-10672

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2006-0341	FDA-2006-0341-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; MULTIHANCE	7/11/2006	null date	7/11/2006		E6-10796
FDA	FDA-2006-0342	FDA-2006-0342-0001	Implantation or Injectable Dosage Form New Animal Drugs; Hyaluronate Sodium Injection	7/12/2006	null date	7/12/2006		E6-10879
FDA	FDA-2006-0343	FDA-2006-0343-0001	New Animal Drugs for Use in Animal Feeds; Melengestrol, Lasalocid, and Tylosin	12/1/2006	null date	12/1/2006		E6-10878
FDA	FDA-2006-0344	FDA-2006-0344-0001	Oral Dosage Form New Animal Drugs; Clindamycin Capsules and Tablets	7/12/2006	null date	7/12/2006		E6-10877
FDA	FDA-2006-0345	FDA-2006-0345-0001	Implantation or Injectable Dosage Form New Animal Drugs; Furosemide	7/13/2006	null date	7/13/2006		E6-10974
FDA	FDA-2006-0346	FDA-2006-0346-0001	Implantation or Injectable Dosage Form New Animal Drugs; Mepivacaine	7/13/2006	null date	7/13/2006		E6-10970
FDA	FDA-2006-0347	FDA-2006-0347-0001	New Animal Drugs; Ceftiofur	7/13/2006	null date	7/13/2006		E6-10973
FDA	FDA-2006-0348	FDA-2006-0348-0001	Guidance for Industry on Providing Regulatory Submissions to the Center for Biologics Evaluation and Research in Electronic Format--Lot Release Protocols; Availability	7/13/2006	null date	7/13/2006		E6-11040
FDA	FDA-2006-0349	FDA-2006-0349-0001	New Animal Drugs; Ceftiofur	7/13/2006	null date	7/13/2006		E6-10972
FDA	FDA-2006-0350	FDA-2006-0350-0001	Oral Dosage Form New Animal Drugs; Clindamycin Liquid	7/13/2006	null date	7/13/2006		E6-10971
FDA	FDA-2006-0351	FDA-2006-0351-0001	Medical Devices; Anesthesiology Devices; Neurological Devices; Denial of Request for Change in Classification of Breathing Frequency Monitor and Electroencephalograph	7/14/2006	null date	7/14/2006		E6-11115
FDA	FDA-2006-0352	FDA-2006-0352-0001	Determination That PHENERGAN (Promethazine Hydrochloride) Tablets, 12.5 Milligrams and 50 Milligrams, Were Not Withdrawn From Sale for Reasons of Safety or Effectiveness	7/14/2006	null date	7/14/2006		E6-11072
FDA	FDA-2006-0353	FDA-2006-0353-0001	Oral Dosage Form New Animal Drugs; Ivermectin Paste	7/14/2006	null date	7/14/2006		E6-11073
FDA	FDA-2006-0354	FDA-2006-0354-0001	Request for Nominations for Voting Members on a Public Advisory Committee; Pediatric Advisory Committee	7/18/2006	null date	7/18/2006		06-06276
FDA	FDA-2006-0355	FDA-2006-0355-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Draft Guidance: Emergency Use Authorization of Medical Products	7/18/2006	null date	7/18/2006		E6-11287
FDA	FDA-2006-0356	FDA-2006-0356-0001	Guidance on Useful Written Consumer Medication Information; Availability	7/18/2006	null date	7/18/2006		E6-11329

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Food and Drug Administration (FDA)								
FDA	FDA-2006-0357	FDA-2006-0357-0001	Joint Meeting of the Endocrinologic and Metabolic Drugs Advisory Committee and the Advisory Committee for Pharmaceutical Science; Notice of Meeting	7/19/2006	null date	7/19/2006		E6-11471
FDA	FDA-2006-0358	FDA-2006-0358-0001	Listing of Color Additives Exempt From Certification; Mica-Based Pearlescent Pigments	7/20/2006	null date	7/20/2006		E6-11536
FDA	FDA-2006-0359	FDA-2006-0359-0001	Advisory Committee for Reproductive Health Drugs; Notice of Meeting	7/20/2006	null date	7/20/2006		E6-11538
FDA	FDA-2006-0360	FDA-2006-0360-0001	Psychopharmacologic Drugs Advisory Committee; Notice of Meeting	7/20/2006	null date	7/20/2006		E6-11537
FDA	FDA-2006-0361	FDA-2006-0361-0001	Draft Manufactured Food Regulatory Program Standards; Availability	7/20/2006	null date	7/20/2006		E6-11539
FDA	FDA-2006-0362	FDA-2006-0362-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Food Labeling; Notification Procedures for Statements on Dietary Supplements	7/24/2006	null date	7/24/2006		E6-11642
FDA	FDA-2006-0363	FDA-2006-0363-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Food and Drug Administration Public Health Notification (formerly known as Safety Alert/Public Health Advisory) Readership Survey	7/24/2006	null date	7/24/2006		E6-11644
FDA	FDA-2006-0364	FDA-2006-0364-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Survey of Need for Online Medical Device Survey	7/24/2006	null date	7/24/2006		E6-11640
FDA	FDA-2006-0365	FDA-2006-0365-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Guidance for Industry on Continuous Marketing Applications: Pilot 2--Scientific Feedback and Interactions During Development of Fast Track Products Under the Prescription Drug User Fee Act	7/24/2006	null date	7/24/2006		E6-11643
FDA	FDA-2006-0366	FDA-2006-0366-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Bar Code Label Requirement for Human Drug and Biological Products	7/24/2006	null date	7/24/2006		E6-11641
FDA	FDA-2006-0367	FDA-2006-0367-0001	Obstetrics and Gynecology Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting	7/25/2006	null date	7/25/2006		E6-11773

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AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2006-0368	FDA-2006-0368-0001	Medical Device Regulations; Addresses; Technical Amendment	7/25/2006	null date	7/25/2006		E6-11777
FDA	FDA-2006-0369	FDA-2006-0369-0001	Anti-Infective Drugs Advisory Committee; Notice of Meeting	7/25/2006	null date	7/25/2006		E6-11772
FDA	FDA-2006-0370	FDA-2006-0370-0001	Nonprescription Drugs Advisory Committee; Notice of Meeting	7/25/2006	null date	7/25/2006		E6-11774
FDA	FDA-2006-0371	FDA-2006-0371-0002	Food Labeling; Guidelines for Voluntary Nutrition Labeling of Raw Fruits, Vegetables, and Fish; Correction	8/17/2006	null date	8/17/2006		06-06957
FDA	FDA-2006-0371	FDA-2006-0371-0001	Food Labeling; Guidelines for Voluntary Nutrition Labeling of Raw Fruits, Vegetables, and Fish	7/25/2006	null date	7/25/2006		06-06436
FDA	FDA-2006-0372	FDA-2006-0372-0001	General and Plastic Surgery Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting	7/25/2006	null date	7/25/2006		E6-11775
FDA	FDA-2006-0373	FDA-2006-0373-0001	Over-the-Counter Drug Products; Safety and Efficacy Review; Additional Sunscreen Ingredient	7/26/2006	null date	7/26/2006		E6-11874
FDA	FDA-2006-0374	FDA-2006-0374-0001	Medical Devices; Immunology and Microbiology Devices; Classification of Fecal Calprotectin Immunological Test Systems	7/27/2006	null date	7/27/2006		E6-11975
FDA	FDA-2006-0375	FDA-2006-0375-0001	Guidance for Industry and Food and Drug Administration Staff; Class II Special Controls Guidance Document: Fecal Calprotectin Immunological Test Systems; Availability	7/27/2006	null date	7/27/2006		E6-11974
FDA	FDA-2006-0376	FDA-2006-0376-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Guidance for Industry on Special Protocol Assessment	7/31/2006	null date	7/31/2006		E6-12158
FDA	FDA-2006-0377	FDA-2006-0377-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; FDA Survey of Physicians Perceptions of the Impact of Early Risk Communication About Medical Products	7/31/2006	null date	7/31/2006		E6-12159
FDA	FDA-2006-0378	FDA-2006-0378-0001	Draft Manufactured Food Regulatory Program Standards; Availability; Correction	7/31/2006	null date	7/31/2006		E6-12179
FDA	FDA-2006-0379	FDA-2006-0379-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Establishing and Maintaining a List of United States Dairy Product Manufacturers/Processors With Interest in Exporting to Chile	7/31/2006	null date	7/31/2006		E6-12160

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Food and Drug Administration (FDA)								
FDA	FDA-2006-0380	FDA-2006-0380-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Guidance for Clinical Trial Sponsors: Establishment and Operation of Clinical Trial Data Monitoring Committees	7/31/2006	null date	7/31/2006		E6-12157
FDA	FDA-2006-0381	FDA-2006-0381-0001	Oncologic Drugs Advisory Committee; Notice of Meeting	8/1/2006	null date	8/1/2006		E6-12270
FDA	FDA-2006-0382	FDA-2006-0382-0001	Cold, Cough, Allergy, Bronchodilator, and Antiasthmatic Drug Products for Over-the-Counter Human Use; Amendment of Monograph for OTC Nasal Decongestant Drug Products	8/1/2006	null date	8/1/2006	0910-AF34	E6-12265
FDA	FDA-2006-0383	FDA-2006-0383-0001	Memorandum of Understanding Between the U.S. Food and Drug Administration, Department of Health and Human Services and the Centers for Disease Control and Prevention	8/1/2006	null date	8/1/2006		06-06603
FDA	FDA-2006-0384	FDA-2006-0384-0001	Cardiovascular and Renal Drugs Advisory Committee; Notice of Meeting	8/1/2006	null date	8/1/2006		E6-12269
FDA	FDA-2006-0385	FDA-2006-0385-0001	Current Good Manufacturing Practice, Quality Control Procedures, Quality Factors, Notification Requirements, and Records and Reports for the Production of Infant Formula; Reopening of the Comment Period	8/1/2006	9/15/2006	8/1/2006	0910-AA04	E6-12268
FDA	FDA-2006-0386	FDA-2006-0386-0001	Meeting to Present Work-In-Progress on a Method for RankingFeed Contaminants According to the Relative Risks They Pose to Animal and Public Health; Part 1: Health Consequence Scoring for Feed Contaminants	8/1/2006	null date	8/1/2006		E6-12266
FDA	FDA-2006-0387	FDA-2006-0387-0001	Medical Device User Fee Rates for Fiscal Year 2007	8/2/2006	null date	8/2/2006		E6-12394
FDA	FDA-2006-0388	FDA-2006-0388-0001	Animal Drug User Fee Rates and Payment Procedures for Fiscal Year 2007	8/2/2006	null date	8/2/2006		E6-12396
FDA	FDA-2006-0389	FDA-2006-0389-0001	Prescription Drug User Fee Rates for Fiscal Year 2007	8/2/2006	null date	8/2/2006		E6-12397
FDA	FDA-2006-0390	FDA-2006-0390-0001	Transmissible Spongiform Encephalopathies Advisory Committee; Notice of Meeting	8/3/2006	null date	8/3/2006		E6-12567
FDA	FDA-2006-0391	FDA-2006-0391-0001	National Mammography Quality Assurance Advisory Committee; Notice of Meeting	8/3/2006	null date	8/3/2006		E6-12569
FDA	FDA-2006-0392	FDA-2006-0392-0001	New Animal Drugs; Change of Sponsor; Isoflurane	8/3/2006	null date	8/3/2006		E6-12570
FDA	FDA-2006-0393	FDA-2006-0393-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; CLINACOX	8/3/2006	null date	8/3/2006		E6-12572

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2006-0394	FDA-2006-0394-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; CYMBALTA	8/3/2006	null date	8/3/2006		E6-12574
FDA	FDA-2006-0395	FDA-2006-0395-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; IPLEX	8/3/2006	null date	8/3/2006		E6-12571
FDA	FDA-2006-0396	FDA-2006-0396-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; EMEND	8/3/2006	10/2/2006	8/3/2006		E6-12573
FDA	FDA-2006-0397	FDA-2006-0397-0001	Oral Dosage Form New Animal Drugs; Kanamycin, Bismuth Subcarbonate, Activated Attapulgite	8/3/2006 12:00	null date	8/3/2006		E6-12568
FDA	FDA-2006-0398	FDA-2006-0398-0001	Reports and guidance documents; availability, etc.: International Conference on Harmonisation--	8/8/2006	null date	8/8/2006		E6-12806
FDA	FDA-2006-0399	FDA-2006-0399-0001	Practice and procedure: Electronic submissions gateway	8/8/2006	null date	8/8/2006		E6-12808
FDA	FDA-2006-0400	FDA-2006-0400-0001	Reports and guidance documents; availability, etc.: InternationalConference on Harmonisation--	8/1/2006	null date	8/8/2006		E6-12807
FDA	FDA-2006-0401	FDA-2006-0401-0001	Animal drugs, feeds, and related products: Oxytetracycline	8/8/2006	null date	8/8/2006		E6-12862
FDA	FDA-2006-0402	FDA-2006-0402-0001	Meetings: Science Advisory Board	8/8/2006	null date	8/8/2006		E6-12863
FDA	FDA-2006-0403	FDA-2006-0403-0001	Antiviral Drugs Advisory Committee; Notice of Meeting	8/9/2006	null date	8/9/2006		E6-12890
FDA	FDA-2006-0404	FDA-2006-0404-0001	Food and Drug Administration-Regulated Products Containing Nanotechnology Materials; Public Meeting	1/1/2006	null date	1/1/2006		06-06867
FDA	FDA-2006-0405	FDA-2006-0405-0001	Industry Exchange Workshop on Food and Drug Administration Clinical Trial Requirements; Public Workshop	8/11/2006	null date	8/11/2006		E6-13114
FDA	FDA-2006-0406	FDA-2006-0406-0001	Unique Device Identification; Request for Comments	8/8/2006	null date	8/11/2006		06-06870
FDA	FDA-2006-0407	FDA-2006-0407-0001	Draft Guidance for Industry on an Amendment Involving Donor Deferral for Transfusion in France Since 1980 to ``Guidance for Industry: Revised Preventive Measures to Reduce the Possible Risk of Transmission of Creutzfeldt-Jakob Disease (CJD) and Variant Creutzfeldt-Jakob Disease (vCJD) by Blood and Blood Products; Availability	8/14/2006	null date	8/14/2006		E6-13234
FDA	FDA-2006-0408	FDA-2006-0408-0001	Guidance for Industry on Implementing a Collection Program for Source Plasma Containing Disease-Associated and Other Immunoglobulin G (IgG) Antibodies; Availability	8/14/2006	8/14/2006	8/14/2006		E6-13233

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2006-0409	FDA-2006-0409-0001	Psychopharmacologic Drugs Advisory Committee; Amendment of Notice	8/17/2006	null date	8/17/2006		E6-13502
FDA	FDA-2006-0410	FDA-2006-0410-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; MedWatch--The Food and Drug Administration Safety Information and Adverse Event Reporting Program; Proposal to Survey MedWatch Partners Organizations	8/17/2006	null date	8/17/2006		E6-13503
FDA	FDA-2006-0411	FDA-2006-0411-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Requirements for Collection of Data Relating to the Prevention of Medical Gas Mixups at Health Care Facilities--Survey	8/17/2006	null date	8/17/2006		E6-13565
FDA	FDA-2006-0412	FDA-2006-0412-0001	Advisory Committee for Pharmaceutical Science; Notice of Meeting	8/17/2006	null date	8/17/2006		E6-13506
FDA	FDA-2006-0413	FDA-2006-0413-0001	Heparin Catheter Lock-Flush Solutions; Transfer of Primary Responsibility from Center for Drug Evaluation and Research to Center for Devices and Radiological Health	8/17/2006	null date	8/17/2006		E6-13509
FDA	FDA-2006-0414	FDA-2006-0414-0001	Preparation for International Conference on Harmonization Meetings in Chicago, Illinois; Public Meeting	8/17/2006	null date	8/17/2006		E6-13505
FDA	FDA-2006-0415	FDA-2006-0415-0001	Draft Guidance for Industry; Animal Drug User Fees: Fees Exceed Costs Waivers and Reductions; Availability	8/17/2006	null date	8/17/2006		E6-13507
FDA	FDA-2006-0416	FDA-2006-0416-0001	Agency information collection activities; proposals, submissions, and approvals	8/18/2006	null date	8/18/2006		E6-13609
FDA	FDA-2006-0417	FDA-2006-0417-0001	Food Additives Permitted for Direct Addition to Food for Human Consumption; Bacteriophage Preparation	8/18/2006	8/18/2006	8/18/2006		E6-13621
FDA	FDA-2006-0418	FDA-2006-0418-0001	Molecular Methods in Immunohematology; Public Workshop	8/21/2006	null date	8/21/2006		E6-13695
FDA	FDA-2006-0419	FDA-2006-0419-0003	Index of Legally Marketed Unapproved New Animal Drugs for Minor Species		null date	12/6/2007	0910-AF67	E7-23580
FDA	FDA-2006-0419	FDA-2006-0419-0001	Index of Legally Marketed Unapproved New Animal Drugs for Minor Species	8/22/2006	12/20/2006	8/22/2006	0910-AF67	06-07070
FDA	FDA-2006-0419	FDA-2006-0419-0002	Index of Legally Marketed Unapproved New Animal Drugs for Minor Species; Extension of Comment Period	10/2/2006	null date	10/2/2006	0910-AF67	E6-16208
FDA	FDA-2006-0420	FDA-2006-0420-0001	Veterinary Medicine Advisory Committee; Notice of Meeting	8/22/2006	null date	8/22/2006		E6-13818

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Food and Drug Administration (FDA) -- 1547 Documents

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Food and Drug Administration (FDA)								
FDA	FDA-2006-0421	FDA-2006-0421-0001	Orthopaedic and Rehabilitation Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting	8/22/2006	null date	8/22/2006		E6-13823
FDA	FDA-2006-0422	FDA-2006-0422-0001	Memorandum of Understanding Between the U.S. Food and Drug Administration, the National Cancer Institute, and the National Institute of Standards and Technology	8/24/2006	null date	8/24/2006		06-07127
FDA	FDA-2006-0423	FDA-2006-0423-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Inspection by Accredited Persons Program Under the Medical Device User Fee and Modernization Act of 2002	8/24/2006	null date	8/24/2006		E6-14056
FDA	FDA-2006-0424	FDA-2006-0424-0001	Food Safety and Security Monitoring Project--Radiological Health; Announcement Type: Cooperative Agreements Under a Limited Competition; Funding Opportunity Number: Request for Applications: RFA-FDA-ORA-2006-4; Catalog of Federal Domestic Assistance Number: 93.448	8/24/2006	null date	8/24/2006		06-07124
FDA	FDA-2006-0425	FDA-2006-0425-0001	National Center for Natural Products Research, University of Mississippi; Single Source Cooperative Agreement; Catalog of Federal Domestic Assistance Number 93.103; Request for Application	8/25/2006	null date	8/25/2006		E6-14109
FDA	FDA-2006-0426	FDA-2006-0426-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Medicated Feed Mill License Application--Extension	8/25/2006	null date	8/25/2006		E6-14076
FDA	FDA-2006-0427	FDA-2006-0427-0001	Allergenic Products Advisory Committee; Notice of Meeting	8/29/2006	null date	8/29/2006		E6-14295
FDA	FDA-2006-0428	FDA-2006-0428-0001	Clinical Pharmacology Subcommittee of the Advisory Committee for Pharmaceutical Science; Notice of Meeting	8/29/2006	null date	8/29/2006		E6-14296
FDA	FDA-2006-0429	FDA-2006-0429-0001	Cardiovascular and Renal Drugs Advisory Committee; Amendment of Notice	8/29/2006	null date	8/29/20061		E6-14294
FDA	FDA-2006-0430	FDA-2006-0430-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; User Fee Cover Sheet; Form FDA 3397	8/29/2006	null date	8/29/2006		E6-14266

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2006-0431	FDA-2006-0431-0002	Draft Guidance for Institutional Review Boards, Clinical Investigators, and Sponsors; Exception from Informed Consent Requirements for Emergency Research	8/29/2006	null date	8/29/2006		E6-14262
FDA	FDA-2006-0431	FDA-2006-0431-0001	Conduct of Emergency Clinical Research; Public Hearing	11/27/2006	null date	8/29/2006		E6-14264
FDA	FDA-2006-0432	FDA-2006-0432-0002	Requirements for Foreign and Domestic Establishment Registration and Listing for Human Drugs, Including Drugs That Are Regulated Under a Biologics License Application, and Animal Drugs; Public Meeting; Extension of Comment Period	10/31/2006	1/26/2007	10/31/2006	0910-AA49	E6-18310
FDA	FDA-2006-0432	FDA-2006-0432-0003	Requirements for Foreign and Domestic Establishment Registration and Listing for Human Drugs, Including Drugs That Are Regulated Under a Biologics License Application, and Animal Drugs; Reopening of Comment Period	2/8/2007	2/26/2007	2/8/2007	0910-AA49	E7-02123
FDA	FDA-2006-0432	FDA-2006-0432-0001	Requirements for Foreign and Domestic Establishment Registration and Listing for Human Drugs, Including Drugs that are Regulated Under a Biologics License Application, and Animal Drugs	8/29/2006	2/26/2007	8/29/2006	0910-AA49	06-07172
FDA	FDA-2006-0433	FDA-2006-0433-0001	Skin Bleaching Drug Products For Over-the-Counter Human Use; Proposed Rule	12/27/2006	null date	8/29/006	0910-AF53	E6-14263
FDA	FDA-2006-0434	FDA-2006-0434-0001	Psychopharmacologic Drugs Advisory Committee; Cancellation	8/29/2006	null date	8/29/2006		E6-14293
FDA	FDA-2006-0435	FDA-2006-0435-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Medical Device User Fee and Modernization Act Small Business Qualification Certification (Form FDA 3602)	8/29/2006	8/29/2006	8/29/2006		E6-14267
FDA	FDA-2006-0436	FDA-2006-0436-0001	Advisory Committee on Special Studies Relating to the Possible Long-Term Health Effects of Phenoxy Herbicides and Contaminants (Ranch Hand Advisory Committee); Notice of Meeting	8/30/2006	8/30/2006	8/30/2006		E6-14371
FDA	FDA-2006-0437	FDA-2006-0437-0001	Anti-Infective Drugs Advisory Committee Meeting; Amendment of Notice	8/31/2006	null date	8/31/2006		06-07310
FDA	FDA-2006-0438	FDA-2006-0438-0001	Nutrition Labeling of Dietary Supplements; Technical Amendment	8/31/2006	null date	8/31/2006		06-07306

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Food and Drug Administration (FDA)								
FDA	FDA-2006-0439	FDA-2006-0439-0001	Certain Other Dosage Form New Animal Drugs; Gentamicin Sulfate Intrauterine Solution	8/31/2006	null date	8/31/2006		06-07307
FDA	FDA-2006-0440	FDA-2006-0440-0001	Implantation or Injectable Dosage Form New Animal Drugs; Lincomycin	9/1/2006	null date	9/1/2006		E6-14509
FDA	FDA-2006-0441	FDA-2006-0441-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Food Additive Petitions	9/1/2006	null date	9/1/2006		E6-14510
FDA	FDA-2006-0442	FDA-2006-0442-0001	Oral Dosage Form New Animal Drugs; Carprofen	9/1/2006	null date	9/1/2006		E6-14508
FDA	FDA-2006-0443	FDA-2006-0443-0001	[Docket No. 2004N-0234]	9/1/2006	null date	9/1/2006		E6-14549
FDA	FDA-2006-0444	FDA-2006-0444-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Guidance on Informed Consent for In Vitro Diagnostic Device Studies Using Leftover Human Specimens That Are Not Identifiable	9/6/2006	null date	9/6/2006		E6-14671
FDA	FDA-2006-0445	FDA-2006-0445-0001	New Animal Drugs For Use in Animal Feeds; Amprolium	9/6/2006	null date	9/6/2006		E6-14673
FDA	FDA-2006-0446	FDA-2006-0446-0001	Draft Guidance for Industry, Clinical Laboratories, and FDA Staff on In Vitro Diagnostic Multivariate Index Assays; Availability	9/7/2006	null date	9/7/2006		06-07499
FDA	FDA-2006-0447	FDA-2006-0447-0001	Draft Guidance for Industry and Food and Drug Administration Staff; Commercially Distributed Analyte Specific Reagents (ASRs): Frequently Asked Questions; Availability	9/7/2006	null date	9/7/2006		06-07500
FDA	FDA-2006-0447	FDA-2006-0447-0002	Draft Guidance for Industry and Food and Drug Administration Staff; Commercially Distributed Analyte Specific Reagents (ASRs): Frequently Asked Questions; Availability; Extension of Comment Period	11/28/2006	null date	11/28/2006		E6-20030
FDA	FDA-2006-0448	FDA-2006-0448-0001	New Animal Drugs; Zilpaterol	9/8/2006	null date	9/8/2006		E6-14899
FDA	FDA-2006-0449	FDA-2006-0449-0001	Risk Communication on Medical Devices: Sharing Perspectives	9/8/2006	null date	9/8/2006		E6-14852
FDA	FDA-2006-0450	FDA-2006-0450-0001	New Animal Drugs For Use in Animal Feed; Oxytetracycline	9/8/2006	null date	9/8/2006		E6-14898
FDA	FDA-2006-0451	FDA-2006-0451-0001	Draft Guidance for Industry on Drug Interaction Studies--Study Design, Data Analysis, and Implications for Dosing and Labeling; Availability	9/12/2006	null date	9/12/2006		E6-15058

