interest is preserved in properties acquired with public funds. The rule further ensures compliance with all other Federal statutes applicable to the expenditure of Federal funds when acquiring real property.

Respondents: Head Start and Early Head Start grantees and delegate agencies.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Regulation	200	1	41	8,200.

Estimated Total Annual Burden Hours: 8,200

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Administration, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. All requests should be identified by the title of the information collection. E-mail address: *infocollection@acf.hhs.gov.*

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register.** Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, Attn: Desk Officer for ACF, E-mail address: *Katherine_T._Astrich@omb.eop.gov.*

Dated: June 8, 2006.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 06–5437 Filed 6–14–06; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2006F–0225]

Georgia-Pacific Resins, Inc.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Georgia-Pacific Resins, Inc., has filed a petition proposing that the food additive regulations be amended to provide for the safe use of glycerol ester of tall oil rosin to adjust the density of citrus oils used in the preparation of beverages and to provide for the use of steam stripping as a purification method for producing glycerol ester of wood rosin, gum rosin, or tall oil rosin.

FOR FURTHER INFORMATION CONTACT: Clarence W. Murray III, Center for Food Safety and Applied Nutrition (HFS–265), Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740–3835, 301–436–1311.

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (sec. 409(b)(5) (21 U.S.C. 348(b)(5))), notice is given that a food additive petition (FAP 6A4765) has been filed by Georgia-Pacific Resins, Inc., P.O. Box 105734, Atlanta, GA 30348. The petition proposes to amend the food additive regulations in § 172.735 *Glycerol ester of wood or gum rosin* (21 CFR 172.735) to provide for the following: (1) The safe use of glycerol ester of tall oil rosin to adjust the density of citrus oils used in the preparation of beverages; and (2) the use of steam stripping as a purification method for producing glycerol ester of wood rosin, gum rosin, or tall oil rosin.

The agency has determined under 21 CFR 25.32(k) that this action is of a type that does not individually or cumulatively have a significant effect on the human environment. Therefore, neither an environmental assessment nor an environmental impact statement is required.

Dated: May 24, 2006.

Laura M. Tarantino,

Director, Office of Food Additive Safety, Center for Food Safety and Applied Nutrition.

[FR Doc. E6–9319 Filed 6–14–06; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2005D–0200]

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

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of availability.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of a guidance document for industry (#177) entitled ''Specifications: Test Procedures and Acceptance Criteria for New Biotechnological/Biological Veterinary Medicinal Products'' (VICH GL40). This guidance has been developed for veterinary use by the International Cooperation on Harmonization of Technical Requirements for Registration of Veterinary Medicinal Products (VICH). This VICH guidance document provides general principles through recommendations on the setting and justification, to the extent possible, of a uniform set of international specifications for biotechnological and biological products to support new marketing applications.

DATES: Submit written or electronic comments at any time.

ADDRESSES: Submit written requests for single copies of the guidance to the Communications Staff (HFV–12), Center for Veterinary Medicine (CVM), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855. Send one self-addressed adhesive label to assist that office in processing your requests. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the guidance document.