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Food and Drug Administration (FDA)								
FDA	FDA-2006-0452	FDA-2006-0452-0001	Cooperative Agreement to Support the Shellfish and Seafood Safety Assistance Project; Announcement Type: Single Source Application; Agency Funding Opportunity Number: RFA-FDA-CFSAN-2006-1	9/13/2006	null date	9/13/2006		E6-15102
FDA	FDA-2006-0453	FDA-2006-0453-0001	New Animal Drugs for Use in Animal Feeds; Chlortetracycline	9/13/2006	null date	9/13/2006		E6-15103
FDA	FDA-2006-0454	FDA-2006-0454-0001	Memorandum of Understanding Between the United States Food and Drug Administration and the National Cancer Institute	9/14/2006	null date	9/14/2006		06-07630
FDA	FDA-2006-0455	FDA-2006-0455-0001	Transmissible Spongiform Encephalopathies Advisory Committee; Amendment of Notice	9/15/2006	null date	9/15/2006		E6-15283
FDA	FDA-2006-0456	FDA-2006-0456-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; AVASTIN	9/20/2006	null date	9/20/2006		E6-15555
FDA	FDA-2006-0457	FDA-2006-0457-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; APTIVUS	9/20/2006	null date	9/20/2006		E6-15553
FDA	FDA-2006-0458	FDA-2006-0458-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; FUZEON	9/20/2006	null date	9/20/2006		E6-15554
FDA	FDA-2006-0459	FDA-2006-0459-0001	Draft Guidance for Industry on Public Availability of Labeling Changes in ``Changes Being Effected Supplements; Availability	9/20/2006	null date	9/20/2006		06-07983
FDA	FDA-2006-0460	FDA-2006-0460-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; CYDECTIN	9/20/2006	null date	9/20/2006		06-07800
FDA	FDA-2006-0461	FDA-2006-0461-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; MACUGEN	9/20/2006	null date	9/20/2006		E6-15556
FDA	FDA-2006-0462	FDA-2006-0462-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; MYCAMINE--New Drug Application 21-754	9/20/2006	null date	9/20/2006		06-07985
FDA	FDA-2006-0463	FDA-2006-0463-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Mammography Quality Standards Act Requirements	9/22/2006	null date	9/22/2006		06-08027
FDA	FDA-2006-0464	FDA-2006-0464-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Export of Medical Devices-Foreign Letters of Approval	9/22/2006	null date	9/22/2006		06-08026
FDA	FDA-2006-0465	FDA-2006-0465-0001	Unique Device Identification; Notice of Public Meeting	9/22/2006	null date	9/22/2006		06-07969
FDA	FDA-2006-0466	FDA-2006-0466-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Environmental Impact Considerations	9/22/2006	9/22/2006	9/22/2006		06-08025

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Food and Drug Administration (FDA)								
FDA	FDA-2006-0467	FDA-2006-0467-0001	Neurological Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting	9/22/2006	null date	9/22/2006		06-08114
FDA	FDA-2006-0468	FDA-2006-0468-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Records and Reports Concerning Experience With Approved New Animal Drugs	9/22/2006	null date	9/22/2006		06-08023
FDA	FDA-2006-0469	FDA-2006-0469-0003	Medical Devices; Reprocessed Single-Use Devices; Requirement for Submission of Validation Data; Withdrawal	1/12/2007	null date	1/12/2007		07-00105
FDA	FDA-2006-0469	FDA-2006-0469-0001	Medical Devices; Reprocessed Single-Use Devices; Requirement for Submission of Validation Data; Companion to Direct Final Rule	9/25/2006	12/11/2006	9/25/2006		06-08165
FDA	FDA-2006-0469	FDA-2006-0469-0002	Medical Devices; Reprocessed Single-Use Devices; Requirement for Submission of Validation Data	9/25/2006	12/11/2006	9/25/2006		06-08166
FDA	FDA-2006-0470	FDA-2006-0470-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; CLOLAR	5/1/2006	null date	9/25/2006		06-08115
FDA	FDA-2006-0471	FDA-2006-0471-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; KETEK	9/26/2006	null date	9/26/2006		E6-15690
FDA	FDA-2006-0472	FDA-2006-0472-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Draft Guidance for Industry and Food and Drug Administration Staff: Recommendations for Clinical Laboratory Improvement Amendments of 1988 Waiver Applications; Availability	9/26/2006	null date	9/26/2006		E6-15693
FDA	FDA-2006-0473	FDA-2006-0473-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Procedures for the Safe and Sanitary Processing and Importing of Fish and Fishery Products	9/26/2006	null date	9/26/2006		E6-15694
FDA	FDA-2006-0474	FDA-2006-0474-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; MYCAMINE--New Drug Application 21-506	9/26/2006	null date	9/26/2006		E6-15767
FDA	FDA-2006-0475	FDA-2006-0475-0001	Medical Devices; Availability of Safety and Effectiveness Summaries for Premarket Approval Applications	9/26/2006	null date	9/26/2006		E6-15755
FDA	FDA-2006-0476	FDA-2006-0476-0001	New Animal Drugs for Use in Animal Feeds; Lasalocid	9/26/2006	null date	9/26/2006		06-08261

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2006-0477	FDA-2006-0477-0001	Food and Drug Administration-Regulated Products Containing Nanotechnology Materials; Public Meeting	6/1/2006	null date	6/1/2006		E6-08242
FDA	FDA-2006-0478	FDA-2006-0478-0001	Oral Dosage Form New Animal Drugs; Amprolium Solution	9/27/2006	null date	9/27/2006		06-08275
FDA	FDA-2006-0479	FDA-2006-0479-0001	Determination of Regulatory ReviewPeriod for Purposes of Patent Extension; LYRICA (New Drug Application 21-446)	9/27/2006	null date	9/27/2006		E6-15908
FDA	FDA-2006-0480	FDA-2006-0480-0001	Oral Dosage Form New Animal Drugs; Neomycin	9/28/2006	null date	9/28/2006		E6-15889
FDA	FDA-2006-0481	FDA-2006-0481-0001	Ophthalmic and Topical Dosage Form New Animal Drugs; Gentamicin Sulfate, Betamethasone Valerate, Clotrimazole Ointment	9/28/2006	null date	9/28/2006		E6-15888
FDA	FDA-2006-0482	FDA-2006-0482-0001	Review of Agreements, Guidances, and Practices Specific to Assignment of Combination Products in Compliance With the Medical Device User Fee and Modernization Act of 2002; Request for Comments	9/28/2006	null date	9/28/2006		E6-15967
FDA	FDA-2006-0483	FDA-2006-0483-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; PLENAXIS	9/29/2006	null date	9/29/2006		E6-15969
FDA	FDA-2006-0484	FDA-2006-0484-0001	Guidances on Providing Regulatory Submissions in Electronic Format; Withdrawal of Guidances	9/29/2006	null date	9/29/2006		E6-15966
FDA	FDA-2006-0485	FDA-2006-0485-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; LYRICA (New Drug Application 21-723)	9/29/2006	null date	9/29/2006		E6-15962
FDA	FDA-2006-0486	FDA-2006-0486-0001	Draft Guidance for Industry on Characterization and Qualification of Cell Substrates and Other Biological Starting Materials Used in the Production of Viral Vaccines for the Prevention and Treatment of Infectious Diseases; Availability	9/29/2006	null date	9/29/2006		E6-15963
FDA	FDA-2006-0487	FDA-2006-0487-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; TARCEVA	9/29/2006	null date	9/29/2006		E6-15987
FDA	FDA-2006-0488	FDA-2006-0488-0001	Implantation or Injectable Dosage Form New Animal Drugs; Tulathromycin	9/29/2006	null date	9/29/2006		E6-15965
FDA	FDA-2006-0489	FDA-2006-0489-0001	Guidance for Industry on Quality Systems Approach to Pharmaceutical Current Good Manufacturing Practice Regulations; Availability	10/2/2006	null date	10/2/2006		E6-16215

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Food and Drug Administration (FDA) -- 1547 Documents

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Food and Drug Administration (FDA)								
FDA	FDA-2006-0490	FDA-2006-0490-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; NATRECOR	10/2/2006	null date	10/2/2006		E6-16091
FDA	FDA-2006-0491	FDA-2006-0491-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Proposed Collection; Comment Request; Guidance for Industry on Submitting and Reviewing Complete Responses to Clinical Holds	10/2/2006	null date	10/2/2006		E6-16225
FDA	FDA-2006-0492	FDA-2006-0492-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; BYETTA	2/1/2006	null date	10/2/2006		E6-16086
FDA	FDA-2006-0493	FDA-2006-0493-0001	Request for Nominations for Voting and Nonvoting Consumer Representative Members on Public Advisory Committees and Panels	10/2/2006	null date	10/2/2006		E6-16216
FDA	FDA-2006-0494	FDA-2006-0494-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Postmarket Surveillance	10/2/2006	null date	10/2/2006		E6-16231
FDA	FDA-2006-0495	FDA-2006-0495-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; DRAXXIN	10/2/2006	null date	10/2/2006		E6-16087
FDA	FDA-2006-0496	FDA-2006-0496-0001	Immunology Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting	10/3/2006	null date	10/3/2006		E6-16319
FDA	FDA-2006-0497	FDA-2006-0497-0001	Guidance for Industry on Bar Code Label Requirements--Questions and Answers; Availability	10/5/2006	null date	10/5/2006		E6-16436
FDA	FDA-2006-0498	FDA-2006-0498-0001	Request for Nominations for Nonvoting Members Representing Industry Interests on Public Advisory Panels or Committees	10/5/2006	null date	10/5/2006		E6-16438
FDA	FDA-2006-0499	FDA-2006-0499-0001	Workshop on Sex Differences and the Food and Drug Administration Critical Path Initiative	10/10/2006	null date	10/10/2006		E6-16605
FDA	FDA-2006-0500	FDA-2006-0500-0001	Oral Dosage Form New Animal Drugs; Omeprazole	10/10/2006	null date	10/10/2006		E6-16604
FDA	FDA-2006-0501	FDA-2006-0501-0001	Request for Nominations for Voting Members on Public Advisory Panels or Committees	11/1/2006	null date	10/11/2006		E6-16679
FDA	FDA-2006-0502	FDA-2006-0502-0001	Redetermination of Regulatory Review Period for Purposes of Patent Extension; BONIVA; Correction	10/11/2006	null date	10/11/2006		E6-16816
FDA	FDA-2006-0503	FDA-2006-0503-0001	Recordkeeping Requirements for Human Food and Cosmetics Manufactured From, Processed With, or Otherwise Containing, Material From Cattle	10/11/2006	null date	10/11/2006	0910-AF48	E6-16830

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2006-0504	FDA-2006-0504-0001	Guidance for Industry on Investigating Out-of-Specification Test Results for Pharmaceutical Production; Availability	10/12/2006	null date	10/12/2006		E6-16838
FDA	FDA-2006-0505	FDA-2006-0505-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Food Labeling; Trans Fatty Acids in Nutrition Labeling	10/12/2006	null date	10/12/2006		E6-16840
FDA	FDA-2006-0506	FDA-2006-0506-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Interstate Shellfish Dealers Certificate	10/13/2006	null date	10/13/2006		E6-16953
FDA	FDA-2006-0507	FDA-2006-0507-0001	Memorandum of Understanding Between the Food and Drug Administration, and Duke University for the Cardiac Safety Research Consortium	10/16/2006	null date	10/16/2006		06-08708
FDA	FDA-2006-0508	FDA-2006-0508-0001	Debarment orders: Butkovitz, Anne L.	10/17/2006	null date	10/17/2006		E6-17178
FDA	FDA-2006-0509	FDA-2006-0509-0001	Summaries of Medical and Clinical Pharmacology Reviews of Pediatric Studies; Availability	10/18/2006	null date	10/18/2006		E6-17284
FDA	FDA-2006-0510	FDA-2006-0510-0001	Guidance for Industry on Fixed Dose Combinations, Co-Packaged Drug Products, and Single-Entity Versions of Previously Approved Antiretrovirals for the Treatment of HIV; Availability	10/18/2006	null date	10/18/2006		E6-17324
FDA	FDA-2006-0511	FDA-2006-0511-0001	Guidance for Industry: Biological Product Deviation Reporting for Blood and Plasma Establishments; Availability	10/19/2006	null date	10/19/2006		E6-17378
FDA	FDA-2006-0512	FDA-2006-0512-0001	Guidance for Industry: Biological Product Deviation Reporting for Licensed Manufacturers of Biological Products Other than Blood and Blood Components; Availability	10/19/2006	null date	10/19/2006		E6-17374
FDA	FDA-2006-0513	FDA-2006-0513-0001	Neurological Devices Panel of the Medical Devices Advisory Committee; Notice of Postponement of Meeting	10/20/2006	null date	10/20/2006		06-08788
FDA	FDA-2006-0514	FDA-2006-0514-0001	Circulatory System Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting	10/20/2006	null date	10/20/2006		E6-17519
FDA	FDA-2006-0515	FDA-2006-0515-0001	Arthritis Advisory Committee; Notice of Meeting	10/20/2006	null date	10/20/2006		06-08787
FDA	FDA-2006-0516	FDA-2006-0516-0001	Agency information collection activities; proposals, submissions, and approvals	10/24/2006	null date	10/24/2006		E6-17720

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Food and Drug Administration (FDA) -- 1547 Documents

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Food and Drug Administration (FDA)								
FDA	FDA-2006-0517	FDA-2006-0517-0001	Reports and guidance documents; availability, etc.: Portable invasive blood glucose monitoring systems; total product life cycle	10/24/2006	null date	10/24/2006		E6-17757
FDA	FDA-2006-0518	FDA-2006-0518-0001	Agency information collection activities; proposals, submissions, and approvals	10/24/2006	null date	10/24/2006		E6-17718
FDA	FDA-2006-0519	FDA-2006-0519-0001	Medical devices: Premarket notification exemptions; Class II devices--	10/24/2006	null date	10/24/2006		E6-17729
FDA	FDA-2006-0520	FDA-2006-0520-0001	Reports and guidance documents; availability, etc.: Global Harmonization Task Force Study Groups; proposed and final documents	10/24/2006	null date	10/24/2006		E6-17727
FDA	FDA-2006-0521	FDA-2006-0521-0002	Conventional Foods Being Marketed as ``Functional Foods; Extension of Comment Period	1/8/2007	3/5/2007	1/8/2007		E7-00047
FDA	FDA-2006-0521	FDA-2006-0521-0001	Conventional Foods Being Marketed as ``Functional Foods; Public Hearing; Request for Comments	10/25/2006	3/5/2007	10/25/2006		06-08895
FDA	FDA-2006-0522	FDA-2006-0522-0001	Safe Foods Corporation; Filing of Food Additive Petition	10/25/2006	null date	10/25/2006		E6-17834
FDA	FDA-2006-0522	FDA-2006-0522-0002	Secondary Direct Food Additives Permitted in Food for Human Consumption	11/29/2007	12/31/2007	11/29/20077		E7-23182
FDA	FDA-2006-0523	FDA-2006-0523-0001	Draft Guidance for Industry and Food and Drug Administration Staff; Annual Reports for Approved Premarket Approval Applications; Availability	10/26/2006	null date	10/26/2006		E6-17908
FDA	FDA-2006-0524	FDA-2006-0524-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget; Extension of Expiration Date for MedWatch (Food and Drug Administration Medical Products Reporting Program) Form	10/26/2006	null date	10/26/2006		E6-17907
FDA	FDA-2006-0525	FDA-2006-0525-0001	Pediatric Advisory Committee; Notice of Meeting	10/26/2006	null date	10/26/2006		E6-17965
FDA	FDA-2006-0526	FDA-2006-0526-0001	Dental Products Panel of the Medical Devices Advisory Committee; Notice of Meeting	10/26/2006	null date	10/26/2006		E6-17932
FDA	FDA-2006-0527	FDA-2006-0527-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Orphan Drugs	10/30/2006	null date	10/30/2006		E6-18067
FDA	FDA-2006-0528	FDA-2006-0528-0001	Draft Guidance for Industry; Blue Bird Medicated Feed Labels; Availability	10/30/2006	null date	10/30/2006		E6-18148

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Food and Drug Administration (FDA)								
FDA	FDA-2006-0529	FDA-2006-0529-0001	Guidance for Industry on Implementation of Acceptable Full-Length Donor History Questionnaire and Accompanying Materials for Use in Screening Donors of Blood and Blood Components; Availability	10/30/2006	null date	10/30/2006		E6-18068
FDA	FDA-2006-0530	FDA-2006-0530-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Biological Products: Reporting of Biological Product Deviations in Manufacturing; Forms FDA 3486 and 3486A	10/31/2006	null date	10/31/2006		E6-18313
FDA	FDA-2006-0531	FDA-2006-0531-0001	Vaccines and Related Biological Products Advisory Committee; Notice of Meeting	10/31/2006	null date	10/31/2006		E6-18314
FDA	FDA-2006-0532	FDA-2006-0532-0001	General and Plastic Surgery Devices; Reclassification of the Absorbable Hemostatic Device	10/31/2006	1/29/2007	10/31/2006		E6-18324
FDA	FDA-2006-0532	FDA-2006-0532-0002	General and Plastic Surgery Devices; Reclassification of the Absorbable Hemostatic Device; Reopening of Comment Period	5/8/2007	6/7/2007	5/8/2007		E7-08784
FDA	FDA-2006-0533	FDA-2006-0533-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Administrative Detention and Banned Medical Devices	10/31/2006	null date	10/31/2006		E6-18190
FDA	FDA-2006-0534	FDA-2006-0534-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Investigational Device Exemptions Reports and Records	10/31/2006	null date	10/31/2006		E6-18200
FDA	FDA-2006-0535	FDA-2006-0535-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Medical Device User Fee and Modernization Act Small Business Qualification Certification (Form FDA 3602)	10/31/2006	null date	10/31/2006		E6-18198
FDA	FDA-2006-0536	FDA-2006-0536-0001	Draft Guidance for Industry and Food and Drug Administration Staff; Class II Special ControlsGuidance Document: Absorbable Hemostatic Device; Availability	10/31/2006	null date	10/31/2006		E6-18318
FDA	FDA-2006-0537	FDA-2006-0537-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Infectious Disease Issues in Xenotransplantation	10/31/2006	null date	10/31/2006		E6-18203
FDA	FDA-2006-0538	FDA-2006-0538-0001	Marketed Unapproved Drugs; Public Workshop	11/1/2006	null date	11/1/2006		E6-17959

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Food and Drug Administration (FDA)								
FDA	FDA-2006-0539	FDA-2006-0539-0001	Cellular, Tissue and Gene Therapies Advisory Committee; Notice of Meeting	11/2/2006	null date	11/2/2006		E6-18472
FDA	FDA-2006-0540	FDA-2006-0540-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; General Licensing Provisions: Biologics License Application, Changes to an Approved Application, Labeling, Revocation and Suspension, Postmarketing Studies Status Reports, and Forms FDA 356h and 2567	11/2/2006	null date	11/2/2006		E6-18445
FDA	FDA-2006-0541	FDA-2006-0541-0001	Guidance for Industry: Questions and Answers Regarding Food Allergens, Including the Food Allergen Labeling and Consumer Protection Act of 2004 (Edition 4); Availability	11/2/2006	null date	11/2/2006		E6-18443
FDA	FDA-2006-0542	FDA-2006-0542-0001	Pediatric Oncology Subcommittee of the Oncologic Drugs Advisory Committee; Notice of Meeting	11/2/2006	null date	11/2/2006		E6-18442
FDA	FDA-2006-0543	FDA-2006-0543-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Substantial Evidence of Effectiveness of New Animal Drugs	11/2/2006	null date	11/2/2006		E6-18432
FDA	FDA-2006-0544	FDA-2006-0544-0001	Implantation or Injectable Dosage Form New Animal Drugs; Glycopyrrolate	11/2/2006	null date	11/2/2006		E6-18444
FDA	FDA-2006-0545	FDA-2006-0545-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Premarket Notification	11/3/2006	null date	11/3/2006		E6-18553
FDA	FDA-2006-0546	FDA-2006-0546-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Reporting and Recordkeeping Requirements and Availability of Sample Electronic Products for Manufacturers and Distributors of Electronic Products	11/3/2006	null date	11/3/2006		E6-18559
FDA	FDA-2006-0547	FDA-2006-0547-0001	Food and Drug Administration Modernization Act of 1997: Modifications to the List of Recognized Standards, Recognition List Number: 016	11/3/2006	null date	11/3/2006		E6-18604
FDA	FDA-2006-0548	FDA-2006-0548-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Medical Device User Fee Cover Sheet	11/3/2006	null date	11/3/2006		E6-18557

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Food and Drug Administration (FDA)								
FDA	FDA-2006-0549	FDA-2006-0549-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Inspection by Accredited Persons Program Under the Medical Device User Fee and Modernization Act of 2002	11/3/2006	null date	11/3/2006		E6-18603
FDA	FDA-2006-0550	FDA-2006-0550-0001	Implantation or Injectable Dosage Form New Animal Drugs; Lincomycin; Correction	11/17/2006	null date	11/7/2006		E6-18679
FDA	FDA-2006-0551	FDA-2006-0551-0001	New Animal Drugs for Use in Animal Feeds; Bambermycins	11/7/2006	null date	11/7/2006		E6-18680
FDA	FDA-2006-0552	FDA-2006-0552-0001	Oral Dosage Form New Animal Drugs; Ivermectin, Pyrantel, and Praziquantel Tablets	11/7/2006	null date	11/7/2006		E6-18684
FDA	FDA-2006-0553	FDA-2006-0553-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Guidance for Industry on How to Use E-Mail to Submit a Notice of Intent to Slaughter for Human Food Purposes	11/8/2006	null date	11/8/2006		E6-18896
FDA	FDA-2006-0554	FDA-2006-0554-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Guidance For Industry on How to Use E-Mail to Submit a Request for a Meeting or Teleconference to the Office Of New Animal Drug Evaluation	11/8/2006	null date	11/8/2006		E6-18911
FDA	FDA-2006-0555	FDA-2006-0555-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Guidance for Industry on How To Use E-Mail To Submit A Study Protocol	11/8/2006	null date	11/8/2006		E6-18908
FDA	FDA-2006-0556	FDA-2006-0556-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Guidance for Industry on How to Use E-Mail to Submit Information to the Center for Veterinary Medicine	11/8/2006	null date	11/8/2006		E6-18901
FDA	FDA-2006-0557	FDA-2006-0557-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Guidance on Reagents for Detection of Specific Novel Influenza A Viruses	11/9/2006	null date	11/9/2006		E6-19045
FDA	FDA-2006-0558	FDA-2006-0558-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Guidance for Industry on How to Use E-Mail to Submit a Notice of Final Disposition of Animals Not Intended for Immediate Slaughter	11/9/2006	null date	11/9/2006		E6-19044

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2006-0559	FDA-2006-0559-0001	Distribution of Blood Derivatives by Registered Blood Establishments That Qualify as Health Care Entities; Prescription Drug Marketing Act of 1987; Prescription Drug Amendments of 1992; Delay of Applicability Date	11/13/2006	null date	11/13/2006	0905-AC81	E6-18892
FDA	FDA-2006-0560	FDA-2006-0560-0001	New Animal Drugs for Use in Animal Feeds; Monensin	11/14/2006	null date	11/14/2006		E6-19203
FDA	FDA-2006-0561	FDA-2006-0561-0001	Draft Guidance for Industry: Protocols for the Conduct of Method Transfer Studies for Type C Medicated Feed Assay Methods; Availability	11/14/2006	null date	11/14/2006		E6-19204
FDA	FDA-2006-0562	FDA-2006-0562-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Medicated Feed Mill License Application	11/14/2006	null date	11/14/2006		E6-19152
FDA	FDA-2006-0563	FDA-2006-0563-0001	Draft Voluntary National Retail Food Regulatory Program Standards; Availability	11/14/2006	null date	11/14/2006		E6-19195
FDA	FDA-2006-0564	FDA-2006-0564-0001	Psychopharmacologic Drugs Advisory Committee; Notice of Meeting	11/15/2006	null date	11/15/2006		E6-19248
FDA	FDA-2006-0565	FDA-2006-0565-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Food Additive Petitions	11/15/2006	null date	11/15/2006		E6-19201
FDA	FDA-2006-0566	FDA-2006-0566-0001	Joint Meeting of the Anti-Infective Drugs Advisory Committee and the Drug Safety and Risk Management Advisory Committee; Notice of Meeting	11/15/2006	null date	11/15/2006		E6-19249
FDA	FDA-2006-0567	FDA-2006-0567-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Agreement for Shipment of Devices for Sterilization	11/15/2006	null date	11/15/2006		E6-19283
FDA	FDA-2006-0568	FDA-2006-0568-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Medical Device User Fee and Modernization Act Small Business Qualification Certification (Form FDA 3602)	11/15/2006	null date	11/15/2006		E6-19285
FDA	FDA-2006-0569	FDA-2006-0569-0001	Clinical Chemistry and Clinical Toxicology Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting	11/17/2006	null date	11/17/2006		E6-19492
FDA	FDA-2006-0570	FDA-2006-0570-0001	New Animal Drugs for Use in Animal Feeds; Lasalocid	11/21/2006	null date	11/21/2006		E6-19614

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Food and Drug Administration (FDA) -- 1547 Documents

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Food and Drug Administration (FDA)								
FDA	FDA-2006-0571	FDA-2006-0571-0001	Electronic Submission of Regulatory Information, and Creating an Electronic Platform for Enhanced Information Management; Public Hearing	11/21/2006	null date	11/21/2006		06-09313
FDA	FDA-2006-0572	FDA-2006-0572-0001	Oral Dosage Form New Animal Drugs; Ivermectin Paste	11/21/2006	null date	11/21/2006		E6-19616
FDA	FDA-2006-0573	FDA-2006-0573-0001	New Animal Drugs For Use in Animal Feeds; Ractopamine	11/21/2006	null date	11/21/2006		E6-19615
FDA	FDA-2006-0574	FDA-2006-0574-0001	Draft Guidance for Industry on Sinusitis: Designing Clinical Development Programs of Nonantimicrobial Drugs for Treatment; Availability	11/22/2006	null date	11/22/2006		E6-19689
FDA	FDA-2006-0575	FDA-2006-0575-0001	Guidance for Industry and Food and Drug Administration Staff; Saline, Silicone Gel, and Alternative Breast Implants; Availability	11/22/2006	null date	11/22/2006		06-09325
FDA	FDA-2006-0576	FDA-2006-0576-0001	Guidance for Industry, Food and Drug Administration Staff, Eye Care Professionals, and Consumers; Decorative, Non-Corrective Contact Lenses; Availability	11/24/2006	null date	11/24/2006		E6-19887
FDA	FDA-2006-0577	FDA-2006-0577-0001	Medical Devices Dispute Resolution Panel of the Medical Devices Advisory Committee; Notice of Meeting	1/24/2006	null date	1/24/2006		E6-19895
FDA	FDA-2006-0578	FDA-2006-0578-0001	Food Defense Workshop; Public Workshop	11/24/2006	null date	11/24/2006		E6-19886
FDA	FDA-2006-0579	FDA-2006-0579-0001	Guidance for Industry: Lead in Candy Likely to Be Consumed Frequently by Small Children; Recommended Maximum Level and Enforcement Policy, Availability; and Supporting Document: Supporting Document for Maximum Recommended Level for Lead in Candy Likely to Be Consumed Frequently By Small Children; Availability	11/24/2006	null date	11/24/2006		E6-19809
FDA	FDA-2006-0580	FDA-2006-0580-0001	Guidance for Industry: Gene Therapy Clinical Trials--Observing Subjects for Delayed Adverse Events; Availability	11/28/2006	null date	11/28/2006		E6-20129
FDA	FDA-2006-0581	FDA-2006-0581-0001	Draft Guidance for Industry, Clinical Laboratories, and Food and Drug Administration Staff on In Vitro Diagnostic Multivariate Index Assays; Availability; Extension of Comment Period	11/28/2006	null date	11/28/2006		E6-20032
FDA	FDA-2006-0582	FDA-2006-0582-0001	Improving Patient Safety by Enhancing the Container Labeling for Parenteral Infusion Drug Products; Public Meeting	11/28/2006	null date	11/28/2006		E6-20035

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Food and Drug Administration (FDA) -- 1547 Documents

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Food and Drug Administration (FDA)								
FDA	FDA-2006-0583	FDA-2006-0583-0001	Training Program for Regulatory Project Managers; Information Available to Industry	11/28/2006	null date	11/28/2006		E6-20041
FDA	FDA-2006-0584	FDA-2006-0584-0001	Oral Dosage Form New Animal Drugs; Neomycin	11/28/2006	null date	11/28/2006		E6-20126
FDA	FDA-2006-0585	FDA-2006-0585-0001	Transmissible Spongiform Encephalopathies Advisory Committee; Notice of Meeting	11/29/2006	null date	11/29/2006		E6-20251
FDA	FDA-2006-0586	FDA-2006-0586-0001	New Animal Drugs; Change of Sponsors Name	11/29/2006	null date	11/29/2006		E6-20250
FDA	FDA-2006-0587	FDA-2006-0587-0001	Agency Information Collection Activities; Announcement of Officeof Management and Budget Approval; Experimental Study of Qualified Health Claims: Consumer Inferences About Monounsaturated Fatty Acids From Olive Oil, EPA and DHA Omega-3 Fatty Acids, and Green Tea	11/29/2006	null date	11/29/2006		E6-20200
FDA	FDA-2006-0588	FDA-2006-0588-0001	Blood Products Advisory Committee; Notice of Meeting	11/30/2006	null date	11/30/2006		E6-20265
FDA	FDA-2006-0589	FDA-2006-0589-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Food Labeling; Notification Procedures for Statements on Dietary Supplements	12/1/2006	null date	12/1/2006		E6-20307
FDA	FDA-2006-0590	FDA-2006-0590-0001	Oral Dosage Form New Animal Drugs; Sulfamethazine Soluble Powder	12/4/2006	null date	12/4/2006		E6-20404
FDA	FDA-2006-0591	FDA-2006-0591-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Human Tissue Intended for Transplantation	12/4/2006	null date	12/4/2006		E6-20477
FDA	FDA-2006-0592	FDA-2006-0592-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Cosmetic Labeling Regulations	12/4/2006	null date	12/4/2006		E6-20478
FDA	FDA-2006-0593	FDA-2006-0593-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Substances Prohibited from Use in Animal Food or Feed; Animal Proteins Prohibited in Ruminant Feed	12/4/2006	null date	12/4/2006		E6-20476
FDA	FDA-2006-0594	FDA-2006-0594-0001	New Animal Drugs For Use in Animal Feeds; Florfenicol	12/4/2006	null date	12/4/2006		E6-20398
FDA	FDA-2006-0595	FDA-2006-0595-0001	Notice of Approval of Original Abbreviated New Animal Drug Application; Pyrantel Pamoate Suspension	12/4/2006	null date	12/4/2006		E6-20399

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Food and Drug Administration (FDA)								
FDA	FDA-2006-0596	FDA-2006-0596-0001	Joint Meeting of the Anti-Infective Drugs Advisory Committee and the Drug Safety and Risk Management Advisory Committee; Amendment of Notice	12/5/2006	null date	12/5/2006		E6-20538
FDA	FDA-2006-0597	FDA-2006-0597-0001	Neurological Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting	12/6/2006	null date	12/6/2006		E6-20552
FDA	FDA-2006-0598	FDA-2006-0598-0003	Use of Ozone-Depleting Substances; Removal of Essential Use Designations; Confirmation of Effective Date	4/27/2007	null date	4/27/2007		E7-08043
FDA	FDA-2006-0598	FDA-2006-0598-0001	Use of Ozone-Depleting Substances; Removal of Essential Use Designations; Companion Document to Direct Final Rule	12/7/2006	2/20/2007	12/7/2006	0910-AF93	E6-20796
FDA	FDA-2006-0598	FDA-2006-0598-0002	Use of Ozone-Depleting Substances; Removal of Essential Use Designations	12/7/2006	2/20/2007	12/7/2006	0910-AF93	E6-20797
FDA	FDA-2006-0599	FDA-2006-0599-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Establishing and Maintaining a List of United States Dairy Product Manufacturers/ Processors With Interest in Exporting to Chile	12/7/2006	null date	12/7/2006		E6-20704
FDA	FDA-2006-0600	FDA-2006-0600-0001	Withdrawal of Federal Register Notice	12/7/2006	null date	12/7/2006		E6-20705
FDA	FDA-2006-0601	FDA-2006-0601-0001	Color Additive Certification; Increase in Fees for Certification Services	12/7/2006	2/5/2007	12/7/2006		E6-20800
FDA	FDA-2006-0602	FDA-2006-0602-0001	Oral Dosage Form New Animal Drugs; Lincomycin and Spectinomycin Powder	12/8/2006	null date	12/8/2006		E6-20929
FDA	FDA-2006-0603	FDA-2006-0603-0001	Oral Dosage Form New Animal Drugs; Oxytetracycline Powder	12/8/2006	null date	12/8/2006		E6-20928
FDA	FDA-2006-0604	FDA-2006-0604-0001	Medical Devices Dispute Resolution Panel of the Medical Devices Advisory Committee; Amendment of Notice	12/12/2006	null date	12/12/2006		E6-21020
FDA	FDA-2006-0605	FDA-2006-0605-0001	Over-the-Counter Human Drugs; Labeling Requirements; Proposed Rule	12/12/2006	4/11/2007	12/12/2006	0910-AD47	E6-21019
FDA	FDA-2006-0606	FDA-2006-0606-0001	New Animal Drugs For Use in Animal Feeds; Tylosin	12/12/2006	null date	12/12/2006		E6-21021
FDA	FDA-2006-0607	FDA-2006-0607-0001	Industry Exchange Workshop on Food and Drug Administration Clinical Trial Requirements; Public Workshop	12/13/2006	null date	12/13/2006		E6-21138
FDA	FDA-2006-0608	FDA-2006-0608-0001	Supplements and Other Changes to Approved New Animal Drug Applications	12/13/2006	null date	12/13/2006	0910-AF59	E6-21133
FDA	FDA-2006-0609	FDA-2006-0609-0001	Food Labeling: Nutrition Labeling of Dietary Supplements on a ``Per Day Basis	12/13/2006	null date	12/13/2006		06-09657

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Food and Drug Administration (FDA)								
FDA	FDA-2006-0610	FDA-2006-0610-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Requirements for Submission of Labeling for Human Prescription Drugs and Biologics in Electronic Format	12/13/2006	null date	12/13/2006		E6-21132
FDA	FDA-2006-0611	FDA-2006-0611-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Postmarket Surveillance	12/13/2006	null date	12/13/2006		E6-21167
FDA	FDA-2006-0612	FDA-2006-0612-0001	Expanded Access to Investigational Drugs for Treatment Use	1/16/2007	null date	12/14/2006	0910-AF14	06-09684
FDA	FDA-2006-0613	FDA-2006-0613-0001	Charging for Investigational Drugs	12/14/2006	3/14/2007	12/14/2006	0910-AF13	06-09685
FDA	FDA-2006-0614	FDA-2006-0614-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Proposed Experimental Study of Trans Fat Claims on Foods	12/15/2006	null date	12/15/2006		E6-21317
FDA	FDA-2006-0615	FDA-2006-0615-0001	Drug Products Containing Quinine; Enforcement Action Dates	12/15/2007	null date	12/15/2007		06-09713
FDA	FDA-2006-0616	FDA-2006-0616-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Registration of Food Facilities Under the Public Health Security and Bioterrorism Preparedness and Response Act of 2002	12/15/2006	null date	12/15/2006		E6-21375
FDA	FDA-2006-0617	FDA-2006-0617-0001	International Drug Scheduling; Convention on Psychotropic Substances; Single Convention on Narcotic Drugs; World Health Organization Scheduling Recommendations for Dronabinol and its Stereoisomers, and Oripavine	12/15/2006	null date	12/15/2006		E6-21318
FDA	FDA-2006-0618	FDA-2006-0618-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Manufactured Food Regulatory Program Standards	12/18/2006	null date	12/18/2006		E6-21472
FDA	FDA-2006-0619	FDA-2006-0619-0002	Agency information collection activities; proposals, submissions, and approvals; correction	1/18/2007	null date	1/18/2007		Z6-21486

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Food and Drug Administration (FDA) -- 1547 Documents

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Food and Drug Administration (FDA)								
FDA	FDA-2006-0619	FDA-2006-0619-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Experimental Study of Possible Footnotes and Cueing Schemes to Help Consumers Interpret Quantitative Trans Fat Disclosure on the Nutrition Facts Panel	12/18/2006	null date	12/18/2006		E6-21486
FDA	FDA-2006-0620	FDA-2006-0620-0001	Medical Devices; Patient Examination and Surgeons Gloves; Test Procedures and Acceptance Criteria	12/19/2006	null date	12/19/2006		E6-21591
FDA	FDA-2006-0620	FDA-2006-0620-0002	Medical Devices; Patient Examination and Surgeons Gloves; Test Procedures and Acceptance Criteria; Correction	1/19/2007	null date	1/19/2007		E7-00682
FDA	FDA-2006-0621	FDA-2006-0621-0001	Marketed Unapproved Drugs; Public Workshop; Change of Meeting Location and Time	12/20/2006	null date	12/20/2006		E6-21738
FDA	FDA-2006-0622	FDA-2006-0622-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Product Jurisdiction: Assignment of Agency Component for Review of Premarket Applications	12/20/2006	null date	12/20/2006		E6-21636
FDA	FDA-2006-0623	FDA-2006-0623-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Prior Notice of Imported Food Under the Public Health Security and Bioterrorism Preparedness and Response Act of 2002	12/20/2006	null date	12/20/2006		E6-21737
FDA	FDA-2006-0624	FDA-2006-0624-0001	Uniform Compliance Date for Food Labeling Regulations	12/21/2006	3/6/2007			E6-21902
FDA	FDA-2006-0625	FDA-2006-0625-0001	Guidance for Industry and Food and Drug Administration Staff; Procedures for Handling Post-Approval Studies Imposed by Premarket Approval Application Order; Availability	12/21/2006	null date	12/21/2006		E6-21901
FDA	FDA-2006-0626	FDA-2006-0626-0001	Implantation or Injectable Dosage Form New Animal Drugs; Gentamicin	12/22/2006	null date	12/22/2006		E6-21951
FDA	FDA-2006-0627	FDA-2006-0627-0001	Advisory Committee for Reproductive Health Drugs; Notice of Meeting	12/22/2006	null date	12/22/2006		E6-21949

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Food and Drug Administration (FDA)								
FDA	FDA-2006-0628	FDA-2006-0628-0001	Guidance for Clinical Investigators, Institutional Review Boards, and Sponsors; Process for Handling Referrals to Food and Drug Administration Under 21 CFR 50.54: Additional Safeguards for Children in Clinical Investigations	12/22/2006	null date	12/22/2006		E6-21950
FDA	FDA-2006-0629	FDA-2006-0629-0001	Medical Device Regulations; Disqualification of a Clinical Investigator; Technical Amendment	12/22/2006	null date	12/22/2006		E6-21952
FDA	FDA-2006-0630	FDA-2006-0630-0001	Neurological Devices Panel of the Medical Devices Advisory Committee; Amendment of Notice	12/26/2006	null date	12/26/2006		E6-21995
FDA	FDA-2006-0631	FDA-2006-0631-0001	Medical Devices; Exemptions from Premarket Notification; Class II Devices	12/26/2006	null date	12/26/2006		E6-22072
FDA	FDA-2006-0632	FDA-2006-0632-0001	Internal Analgesic, Antipyretic, and Antirheumatic Drug Products for Over-the-Counter Human Use; Proposed Amendment of the Tentative Final Monograph; Required Warnings and Other Labeling	12/26/2006	5/25/2007	12/26/2006	0910-AF36	E6-21855
FDA	FDA-2006-0633	FDA-2006-0633-0001	Advisory Committees; Tentative Schedule of Meetings for 2007	12/29/2006	null date	12/29/2006		E6-22389
FDA	FDA-2007-0001	FDA-2007-0001-0001	Draft Animal Cloning Risk Assessment; Proposed Risk Management Plan; Draft Guidance for Industry; Availability	1/3/2007	null date	1/3/2007		06-09927
FDA	FDA-2007-0002	FDA-2007-0002-0001	Draft Guidance for Industry and Food and Drug Administration Staff; Radio-Frequency Wireless Technology in Medical Devices; Availability	1/3/2007	null date	1/3/2007		E6-22449
FDA	FDA-2007-0003	FDA-2007-0003-0001	Advisory Committees; Filing of Annual Reports	1/3/2007	null date	1/3/2007		E6-22450
FDA	FDA-2007-0004	FDA-2007-0004-0001	Animal drugs, feeds, and related products: Dexmedetomidine	1/4/2007	null date	1/4/2007		E6-22508
FDA	FDA-2007-0005	FDA-2007-0005-0001	Animal drugs, feeds, and related products: Atipamezole	1/4/2007	null date	1/4/2007		E6-22515
FDA	FDA-2007-0006	FDA-2007-0006-0001	Animal drugs, feeds, and related products: Dirlotapide solution	1/4/2007	null date	1/4/2007		E6-22542
FDA	FDA-2007-0007	FDA-2007-0007-0001	Animal drugs, feeds, and related products: Chlorhexidine	1/4/2007	null date	1/4/2007		E6-22514
FDA	FDA-2007-0008	FDA-2007-0008-0001	Reports and guidance documents; availability, etc.: Best Pharmaceuticals for Children Act--	1/4/2007	null date	1/4/2007		E6-22517
FDA	FDA-2007-0009	FDA-2007-0009-0001	Animal drugs, feeds, and related products: Florfenicol	1/4/2007	null date	1/4/2007		E6-22516
FDA	FDA-2007-0010	FDA-2007-0010-0001	Animal drugs, feeds, and related products: Doxapram	1/4/2007	null date	1/4/2007		E6-22510

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0011	FDA-2007-0011-0001	Animal drugs, feeds, and related products: Clomipramine tablets	1/4/2007	null date	1/4/2007		E6-22509
FDA	FDA-2007-0012	FDA-2007-0012-0001	Supplements and Other Changes to an Approved Application; Public Meeting	1/5/2007	null date	1/5/2007		E6-22588
FDA	FDA-2007-0013	FDA-2007-0013-0001	Food Labeling: Nutrient Content Claims, General Principles; Health Claims, General Requirements and Other Specific Requirements for Individual Health Claims; Withdrawal in Part	1/5/2007	null date	1/5/2007		E6-21996
FDA	FDA-2007-0014	FDA-2007-0014-0001	Food Labeling: Health Claims; Calcium and Osteoporosis, and Calcium, Vitamin D, and Osteoporosis	1/5/2007	3/21/2007	1/5/2007		E6-22573
FDA	FDA-2007-0015	FDA-2007-0015-0001	New Animal Drugs For Use in Animal Feeds; Monensin	1/8/2007	null date	1/8/2007		E7-0004
FDA	FDA-2007-0016	FDA-2007-0016-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Threshold of Regulation for Substances Used in Food- Contact Articles	1/8/2007	null date	1/8/2007		E7-00006
FDA	FDA-2007-0017	FDA-2007-0017-0001	International Conference on Harmonisation; Draft Guidance on E15 Terminology in Pharmacogenomics; Availability	1/8/2007	null date	1/8/2007		E7-00005
FDA	FDA-2007-0018	FDA-2007-0018-0001	Meetings: Medical Devices 101: An Educational Forum; public workshop	1/9/2007	null date	1/9/2007		E7-00092
FDA	FDA-2007-0019	FDA-2007-0019-0001	Meetings: In vitro diagnostic multivariate index assays	1/9/2007	null date	1/9/2007		E7-00093
FDA	FDA-2007-0020	FDA-2007-0020-0001	Vaccines and Related Biological Products Advisory Committee; Notice of Meeting	9/1/2007	null date	9/1/2007		07-00028
FDA	FDA-2007-0021	FDA-2007-0021-0001	Medical Devices; Immunology and Microbiology Devices; Classification of Quality Control Material for Cystic Fibrosis Nucleic Acid Assays	1/10/2007	null date	1/10/2007		E7-00119
FDA	FDA-2007-0022	FDA-2007-0022-0001	New Animal Drugs; Change of Sponsor	1/10/2007	null date	1/10/2007		E7-00118
FDA	FDA-2007-0023	FDA-2007-0023-0001	Guidance for Industry and Food and Drug Administration Staff; Class II Special Controls Guidance Document: Quality Control Material for Cystic Fibrosis Nucleic Acid Assays; Availability	1/10/2007	null date	1/10/2007		E7-00120
FDA	FDA-2007-0024	FDA-2007-0024-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Infant Formula Requirements	1/12/2007	null date	1/12/2007		E7-00331

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0025	FDA-2007-0025-0002	Use of Materials Derived From Cattle in Medical Products Intended for Use in Humans and Drugs Intended for Use in Ruminants; Reopening of the Comment Period	3/30/2007	5/14/2007	3/30/2007	0910-AF54	E7-05894
FDA	FDA-2007-0025	FDA-2007-0025-0001	Use of Materials Derived from Cattle in Medical Products Intended for Use in Humans and Drugs Intended for Use in Ruminants	1/12/2007	5/14/2007	1/12/2007	0910-AF54	E6-22329
FDA	FDA-2007-0026	FDA-2007-0026-0001	Prescription Drug User Fee Act; Public Meeting	1/16/2007	null date	1/16/2007		07-00122
FDA	FDA-2007-0027	FDA-2007-0027-0001	Draft Guidance for Industry: Minimally Manipulated, Unrelated, Allogeneic Placental/Umbilical Cord Blood Intended for Hematopoietic Reconstitution in Patients with Hematological Malignancies; Availability	1/17/2007	null date	1/17/2007		E7-00549
FDA	FDA-2007-0028	FDA-2007-0028-0001	Orthopedic Devices; Reclassification of Non-Invasive Bone Growth Stimulator	1/17/2007	4/17/2020	1/17/2007		E7-00476
FDA	FDA-2007-0029	FDA-2007-0029-0001	Sentinel Network To Promote Medical Product Safety; Public Meeting	1/1/8/2007	null date	1/18/2007		07-00141
FDA	FDA-2007-0030	FDA-2007-0030-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Submission of Petitions: Food Additive, Color Additive (Including Labeling), and Generally Recognized as Safe Affirmation; Electronic Submission Using Food and Drug Administration Forms 3503 and 3504	1/19/2007	null date	1/19/2007		E7-00681
FDA	FDA-2007-0031	FDA-2007-0031-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Guidance for Industry and Food and Drug Administration Staff on Class II Special Controls Guidance Document: Automated Blood Cell Separator Device Operating by Centrifugal or Filtration Separation Principle	1/22/2007	null date	1/22/2007		E7-00804
FDA	FDA-2007-0032	FDA-2007-0032-0001	Orthopaedic and Rehabilitation Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting	1/23/2007	null date	1/23/2007		E7-00946
FDA	FDA-2007-0033	FDA-2007-0033-0001	Hydrogen Peroxide Solution for Control of Various Fungal and Bacterial Diseases in Fish; Availability of Data	1/23/2007	null date	1/23/2007		E7-00947
FDA	FDA-2007-0034	FDA-2007-0034-0001	Food Labeling; Gluten-Free Labeling of Foods	1/23/2007	4/23/2007	1/23/2007	0910-ZA26	E7-00843

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0035	FDA-2007-0035-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Bar Code Label Requirement for Human Drug and Biological Products	1/23/2007	null date	1/23/2007		E7-00916
FDA	FDA-2007-0036	FDA-2007-0036-0001	Guidance for Industry: Certain Human Cells, Tissues, and Cellular and Tissue-Based Products Recovered From Donors Who Were Tested for Communicable Diseases Using Pooled Specimens or Diagnostic Tests; Availability	1/24/2007	null date	1/24/2007		E7-00978
FDA	FDA-2007-0037	FDA-2007-0037-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Adoption of the Food and Drug Administration Food Code by Local, State, and Tribal Governments	1/26/2007	null date	1/26/2007		E7-01231
FDA	FDA-2007-0038	FDA-2007-0038-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Human Cells, Tissues, and Cellular and Tissue-Based Products: Establishment Registration and Listing; Form Food and Drug Administration 3356; Eligibility Determination for Donors; and Current Good Tissue Practice	1/26/2007	null date	1/26/2007		E7-01196
FDA	FDA-2007-0039	FDA-2007-0039-0001	Medical Devices; Availability of Safety and Effectiveness Summaries for Premarket Approval Applications	1/26/2007	null date	1/26/2007		E7-01199
FDA	FDA-2007-0040	FDA-2007-0040-0001	Indevus Pharmaceuticals, Inc.; Withdrawal of Approval of a New Drug Application	1/30/2007	null date	1/30/2007		E7-01414
FDA	FDA-2007-0041	FDA-2007-0041-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Biological Products: Reporting of Biological Product Deviations in Manufacturing; Forms FDA 3486 and 3486A	1/30/2007	null date	1/30/2007		E7-01415
FDA	FDA-2007-0042	FDA-2007-0042-0001	James T. Kimball; Denial of Hearing; Final Debarment Order	1/30/2007	null date	1/30/2007		E7-01416
FDA	FDA-2007-0043	FDA-2007-0043-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Interstate Shellfish Dealers Certificate	1/31/2007	null date	1/31/2007		E7-01549

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0044	FDA-2007-0044-0001	Memorandum of Understanding Between the United States Food and Drug Administration and the Veterans Health Administration	1/31/2007	null date	1/31/2007		07-00421
FDA	FDA-2007-0045	FDA-2007-0045-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Infectious Disease in Xenotransplantation	2/1/2007	null date	2/1/2007		E7-01550
FDA	FDA-2007-0046	FDA-2007-0046-0001	Guidance for Industry; Class II Special Controls Guidance Document: Cord Blood Processing System and Storage Container; Availability	2/1/2007	null date	2/1/2007		E7-01568
FDA	FDA-2007-0047	FDA-2007-0047-0001	University of Arkansas/Food and Drug Administration Food Labeling Workshop; Public Workshop	2/1/2007	null date	2/1/2007		E7-01570
FDA	FDA-2007-0048	FDA-2007-0048-0001	Medical Devices; Hematology and Pathology Devices; Classification of Cord Blood Processing System and Storage Container	2/1/2007	null date	2/1/2007		E7-01566
FDA	FDA-2007-0049	FDA-2007-0049-0001	Regulatory Site Visit Training Program	2/1/2007	null date	2/1/2007		E7-01576
FDA	FDA-2007-0050	FDA-2007-0050-0001	Determination That SUSTIVA (Efavirenz) 300-Milligram Tablets Were Not Withdrawn From Sale for Reasons of Safety or Effectiveness	2/2/2007	null date	2/2/2007		E7-01748
FDA	FDA-2007-0051	FDA-2007-0051-0001	New Animal Drugs for Use in Animal Feeds; Lasalocid	2/2/2007	null date	2/2/2007		E7-01684
FDA	FDA-2007-0052	FDA-2007-0052-0001	Memorandum of Understanding Between the Food and Drug Administration, Duke University and Duke University Health System, Inc.	2/2/2007	null date	2/2/2007		07-00454
FDA	FDA-2007-0053	FDA-2007-0053-0001	Agency Emergency Processing Under Office of Management and Budget Review; Toll-Free Number for Reporting Adverse Events on Labeling for Human Drug Products; Labeling Comprehension Study	2/2/2007	null date	2/2/2007		E7-01674
FDA	FDA-2007-0054	FDA-2007-0054-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; General Licensing Provisions: Biologics License Application, Changes to an Approved Application, Labeling, Revocation and Suspension, Postmarketing Studies Status Reports, and Forms FDA 456h and 2567	2/2/2007	null date	2/2/2007		E7-01741
FDA	FDA-2007-0055	FDA-2007-0055-0001	Report on the Performance of Drug and Biologics Firms in Conducting Postmarketing Commitment Studies; Availability	2/2/2007	null date	2/2/2007		E7-01749

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Food and Drug Administration (FDA)								
FDA	FDA-2007-0056	FDA-2007-0056-0001	Food Defense Workshop; Public Workshop	2/6/2007	null date	2/6/2007		E7-01865
FDA	FDA-2007-0057	FDA-2007-0057-0001	Global Harmonization Task Force, Study Groups 1, 2, and 4; New Proposed and Final Documents; Availability	2/6/2007	null date	2/6/2007		E7-01864
FDA	FDA-2007-0058	FDA-2007-0058-0001	New Animal Drugs; Hydrogen Peroxide	2/6/2007	null date	2/6/2007		E7-01848
FDA	FDA-2007-0059	FDA-2007-0059-0001	Food Labeling: Health Claims; Soluble Fiber From Certain Foods and Risk of Coronary Heart Disease	2/6/2007	4/23/2007	2/6/2007	0910-AF94	E7-01849
FDA	FDA-2007-0060	FDA-2007-0060-0001	Vaccines and Related Biological Products Advisory Committee; Notice of Meeting	7/1/2007	null date	7/1/2007		E7-01899
FDA	FDA-2007-0061	FDA-2007-0061-0001	Anesthetic and Life Support Drugs Advisory Committee; Notice of Meeting	2/7/2007	null date	2/7/2007		E7-01991
FDA	FDA-2007-0062	FDA-2007-0062-0001	Antiviral Drugs Advisory Committee; Notice of Meeting	2/7/2007	null date	2/7/2007		E7-01900
FDA	FDA-2007-0063	FDA-2007-0063-0001	Guidance for Industry on User Fee Waivers for Fixed Dose Combination and Co-Packaged Human Immunodeficiency Virus Drugs for the Presidents Emergency Plan for Acquired Immunodeficiency Syndrome Relief; Availability	2/8/2007	null date	2/8/2007		E7-02124
FDA	FDA-2007-0064	FDA-2007-0064-0001	Circulatory System Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting	8/1/2007	null date	2/8/2007		E7-02122
FDA	FDA-2007-0065	FDA-2007-0065-0001	Ophthalmic and Topical Dosage Form New Animal Drugs; Gentamicin and Betamethasone Spray	2/8/2007	null date	2/8/2007		E7-02121
FDA	FDA-2007-0066	FDA-2007-0066-0001	Ophthalmic and Topical Dosage Form New Animal Drugs; Ivermectin Topical Solution	2/12/2007	null date	2/12/2007		E7-02368
FDA	FDA-2007-0067	FDA-2007-0067-0001	Oral Dosage Form New Animal Drugs; Fluoxetine	2/12/2007	null date	2/12/2007		E7-02172
FDA	FDA-2007-0068	FDA-2007-0068-0001	Voluntary Self Inspection of Medicated Feed Manufacturing Facilities; Draft Compliance Policy Guide; Availability	2/12/2007	null date	2/12/2007		E7-02232
FDA	FDA-2007-0069	FDA-2007-0069-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Food Labeling; Notification Procedures for Statements on Dietary Supplements	2/14/2007	null date	2/14/2007		E7-02480

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Food and Drug Administration (FDA)								
FDA	FDA-2007-0070	FDA-2007-0070-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Administrative Procedures for the Clinical Laboratory Improvement Amendments of 1998 Categorization	2/14/2007	null date	2/14/2007		E7-02468
FDA	FDA-2007-0071	FDA-2007-0071-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Agreement for Shipment of Devices for Sterilization	2/14/2007	null date	2/14/2007		E7-02467
FDA	FDA-2007-0072	FDA-2007-0072-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Guidance for Industry on How to Use E-mail to Submit Information to the Center for Veterinary Medicine	2/14/2007	null date	2/14/2007		E7-02470
FDA	FDA-2007-0073	FDA-2007-0073-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; How to Use E-mail to Submit a Notice of Final Disposition of Animals Not Intended for Immediate Slaughter	2/14/2007	null date	2/14/2007		E7-02485
FDA	FDA-2007-0074	FDA-2007-0074-0001	Agency Information Collection Activities: Submission for Office of Management and Budget Review; Comment Request; Export of Medical Devices-Foreign Letters of Approval	2/14/2007	null date	2/14/2007		E7-02489
FDA	FDA-2007-0075	FDA-2007-0075-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; User Fee Cover Sheet; Form FDA 3397	2/14/2007	null date	2/14/2007		E7-02469
FDA	FDA-2007-0076	FDA-2007-0076-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Substantial Evidence of Effectiveness of New Animal Drugs	2/14/2007	null date	2/14/2007		E7-02497

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AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0077	FDA-2007-0077-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; General Licensing Provisions: Biologics License Application, Changes to an Approved Application, Labeling, Revocation and Suspension, Postmarketing Studies Status Reports, and Forms FDA 356h and 2567; Correction	2/15/2007	null date	2/15/2007		E7-02576
FDA	FDA-2007-0078	FDA-2007-0078-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Guidance for Industry on How to Use E-Mail to Submit a Request for a Meeting or Teleconference to the Office Of New Animal Drug Evaluation	2/15/2007	null date	2/15/2007		E7-02579
FDA	FDA-2007-0079	FDA-2007-0079-0001	Sentinel Network To Promote Medical Product Safety; Public Meeting		null date	2/15/2007		07-00710
FDA	FDA-2007-0080	FDA-2007-0080-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Mammography Quality Standards Act Requirements	2/15/2007	null date	2/15/2007		E7-02578
FDA	FDA-2007-0081	FDA-2007-0081-0001	Draft Guidance for Industry on Developing Products for Weight Management; Availability	2/15/2007	null date	2/15/2007		E7-02581
FDA	FDA-2007-0082	FDA-2007-0082-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Guidance for Industry on How To Use E-Mail To Submit a Study Protocol	2/15/2007	null date	2/15/2007		E7-02577
FDA	FDA-2007-0083	FDA-2007-0083-0001	Implantation or Injectable Dosage Form New Animal Drugs; Trenbolone Acetate and Estradiol		null date			E7-02580
FDA	FDA-2007-0084	FDA-2007-0084-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Medical Devices; Exception From General Requirements for Informed Consent	2/16/2007	null date	2/16/2007		E7-02794
FDA	FDA-2007-0085	FDA-2007-0085-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; TYGACIL	2/16/2007	null date	2/16/2007		E7-02805
FDA	FDA-2007-0086	FDA-2007-0086-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Label Comprehension Study	2/16/2007	null date	2/16/2007		E7-02716

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Food and Drug Administration (FDA)								
FDA	FDA-2007-0087	FDA-2007-0087-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Guidance for Industry on How to Use E-Mail to Submit a Notice of Intent to Slaughter for Human Food Purposes	2/16/2007	null date	2/16/2007		E7-02710
FDA	FDA-2007-0088	FDA-2007-0088-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Establishing and Maintaining a List of United States Dairy Product Manufacturers/Processors With Interest in Exporting to Chile	2/16/2007	null date	2/16/2007		E7-02708
FDA	FDA-2007-0089	FDA-2007-0089-0001	Control of Communicable Diseases; Restrictions on African Rodents, Prairie Dogs, and Certain Other Animals	2/21/2007	3/23/2007	2/21/2007	0910-ZA21	E7-02857
FDA	FDA-2007-0090	FDA-2007-0090-0001	Insect Repellent-Sunscreen Drug Products for Over-the-Counter Human Use; Request for Information and Comments	2/22/2007	null date	2/22/2007	0910-AF43	E7-02890
FDA	FDA-2007-0091	FDA-2007-0091-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; LEVEMIR	2/22/2007	null date	2/22/2007		E7-03001
FDA	FDA-2007-0092	FDA-2007-0092-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; BARACLUDE	2/23/2007	null date	2/23/2007		07-03042
FDA	FDA-2007-0093	FDA-2007-0093-0001	Withdrawal of Approval of 128 Suitability Petitions	2/23/2007	null date	2/23/2007		07-03043
FDA	FDA-2007-0094	FDA-2007-0094-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; AMITIZA	2/23/2007	null date	2/23/2007		E7-03128
FDA	FDA-2007-0095	FDA-2007-0095-0001	Substances Approved for Use in the Preparation of Meat and Poultry Products; Announcement of Effective Date	2/23/2007	null date	2/23/2007		07-00801
FDA	FDA-2007-0096	FDA-2007-0096-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; EXJADE	3/1/2007	null date	2/23/2007		E7-03041
FDA	FDA-2007-0097	FDA-2007-0097-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; S8 OVER-THE-WIRE SYSTEM	3/1/2007	null date	3/1/2007		07-03127
FDA	FDA-2007-0098	FDA-2007-0098-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Reporting and Recordkeeping Requirements and Availability of Sample Electronic Products for Manufacturers and Distributors of Electronic Products	2/27/2007	null date	2/27/2007		E7-03258

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0099	FDA-2007-0099-0001	Draft Guidance for Industry and Food and Drug Administration Staff; Class II Special Controls Guidance Document; Oxygen Pressure Regulators and Oxygen Conserving Devices; Availability	2/27/2007	null date	2/27/2007		E7-03254
FDA	FDA-2007-0100	FDA-2007-0100-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Food Labeling Regulations	2/27/2007	null date	2/27/2007		07-03257
FDA	FDA-2007-0101	FDA-2007-0101-0001	Draft Guidance for Industry on Complementary and Alternative Medicine Products and Their Regulation by the Food and Drug Administration; Availability	2/27/2007	null date	2/27/2007		E7-03259
FDA	FDA-2007-0102	FDA-2007-0102-0001	Medical Devices; Anesthesiology Devices; Oxygen Pressure Regulators and Oxygen Conserving Devices	2/27/2007	5/29/2007	2/27/2007		E7-03253
FDA	FDA-2007-0103	FDA-2007-0103-0001	Safety of Fresh Produce; Public Hearings; Request for Comments	2/27/2007	null date	2/27/2007		07-00891
FDA	FDA-2007-0104	FDA-2007-0104-0001	Draft Guidance for Industry on Advisory Committee Meetings: Preparation and Public Availability of Information Given to Advisory Committee Members; Availability	2/28/2007	null date	2/28/2007		07-00887
FDA	FDA-2007-0105	FDA-2007-0105-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Premarket Notification	2/28/2007	null date	2/28/2007		E7-03444
FDA	FDA-2007-0106	FDA-2007-0106-0001	Guidance for Industry: Eligibility Determination for Donors of Human Cells, Tissues, and Cellular and Tissue-Based Products; Availability	2/28/2007	null date	2/28/2007		E7-03445
FDA	FDA-2007-0107	FDA-2007-0107-0001	The Essentials of Food and Drug Administration Medical Device Regulations: A Primer for Manufacturers and Suppliers; Public Seminar	3/1/2007	null date	3/1/2007		E7-03619
FDA	FDA-2007-0108	FDA-2007-0108-0001	Implantation or Injectable Dosage Form New Animal Drugs; Trenbolone and Estradiol	3/1/2007	null date	3/1/2007		E7-03620
FDA	FDA-2007-0109	FDA-2007-0109-0001	New Animal Drugs For Use in Animal Feeds; Monensin	3/1/2007	null date	3/1/2007		07-03621
FDA	FDA-2007-0110	FDA-2007-0110-0001	New Animal Drugs; Maropitant	3/1/2007	null date	3/1/2007		E7-03402
FDA	FDA-2007-0111	FDA-2007-0111-0001	New Animal Drugs For Use in Animal Feeds; Zilpaterol	3/1/2007	null date	3/1/2007		E7-03615

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0112	FDA-2007-0112-0001	Determination That LAMICTAL (Lamotrigine) Tablets, 50 Milligrams and 250 Milligrams, Were Not Withdrawn From Sale for Reasons of Safety or Effectiveness	3/5/2007	null date	3/5/2007		E7-03713
FDA	FDA-2007-0113	FDA-2007-0113-0001	Joint Meeting of the Anti-Infective Drugs Advisory Committee and the Pediatric Advisory Committee; Notice of Meeting	3/5/2007	null date	3/5/2007		E7-03720
FDA	FDA-2007-0114	FDA-2007-0114-0001	Advisory Committee: Change of Name and Function	3/5/2007	null date	3/5/2007		E7-03716
FDA	FDA-2007-0115	FDA-2007-0115-0001	Cellular, Tissue, and Gene Therapies Advisory Committee; Notice of Meeting	3/5/2007	null date	3/5/2007		E7-03712
FDA	FDA-2007-0116	FDA-2007-0116-0001	Arthritis Advisory Committee; Notice of Meeting	3/5/2007	null date	3/5/2007		E7-03722
FDA	FDA-2007-0117	FDA-2007-0117-0001	Manufacturing Subcommittee of the Advisory Committee for Pharmaceutical Science and Clinical Pharmacology (Formerly Advisory Committee for Pharmaceutical Science); Notice of Meeting	3/5/2007	null date	3/5/2007		E7-03717
FDA	FDA-2007-0118	FDA-2007-0118-0001	Cardiovascular and Renal Drugs Advisory Committee; Notice of Meeting	3/5/2007	null date	3/5/2007		E7-03721
FDA	FDA-2007-0119	FDA-2007-0119-0001	Guidance for Industry on Orally Inhaled and Intranasal Corticosteroids: Evaluation of the Effects on Growth in Children; Availability	3/6/2007	null date	3/6/2007		E7-03807
FDA	FDA-2007-0120	FDA-2007-0120-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Procedures for the Safe and Sanitary Processing and Importing of Fish and Fishery Products	3/7/2007	null date	3/7/2007		E7-03915
FDA	FDA-2007-0121	FDA-2007-0121-0001	Guidance on Drug Safety Information--Food and Drug Administrations Communication to the Public; Availability	3/7/2007	null date	3/7/2007		07-01048
FDA	FDA-2007-0122	FDA-2007-0122-0001	Food and Color Additives and Generally Recognized As Safe Substances; Technical Amendments		null date	3/8/2007		E7-04104
FDA	FDA-2007-0123	FDA-2007-0123-0001	New Animal Drugs for Use in Animal Feeds; Melengestrol, Ractopamine, and Monensin		null date	3/8/2007		E7-04100
FDA	FDA-2007-0124	FDA-2007-0124-0001	Ophthalmic and Topical Dosage Form New Animal Drugs; Imidacloprid and Moxidectin		null date	3/9/2007		E7-04226
FDA	FDA-2007-0125	FDA-2007-0125-0001	Oral Dosage Form New Animal Drugs; Oxfendazole Suspension		null date	3/9/2007		E7-04205

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0126	FDA-2007-0126-0001	Guidance for Industry: Animal Drug User Fees; Fees Exceed Costs Waiver/Reduction; Availability		null date	3/9/2007		E7-04322
FDA	FDA-2007-0127	FDA-2007-0127-0001	Oral Dosage Form New Animal Drugs; Fenbendazole Paste		null date	3/9/2007		E7-04204
FDA	FDA-2007-0128	FDA-2007-0128-0001	Implantation or Injectable Dosage Form New Animal Drugs; Enrofloxacin		null date	3/9/2007		E7-04206
FDA	FDA-2007-0129	FDA-2007-0129-0001	Immune Globulins for Primary Immune Deficiency Diseases: Antibody Specificity, Potency and Testing; Public Workshop	3/12/2007	null date	3/12/2007		E7-04313
FDA	FDA-2007-0130	FDA-2007-0130-0001	Draft Final Guidance for Industry: Guide to Minimize Food Safety Hazards for Fresh-Cut Fruits and Vegetables; Availability; Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request	3/13/2007	null date	3/13/2007		E7-04446
FDA	FDA-2007-0130	FDA-2007-0130-0002	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Guide to Minimize Food Safety Hazards for Fresh-Cut Fruits and Vegetables	10/19/2007	null date	10/19/2007		E7-20632
FDA	FDA-2007-0131	FDA-2007-0131-0001	Electronic Case Report Form Submission; Notice of Pilot Project		null date	3/1/2007		E7-04451
FDA	FDA-2007-0132	FDA-2007-0132-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Control of Communicable Diseases; Restrictions on African Rodents, Prairie Dogs, and Certain Other Animals	3/13/2007	null date	3/13/2007		E7-04450
FDA	FDA-2007-0133	FDA-2007-0133-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Recordkeeping Requirements for Human Food and Cosmetics Manufactured From, Processed With, or Otherwise Containing, Material from Cattle	3/13/2007	null date	3/13/2007		E7-04455
FDA	FDA-2007-0134	FDA-2007-0134-0001	Animal Drug User Fee Act; Public Meeting		null date	3/13/2007		E7-04452
FDA	FDA-2007-0135	FDA-2007-0135-0001	Guidance for Industry and Food and Drug Administration Staff; Statistical Guidance on Reporting Results from Studies Evaluating Diagnostic Tests; Availability	3/13/2007	null date	3/13/2007		E7-04453

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0136	FDA-2007-0136-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Food Labeling; Trans Fatty Acids in Nutrition Labeling	3/13/2007	null date	3/13/2007		E7-04454
FDA	FDA-2007-0137	FDA-2007-0137-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Experimental Evaluation of Variations in Content and Format of the Brief Summary in Direct-to-Consumer Print Advertisements for Prescription Drugs	3/14/2007	null date	3/14/2007		E7-04556
FDA	FDA-2007-0138	FDA-2007-0138-0001	Temperature-Indicating Devices; Thermally Processed Low-Acid Foods Packaged in Hermetically Sealed Containers	3/14/2007	6/12/2007	3/14/2007		07-01172
FDA	FDA-2007-0139	FDA-2007-0139-0001	Medical Devices Regulated by the Center for Biologics Evaluation and Research; Availability of Summaries of Safety and Effectiveness Data for Premarket Approval Applications	3/15/2007	null date	3/15/2007		E7-04677
FDA	FDA-2007-0140	FDA-2007-0140-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Substances Prohibited From Use in Animal Food or Feed; Animal Proteins Prohibited in Ruminant Feed	3/15/2007	null date	3/15/2007		E7-04685
FDA	FDA-2007-0141	FDA-2007-0141-0001	Advisory Committee for Pharmaceutical Science and Clinical Pharmacology (formerly called Advisory Committee for Pharmaceutical Science); Notice of Meeting	3/16/2007	null date	3/16/2007		E7-04797
FDA	FDA-2007-0142	FDA-2007-0142-0001	Anti-Infective Drugs Advisory Committee; Notice of Meeting		null date	3/16/2007		07-04860
FDA	FDA-2007-0143	FDA-2007-0143-0001	Food Substances Affirmed as Generally Recognized as Safe in Feed and Drinking Water of Animals: 25-Hydroxyvitamin D3	3/16/2007	null date	3/16/2007		E7-04796
FDA	FDA-2007-0144	FDA-2007-0144-0001	Draft Guidance for Industry on Indexing Structured Product Labeling; Availability	3/19/2007	null date	3/19/2007		E7-04881
FDA	FDA-2007-0145	FDA-2007-0145-0001	Pediatric Advisory Committee; Notice of Meeting	3/19/2007	null date	3/19/2007		E7-04877

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Food and Drug Administration (FDA) -- 1547 Documents

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Food and Drug Administration (FDA)								
FDA	FDA-2007-0146	FDA-2007-0146-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Survey of Health Care Professionals on the Food Safety and Nutrition Information That They Provide to Pregnant Women	3/20/2007	null date	3/20/2007		E7-05046
FDA	FDA-2007-0147	FDA-2007-0147-0001	National Antimicrobial Resistance Monitoring System Program Subcommittee ofthe Science Advisory Board to the Food and Drug Administration; Notice of Public Meeting	3/21/2007	null date	3/21/2007		E7-05153
FDA	FDA-2007-0148	FDA-2007-0148-0001	Medical Devices Dispute Resolution Panel of the Medical Devices Advisory Committee; Notice of Meeting	3/21/2007	null date	3/21/2007		07-05152
FDA	FDA-2007-0149	FDA-2007-0149-0001	Pulmonary-Allergy Drugs Advisory Committee; Notice of Meeting		null date	3/22/2007		07-05194
FDA	FDA-2007-0150	FDA-2007-0150-0001	Request for Nominations for Voting Members on Public Advisory Committees		null date	3/22/2007		E7-05193
FDA	FDA-2007-0151	FDA-2007-0151-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Threshold of Regulation for Substances Used in Food-Contact Articles	3/22/2007	null date	3/22/2007		E7-05196
FDA	FDA-2007-0152	FDA-2007-0152-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Generic Food and Drug Administration Rapid Response Surveys	3/22/2007	null date	3/22/2007		E7-05195
FDA	FDA-2007-0153	FDA-2007-0153-0001	Draft Guidance for the Public, FDA Advisory Committee Members, and FDA Staff on Procedures for Determining Conflict of Interest and Eligibility for Participation in FDA Advisory Committees; Availability	3/23/2007	null date	3/23/2007		07-01459
FDA	FDA-2007-0154	FDA-2007-0154-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Infant Formula Requirements	3/26/2007	null date	3/26/2007		E7-05470
FDA	FDA-2007-0155	FDA-2007-0155-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; NOVOLOG		null date	3/26/2007		E7-05445
FDA	FDA-2007-0156	FDA-2007-0156-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; EMTRIVA		null date	3/26/2007		E7-05446
FDA	FDA-2007-0157	FDA-2007-0157-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; PREVICOX		null date	3/26/2007		E7-05443

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Food and Drug Administration (FDA)								
FDA	FDA-2007-0158	FDA-2007-0158-0001	Determination That DURICEF (Cefadroxil USP) Tablets, 1 Gram, and Capsules, 500 Milligrams, Were Not Withdrawn From Sale for Reasons of Safety or Effectiveness	3/26/2007	null date	3/26/2007		E7-05415
FDA	FDA-2007-0159	FDA-2007-0159-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; REVLIMID		null date	3/26/2007		E7-05439
FDA	FDA-2007-0160	FDA-2007-0160-0001	Orthopaedic and Rehabilitation Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting	3/26/2007	null date	3/26/2007		E7-05469
FDA	FDA-2007-0161	FDA-2007-0161-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; ONYX LES		null date	3/26/2007		E7-05444
FDA	FDA-2007-0162	FDA-2007-0162-0001	Draft Guidance for Industry and Food and Drug Administration Staff; Modifications to Devices Subject to Premarket Approval--The Premarket Approval Supplement Decision-Making Process; Availability	3/27/2007	null date	3/27/2007		E7-05572
FDA	FDA-2007-0163	FDA-2007-0163-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; A180		null date	3/27/2007		E7-05504
FDA	FDA-2007-0164	FDA-2007-0164-0001	Endocrinologic and Metabolic Drugs Advisory Committee; Notice of Meeting		null date	3/27/2007		E7-05506
FDA	FDA-2007-0165	FDA-2007-0165-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Focus Groups as Used by the Food and Drug Administration	3/27/2007	null date	3/27/2007		E7-05505
FDA	FDA-2007-0166	FDA-2007-0166-0001	Industry Exchange Workshop on Food and Drug Administration Clinical Trial Requirements; Public Workshop	3/28/2007	null date	3/28/2007		E7-05633
FDA	FDA-2007-0167	FDA-2007-0167-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Preparing a Claim of Categorical Exclusion or an Environmental Assessment for Submission to the Center for Food Safety and Applied Nutrition	3/28/2007	null date	3/28/2007		E7-05634
FDA	FDA-2007-0168	FDA-2007-0168-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; INFUSE BONE GRAFT/LT-CAGE LUMBAR TAPERED FUSION DEVICE	3/28/2007	null date	3/28/2007		E7-05635
FDA	FDA-2007-0169	FDA-2007-0169-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; RETEVASE		null date	3/29/2007		E7-05736
FDA	FDA-2007-0170	FDA-2007-0170-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; VAPRISOL		null date	3/29/2007		E7-05737

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Food and Drug Administration (FDA)								
FDA	FDA-2007-0171	FDA-2007-0171-0001	Workshop to Discuss Development of a Womens Health Information Sharing Network		null date	3/29/2007		07-01546
FDA	FDA-2007-0172	FDA-2007-0172-0001	Meeting to Present Work-in-Progress on a Method for Ranking Feed Contaminants According to the Relative Risks They Pose to Animal and Public Health; Part 2: Exposure Scoring for Feed Contaminants; Public Meeting	3/29/2007	null date	3/29/2007		E7-05820
FDA	FDA-2007-0173	FDA-2007-0173-0001	Laxative Drug Products for Over-the-Counter Human Use; Psyllium Ingredients in Granular Dosage Forms		null date	3/29/2007	0910-AF38	07-05740
FDA	FDA-2007-0174	FDA-2007-0174-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Mental Models Study of Food Bioterrorism Risk Awareness	3/30/2007	null date	3/30/2007		07-01577
FDA	FDA-2007-0175	FDA-2007-0175-0001	Draft Guidance for Industry and Review Staff on Target Product Profile--A Strategic Development Process Tool; Availability	3/30/2007	null date	3/30/2007		E7-05949
FDA	FDA-2007-0176	FDA-2007-0176-0001	New Drugs Exempted From Prescription-Dispensing Requirements; Technical Amendment		null date	3/30/2007		E7-05895
FDA	FDA-2007-0177	FDA-2007-0177-0001	The 10th Annual Food and Drug Administration-Orange County Regulatory Affairs Educational Conference	2/1/2007	null date	2/1/2007		07-06052
FDA	FDA-2007-0178	FDA-2007-0178-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; RANEXA		null date	4/2/2007		E7-06061
FDA	FDA-2007-0179	FDA-2007-0179-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; KEPIVANCE	null date	null date	4/2/2007		E7-06053
FDA	FDA-2007-0180	FDA-2007-0180-0001	Electronic Distribution of Prescribing Information for Prescription Drug Products; Public Hearing; Request for Comments	4/2/2007	null date	4/2/2007		07-01604
FDA	FDA-2007-0181	FDA-2007-0181-0001	Reports and guidance documents; availability, etc.: Herpes simplex virus types 1 and 2 serological assays; Class II special controls	4/3/2007	null date	4/3/2007		E7-06168
FDA	FDA-2007-0182	FDA-2007-0182-0001	Meetings: Oncologic Drugs Advisory Committee		null date	4/3/2007		E7-06171
FDA	FDA-2007-0183	FDA-2007-0183-0001	Medical devices: Immunology and microbiology devices--		null date	4/3/2007		E7-06167
FDA	FDA-2007-0184	FDA-2007-0184-0001	Medical devices: Premarket approval applications, list; safety and effectiveness summaries availability	4/3/2007	null date	4/3/2007		E7-06166

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0185	FDA-2007-0185-0001	Reports andguidance documents; availability, etc.: Animal cloning risk assessment plan; industry guidance	4/3/2007	null date	4/3/2007		E7-06170
FDA	FDA-2007-0186	FDA-2007-0186-0001	New Animal Drugs for Use in Animal Feeds; Melengestrol and Lasalocid		null date	4/4/2007		E7-06180
FDA	FDA-2007-0187	FDA-2007-0187-0001	Preparation for International Conference on Harmonisation Meetings in Brussels, Belgium; Public Meeting	4/4/2007	null date	4/4/2007		07-01633
FDA	FDA-2007-0188	FDA-2007-0188-0001	Oral Dosage Form New Animal Drugs; Praziquantel and Pyrantel		null date	4/4/2007		E7-06181
FDA	FDA-2007-0189	FDA-2007-0189-0001	Irradiation in the Production, Processing and Handling of Food	4/4/2007	7/3/2007	4/4/2007	0910-ZA29	07-01636
FDA	FDA-2007-0190	FDA-2007-0190-0001	Manufacturing Subcommittee of the Advisory Committee for Pharmaceutical Science and Clinical Pharmacology (formerly called Advisory Committee for Pharmaceutical Science); Notice of Meeting; Cancellation	4/5/2007	null date	4/5/2007		E7-06288
FDA	FDA-2007-0191	FDA-2007-0191-0001	Advisory Committee for Pharmaceutical Science and Clinical Pharmacology (formerly called Advisory Committee for Pharmaceutical Science); Notice of Meeting; Cancellation	4/5/2007	null date	4/5/2007		E7-06283
FDA	FDA-2007-0192	FDA-2007-0192-0001	Use of Medication Guides to Distribute Drug Risk Information to Patients; Public Hearing		null date	4/9/2007		E7-06506
FDA	FDA-2007-0193	FDA-2007-0193-0001	Draft Guidance for Industry on Orally Disintegrating Tablets; Availability		null date	4/9/2007		E7-06509
FDA	FDA-2007-0194	FDA-2007-0194-0001	Blood Products Advisory Committee; Notice of Meeting		null date	4/9/2007		E7-06594
FDA	FDA-2007-0195	FDA-2007-0195-0001	Irradiation in the Production, Processing and Handling of Food		null date	4/9/2007		E7-06646
FDA	FDA-2007-0196	FDA-2007-0196-0001	Draft Guidance for Clinical Investigators, Sponsors, and Investigational Review Boards on Adverse Event Reporting--Improving Human Subject Protection; Availability	4/9/2007	null date	4/9/2007		E7-06595
FDA	FDA-2007-0197	FDA-2007-0197-0001	Draft Guidance for Industry on the Content and Format of the Dosage and Administration Section of Labeling for Human Prescription Drug and Biological Products; Availability	4/9/2007	null date	4/9/2007		E7-06508
FDA	FDA-2007-0198	FDA-2007-0198-0001	Medical Devices; Technical Amendment		null date	4/9/2007		E7-06290
FDA	FDA-2007-0199	FDA-2007-0199-0001	General Hospital and Personal Use Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting	4/9/2007	null date	4/9/2007		E7-06645

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0200	FDA-2007-0200-0001	Trimethobenzamide Hydrochloride Suppositories; Withdrawal of Approval		null date	4/9/2007		E7-06593
FDA	FDA-2007-0201	FDA-2007-0201-0001	Durand-Wayland, Inc.; Filing of Food Additive Petition		null date	4/11/2007		E7-06765
FDA	FDA-2007-0202	FDA-2007-0202-0001	Ophthalmic and Topical Dosage Form New Animal Drugs; Mupirocin Ointment		null date	4/11/2007		E7-06828
FDA	FDA-2007-0203	FDA-2007-0203-0001	Anti-Infective Drugs Advisory Committee Meeting; Notice of Meeting; Cancellation		null date	4/12/2007		07-01825
FDA	FDA-2007-0204	FDA-2007-0204-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Adoption of the Food and Drug Administration Food Code by Local, State, and Tribal Governments	4/13/2007	null date	4/13/2007		E7-06983
FDA	FDA-2007-0205	FDA-2007-0205-0001	Supplements and Other Changes to an Approved Application; Public Meeting; Reopening of Comment Period		null date	4/13/2007		07-06985
FDA	FDA-2007-0206	FDA-2007-0206-0001	Vaccines and Related Biological Products Advisory Committee; Notice of Meeting		null date	4/16/2007		E7-07090
FDA	FDA-2007-0207	FDA-2007-0207-0001	Medical Device User Fee and Modernization Act; Public Meeting		null date	4/18/2007		07-01919
FDA	FDA-2007-0208	FDA-2007-0208-0001	Memorandum of Understanding Between the National Cancer Institute and the Food and Drug Administration	4/18/2007	null date	4/18/2007		07-01921
FDA	FDA-2007-0209	FDA-2007-0209-0001	Withdrawal of Approval of New Animal Drug Applications; Pyrantel; Tylosin; Tylosin and Sulfamethazine	4/19/2007	null date	4/19/2007		07-07461
FDA	FDA-2007-0210	FDA-2007-0210-0001	New Animal Drugs For Use in Animal Feed; Withdrawal of Approval of NADAs; Pyrantel; Tylosin; Tylosin and Sulfamethazine	4/19/2007	null date	4/19/2007		E7-07460
FDA	FDA-2007-0211	FDA-2007-0211-0001	Implantation or Injectable Dosage Form New Animal Drugs; Withdrawal of Approval of NADAs; Estradiol Benzoate	4/19/2007	null date	4/19/2007		E7-07458
FDA	FDA-2007-0212	FDA-2007-0212-0001	Withdrawal of Approval of New Animal Drug Applications; Estradiol Benzoate		null date	4/19/2007		07-01941
FDA	FDA-2007-0213	FDA-2007-0213-0001	Preparation for International Conference on Harmonization Meetings in Brussels, Belgium; Public Meeting	4/20/2007	null date	4/20/2007		07-01952
FDA	FDA-2007-0214	FDA-2007-0214-0001	Oral Dosage Form New Animal Drugs; Clindamycin Solution		null date	4/20/2007		E7-07472
FDA	FDA-2007-0215	FDA-2007-0215-0001	New Animal Drugs; Florfenicol		null date	4/20/2007		E7-07475
FDA	FDA-2007-0216	FDA-2007-0216-0001	Oral Dosage Form New Animal Drugs; Dexmedetomidine; Technical Amendment		null date	4/20/2007		E7-07594

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0217	FDA-2007-0217-0001	Antiviral Drugs Advisory Committee; Amendment of Notice		null date	4/23/2007		07-02001
FDA	FDA-2007-0218	FDA-2007-0218-0001	Guidance for Industry and Food and Drug Administration Staff; Class II Special Controls Guidance Document: Computerized Labor Monitoring Systems; Availability	4/24/2007	null date	4/24/2007		E7-07700
FDA	FDA-2007-0219	FDA-2007-0219-0001	Medical Devices; Obstetrical and Gynecological Devices; Classification of Computerized Labor Monitoring System	4/24/2007	null date	4/24/2007		E7-07702
FDA	FDA-2007-0220	FDA-2007-0220-0001	Summaries of Medical and Clinical Pharmacology Reviews of Pediatric Studies; Availability		null date	4/24/2007		E7-07717
FDA	FDA-2007-0221	FDA-2007-0221-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Human Tissue Intended for Transplantation	4/25/2007	null date	4/25/2007		E7-07815
FDA	FDA-2007-0222	FDA-2007-0222-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Submission of Petitions: Food Additive, Color Additive (Including Labeling), and Generally Recognized as Safe Affirmation; Electronic Submission Using Food and Drug Administration Forms 3503 and 3504	4/25/2007	null date	4/25/2007		E7-07813
FDA	FDA-2007-0223	FDA-2007-0223-0001	Canned Pacific Salmon Deviating From Identity Standard; Temporary Permit for Market Testing		null date	4/27/2007		E7-08039
FDA	FDA-2007-0224	FDA-2007-0224-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Human Cells, Tissues, and Cellular and Tissue-Based Products: Establishment Registration and Listing; Form Food and Drug Administration 3356; Eligibility Determination for Donors; and Current Good Tissue Practice	4/27/2007	null date	4/27/2007		E7-08038
FDA	FDA-2007-0225	FDA-2007-0225-0001	Guidance for Industry: Protocols for the Conduct of Method Transfer Studies for Type C Medicated Feed Assay Methods; Availability	4/27/2007	null date	4/27/2007		E7-08042
FDA	FDA-2007-0226	FDA-2007-0226-0001	Oral Dosage Form New Animal Drugs; Diclazuril		null date	4/27/2007		E7-0804
FDA	FDA-2007-0227	FDA-2007-0227-0001	Medical Device User Fee and Modernization Act; Public Meeting; Correction		null date	4/27/2007		07-02085

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0228	FDA-2007-0228-0001	Iceberg Water Deviating From Identity Standard; Temporary Permit for Market Testing		null date	4/27/2007		E7-08040
FDA	FDA-2007-0229	FDA-2007-0229-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Applications for Food and Drug Administration Approval to Market a New Drug: Patent Submission and Listing Requirements and Application of 30-Month Stays on Approval of Abbreviated New Drug Applications Certifying That a Patent Claiming a Drug Is Valid or Will Not Be Infringed	4/30/2007	null date	4/30/2007		E7-08141
FDA	FDA-2007-0230	FDA-2007-0230-0001	Defining and Implementing Quality in Clinical Investigations: From Design to Completion; Public Workshop; Request for Comments	4/30/2007	null date	4/30/2007		E7-08137
FDA	FDA-2007-0231	FDA-2007-0231-0001	Memorandum of Understanding Between the National Cancer Institute and the Food and Drug Administration	4/30/2007	null date	4/30/2007		07-02106
FDA	FDA-2007-0232	FDA-2007-0232-0001	Oral Dosage Form New Animal Drugs; Fenbendazole Paste		null date	5/2/2007		E7-08391
FDA	FDA-2007-0233	FDA-2007-0233-0001	New Animal Drugs; Change of Sponsors Address		null date	5/2/2007		E7-08322
FDA	FDA-2007-0234	FDA-2007-0234-0001	Guidance for Industry on Testing of Glycerin for Diethylene Glycol; Availability	lldate	null date	5/2/2007		E7-08389
FDA	FDA-2007-0235	FDA-2007-0235-0001	Pediatric Oncology Subcommittee of the Oncologic Drugs Advisory Committee; Notice of Meeting	5/7/2007	null date	5/7/2007		E7-08656
FDA	FDA-2007-0236	FDA-2007-0236-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Food and Drug Administration Survey of Current Manufacturing Practices in the Food Industry	5/8/2007	null date	5/8/2007		E7-08783
FDA	FDA-2007-0237	FDA-2007-0237-0001	Draft Guidance for Industry and Food and Drug Administration Staff; Class II Special Controls Guidance Document: Absorbable Hemostatic Device; Availability; Reopening of Comment Period	5/8/2007	null date	5/8/2007		E7-08780
FDA	FDA-2007-0238	FDA-2007-0238-0001	Voluntary Self-Inspection of Medicated Feed Manufacturing Facilities; Draft Compliance Policy Guide; Availability; Reopening of Comment Period	5/8/2007	null date	5/8/2007		E7-08781

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Food and Drug Administration (FDA)								
FDA	FDA-2007-0239	FDA-2007-0239-0001	Guidance for Industry: Analytical Methods Description for Type C Medicated Feeds; Availability	9/1/2007	null date	5/9/2007		E7-08808
FDA	FDA-2007-0240	FDA-2007-0240-0001	New Animal Drugs; Change of Sponsors Name and Address		null date	5/9/2007		E7-08870
FDA	FDA-2007-0241	FDA-2007-0241-0001	Guidance for Industry and Food and Drug Administration Staff; Class II Special Controls Guidance Document: Gene Expression Profiling Test System for Breast Cancer Prognosis; Availability	5/9/2007	null date	5/9/2007		E7-08872
FDA	FDA-2007-0242	FDA-2007-0242-0001	Medical Devices; Immunology and Microbiology Devices; Classification of Gene Expression Profiling Test System for Breast Cancer Prognosis	5/9/2007	null date	5/9/2007		E7-08871
FDA	FDA-2007-0243	FDA-2007-0243-0001	Certain Other Dosage Form New Animal Drugs; Oxytetracycline		null date	5/9/2007		E7-08869
FDA	FDA-2007-0244	FDA-2007-0244-0001	Circulatory System Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting	null date	null date	5/10/2007		E7-09054
FDA	FDA-2007-0245	FDA-2007-0245-0001	Vaccines and Related Biological Products Advisory Committee; Amendment of Notice	null date	null date	5/10/2007		E7-09053
FDA	FDA-2007-0246	FDA-2007-0246-0001	Draft Guidance for Industry on Protecting the Rights, Safety, and Welfare of Study Subjects--Supervisory Responsibilities of Investigators; Availability	5/10/2007	null date	5/10/2007		E7-09055
FDA	FDA-2007-0247	FDA-2007-0247-0001	Guidance for Industry on Computerized Systems Used in Clinical Investigations; Availability	2:00:00 null date	null date	5/10/2007		E7-09056
FDA	FDA-2007-0248	FDA-2007-0248-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Hazard Analysis and Critical Control Point Procedures for the Safe and Sanitary Processing and Importing of Juice	5/14/2007	null date	5/14/2007		E7-09220
FDA	FDA-2007-0249	FDA-2007-0249-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Food Labeling Regulations	5/14/2007	null date	5/14/2007		E7-09219
FDA	FDA-2007-0250	FDA-2007-0250-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Information Program on Clinical Trials for Serious or Life-Threatening Diseases: Maintaining a Data Bank	5/14/2007	null date	5/14/2007		E7-09221

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AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0251	FDA-2007-0251-0001	Determination That MEPRON (Atovaquone) Tablets, 250 milligrams, Were Not Withdrawn From Sale for Reasons of Safety or Effectiveness	5/16/2007	null date	5/16/2007		E7-09348
FDA	FDA-2007-0252	FDA-2007-0252-0001	Draft Guidance for Industry and Review Staff on Labeling for Human Prescription Drugs--Determining Established Pharmacologic Class for Use in the Highlights of Prescribing Information; Availability	5/16/2007	null date	5/16/2007		E7-09347
FDA	FDA-2007-0253	FDA-2007-0253-0001	Agency Information Collection Activities; Announcementof Office of Management and Budget Approval; Cosmetic Labeling Regulations	5/16/2007	null date	5/16/2007		E7-09436
FDA	FDA-2007-0254	FDA-2007-0254-0001	Guidance for Industry on Clinical Trial Endpoints for the Approval of Cancer Drugs and Biologics; Availability	5/16/2007	null date	5/16/2007		E7-09345
FDA	FDA-2007-0255	FDA-2007-0255-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Administrative Procedures for the Clinical Laboratory Improvement Amendments of 1988 Categorization	5/16/2007	null date	5/16/2007		E7-09435
FDA	FDA-2007-0256	FDA-2007-0256-0001	Implantation or Injectable Dosage Form New Animal Drugs; Ivermectin and Clorsulon		null date	5/17/2007		E7-09517
FDA	FDA-2007-0257	FDA-2007-0257-0001	Implantation or Injectable Dosage Form New Animal Drugs; Ivermectin		null date	5/17/2007		E7-09515
FDA	FDA-2007-0258	FDA-2007-0258-0001	Oral Dosage Form New Animal Drugs; Pimobendan		null date	5/17/2007		E7-09516
FDA	FDA-2007-0259	FDA-2007-0259-0001	Implantation or Injectable Dosage Form New Animal Drugs; Butorphanol		null date	5/18/2007		E7-09557
FDA	FDA-2007-0260	FDA-2007-0260-0001	Oral Dosage Form New Animal Drugs; Phenylbutazone Powder		null date	5/18/2007		E7-09559
FDA	FDA-2007-0261	FDA-2007-0261-0001	New Animal Drugs; Change of Sponsors Address		null date	5/18/2007		E7-09555
FDA	FDA-2007-0262	FDA-2007-0262-0001	International Cooperation on Harmonisation of Technical Requirements for Registration of Veterinary Medicinal Products (VICH); Draft Guidance for Industry on Target Animal Safety for Veterinary Pharmaceutical Products, VICH GL43, Request for Comments; Availability	5/18/2007	null date	5/18/2007		E7-09592

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Food and Drug Administration (FDA)								
FDA	FDA-2007-0263	FDA-2007-0263-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; GALILEO INTRAVASCULAR RADIOTHERAPY SYSTEM	5/21/2007	null date	5/21/2007		E7-09720
FDA	FDA-2007-0264	FDA-2007-0264-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; NOXAFIL		null date	5/21/2007		E7-09730
FDA	FDA-2007-0265	FDA-2007-0265-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Guidance for Industry on Continuous Marketing Applications: Pilot 2--Scientific Feedback and Interactions During Development of Fast Track Products Under the Prescription Drug User Fee Act	5/21/2007	null date	5/21/2007		E7-09709
FDA	FDA-2007-0266	FDA-2007-0266-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; IRESSA		null date	5/21/2007		E7-09733
FDA	FDA-2007-0267	FDA-2007-0267-0001	Science Board to the Food and Drug Administration; Notice of Meeting	5/21/2007	null date	5/21/2007		E7-09737
FDA	FDA-2007-0268	FDA-2007-0268-0001	Food and Drug Administration Modernization Act of 1997: Modifications to the List of Recognized Standards, Recognition List Number: 017	5/21/2007	null date	5/21/2007		E7-09718
FDA	FDA-2007-0269	FDA-2007-0269-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; FOSRENOL		null date	5/22/2007		E7-09787
FDA	FDA-2007-0270	FDA-2007-0270-0001	Implementation of Risk Minimization Action Plans (RiskMAPs) to Support Quality Use of Pharmaceuticals: Opportunities and Challenges; Public Workshop	5/23/2007	null date	5/23/2007		07-02574
FDA	FDA-2007-0271	FDA-2007-0271-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; BEXTRA	null date	null date	5/23/2007		E7-09957
FDA	FDA-2007-0272	FDA-2007-0272-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; ELAPRASE	null date	null date	5/23/2007		E7-09951
FDA	FDA-2007-0273	FDA-2007-0273-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; GARDASIL	null date	null date	5/23/2007		E7-09950
FDA	FDA-2007-0274	FDA-2007-0274-0001	Determination That Protamine Sulfate Injection and 26 Other Drug Products Were Not Withdrawn From Sale for Reasons of Safety or Effectiveness	5/23/2007	null date	5/23/2007		E7-09962
FDA	FDA-2007-0275	FDA-2007-0275-0001	Determination That ESTROSTEP 21 (Ethinyl Estradiol and Norethindrone Acetate) Tablets Were Not Withdrawn From Sale for Reasons of Safety or Effectiveness	5/23/2007	null date	5/23/2007		E7-09949

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0276	FDA-2007-0276-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; ORENCIA		null date	5/23/2007		E7-09945
FDA	FDA-2007-0277	FDA-2007-0277-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; SOMAVERT		null date	5/24/2007		E7-10052
FDA	FDA-2007-0278	FDA-2007-0278-0001	Determination of Regulatory Review Periods for Purposes of Patent Extension; SPRYCEL--New Drug Applications 21-986 and 22-072	5/25/2007	null date	5/25/2007		E7-10089
FDA	FDA-2007-0279	FDA-2007-0279-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Health and Diet Survey	5/25/2007	null date	5/25/2007		E7-10086
FDA	FDA-2007-0280	FDA-2007-0280-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; PREZISTA	null date	null date	5/25/2007		E7-10147
FDA	FDA-2007-0281	FDA-2007-0281-0001	Draft Guidance for Industry on Complementary and Alternative Medicine Products and Their Regulation by the Food and Drug Administration; Availability	5/25/2007	null date	5/25/2007		07-02610
FDA	FDA-2007-0282	FDA-2007-0282-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; MYOZYME		null date	5/25/2007		E7-10087
FDA	FDA-2007-0283	FDA-2007-0283-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; KDR 401 and 403 PACEMAKERS	5/25/2007	null date	5/25/2007		E7-10127
FDA	FDA-2007-0284	FDA-2007-0284-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Orphan Drugs	5/29/2007	null date	5/29/2007		E7-10271
FDA	FDA-2007-0284	FDA-2007-0284-0002	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Orphan Drugs	8/29/2007	null date	8/29/2007		E7-17094
FDA	FDA-2007-0285	FDA-2007-0285-0001	Timed-Release Drug Products Containing Guaifenesin; Enforcement Action Dates	null date	null date	5/29/2007		E7-10266
FDA	FDA-2007-0286	FDA-2007-0286-0001	Joint Meeting of the Gastrointestinal Drugs Advisory Committee and the Drug Safety and Risk Management Advisory Committee; Notice of Meeting	5/29/2007	null date	5/29/2007		E7-10270
FDA	FDA-2007-0287	FDA-2007-0287-0001	Orthopaedic and Rehabilitation Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting	5/29/2007	null date	5/29/2007		E7-10267
FDA	FDA-2007-0288	FDA-2007-0288-0001	Interim Melamine and Melamine Analogues Safety/Risk Assessment; Availability		null date	5/30/2007		07-02679

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0289	FDA-2007-0289-0001	Guidance for Industry: Chemistry, Manufacturing, and Control Changes to an Approved New Animal Drug Application or Abbreviated New Animal Drug Application		null date	5/31/2007		E7-10515
FDA	FDA-2007-0290	FDA-2007-0290-0001	Draft Guidances for Industry Describing Product-Specific Bioequivalence Recommendations; Availability		null date	5/31/2007		07-10491
FDA	FDA-2007-0291	FDA-2007-0291-0001	Draft Guidance for Industry on Bioequivalence Recommendations for Specific Products		null date	5/31/2007		E7-10492
FDA	FDA-2007-0292	FDA-2007-0292-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; X-STOP INTERSPINOUS PROCESS DECOMPRESSION SYSTEM		null date	6/1/2007		E7-10618
FDA	FDA-2007-0293	FDA-2007-0293-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; GEM 21S GROWTH-FACTOR ENHANCED MATRIX	6/1/2007	null date	6/1/2007		E7-10633
FDA	FDA-2007-0294	FDA-2007-0294-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; PHAKIC INTRAOCULAR LENSES	6/1/2007	null date	6/1/2007		07-10631
FDA	FDA-2007-0295	FDA-2007-0295-0001	Guidance for Industry: Clinical Data Needed to Support the Licensure of Pandemic Influenza Vaccines; Availability	6/1/2007	null date	6/1/2007		E7-10499
FDA	FDA-2007-0296	FDA-2007-0296-0001	Guidance for Industry: Clinical Data Needed to Support the Licensure of Seasonal Inactivated Influenza Vaccines; Availability	6/1/2007	null date	6/1/2007		E7-10497
FDA	FDA-2007-0297	FDA-2007-0297-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; ZILMAX	null date	null date	6/1/2007		E7-10602
FDA	FDA-2007-0298	FDA-2007-0298-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Registration of Food Facilities Under the Public Health Security and Bioterrorism Preparedness and Response Act of 2002	6/1/2007	null date	6/1/2007		E7-10617
FDA	FDA-2007-0299	FDA-2007-0299-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; LANTUS		null date	6/1/2007		E7-10632
FDA	FDA-2007-0300	FDA-2007-0300-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; NAMENDA	5/1/2007	null date	6/5/2007		E7-10730
FDA	FDA-2007-0301	FDA-2007-0301-0001	Guidance for Industry: Refrigerated Carrot Juice and Other Refrigerated Low-Acid Juices; Availability	6/5/2007	null date	6/5/2007		7-10792
FDA	FDA-2007-0302	FDA-2007-0302-0001	Advisory Committee; Risk Communication Advisory Committee; Establishment	6/5/2007	null date	6/5/2007		E7-10740

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0303	FDA-2007-0303-0001	New Animal Drugs; Change of Sponsors Address	6/5/2007	null date	6/5/2007		E7-10771
FDA	FDA-2007-0304	FDA-2007-0304-0001	Draft Guidance for Industry on Providing Regulatory Submissions in Electronic Format--Receipt Date; Availability	6/5/2007	null date	6/5/2007		E7-10780
FDA	FDA-2007-0305	FDA-2007-0305-0001	Request for Nominations for Voting Members on a Public Advisory Committee; Risk Communication Advisory Committee	6/5/2007	null date	6/5/2007		E7-10737
FDA	FDA-2007-0306	FDA-2007-0306-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Prior Notice of Imported Food Under the Public Health Security and Bioterrorism Preparedness and Response Act of 2002	6/5/2007	null date	6/5/2007		E7-10785
FDA	FDA-2007-0307	FDA-2007-0307-0001	Advisory Committee Information Hotline		null date	6/5/2007		E7-10738
FDA	FDA-2007-0308	FDA-2007-0308-0001	Joint Meeting of the Endocrinologic and Metabolic Drugs Advisory Committee and the Drug Safety and Risk Management Advisory Committee; Notice of Meeting	6/6/2007	null date	6/6/2007		E7-10850
FDA	FDA-2007-0309	FDA-2007-0309-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; RAPLON		null date	6/6/2007		E7-10853
FDA	FDA-2007-0310	FDA-2007-0310-0001	Implantation or Injectable Dosage Form New Animal Drugs; Spectinomycin Sulfate		null date	6/1/2007		E7-10801
FDA	FDA-2007-0311	FDA-2007-0311-0001	Determination That CEFOTAN (Cefotetan Disodium For Injection), Equivalent 1 Gram Base/Vial and 2 Grams Base/Vial, Was Not Withdrawn From Sale for Reasons of Safety or Effectiveness	6/7/2007	null date	6/7/2007		E7-10959
FDA	FDA-2007-0312	FDA-2007-0312-0001	Guidance for Industry and Food and Drug Administration Staff; Assayed and Unassayed Quality Control Material; Availability	6/7/2007	null date	6/7/2007		E7-10996
FDA	FDA-2007-0313	FDA-2007-0313-0001	Medical Devices; Availability of Safety and Effectiveness Summaries for Premarket Approval Applications	6/7/2007	null date	6/7/2007		E7-11002
FDA	FDA-2007-0314	FDA-2007-0314-0001	Science Board to the Food and Drug Administration; Amendment of Notice		null date	6/7/2007		07-02829
FDA	FDA-2007-0315	FDA-2007-0315-0001	Determination of Regulatory Review Period for Purposes of Patent Extension; CHANTIX	7/1/2007	null date	6/7/2007		E7-10915

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0316	FDA-2007-0316-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Substantiation for Dietary Supplement Claims Made Under the Federal Food, Drug, and Cosmetic Act	6/7/2007	null date	6/7/2007		E7-10911
FDA	FDA-2007-0317	FDA-2007-0317-0001	Draft Guidance for Industry on Malaria: Developing Drug and Nonvaccine Biological Products for Treatment and Prophylaxis; Availability	6/7/2007	null date	6/7/2007		E7-11001
FDA	FDA-2007-0318	FDA-2007-0318-0001	Request for Notification From Industry Organizations Interested in Participating in Selection Process for Nonvoting Industry Representatives on Public Advisory Committees and Request for Nominations for Nonvoting Industry Representatives on Public Advisory Committees	6/8/2007	null date	6/8/2007		E7-11065
FDA	FDA-2007-0319	FDA-2007-0319-0002	Use of Ozone-Depleting Substances; Removal of Essential-Use Designations; Public Meeting	7/9/2007	8/10/2007	7/9/2007	0910-AF93	E7-13300
FDA	FDA-2007-0319	FDA-2007-0319-0001	Use of Ozone-Depleting Substances; Removal of Essential-Use Designations	6/11/2007	8/10/2007	6/11/2007	0910-AF93	07-02883
FDA	FDA-2007-0319	FDA-2007-0319-0003	Use of Ozone-Depleting Substances; Removal of Essential-Use Designations; Extension of Comment Period	8/7/2007	9/10/2007	8/7/2007	0910-AF93	E7-15372
FDA	FDA-2007-0320	FDA-2007-0320-0001	Request for Notification From Industry Organizations Interested in Participating in Selection Process for Nonvoting Industry Representatives on FoodSafety Public Advisory Committee and Request for Nominations for Nonvoting Industry Representatives on Food Safety Public Advisory Committee	6/11/2007	null date	6/11/2007		E7-11141
FDA	FDA-2007-0321	FDA-2007-0321-0001	Guidance for Industry and Food and Drug Administration Staff; Class II Special Controls Guidance Document: Intervertebral Body Fusion Device; Availability	6/12/2007	null date	6/12/2007		E7-11235
FDA	FDA-2007-0322	FDA-2007-0322-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Adverse Event Pilot Program for Medical Products (Formally Medical Device Adverse Event Reporting Program)	6/13/2007	null date	6/13/2007		E7-11400

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0323	FDA-2007-0323-0001	Agency Information Collection Activities; Proposed Collection; CommentRequest; Animal Drug User Fees and Fee Waivers and Reductions	6/14/2007	null date	6/14/2007		E7-11425
FDA	FDA-2007-0324	FDA-2007-0324-0001	Otsuka Pharmaceutical Co., Ltd.; Withdrawal of Approval of a New Drug Application	null date	null date	6/14/2007		E7-11427
FDA	FDA-2007-0325	FDA-2007-0325-0001	Oncologic Drugs Advisory Committee; Notice of Meeting		null date	6/14/2007		E7-11496
FDA	FDA-2007-0326	FDA-2007-0326-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Label Comprehension Study	6/15/2007	null date	6/15/2007		E7-11528
FDA	FDA-2007-0327	FDA-2007-0327-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Animal Drug User Fee Cover Sheet, FDA Form 3546	6/15/2007	null date	6/15/2007		E7-11527
FDA	FDA-2007-0328	FDA-2007-0328-0001	New Animal Drugs for Use in Animal Feeds; Lincomycin		null date	6/18/2007		E7-11611
FDA	FDA-2007-0329	FDA-2007-0329-0001	Licensure of Apheresis Blood Products; Public Workshop		null date	6/18/2007		E7-11615
FDA	FDA-2007-0330	FDA-2007-0330-0001	Anthrax Vaccines--Bridging Correlates of Protection in Animals to Immunogenicity in Humans; Public Workshop	6/18/2007	null date	6/18/2007		E7-11613
FDA	FDA-2007-0331	FDA-2007-0331-0001	Guidance for Industry and Food and Drug Administration Staff; Pharmacogenetic Tests and Genetic Tests for Heritable Markers; Availability	6/19/2007	null date	6/19/2007		E7-11817
FDA	FDA-2007-0332	FDA-2007-0332-0001	Human Cells, Tissues, and Cellular and Tissue-Based Products; Donor Screening and Testing, and Related Labeling	6/19/2007	null date	6/19/2007		E7-11795
FDA	FDA-2007-0333	FDA-2007-0333-0001	Cellular, Tissue, and Gene Therapies Advisory Committee; Notice of Meeting		null date	6/19/2007		E7-11728
FDA	FDA-2007-0334	FDA-2007-0334-0001	Science Board to the Food and Drug Administration; Amendment of Notice; Correction		null date	6/19/2007		E7-11727
FDA	FDA-2007-0335	FDA-2007-0335-0001	Listing of Color Additives Subject to Certification; D&C Black No. 3	7/19/2007	7/19/2007	6/19/2007		E7-11801
FDA	FDA-2007-0336	FDA-2007-0336-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Medical Devices Third-Party Review Under the Food and Drug Administration Modernization Act	6/21/2007	null date	6/21/2007		E7-11981

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Food and Drug Administration (FDA)								
FDA	FDA-2007-0337	FDA-2007-0337-0001	Guidance for Industry: Informed Consent Recommendations for Source Plasma Donors Participating in Plasmapheresis and Immunization Programs; Availability	6/21/2007	null date	6/21/2007		E7-11997
FDA	FDA-2007-0338	FDA-2007-0338-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Preparing a Claim of Categorical Exclusion or an Environmental Assessment for Submission to the Center for Food Safety and Applied Nutrition	6/21/2007	null date	6/21/2007		E7-11969
FDA	FDA-2007-0339	FDA-2007-0339-0001	International Cooperation on Harmonisation of Technical Requirements for Approval of Veterinary Medicinal Products (VICH); Revised Draft Guidance for Industry on Pharmacovigilance of Veterinary Medicinal Products: Controlled List of Terms (VICH GL30); Request for Comments; Availability	6/21/2007	null date	6/21/2007		E7-11996
FDA	FDA-2007-0340	FDA-2007-0340-0001	Draft Guidance for Industry on Use of the Computer Crossmatch; Availability		null date	6/21/2007		E7-11998
FDA	FDA-2007-0341	FDA-2007-0341-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Information From United States Processors That Export to the European Community	6/21/2007	null date	6/21/2007		E7-11980
FDA	FDA-2007-0342	FDA-2007-0342-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Guidance for Industry on Special Protocol Assessment	6/22/2007	null date	6/22/2007		E7-12056
FDA	FDA-2007-0343	FDA-2007-0343-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Infant Formula Requirements	6/22/2007	null date	6/22/2007		E7-12057
FDA	FDA-2007-0344	FDA-2007-0344-0001	Current Good Manufacturing Practice in Manufacturing, Packaging, Labeling, or Holding Operations for Dietary Supplements	6/25/2007	null date	6/25/2007	0910-AB88	07-03039
FDA	FDA-2007-0345	FDA-2007-0345-0001	Petition to Request an Exemption From 100 Percent Identity Testing of Dietary Ingredients: Current Good Manufacturing Practice in Manufacturing, Packaging, Labeling, or Holding Operations for Dietary Supplements	6/25/2007	null date	6/25/2007	0910-AB88	07-03038

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Food and Drug Administration (FDA) -- 1547 Documents

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Food and Drug Administration (FDA)								
FDA	FDA-2007-0345	FDA-2007-0345-0002	Petition to Request an Exemption From 100 Percent Identity Testing of Dietary Ingredients: Current Good Manufacturing Practice in Manufacturing, Packaging, Labeling, or Holding Operations for Dietary Supplements; Extension of Comment Period	9/17/2007	10/24/2007	9/17/2007	0910-AB88	E7-18293
FDA	FDA-2007-0346	FDA-2007-0346-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Adoption of Food and Drug Administration Food Code by Local, State and Tribal Governments	6/28/2007	null date	6/28/2007		E7-12499
FDA	FDA-2007-0347	FDA-2007-0347-0001	Request for Notification From Industry Organizations Interested in Participating in the Selection Process for a Nonvoting Industry Representative on the Allergenic Products Advisory Committee and Request for Nominations for a Nonvoting Industry Representative on the Allergenic Products Advisory Committee	6/28/2007	null date	6/28/2007		E7-12527
FDA	FDA-2007-0348	FDA-2007-0348-0001	Dermatologic and Ophthalmic Drugs Advisory Committee and the Drug Safety and Risk Management Advisory Committee; Notice of Meeting	6/28/2007	null date	6/28/2007		E7-12501
FDA	FDA-2007-0349	FDA-2007-0349-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Institutional Review Boards	6/28/2007	null date	6/28/2007		E7-12496
FDA	FDA-2007-0350	FDA-2007-0350-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Pre-market Approval of Medical Devices	6/28/2007	null date	6/28/2007		E7-12502
FDA	FDA-2007-0351	FDA-2007-0351-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; General Licensing Provisions: Biologics License Application, Changes to an Approved Application, Labeling, Revocation and Suspension, Postmarketing Studies Status Reports, and Forms FDA 356h and 2567	6/28/2007	null date	6/28/2007		E7-12497
FDA	FDA-2007-0352	FDA-2007-0352-0001	Menley and James Laboratories, Inc. et al.; Proposal to Withdraw Approval of Six New Drug Applications; Opportunity for a Hearing	6/28/2007	null date	6/28/2007		E7-12494

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Food and Drug Administration (FDA)								
FDA	FDA-2007-0353	FDA-2007-0353-0001	General and Plastic Surgery Devices; Reclassification of the Tissue Adhesive for Topical Approximation of Skin Device	7/3/2007	9/4/2007	7/3/2007		E7-12797
FDA	FDA-2007-0354	FDA-2007-0354-0001	Draft Guidance for Industry on Integrated Summaries of Effectiveness and Safety: Location Within the Common Technical Document; Availability	7/3/2007	null date	7/3/2007		E7-12792
FDA	FDA-2007-0355	FDA-2007-0355-0001	Neurological Devices; Denial of Request for Change in Classification of Cutaneous Electrode		null date	7/3/2007		E7-12882
FDA	FDA-2007-0356	FDA-2007-0356-0001	Medical Devices; Cardiovascular Devices; Denial of Request for Change in Classification of Impedance Plethysmograph	7/3/2007	null date	7/3/2007		E7-12883
FDA	FDA-2007-0357	FDA-2007-0357-0001	Program Priorities in the Center for Food Safety and Applied Nutrition; Request for Comments		null date	7/3/2007		E7-12884
FDA	FDA-2007-0358	FDA-2007-0358-0001	Medical Devices; General Hospital and Personal Use Devices; Classification of the Filtering Facepiece Respirator for Use by the General Public in Public Health Medical Emergencies; Availability	7/3/2007	null date	7/3/2007		E7-12790
FDA	FDA-2007-0359	FDA-2007-0359-0001	Request for Nominations for Voting Members on Public Advisory Panels or Committees		null date	7/3/2007		E7-12799
FDA	FDA-2007-0360	FDA-2007-0360-0001	Draft Guidance for Industry and Food and Drug Administration Staff; Class II Special Controls Guidance Document: Tissue Adhesive for the Topical Approximation of Skin; Availability	7/3/2007	null date	7/3/2007		E7-12795
FDA	FDA-2007-0361	FDA-2007-0361-0001	Clinical Studies of Safety and Effectiveness of Orphan Products; Availability of Grants; Request for Applications: RFA-FD08-001; Research Project Grants (R01); Catalog of Federal Domestic Assistance Number: 93.103	7/3/2007	null date	7/3/2007		E7-12881
FDA	FDA-2007-0362	FDA-2007-0362-0001	Medical Devices; General Hospital and Personal Use Devices; Classification of the Filtering Facepiece Respirator for Use by the General Public in Public Health Medical Emergencies	7/3/2007	null date	7/3/2007		E7-12789
FDA	FDA-2007-0363	FDA-2007-0363-0001	New Animal Drugs; Change of Sponsors Name; Liquid Crystalline Trypsin, Peru Balsam, Castor Oil	5/1/2007	null date	7/5/2007		E7-13010

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0364	FDA-2007-0364-0001	Medical Devices: The Mammography Quality Standards Act of 1992 and Subsequent Mammography Quality Standards Reauthorization Act and Amendments; Inspection Fees	7/6/2007	null date	7/6/2007		E7-13044
FDA	FDA-2007-0365	FDA-2007-0365-0001	Cooperative Agreement to Support the Joint Institute for Food Safety and Applied Nutrition	7/6/2007	null date	7/6/2007		E7-13046
FDA	FDA-2007-0366	FDA-2007-0366-0001	Guidance for Industry on ANDAs: Pharmaceutical Solid Polymorphism; Chemistry, Manufacturing, and Controls Information; Availability	7/9/2007	null date	7/9/2007		E7-13171
FDA	FDA-2007-0367	FDA-2007-0367-0001	Nippon Oil Corp.; Filing of Color Additive Petition	7/9/2007	null date	7/9/2007		07-13161
FDA	FDA-2007-0368	FDA-2007-0368-0001	Draft Guidance for Industry: Preparation of Investigational Device Exemptions and Investigational New Drug Applications for Products Intended to Repair or Replace Knee Cartilage; Availability	7/9/2007	null date	7/9/2007		E7-13162
FDA	FDA-2007-0369	FDA-2007-0369-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Presubmission Conferences, New Animal Drug Applications and Supporting Regulations and Guidance 152, and Form FDA 356V	7/9/2007	null date	7/9/2007		E7-13195
FDA	FDA-2007-0370	FDA-2007-0370-0001	Otsuka Pharmaceutical Co., Ltd.; Withdrawal of Approval of a New Drug Application; Correction	9/1/2007	null date	7/9/2007		E7-13160
FDA	FDA-2007-0371	FDA-2007-0371-0001	Agency Information Collection Activities; Proposed Collection; Comment Request, Patent Term Restoration, Due Diligence Petitions, Filing, Format, and Content of Petitions	7/9/2007	null date	7/9/2007		E7-13269
FDA	FDA-2007-0372	FDA-2007-0372-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Procedures for the Safe and Sanitary Processing and Importing of Fish and Fishery Products	7/9/2007	null date	7/9/2007		E7-13153
FDA	FDA-2007-0373	FDA-2007-0373-0001	Draft Guidance for Industry: Evidence-Based Review System for the Scientific Evaluation of Health Claims; Availability	7/9/2007	9/7/2007	7/9/2007		E7-13274

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0374	FDA-2007-0374-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Medical Devices: Current Good Manufacturing Practice Quality System Regulations	7/9/2007	null date	7/9/2007		E7-13152
FDA	FDA-2007-0375	FDA-2007-0375-0001	New Animal Drugs For Use in Animal Feeds; Ivermectin		null date	7/10/2007		E7-13369
FDA	FDA-2007-0376	FDA-2007-0376-0001	Oral Dosage Form New Animal Drugs; Deracoxib		null date	7/10/2007		E7-13372
FDA	FDA-2007-0377	FDA-2007-0377-0001	Determination That ARISTOCORT FORTE Injectable Suspension (Triamcinolone Diacetate), 40 Milligrams per Milliliter, Was Not Withdrawn From Sale for Reasons of Safety or Effectiveness	7/11/2007	null date	7/11/2007		E7-13416
FDA	FDA-2007-0378	FDA-2007-0378-0001	Antiviral Drugs Advisory Committee; Notice of Meeting		null date	7/12/2007		E7-13560
FDA	FDA-2007-0379	FDA-2007-0379-0001	Global Harmonization Task Force, Study Groups 1 and 5; New Proposed and Final Documents; Availability	7/13/2007	null date	7/13/2007		07-13664
FDA	FDA-2007-0380	FDA-2007-0380-0001	International Conference on Harmonisation; Draft Guidance on Q10 Pharmaceutical Quality System; Availability	7/13/2007	null date	7/13/2007		E7-13667
FDA	FDA-2007-0381	FDA-2007-0381-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; FDA Survey of Current Manufacturing Practices in the Food Industry	7/19/2007	null date	7/19/2007		E7-13951
FDA	FDA-2007-0382	FDA-2007-0382-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Threshold of Regulation for Substances Used in Food-Contact Articles	7/19/2007	null date	7/19/2007		E7-14014
FDA	FDA-2007-0383	FDA-2007-0383-0001	Determination That Brethine (Terbutaline Sulfate) Injection Was Not Withdrawn From Sale for Reasons of Safety or Effectiveness	7/19/2007	null date	7/19/2007		E7-13950
FDA	FDA-2007-0384	FDA-2007-0384-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; FDA Survey of Physicians Perceptions of the Impact of Early Risk Communication About Medical Products	7/19/2007	null date	7/19/2007		E7-14015

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0385	FDA-2007-0385-0001	Draft Guidance for Industry and Food and Drug Administration Staff; Pulse Oximeters--Premarket Notification Submissions [510(k)s]; Availability	7/19/2007	null date	7/19/2007		E7-14012
FDA	FDA-2007-0386	FDA-2007-0386-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Experimental Study of Possible Footnotes and Cueing Schemes to Help Consumers Interpret Quantitative Trans Fat Disclosures on the Nutrition Facts Panel	7/19/2007	null date	7/19/2007		E7-14011
FDA	FDA-2007-0387	FDA-2007-0387-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Experimental Study of Trans Fat Claims on Foods	7/19/2007	null date	7/19/2007		E7-14010
FDA	FDA-2007-0388	FDA-2007-0388-0001	Food Additives Permitted in Feed and Drinking Water of Animals; Selenium Yeast	7/19/20007	8/20/2007	7/19/2007		E7-13954
FDA	FDA-2007-0389	FDA-2007-0389-0001	Irradiation in the Production, Processing and Handling of Food		null date	7/19/2007		E7-13947
FDA	FDA-2007-0390	FDA-2007-0390-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Voluntary Registration of Cosmetic Product Establishments	7/19/2007	null date	7/19/2007		E7-14013
FDA	FDA-2007-0391	FDA-2007-0391-0001	Draft Guidance for Industry and Food and Drug Administration Staff; Premarket Notification Submissions for Medical Devices That Include Antimicrobial Agents; Availability	7/19/2007	null date	7/19/2007		E7-13952
FDA	FDA-2007-0392	FDA-2007-0392-0001	Joint Meeting of the Cardiovascular and Renal Drugs Advisory Committee and the Drug Safety and Risk Management Advisory Committee; Notice of Meeting	7/20/2007	null date	7/20/2007		E7-14086
FDA	FDA-2007-0393	FDA-2007-0393-0001	Food Safety and Defense . . . Be ALERT; Public Workshop		null date	7/20/2007		E7-14045
FDA	FDA-2007-0394	FDA-2007-0394-0001	Food Labeling: Use of Symbols to Communicate Nutrition Information, Consideration of Consumer Studies and Nutritional Criteria; Public Hearing; Request for Comments	7/20/2007	null date	7/20/2007		E7-14046
FDA	FDA-2007-0395	FDA-2007-0395-0001	Blood Products Advisory Committee; Notice of Meeting		null date	7/20/2007		E7-14088

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0396	FDA-2007-0396-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Draft Guidance for Industry: Cooperative Manufacturing Arrangements for Licensed Biologics	7/23/2007	null date	7/23/2007		E7-14149
FDA	FDA-2007-0397	FDA-2007-0397-0001	Joint Meeting of the Cardiovascular and Renal Drugs Advisory Committee and the Drug Safety and Risk Management Advisory Committee; Notice of Meeting	7/23/2007	null date	7/23/2007		E7-14151
FDA	FDA-2007-0398	FDA-2007-0398-0001	Request for Notification From Industry Organizations Interested in Participating in Selection Process for Nonvoting Industry Representatives on Public Advisory Panels or Committees and Request for Nonvoting Industry Representatives on Public Advisory Panels or Committees	7/24/2007	null date	7/24/2007		E7-14206
FDA	FDA-2007-0399	FDA-2007-0399-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Color Additive Certification Requests and Recordkeeping	7/24/2007	null date	7/24/2007		E7-14201
FDA	FDA-2007-0400	FDA-2007-0400-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Mental Models Study of Food Terrorism Risk Awareness	7/24/2007	null date	7/24/2007		E7-14200
FDA	FDA-2007-0401	FDA-2007-0401-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Hazard Analysis and Critical Control Point Procedures for the Safe and Sanitary Processing and Importing of Juice	7/25/2007	null date	7/25/2007		E7-14403
FDA	FDA-2007-0402	FDA-2007-0402-0001	Guidance; Emergency Use Authorization of Medical Products; Availability		null date	7/26/2007		07-03661
FDA	FDA-2007-0403	FDA-2007-0403-0001	Draft Guidance for Industry and Food and Drug Administration Staff; In Vitro Diagnostic Multivariate Index Assays; Availability	7/26/2007	null date	7/26/2007		07-03660
FDA	FDA-2007-0404	FDA-2007-0404-0001	Draft Guidance for Industry: Cell Selection Devices for Point of Care Production of Minimally Manipulated Autologous Peripheral Blood Stem Cells; Availability	7/26/2007	null date	7/26/2007		07-03659

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0405	FDA-2007-0405-0001	Circulatory System Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting	7/27/2007	null date	7/27/2007		E7-14600
FDA	FDA-2007-0406	FDA-2007-0406-0001	Advisory Committee; Risk Communication Advisory Committee; Establishment		null date	7/27/2007		E7-14498
FDA	FDA-2007-0407	FDA-2007-0407-0001	Assuring Radiation Protection; Cooperative Agreement; Request for Applications: RFA-FDA-CDRH-07-004; Catalog of Federal Domestic Assistance Number: 93.103	7/30/2007	null date	7/30/2007		E7-14610
FDA	FDA-2007-0408	FDA-2007-0408-0001	Food Substances Affirmed as Generally Recognized as Safe in Feed and Drinking Water of Animals; Ethyl Alcohol Containing Ethyl Acetate	7/31/2007	null date	7/31/2007		E7-14700
FDA	FDA-2007-0409	FDA-2007-0409-0001	New Animal Drugs For Use in Animal Feeds; Ractopamine and Tylosin		null date	7/31/2007		E7-14699
FDA	FDA-2007-0410	FDA-2007-0410-0001	Ophthalmic and Topical Dosage Form New Animal Drugs; Emodepside and Praziquantel		null date	8/2/2007		E7-14945
FDA	FDA-2007-0411	FDA-2007-0411-0001	Animal Drug User Fee Rates and Payment Procedures for Fiscal Year 2008		null date	8/2/2007		07-03782
FDA	FDA-2007-0412	FDA-2007-0412-0001	Implantation or Injectable Dosage Form New Animal Drugs; Oxytetracycline Hydrochloride Injection		null date	8/2/2007		E7-14950
FDA	FDA-2007-0413	FDA-2007-0413-0001	Food Safety and Security Monitoring Project--Radiological Health; Availability of Cooperative Agreements Under a Limited Competition; Request for Applications: FD07-005; Catalog of Federal Domestic Assistance Number: 93.448	8/3/2007	null date	8/3/2007		E7-15061
FDA	FDA-2007-0414	FDA-2007-0414-0001	Medical Devices; General and Plastic Surgery Devices; Classification of Absorbable Poly(hydroxybutyrate) Surgical Suture Produced by Recombinant DNA Technology	8/3/2007	null date	8/3/2007		E7-15064
FDA	FDA-2007-0415	FDA-2007-0415-0001	Guidance for Industry and Food and Drug Administration Staff; ``Class II Special Controls Guidance Document: Absorbable Poly(hydroxybutyrate) Surgical Suture Produced by Recombinant DNA Technology; Availability	8/3/2007	null date	8/3/2007		E7-15063
FDA	FDA-2007-0416	FDA-2007-0416-0001	Determination That Daranide (Dichlorphenamide) Tablets, 50 Milligrams, Were Not Withdrawn From Sale for Reasons of Safety or Effectiveness	8/6/2007	null date	8/6/2007		E7-15230

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AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0417	FDA-2007-0417-0001	Determination That PHOSLO (Calcium Acetate) 667-Milligram Tablet Was Not Withdrawn From Sale for Reasons of Safety or Effectiveness	8/6/2007	null date	8/6/2007		E7-15172
FDA	FDA-2007-0418	FDA-2007-0418-0001	Determination That PHENERGAN (Promethazine Hydrochloride) Suppositories, 12.5 Milligrams and 25 Milligrams, Were Not Withdrawn From Sale for Reasons of Safety or Effectiveness	8/6/2007	null date	8/6/2007		E7-15174
FDA	FDA-2007-0419	FDA-2007-0419-0001	Determination That PREVACID NAPRAPAC (Copackaged Lansoprazole Delayed-Release 15-Milligram Capsules and Naproxen 250-Milligram Tablets) Was Not Withdrawn From Sale for Reasons of Safety or Effectiveness	8/7/2007	null date	8/7/2007		E7-15233
FDA	FDA-2007-0420	FDA-2007-0420-0001	Determination That DEXEDRINE (Dextroamphetamine Sulfate) Oral Solution, 5 Milligrams per 5 Milliliters, Was Not Withdrawn From Sale for Reasons of Safety or Effectiveness	8/7/2007	null date	8/7/2007	(Dextroam	E7-15236
FDA	FDA-2007-0421	FDA-2007-0421-0001	Summaries of Medical and Clinical Pharmacology Reviews of Pediatric Studies; Availability	8/7/2007	null date	8/7/2007		E7-15234
FDA	FDA-2007-0422	FDA-2007-0422-0001	Determination That Methotrexate Injection, USP, Preservative Free, Equivalent to 500 Milligrams Base/20 Milliliters (25 Milligrams/ Milliliter), Was Not Withdrawn From Sale for Reasons of Safety or Effectiveness	8/8/2007	null date	8/8/2007		E7-15490
FDA	FDA-2007-0423	FDA-2007-0423-0001	Cooperative Agreement To Support the National Alliance for Hispanic Health; Notice of Intent To Accept and Consider a Single Source Application; Availability of Funds for Fiscal Year 2007	8/8/2007	null date	8/8/2007		E7-15491
FDA	FDA-2007-0424	FDA-2007-0424-0001	Guidance for Industry: Class II Special Controls Guidance Document: In Vitro Human Immunodeficiency Virus Drug Resistance Genotype Assay; Availability	8/8/2007	null date	8/8/2007		E7-15477
FDA	FDA-2007-0425	FDA-2007-0425-0001	Guidance for Industry: Adequate and Appropriate Donor Screening Tests for Hepatitis B; Hepatitis B Surface Antigen Assays Used to Test Donors of Whole Blood and Blood Components, Including Source Plasma and Source Leukocytes; Availability	8/8/2007	null date	8/8/2007		E7-15472

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Food and Drug Administration (FDA) -- 1547 Documents

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Food and Drug Administration (FDA)								
FDA	FDA-2007-0426	FDA-2007-0426-0001	Medical Devices: Immunology and Microbiology Devices: Classification of In Vitro Human Immunodeficiency Virus Drug Resistance Genotype Assay	8/8/2007	null date	8/8/2007		E7-15475
FDA	FDA-2007-0427	FDA-2007-0427-0001	Determination That MIVACRON (Mivacurium Chloride) Injection Equivalent to 2 Milligrams Base/Milliliter Was Not Withdrawn From Sale for Reasons of Safety or Effectiveness	8/8/2007	null date	8/8/2007		E7-15488
FDA	FDA-2007-0428	FDA-2007-0428-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Requirements for Submission of Labeling for Human Prescription Drugs and Biologics in Electronic Format	8/10/2007	null date	8/10/2007		E7-15614
FDA	FDA-2007-0429	FDA-2007-0429-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Environmental Impact Considerations	8/10/2007	null date	8/10/2007		E7-15612
FDA	FDA-2007-0430	FDA-2007-0430-0001	Implantation or Injectable Dosage Form New Animal Drugs; Ampicillin Sodium		null date	8/13/2007		E7-15761
FDA	FDA-2007-0431	FDA-2007-0431-0001	Preparation for International Cooperation on Cosmetics Regulations Meeting in Brussels, Belgium; Notice of Public Meeting	8/13/2007	null date	8/13/2007		07-03954
FDA	FDA-2007-0432	FDA-2007-0432-0001	Certain Other Dosage Form New Animal Drugs; Formalin		null date	8/13/2007		E7-15763
FDA	FDA-2007-0433	FDA-2007-0433-0001	Preparation for International Conference on Harmonization Meetings in Yokohama, Japan; Public Meeting	8/13/2007	null date	8/13/2007		07-15803
FDA	FDA-2007-0434	FDA-2007-0434-0001	Determination That ORUDIS KT (Ketoprofen) Tablets, 12.5 Milligrams, Were Not Withdrawn From Sale for Reasons of Safety or Effectiveness	8/14/2007	null date	8/14/2007		E7-15843
FDA	FDA-2007-0435	FDA-2007-0435-0001	Listing of Color Additives Subject to Certification; D&C Black No. 3; Confirmation of Effective Date		null date	8/14/2007		07-15831
FDA	FDA-2007-0436	FDA-2007-0436-0001	Cardiovascular and Renal Drugs Advisory Committee; Notice of Meeting		null date	8/14/2007		E7-15834
FDA	FDA-2007-0437	FDA-2007-0437-0001	Guidance for Industry on Exports Under the Food and Drug Administration Export Reform and Enhancement Act of 1996; Availability	8/14/2007	null date	8/14/2007		E7-15840

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0438	FDA-2007-0438-0001	Clinical Development Programs for Human Drugs, Biological Products, and Medical Devices for the Treatment and Prevention of Osteoarthritis; Request for Assistance	8/14/2007	null date	8/14/2007		E7-15844
FDA	FDA-2007-0439	FDA-2007-0439-0001	Workshop to Discuss Development of a Womens Health Information Sharing Network		null date	8/15/2007		E7-15944
FDA	FDA-2007-0440	FDA-2007-0440-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Control of Communicable Diseases; Restrictions on African Rodents, Prairie Dogs, and Certain Other Animals	8/15/2007	null date	8/15/2007		E7-15939
FDA	FDA-2007-0441	FDA-2007-0441-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Current Good Manufacturing Practice Regulations for Type A Medicated Articles	8/16/2007	null date	8/16/2007		E7-16087
FDA	FDA-2007-0442	FDA-2007-0442-0003	Revisions to the Requirements Applicable to Blood, Blood Components and Source Plasma; Correction	9/24/2007	null date	9/24/2007		E7-18799
FDA	FDA-2007-0442	FDA-2007-0442-0002	Revisions to the Requirements Applicable to Blood, Blood Components, and Source Plasma; Companion Document to Direct Final Rule	8/16/2007	10/30/2007	8/16/2007		E7-15942
FDA	FDA-2007-0442	FDA-2007-0442-0001	Revisions to the Requirements Applicable to Blood, Blood Components and Source Plasma	8/16/2007	10/30/2007	8/16/2007		E7-15943
FDA	FDA-2007-0442	FDA-2007-0442-0004	Revisions to the Requirements Applicable to Blood, Blood Components, and Source Plasma; Companion Document to Direct Final Rule; Correction	9/24/2007	10/30/2007	9/24/2007		E7-18802
FDA	FDA-2007-0443	FDA-2007-0443-0001	Joint Meeting of the Nonprescription Drugs Advisory Committee and the Pediatric Advisory Committee; Notice of Meeting	8/16/2007	null date	8/16/2007		E7-16169
FDA	FDA-2007-0444	FDA-2007-0444-0001	Processing Methods for Orthopedic, Cardiovascular, and Skin Allografts; Public Workshop		null date	8/16/2007		E7-16182
FDA	FDA-2007-0445	FDA-2007-0445-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Current Good Manufacturing Practice Regulations for Medicated Feeds	8/16/2007	null date	8/16/2007		E7-16088
FDA	FDA-2007-0446	FDA-2007-0446-0001	Food Labeling: Safe Handling Statements: Labeling of Shell Eggs		null date	8/20/2007	0910-ZA23	E7-16272

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0447	FDA-2007-0447-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Guidance for Industry on Pharmacogenomic Data Submissions; Extension	8/21/2007	null date	8/21/2007		E7-16470
FDA	FDA-2007-0448	FDA-2007-0448-0001	Medical Devices 101: An Educational Forum; Public Workshop		null date	8/21/2007		E7-16375
FDA	FDA-2007-0449	FDA-2007-0449-0001	Food Additives Permitted for Direct Addition to Food for Human Consumption; Glycerol Ester of Tall Oil Rosin	8/22/2007	9/21/2007	8/22/2007		E7-16558
FDA	FDA-2007-0450	FDA-2007-0450-0001	Agency information collection activities; proposals, submissions, and approvals	8/22/2007	null date	8/22/2007		E7-16603
FDA	FDA-2007-0451	FDA-2007-0451-0001	Agency Information Collection Activities: Proposed Collection; Comment Request; Registration of Producers of Drugs and Listing of Drugs in Commercial Distribution	8/24/2007	null date	8/24/2007		E7-16795
FDA	FDA-2007-0452	FDA-2007-0452-0001	``Guidance for Industry: `Lookback for Hepatitis C Virus (HCV): Product Quarantine, Consignee Notification, Further Testing, Product Disposition, and Notification of Transfusion Recipients Based on Donor Test Results Indicating Infection with HCV; Availability	8/24/2007	null date	8/24/2007		E7-16605
FDA	FDA-2007-0453	FDA-2007-0453-0001	Medical Devices Regulated by the Center for Biologics Evaluation and Research; Availability of Summaries of Safety and Effectiveness Data for Premarket Approval Applications	8/24/2007	null date	8/24/2007		E7-16706
FDA	FDA-2007-0454	FDA-2007-0454-0001	Current Good Manufacturing Practice for Blood and Blood Components; Notification of Consignees and Transfusion Recipients Receiving Blood and Blood Components at Increased Risk of Transmitting Hepatitis C Virus Infection (``Lookback)	8/24/2007	null date	8/24/2007	0910-AB76	E7-16607
FDA	FDA-2007-0455	FDA-2007-0455-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Manufactured Food Regulatory Program Standards	8/24/2007	null date	8/24/2007		E7-16708
FDA	FDA-2007-0456	FDA-2007-0456-0001	Sunscreen Drug Products for Over-the-Counter Human Use; Proposed Amendment of Final Monograph	8/27/2007	11/26/2007	8/27/2007	0910-AF43	07-04131

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0456	FDA-2007-0456-0002	Sunscreen Drug Products for Over-The-Counter Human Use; Proposed Amendment of Final Monograph; Extension of Comment Period	11/28/2007	12/26/2007	11/28/2007	0910-AF43	07-05853
FDA	FDA-2007-0457	FDA-2007-0457-0001	Preparation for International Conference on Harmonization Meetings in Yokohama, Japan; Public Meeting; Correction	8/27/2007	null date	8/27/2007		E7-16892
FDA	FDA-2007-0458	FDA-2007-0458-0001	Withdrawal of Approval of a New Animal Drug Application; Bacitracin Zinc		null date	8/29/2007		E7-16985
FDA	FDA-2007-0459	FDA-2007-0459-0001	New Animal Drugs For Use in Animal Feeds; Withdrawal of Approval of a New Animal Drug Application; Bacitracin Zinc	8/28/2007	null date	8/28/2007		E7-16984
FDA	FDA-2007-0460	FDA-2007-0460-0001	Companion to Guidance for Industry on Pharmacogenomic Data; Availability		null date	8/29/2007		E7-17103
FDA	FDA-2007-0461	FDA-2007-0461-0001	Draft Guidance for Industry: Evidence-Based Review System for the Scientific Evaluation of Health Claims; Availability; Correction	8/29/2007	null date	8/29/2007		E7-17038
FDA	FDA-2007-0462	FDA-2007-0462-0001	Organization, functions, and authority delegations: Office of the Commissioner		null date	8/30/2007		07-04259
FDA	FDA-2007-0463	FDA-2007-0463-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Administrative Procedures for the Clinical Laboratory Improvement Amendments of 1998 Categorization; Correction	8/30/2007	null date	8/30/2007		E7-17153
FDA	FDA-2007-0464	FDA-2007-0464-0001	Presidential Interagency Working Group on Import Safety; Public Meeting		null date	8/31/2007		E7-17305
FDA	FDA-2007-0465	FDA-2007-0465-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Medical Devices: Recommended Glossary and Educational Outreach to Support Use of Symbols on Labels and in Labeling of In Vitro Diagnostic Devices Intended for Professional Use	8/31/2007	null date	8/31/2007		E7-17217
FDA	FDA-2007-0466	FDA-2007-0466-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Adverse Event Pilot Program for Medical Products (Formally Medical Device Adverse Event Reporting Program)	9/6/2007	null date	9/6/2007		E7-17562

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0467	FDA-2007-0467-0001	Determination That MILTOWN (Meprobamate) Tablets and Five Other Drug Products Were Not Withdrawn From Sale for Reasons of Safety or Effectiveness	9/6/2007	null date	9/6/2007		E7-17566
FDA	FDA-2007-0468	FDA-2007-0468-0001	Implantation or Injectable Dosage Form New Animal Drugs; Dexmedetomidine		null date	9/7/2007		E7-17696
FDA	FDA-2007-0469	FDA-2007-0469-0001	Implantation or Injectable Dosage Form New Animal Drugs; Etodolac		null date	9/7/2007		E7-17645
FDA	FDA-2007-0470	FDA-2007-0470-0001	``Guidance for Industry: Manufacturing Biological Intermediates and Biological Drug Substances Using Spore-Forming Microorganisms; Availability	9/7/2007	null date	9/7/2007		E7-17709
FDA	FDA-2007-0471	FDA-2007-0471-0001	Memorandum of Understanding Between the Food and Drug Administration and the University System of Maryland	9/7/2007	null date	9/7/2007		07-04404
FDA	FDA-2007-0472	FDA-2007-0472-0001	National Mammography Quality Assurance Advisory Committee; Notice of Meeting	9/11/2007	null date	9/11/2007		E7-17795
FDA	FDA-2007-0473	FDA-2007-0473-0001	Food and Drug Administration Modernization Act of 1997: Modifications to the List of Recognized Standards, Recognition List Number: 018	9/12/2007	null date	9/12/2007		E7-18021
FDA	FDA-2007-0474	FDA-2007-0474-0001	Circulatory System Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting	9/12/2007	null date	9/12/2007		E7-17983
FDA	FDA-2007-0475	FDA-2007-0475-0001	Cardiovascular and Renal Drugs Advisory Committee; Amendment of Notice	9/12/2007	null date	9/12/2007		E7-18031
FDA	FDA-2007-0476	FDA-2007-0476-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Information From United States Processors That Export to the European Community	9/12/2007	null date	9/12/2007		E7-18033
FDA	FDA-2007-0477	FDA-2007-0477-0001	Medical Devices; Availability of Safety and Effectiveness Summaries for Premarket Approval Applications	9/13/2007	null date	9/13/2007		E7-18034
FDA	FDA-2007-0478	FDA-2007-0478-0001	Guidance for Industry and Food and Drug Administration Staff; Commercially Distributed Analyte Specific Reagents: Frequently Asked Questions; Availability	9/14/2007	null date	9/14/2007		E7-18108

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0479	FDA-2007-0479-0001	Draft Guidance for Industry, Clinical Laboratories, and Food and Drug Administration Staff on In Vitro Diagnostic Multivariate Index Assays; Reopening of the Comment Period	9/17/2007	null date	9/17/2007		E7-18221
FDA	FDA-2007-0480	FDA-2007-0480-0001	Food Labeling; Health Claims; Dietary Noncariogenic Carbohydrate Sweeteners and Dental Caries	9/17/2007	12/3/2007	9/17/2007		E7-18196
FDA	FDA-2007-0481	FDA-2007-0481-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Premarket Approval of Medical Devices	9/17/2007	null date	9/17/2007		E7-18222
FDA	FDA-2007-0482	FDA-2007-0482-0001	Use of Ozone-Depleting Substances; Removal of Essential-Use Designation (Epinephrine)	9/20/2007	11/19/2007	9/20/2007	0910-AF92	07-04663
FDA	FDA-2007-0482	FDA-2007-0482-0002	Use of Ozone-Depleting Substances; Removal of Essential-Use Designation (Epinephrine); Public Meeting; Extension of Comment Period	11/8/2007	12/19/2007	11/8/2007	0910-AF92	07-05593
FDA	FDA-2007-0483	FDA-2007-0483-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Medical Devices: Current Good Manufacturing Practice Quality System Regulations	9/20/2007	null date	9/20/2007		E7-18582
FDA	FDA-2007-0484	FDA-2007-0484-0001	Educational Workshops on Current Good Manufacturing Practices; Public Workshops	9/20/2007	null date	9/20/2007		E7-18556
FDA	FDA-2007-0485	FDA-2007-0485-0001	Agency Information Collection Activities: Proposed Collection; Comment Request; Radioactive Drug Research Committees	9/21/2007	null date	9/21/2007		E7-18646
FDA	FDA-2007-0486	FDA-2007-0486-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Pharmaceutical Development Study	9/21/2007	null date	9/21/2007		E7-18641
FDA	FDA-2007-0487	FDA-2007-0487-0001	Information Technology Strategic Planning; Public Meeting	9/21/2007	null date	9/21/2007		07-04692
FDA	FDA-2007-0488	FDA-2007-0488-0001	Draft Guidance for Industry: Microbiological Considerations for Antimicrobial Food Additive Submissions; Availability	9/25/2007	null date	9/24/2007		E7-18816
FDA	FDA-2007-0489	FDA-2007-0489-0001	Implantation or Injectable Dosage Form New Animal Drugs; Tulathromycin	9/26/2007	null date	9/26/2007		E7-18983
FDA	FDA-2007-0490	FDA-2007-0490-0001	Request for Nominations for Voting Members on Public Advisory Committee, Veterinary Medicine Advisory Committee	9/27/2007	null date	9/27/2007		E7-19130

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Food and Drug Administration (FDA)								
FDA	FDA-2007-0491	FDA-2007-0491-0001	Guidance for Industry: Toxicity Grading Scale for Healthy Adult and Adolescent Volunteers Enrolled in Preventive Vaccine Clinical Trials; Availability	9/27/2007	null date	9/27/2007		E7-19155
FDA	FDA-2007-0492	FDA-2007-0492-0001	Joint Meeting of the Nonprescription Drugs Advisory Committee and the Pediatric Advisory Committee; Amendment of Notice	10/1/2007	null date	10/1/2007		E7-19332
FDA	FDA-2007-0493	FDA-2007-0493-0001	Drug Products Containing Hydrocodone; Enforcement Action Dates	10/1/2007	null date	10/1/2007		E7-19340
FDA	FDA-2007-0494	FDA-2007-0494-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Submission of Petitions: Food Additive, Color Additive (Including Labeling), and Generally Recognized as Safe Affirmation; Electronic Submission Using Food and Drug Administration Forms 3503 and 3504	10/2/2007	null date	10/2/2007		E7-19350
FDA	FDA-2007-0495	FDA-2007-0495-0001	Science Board to the Food and Drug Administration; Notice of Meeting	10/2/2007	null date	10/2/2007		E7-19349
FDA	FDA-2007-0496	FDA-2007-0496-0001	Guidance for Industry: Recommended Study Design and Evaluation of Effectiveness Studies for Swine Respiratory Disease Claims; Availability	10/2/2007	null date	10/2/2007		E7-19412
FDA	FDA-2007-0497	FDA-2007-0497-0001	Agency Emergency ProcessingUnder OMB Review; Medical Device User Fee Amendments of 2007; Foreign Small Business Qualification Certification Form FDA 3602A	10/2/2007	null date	10/2/2007		E7-19411
FDA	FDA-2007-0498	FDA-2007-0498-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Human Tissue Intended for Transplantation	10/3/2007	null date	10/3/2007		E7-19457
FDA	FDA-2007-0499	FDA-2007-0499-0001	Nominations for Membership on the Board of Directors of the Reagan-Udall Foundation From Consumer Advocacy Groups, Professional Scientific and Medical Societies, and Industry Trade Organizations	10/3/2007	null date	10/3/2007		07-04882
FDA	FDA-2007-0500	FDA-2007-0500-0001	Electronic Nonclinical Study Data Submission; Notice of Pilot Project	10/3/2007	null date	10/3/2007		E7-19468

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Food and Drug Administration (FDA)								
FDA	FDA-2007-0501	FDA-2007-0501-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Human Cells, Tissues, and Cellular and Tissue-Based Products: Establishment Registration and Listing; Form Food and Drug Administration 3356; Eligibility Determination for Donors; and Current Good Tissue Practice	10/3/2007	null date	10/3/2007		E7-19454
FDA	FDA-2007-0502	FDA-2007-0502-0001	Guidance for Industry and Food and Drug Administration Staff; Biological Indicator Premarket Notification Submissions; Availability	10/4/2007	null date	10/4/2007		E7-19573
FDA	FDA-2007-0503	FDA-2007-0503-0001	Dean Foods Co.; Filing of Food Additive Petition	10/4/2007	null date	10/4/2007		07-19576
FDA	FDA-2007-0504	FDA-2007-0504-0001	Draft Guidance for Industry and Food and Drug Administration Staff; Class II Special Controls Guidance Document: Electrocardiograph Electrodes; Availability	10/4/2007	null date	10/4/2007		E7-19578
FDA	FDA-2007-0505	FDA-2007-0505-0001	Medical Devices; Cardiovascular Devices; Electrocardiograph Electrode; Designation of Special Controls	10/4/2007	1/2/2008	10/4/2007		E7-19580
FDA	FDA-2007-0506	FDA-2007-0506-0001	Behind the Counter Availability of Certain Drugs; Public Meeting	10/4/2007	null date	10/4/2007		E7-19329
FDA	FDA-2007-0507	FDA-2007-0507-0001	Establishing a Docket for the Development of Safety and Effectiveness Assessments of Vaccines Used for Pandemic Influenza; Availability	10/4/2007	null date	10/4/2007		E7-19577
FDA	FDA-2007-0508	FDA-2007-0508-0001	New Animal Drugs; Ractopamine	10/5/2007	null date	10/5/2007		E7-19732
FDA	FDA-2007-0509	FDA-2007-0509-0001	Implantation or Injectable Dosage Form New Animal Drugs; Polysulfated Glycosaminoglycan	10/5/2007	null date	10/5/2007		E7-19729
FDA	FDA-2007-0510	FDA-2007-0510-0001	New Animal Drugs; Florfenicol	10/9/2007	null date	10/9/2007		E7-19853
FDA	FDA-2007-0511	FDA-2007-0511-0001	Menley and James Laboratories, Inc. et al.; Withdrawal of Approval of Six New Drug Applications	10/10/2007	null date	10/10/2007		E7-19865
FDA	FDA-2007-0512	FDA-2007-0512-0001	Medical Device User Fee and Modernization Act; Notice to Public of Web Location of 2008 Proposed Guidance Development; Establishment of a Public Docket	10/10/2007	null date	10/10/2007		E7-19864

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Food and Drug Administration (FDA)								
FDA	FDA-2007-0513	FDA-2007-0513-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Patent Term Restoration, Due Diligence Petitions, Filing, Format, and Content of Petitions	10/11/2007	null date	10/11/2007		E7-20070
FDA	FDA-2007-0514	FDA-2007-0514-0001	Quality System Regulation Educational Forum on Design Controls; Public Workshop	10/11/2007	null date	10/11/2007		E7-20077
FDA	FDA-2007-0515	FDA-2007-0515-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Generic Food and Drug Administration Rapid Response Surveys	10/11/2007	null date	10/11/2007		E7-20067
FDA	FDA-2007-0516	FDA-2007-0516-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Toll-Free Number for Consumer Reporting of Drug Product Side Effects: Comprehension	10/11/2007	null date	10/11/2007		E7-20075
FDA	FDA-2007-0517	FDA-2007-0517-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Institutional Review Boards	10/11/2007	null date	10/11/2007		E7-20063
FDA	FDA-2007-0518	FDA-2007-0518-0001	Draft Guidance for Industry and Food and Drug Administration Staff; Class II Special Controls Guidance Document: Electrocardiograph Electrodes; Availability; Correction	10/12/2007	null date	10/12/2007		E7-20183
FDA	FDA-2007-0519	FDA-2007-0519-0001	Medical Device User Fee Rates for Fiscal Year 2008	10/12/2007	null date	10/12/2007		07-05051
FDA	FDA-2007-0520	FDA-2007-0520-0001	Prescription Drug User Fee Rates for Fiscal Year 2008	10/12/2007	null date	10/12/2007		07-05052
FDA	FDA-2007-0521	FDA-2007-0521-0001	In Vitro Analysis of Cell/Scaffold Medical Products; Public Workshop	10/12/2007	null date	10/12/2007		E7-20191
FDA	FDA-2007-0522	FDA-2007-0522-0001	Pediatric Advisory Committee; Notice of Meeting	10/15/2007	null date	10/15/2007		E7-20302
FDA	FDA-2007-0523	FDA-2007-0523-0001	Draft Guidance for Industry: Questions and Answers Regarding Adverse Event Reporting and Recordkeeping for Dietary Supplements as Required by the Dietary Supplement and Nonprescription Drug Consumer Protection Act; Availability	10/15/2007	null date	10/15/2007		07-05074

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0524	FDA-2007-0524-0001	Draft Guidance for Industry on Postmarketing Adverse Event Reporting for Nonprescription Human Drug Products Marketed Without an Approved Application; Availability	10/15/2007	null date	10/15/2007		07-05073
FDA	FDA-2007-0525	FDA-2007-0525-0001	Draft Guidance for Industry on Antibacterial Drug Products: Use ofNoninferiority Studies to Support Approval; Availability	10/15/2007	null date	10/15/2007		E7-20282
FDA	FDA-2007-0526	FDA-2007-0526-0001	Pediatric Advisory Committee; Notice of Meeting	10/15/2007	null date	10/15/2007		E7-20304
FDA	FDA-2007-0527	FDA-2007-0527-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Focus Groups as Used by the Food and Drug Administration	10/15/2007	null date	10/15/2007		E7-20291
FDA	FDA-2007-0528	FDA-2007-0528-0001	Training Program for Regulatory Project Managers; Information Available to Industry	10/16/2007	null date	10/16/2007		E7-20430
FDA	FDA-2007-0529	FDA-2007-0529-0001	Guidance for Industry on FDA Recommendations for Sampling and Testing Yellow Corn and Dry-Milled Yellow Corn Shipments Intended for Human Food Use for Cry9C Protein Residues; Comments on Possible Withdrawal	10/17/2007	null date	10/17/2007		E7-20379
FDA	FDA-2007-0530	FDA-2007-0530-0001	Revision of the Requirements for Live Vaccine Processing	1/2/2008	null date	10/18/2007		E7-20610
FDA	FDA-2007-0530	FDA-2007-0530-0002	Revision of the Requirements for Live Vaccine Processing; Companion to Direct Final Rule	10/18/2007	1/2/2008	10/18/2007		E7-20609
FDA	FDA-2007-0531	FDA-2007-0531-0001	Science Board to the Food and Drug Administration; Amendment of Notice	10/18/2007	null date	10/18/2007		E7-20550
FDA	FDA-2007-0532	FDA-2007-0532-0001	Peripheral and Central Nervous System Drugs Advisory Committee; Notice of Meeting	10/18/2007	null date	10/18/2007		E7-20511
FDA	FDA-2007-0533	FDA-2007-0533-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Guidance for Industry on Special Protocol Assessment	10/18/2007	null date	10/18/2007		E7-20549
FDA	FDA-2007-0534	FDA-2007-0534-0001	Applications for Food and Drug Administration Application Approval to Market a New Drug; Revision of Postmarketing Reporting Requirements	10/18/2007	null date	10/18/2007		E7-20510
FDA	FDA-2007-0535	FDA-2007-0535-0001	Cardiovascular and Renal Drugs Advisory Committee; Notice of Meeting	10/18/2007	null date	10/18/2007		E7-20512

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0536	FDA-2007-0536-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Information Program on Clinical Trials for Serious or Life-Threatening Diseases: Maintaining a Data Bank	10/19/2007	null date	10/19/2007		E7-20662
FDA	FDA-2007-0537	FDA-2007-0537-0001	Medical Devices; General Hospital and Personal Use Devices; Classification of Remote Medication Management System	10/19/2007	null date	10/19/2007		E7-20633
FDA	FDA-2007-0538	FDA-2007-0538-0001	Guidance for Industry and Food and Drug Administration Staff; Class II Special Controls Guidance Document: Remote Medication Management System; Availability	10/19/2007	null date	10/19/2007		E7-20635
FDA	FDA-2007-0539	FDA-2007-0539-0001	Draft Guidance for Industry on the Use of Mechanical Calibration of Dissolution Apparatus 1 and 2 - Current Good Manufacturing Practice; Availability	10/19/2007	null date	10/19/2007		E7-20664
FDA	FDA-2007-0540	FDA-2007-0540-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Experimental Evaluation of Variations in Content and Format of the Brief Summary in Direct-to- Consumer Print Advertisements for Prescription Drugs	10/22/2007	null date	10/22/2007		E7-20756
FDA	FDA-2007-0541	FDA-2007-0541-0001	Electronic Distribution of Prescribing Information for Prescription Drug Products; Reopening of Comment Period	10/22/2007	null date	10/22/2007		E7-20759
FDA	FDA-2007-0542	FDA-2007-0542-0001	Vaccines and Related Biological Products Advisory Committee; Notice of Meeting	10/23/2007	null date	10/23/2007		E7-20854
FDA	FDA-2007-0543	FDA-2007-0543-0001	Joint Meeting of the Nonprescription Drugs Advisory Committee and the Endocrinologic and Metabolic Drugs Advisory Committee; Notice of Meeting	10/23/2007	null date	10/23/2007		E7-20855
FDA	FDA-2007-0544	FDA-2007-0544-0001	Guidance for Industry, Food and Drug Administration, and Foreign Governments; Fiscal Year 2008 Medical Device User Fee Small Business Qualification and Certification; Availability	10/23/2007	null date	10/23/2007		07-05226
FDA	FDA-2007-0546	FDA-2007-0546-0001	Nonprescription Drugs Advisory Committee; Notice of Meeting	10/24/2007	null date	10/24/2007		07-05249

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0547	FDA-2007-0547-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Guidance for Industry on Continuous Marketing Applications: Pilot--Scientific Feedback and Interactions During Development of Fast Track Products Under the Prescription Drug User Fee Act	10/25/2007	null date	10/25/2007		E7-21056
FDA	FDA-2007-0548	FDA-2007-0548-0001	Immune Correlates of Protection Against Influenza A Viruses in Support of Pandemic Vaccine Development; Public Workshop	10/25/2007	null date	10/25/2007		E7-20981
FDA	FDA-2007-0549	FDA-2007-0549-0001	User Fee Program for Advisory Review of Direct-to-Consumer Television Advertisements for Prescription Drug and Biological Products; Request for Notification of Participation and Number of Advertisements for Review	10/25/2007	null date	10/25/2007		07-05282
FDA	FDA-2007-0550	FDA-2007-0550-0001	Publication of Guidances for Industry Describing Product-Specific Bioequivalence Recommendations	10/25/2007	null date	10/25/2007		E7-21062
FDA	FDA-2007-0551	FDA-2007-0551-0001	Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; CGMP for Blood and Blood Components; Notification of Consignees and Transfusion Recipients Receiving Blood and Blood Components at Increased Risk of Transmitting HCV Infection (``Lookback)	10/25/2007	null date	10/25/2007		E7-21055
FDA	FDA-2007-0552	FDA-2007-0552-0001	New Animal Drugs For Use in Animal Feeds; Change of Sponsor	10/25/2007	null date	10/25/2007		E7-21059
FDA	FDA-2007-0553	FDA-2007-0553-0001	Oral Dosage Form New Animal Drugs; Spinosad	10/25/2007	null date	10/25/2007		E7-21058
FDA	FDA-2007-0554	FDA-2007-0554-0001	Draft Guidance for Industry on Drug-Induced Liver Injury: Premarketing Clinical Evaluation; Availability	10/25/2007	null date	10/25/2007		E7-21060
FDA	FDA-2007-0555	FDA-2007-0555-0001	Draft Guidance for Industry and Food and Drug Administration Staff; In Vitro Diagnostic Device Studies--Frequently Asked Questions; Availability	10/25/2007	null date	10/25/2007		E7-20982
FDA	FDA-2007-0556	FDA-2007-0556-0001	New Animal Drugs; Change of Sponsor	10/25/2007	null date	10/25/2007		E7-21057
FDA	FDA-2007-0557	FDA-2007-0557-0001	Oral Dosage Form New Animal Drugs; Phenylbutazone Paste	10/25/2007	null date	10/25/2007		E7-21054
FDA	FDA-2007-0558	FDA-2007-0558-0001	Draft Guidance for Industry and Food and Drug Administration Staff; Impact-Resistant Lenses: Questions and Answers; Availability	10/26/2007	null date	10/26/2007		E7-21122

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0559	FDA-2007-0559-0001	Exocrine Pancreatic Insufficiency Drug Products; Extension to Obtain Marketing Approval	10/26/2007	null date	10/26/2007		E7-21082
FDA	FDA-2007-0560	FDA-2007-0560-0001	Guidance for Industry: Considerations for Plasmid Deoxyribonucleic Acid Vaccines for Infectious Disease Indications; Availability	10/29/2007	null date	10/29/2007		E7-21266
FDA	FDA-2007-0561	FDA-2007-0561-0001	Draft Guidance for Industry: Blood Establishment Computer System Validation in the Users Facility; Availability	10/29/2007	null date	10/29/2007		E7-21268
FDA	FDA-2007-0562	FDA-2007-0562-0001	Biomin GmbH; Filing of Food Additive Petition	10/30/2007	null date	10/30/2007		E7-21298
FDA	FDA-2007-0563	FDA-2007-0563-0001	Draft Guidance for Industry on Acute Bacterial Sinusitis: Developing Drugs for Treatment; Availability	10/30/2007	null date	10/30/2007		E7-21332
FDA	FDA-2007-0564	FDA-2007-0564-0001	Guidance for Industry on the Role of Human Immunodeficiency Virus Resistance Testing in Antiretroviral Drug Development; Availability	10/31/2007	null date	10/31/2007		E7-21403
FDA	FDA-2007-0565	FDA-2007-0565-0001	Draft Guidance for the Public, Food and Drug Administration Advisory Committee Members, and Food and Drug Administration Staff: Public Availability of Advisory Committee Members Financial Interest Information and Waivers; Availability	10/31/2007	null date	10/31/2007		07-05408
FDA	FDA-2007-0567	FDA-2007-0567-0001	Adolescent Over-the-Counter Drug Product Use; Public Workshop	11/5/2007	null date	11/5/2007		E7-21713
FDA	FDA-2007-0568	FDA-2007-0568-0001	Oncologic Drugs Advisory Committee; Notice of Meeting	11/5/2007	null date	11/5/2007		E7-21630
FDA	FDA-2007-0569	FDA-2007-0569-0001	Circulatory System Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting	11/6/2007	null date	11/6/2007		E7-21779
FDA	FDA-2007-0570	FDA-2007-0570-0001	New Animal Drugs; Ractopamine	11/6/2007	null date	11/6/2007		E7-21816
FDA	FDA-2007-0571	FDA-2007-0571-0001	Implantation or Injectable Dosage Form New Animal Drugs; Ivermectin	11/7/2007	null date	11/7/2007		E7-21839
FDA	FDA-2007-0572	FDA-2007-0572-0001	Lederle Laboratories et al.; Withdrawal of Approval of 73 New Drug Applications and 62 Abbreviated New Drug Applications	11/7/2007	null date	11/7/2007		E7-21886
FDA	FDA-2007-0573	FDA-2007-0573-0001	Requirements for Human Blood and Blood Components Intended for Transfusion or for Further Manufacturing Use	11/8/2007	null date	11/8/2007		E7-21565
FDA	FDA-2007-0574	FDA-2007-0574-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Application for Participation in the Medical Device Fellowship Program	11/9/2007	null date	11/9/2007		E7-21971

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0575	FDA-2007-0575-0001	Medical Devices Regulated by the Center for Biologics Evaluation and Research; Availability of Summaries of Safety and Effectiveness Data for Premarket Approval Applications	11/9/2007	null date	11/9/2007		E7-21986
FDA	FDA-2007-0576	FDA-2007-0576-0001	Agency Emergency Processing Under Office of Management and Budget Review; Orphan Drug Products; Common European Medicines Evaluation Agency/Food and Drug Administration Application Form for Orphan Medicinal Product Designation	11/9/2007	null date	11/9/2007		E7-21988
FDA	FDA-2007-0577	FDA-2007-0577-0001	Obstetrics and Gynecology Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting	11/9/2007	null date	11/9/2007		07-21979
FDA	FDA-2007-0578	FDA-2007-0578-0001	Draft Guidance for Industry on Chronic Obstructive Pulmonary Disease: Developing Drugs for Treatment; Availability	11/9/2007	null date	11/9/2007		E7-21985
FDA	FDA-2007-0579	FDA-2007-0579-0001	New Animal Drugs; Change of Sponsors Address	11/14/2007	null date	11/14/2007		E7-22210
FDA	FDA-2007-0580	FDA-2007-0580-0001	Oral Dosage Form New Animal Drugs; Chlortetracycline Powder	11/14/2007	null date	11/14/2007		E7-22261
FDA	FDA-2007-0581	FDA-2007-0581-0001	ARCH Chemicals, Inc.; Withdrawal of Food Additive Petition FAP 6B4764	11/19/2007	null date	11/19/2007		E7-22536
FDA	FDA-2007-0582	FDA-2007-0582-0001	Agency Information Collection Activities: Submission for Office of Management and Budget Review; Comment Request; Medical Devices Third-Party Review Under the Food and Drug Administration Modernization Act	11/19/2007	null date	11/19/2007		E7-22586
FDA	FDA-2007-0583	FDA-2007-0583-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Voluntary Registration of Cosmetic Product Establishments	11/19/2007	null date	11/19/2007		E7-22588
FDA	FDA-2007-0584	FDA-2007-0584-0001	Draft Guidance for Food and Drug Administration Advisory Committee Members and Food and Drug Administration Staff: Voting Procedures for Advisory Committee Meetings; Availability	11/19/2007	null date	11/19/2007		07-05751
FDA	FDA-2007-0585	FDA-2007-0585-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Animal Drug User Fees and Fee Waivers and Reductions	11/19/2007	null date	11/19/2007		E7-22495

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0586	FDA-2007-0586-0001	Memorandum of Understanding Between the Food and Drug Administration and the Association of American Feed Control Officials	11/19/2007	null date	11/19/2007		07-05748
FDA	FDA-2007-0587	FDA-2007-0587-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Recordkeeping and Records Access Requirements for FoodFacilities	11/19/2007	null date	11/19/2007		E7-22480
FDA	FDA-2007-0588	FDA-2007-0588-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Draft Guidance for Industry: Cooperative Manufacturing Arrangements for Licensed Biologics	11/19/2007	null date	11/19/2007		E7-22489
FDA	FDA-2007-0589	FDA-2007-0589-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Medical Devices: Recommended Glossary and Educational Outreach to Support Use of Symbols on Labels and in Labeling of In Vitro Diagnostic Devices Intended for Professional Use	11/19/2007	null date	11/19/2007		E7-22492
FDA	FDA-2007-0590	FDA-2007-0590-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Current Good Manufacturing Practice Regulations for Medicated Feeds	11/19/2007	null date	11/19/2007		E7-22587
FDA	FDA-2007-0591	FDA-2007-0591-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Animal Drug User Fee Cover Sheet, FDA Form 3546	11/20/2007	null date	11/20/2007		E7-22649
FDA	FDA-2007-0592	FDA-2007-0592-0001	Compliance Policy Guide; Radiofrequency Identification Feasibility Studies andPilot Programs for Drugs; Notice to Extend Expiration Date	11/23/2007	null date	11/23/2007		E7-22818
FDA	FDA-2007-0593	FDA-2007-0593-0001	International Cooperation on Harmonisation of Technical Requirements for Registration of Veterinary Medicinal Products; Revised Guidance for Industry on Impurities in New Veterinary Drug Substances (Revision); Availability	11/23/2007	null date	11/23/2007		E7-22902

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Food and Drug Administration (FDA) -- 1547 Documents

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Food and Drug Administration (FDA)								
FDA	FDA-2007-0594	FDA-2007-0594-0001	Draft Guidance for Industry on Smallpox (Variola) Infection: Developing Drugs for Treatment or Prevention; Availability	11/23/2007	null date	11/23/2007		E7-22884
FDA	FDA-2007-0595	FDA-2007-0595-0001	Memorandum of Understanding Between the Food and Drug Administration and Duke University	11/23/2007	null date	11/23/2007		07-05793
FDA	FDA-2007-0596	FDA-2007-0596-0001	International Cooperation on Harmonisation of Technical Requirements for Registration of Veterinary Medicinal Products; Revised Guidance for Industry on Impurities in New Veterinary Medicinal Products (Revision); Availability	11/23/2007	null date	11/23/2007		E7-22901
FDA	FDA-2007-0597	FDA-2007-0597-0001	International Cooperation on Harmonisation of Technical Requirements for Registration of Veterinary Medicinal Products; Revised Guidance for Industry on Stability Testing of New Veterinary Drug Substances and Medicinal Products (Revision); Availability	11/23/2007	null date	11/23/2007		E7-22900
FDA	FDA-2007-0598	FDA-2007-0598-0001	New Animal Drugs For Use in Animal Feeds; Ractopamine	11/23/2007	null date	11/23/2007		E7-22882
FDA	FDA-2007-0599	FDA-2007-0599-0001	New Animal Drugs For Use in Animal Feeds; Florfenicol	11/26/2007	null date	11/26/2007		E7-22942
FDA	FDA-2007-0600	FDA-2007-0600-0001	Determination That ELOXATIN (Oxaliplatin for Injection), 50 and 100 Milligrams Per Vial, Sterile Lyophilized Powder for Injection, Was Not Withdrawn From Sale for Reasons of Safety or Effectiveness	11/26/2007	null date	11/26/2007		E7-22973
FDA	FDA-2007-0602	FDA-2007-0602-0001	Behind the Counter Availability of Certain Drugs; Public Meeting; Comment Period Clarification	11/27/2020	null date	11/27/2020		E7-23026
FDA	FDA-2007-0603	FDA-2007-0603-0001	New Animal Drugs For Use in Animal Feeds; Fenbendazole	11/27/2007	null date	11/27/2007		E7-22987
FDA	FDA-2007-0604	FDA-2007-0604-0001	Risk Assessment of the Public Health Impact from Foodborne Listeria Monocytogenes in Soft-Ripened Cheese: Request for Comments and for Scientific Data and Information	11/28/2007	null date	11/28/2007		E7-23104
FDA	FDA-2007-0605	FDA-2007-0605-0001	Gastrointestinal Drugs Advisory Committee; Notice of Meeting	11/29/2007	null date	11/29/2007		E7-23177

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0606	FDA-2007-0606-0001	Food Labeling: Use of Symbols to Communicate Nutrition Information, Consideration of Consumer Studies and Nutritional Criteria; Reopening of Comment Period	11/30/2007	null date	11/30/2007		E7-23211
FDA	FDA-2007-0607	FDA-2007-0607-0001	Cold, Cough, Allergy, Bronchodilator, and Antiasthmatic Drug Products for Over-the-Counter Human Use; Final Rule for Over-the-Counter Antitussive Drug Products; Technical Amendment	11/30/2007	null date	11/30/2007	0910-AF33	E7-23207
FDA	FDA-2007-0608	FDA-2007-0608-0001	Medical Devices; Hematology and Pathology Devices: Reclassification of Automated Blood Cell Separator Device Operating by Centrifugal Separation Principle	11/30/2007	null date	11/30/2007		E7-23285
FDA	FDA-2007-0609	FDA-2007-0609-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Registration of Producers of Drugs and Listing of Drugs in Commercial Distribution	11/30/2007	null date	11/30/2007		E7-23275
FDA	FDA-2007-0610	FDA-2007-0610-0001	Guidance for Industry and Food and Drug Administration Staff: Class II Special Controls Guidance Document: Automated Blood Cell Separator Device Operating by Centrifugal or Filtration Separation Principle; Availability	11/30/2007	null date	11/30/2007		E7-23281
FDA	FDA-2007-0611	FDA-2007-0611-0001	General Mills, Inc.; Filing of Food Additive Petition	12/3/2007	null date	12/3/2007		E7-23400
FDA	FDA-2007-0612	FDA-2007-0612-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Current Good Manufacturing Practice Regulations for Type A Medicated Articles	12/3/2007	null date	12/3/2007		E7-23351
FDA	FDA-2007-0613	FDA-2007-0613-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Color Additive Certification Requests and Recordkeeping	12/3/2007	null date	12/3/2007		E7-23352
FDA	FDA-2007-0615	FDA-2007-0615-0001	DSM Nutritional Products, Inc.; Filing of Color Additive Petition	12/4/2007	null date	12/4/2007		E7-23473
FDA	FDA-2007-0616	FDA-2007-0616-0001	Current Good Manufacturing Practice; Amendment of Certain Requirements For Finished Pharmaceuticals; Withdrawal	12/4/2007	null date	12/4/2007		E7-23271

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0617	FDA-2007-0617-0001	Oral Dosage Form New Animal Drugs; Carprofen	12/5/2007	null date	12/5/2007		E7-23516
FDA	FDA-2007-0618	FDA-2007-0618-0001	New Animal Drugs For Use in Animal Feeds; Monensin USP	12/5/2007	null date	12/5/2007		E7-23517
FDA	FDA-2007-0619	FDA-2007-0619-0001	New Animal Drugs For Use in Animal Feeds; Monensin	12/5/2007	null date	12/5/2007		07-23519
FDA	FDA-2007-0620	FDA-2007-0620-0001	Implantation or Injectable Dosage Form New Animal Drugs; Erythromycin	12/7/2007	null date	12/7/2007		E7-23763
FDA	FDA-2007-0621	FDA-2007-0621-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Mental Models Study of Communicating With Health Care Providers About the Risks and Benefits of Prescription Drug Use for Pregnant and Nursing Women With Chronic Conditions	12/11/2007	null date	12/11/2007		E7-23976
FDA	FDA-2007-0622	FDA-2007-0622-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Guidance for Industry on Pharmacogenomic Data Submissions; Extension	12/11/2007	null date	12/11/2007		E7-23996
FDA	FDA-2007-0623	FDA-2007-0623-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Reports of Corrections and Removals	12/11/2007	null date	12/11/2007		E7-23962
FDA	FDA-2007-0624	FDA-2007-0624-0001	Advisory Committees; Filing of Closed Meeting Reports	12/11/2007	null date	12/11/2007		E7-23986
FDA	FDA-2007-0625	FDA-2007-0625-0001	Establishment of Fiscal Year 2008 User Fee Rates for Advisory Review of Direct-to-Consumer Television Advertisements for Prescription Drug and Biological Products	12/11/2007	null date	12/11/2007		E7-24000
FDA	FDA-2007-0626	FDA-2007-0626-0001	Agency Information Collection Activities; Submission forOffice of Management and Budget Review; Comment Request; Radioactive Drug Research Committees	12/11/2007	null date	12/11/2007		E7-23977
FDA	FDA-2007-0627	FDA-2007-0627-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Presubmission Conferences, New Animal Drug Applications and Supporting Regulations and Guidance 152, and Form FDA 356V	12/11/2007	null date	12/11/2007		E7-23998
FDA	FDA-2007-0628	FDA-2007-0628-0001	Drug Safety and Risk Management Advisory Committee; Notice of Meeting	12/11/2007	null date	12/11/2007		E7-24003

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0630	FDA-2007-0630-0001	Agency Emergency Processing Under the Office of Management and Budget Review; Certification to Accompany Drug, Biological Product, and Device Applications or Submissions	12/12/2007	null date	12/12/2007		07-06023
FDA	FDA-2007-0631	FDA-2007-0631-0001	Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Health and Diet Survey	12/13/2007	null date	12/13/2007		E7-24123
FDA	FDA-2007-0632	FDA-2007-0632-0001	Quality System Regulation Educational Forum on Design Controls; Public Workshop; Amendment of Notice	12/13/2007	null date	12/13/2007		07-24144
FDA	FDA-2007-0633	FDA-2007-0633-0001	New Animal Drugs for Use in Animal Feeds; Oxytetracycline	12/13/2007	null date	12/13/2007		E7-24146
FDA	FDA-2007-0634	FDA-2007-0634-0001	New Animal Drugs For Use in Animal Feeds; Ractopamine	12/13/2007	null date	12/13/2007		E7-24145
FDA	FDA-2007-0635	FDA-2007-0635-0001	International Conference on Harmonisation; Draft Guidance on Q4B Evaluation and Recommendation of Pharmacopoeial Texts for Use in the International Conference on Harmonisation Regions; Annex 2 on Test for Extractable Volume of Parenteral Preparations General Chapter; Availability	12/17/2007	null date	12/17/2007		E7-24434
FDA	FDA-2007-0636	FDA-2007-0636-0001	International Conference on Harmonisation; Draft Guidance on Q4B Evaluation and Recommendation of Pharmacopoeial Texts for Use in the ICH Regions; Annex 3 on Test for Particulate Contamination: Subvisible Particles General Chapter; Availability	12/17/2007	null date	12/17/2007		E7-24431
FDA	FDA-2007-0637	FDA-2007-0637-0001	Guidance for Industry and Food and Drug Administration Review Staff: Collection of Platelets by Automated Methods; Availability	12/17/2007	null date	12/17/2007		E7-24385
FDA	FDA-2007-0638	FDA-2007-0638-0001	Over-the-Counter Vaginal Contraceptive and Spermicide Drug Products Containing Nonoxynol 9; Required Labeling	12/19/2007	null date	12/19/2007	0910-AF44	07-06111
FDA	FDA-2007-0639	FDA-2007-0639-0001	Medical Devices; Availability of Safety and Effectiveness Summaries for Premarket Approval Applications	12/19/2007	null date	12/19/2007		E7-24620
FDA	FDA-2007-0640	FDA-2007-0640-0001	Cellular, Tissue and Gene Therapies Advisory Committee; Notice of Meeting	12/19/2007	null date	12/19/2007		E7-24629

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Food and Drug Administration (FDA) -- 1547 Documents

AGENCY	<www.regulations.gov> DOCKET ID	<www.regulations.gov> DOCUMENT ID	DOCUMENT TITLE	COMMENT START	COMMENT END	FR DATE	RIN	FR NUMBER
Food and Drug Administration (FDA)								
FDA	FDA-2007-0641	FDA-2007-0641-0001	Food and Drug Administration Modernization Act of 1997: Modifications to the List of Recognized Standards, Recognition List Number: 019	12/19/2007	null date	12/19/2007		E7-24580
FDA	FDA-2007-0642	FDA-2007-0642-0001	Psychopharmacologic Drugs Advisory Committee; Notice of Meeting	12/19/2007	null date	12/19/2007		E7-24627
FDA	FDA-2007-0643	FDA-2007-0643-0001	Health Claims and Qualified Health Claims; Dietary Lipids and Cancer, Soy Protein and Coronary Heart Disease, Antioxidant Vitamins and Certain Cancers, and Selenium and Certain Cancers; Reevaluation; Opportunity for Public Comment	12/21/2007	null date	12/21/2007		E7-24813
FDA	FDA-2007-0644	FDA-2007-0644-0001	Pulmonary-Allergy Drugs Advisory Committee; Notice of Meeting	12/21/2007	null date	12/21/2007		E7-24812
FDA	FDA-2007-0645	FDA-2007-0645-0001	Agency Emergency Processing Under the Office of Management and Budget Review; Certification to Accompany Drug, Biological Product, and Device Applications or Submissions; Correction	12/26/2007	null date	12/26/2007		E7-24914
FDA	FDA-2007-0646	FDA-2007-0646-0001	DSM Nutritional Products, Inc.; Filing of Color Additive Petition; Correction	12/26/2007	null date	12/26/2007		E7-24911
FDA	FDA-2007-0647	FDA-2007-0647-0001	Maximizing the Public Health Benefit of Adverse Event Collection Throughout a Products Marketed Life Cycle; Public Workshop; Request for Comments	12/26/2007	null date	12/26/2007		E7-24960
FDA	FDA-2007-0648	FDA-2007-0648-0001	New Animal Drugs; Change of Sponsors Name	12/26/2007	null date	12/26/2007		E7-24974
FDA	FDA-2007-0649	FDA-2007-0649-0001	Clinical Trial Design for Community-Acquired Pneumonia; Public Workshop	12/26/2007	null date	12/26/2007		E7-24927
FDA	FDA-2007-0651	FDA-2007-0651-0001	Cellular, Tissue and Gene Therapies Advisory Committee; Notice of Meeting	12/28/2007	null date	12/28/2007		E7-25124
FDA	FDA-2007-0652	FDA-2007-0652-0001	Draft Prescription Drug User Fee Act IV Information Technology Plan; Availability for Comment	12/28/2007	null date	12/28/2007		E7-25310
FDA	FDA-2008-0001	FDA-2008-0001-0001	Draft Guidance for Industry: Questions and Answers Regarding the Labeling of Dietary Supplements as Required by the Dietary Supplement and Nonprescription Drug Consumer Protection Act; Availability	1/2/2008	null date	1/2/2008		07-06266

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Food and Drug Administration (FDA)								
FDA	FDA-2008-0002	FDA-2008-0002-0001	Draft Guidance for Industry on Questions and Answers Regarding the Labeling of Nonprescription Human Drug Products Marketed Without an Approved Application as Required by the Dietary Supplement and Nonprescription Drug Consumer Protection Act; Availability	1/2/2008	null date	1/2/2008		07-06267
FDA	FDA-2008-0003	FDA-2008-0003-0001	Organization, functions, and authority delegations: Center for Food Safety and Applied Nutrition	1/2/2008	null date	1/2/2008		07-06257
FDA	FDA-2008-0004	FDA-2008-0004-0001	Guidance for Industry and Food and Drug Administration; Interactive Review for Medical Device Submissions: 510(k)s, Original PMAs, PMA Supplements, Original BLAs, and BLA Supplements; Availability	1/2/2008	null date	1/2/2008		07-06268
FDA	FDA-2008-0005	FDA-2008-0005-0001	Toll-Free Number for Reporting Adverse Events on Labeling for Human Drug Products	1/3/2008	null date	1/3/2008	0910-AC35	E7-25426
FDA	FDA-2008-0006	FDA-2008-0006-0001	Intramammary Dosage Form New Animal Drugs; Pirlimycin	1/4/2008	null date	1/4/2008		E7-25606
FDA	FDA-2008-0007	FDA-2008-0007-0001	New Animal Drugs For Use in Animal Feed; Semduramicin	1/4/2008	null date	1/4/2008		E7-25605
FDA	FDA-2008-0008	FDA-2008-0008-0001	Request for Comments on the Science and Technology Report; Establishment of Docket; Request for Comments	1/4/2008	null date	1/4/2008		E7-25607
FDA	FDA-2008-0009	FDA-2008-0009-0001	Draft Guidance for Industry on Antibacterial Drug Products: Use of Noninferiority Studies to Support Approval; Availability; Reopening of Comment Period	1/4/2008	null date	1/4/2008		E7-25601
FDA	FDA-2008-0010	FDA-2008-0010-0001	Agency Information Collection Activities; Proposed Collection; Comment Request; Application for Food and Drug Administration Approval to Market a New Drug	1/4/2008	null date	1/4/2008		E7-25593
FDA	FDA-2008-0011	FDA-2008-0011-0001	Meeting Being Planned to Obtain Public Input for Ensuring the Safety of Pet Food	1/7/2008	null date	1/7/2008		E7-25599
FDA	FDA-2008-0012	FDA-2008-0012-0001	Guidance for Industry and Food and Drug Administration Staff; The Review and Inspection of Premarket Approval Applications Under the Bioresearch Monitoring Program; Availability	1/8/2008	null date	1/8/2008		E8-00143

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Food and Drug Administration (FDA)								
FDA	FDA-2008-0013	FDA-2008-0013-0001	Guidance for Industry and Food and Drug Administration Staff; The Review and Inspection of Premarket Approval Application Manufacturing Information and Operations; Availability	1/8/2008	null date	1/8/2008		E8-00126
FDA	FDA-2008-0014	FDA-2008-0014-0001	Draft, Revised Compliance Policy Guide Sec. 575.100 Pesticide Chemical Residues in Food--Enforcement Criteria (CPG 7141.01); Availability	1/8/2008	null date	1/8/2008		E8-00123
FDA	FDA-2008-0015	FDA-2008-0015-0001	Compliance Policy Guide Sec. 555.700 Revocation of Tolerances for Cancelled Pesticides (CPG 7120.29); Withdrawal	1/8/2008	null date	1/8/2008		E8-00127
FDA	FDA-2008-0016	FDA-2008-0016-0001	Memorandum of Understanding Between the Food and Drug Administration and Regents of the University of California	1/9/2008	null date	1/9/2008		08-00030
FDA	FDA-2008-0017	FDA-2008-0017-0001	Determination That INDERAL (Propranolol Hydrochloride) Tablets, 10 Milligrams, 20 Milligrams, and 90 Milligrams Were Not Withdrawn From Sale for Reasons of Safety or Effectiveness	1/9/2008	null date	1/9/2008		E8-00190
FDA	FDA-2008-0018	FDA-2008-0018-0001	International Conference on Harmonisation; Draft Guidance on Q8(R1) Pharmaceutical Development; Availability	1/10/2008	null date	1/10/2008		E8-00213
FDA	FDA-2008-0019	FDA-2008-0019-0001	Anti-Infective Drugs Advisory Committee; Notice of Meeting	1/11/2007	null date	1/11/2007		E8-00343
FDA	FDA-2008-0020	FDA-2008-0020-0001	Kemira Oyi; Filing of Food Additive Petition (Animal Use); Partially Ammoniated Formic Acid	1/11/2007	null date	1/11/2007		E8-00316
	FDA Total Documents	1547						