Subject name and address	Effective date
PAWTUCKET, RI	
PINKERTON, LINDA ONEIDA, NY	1/19/2005
RADOMSKI, JULIENNE LEESBURG, VA	1/19/2005
RAYMOND, JERRY SPRINGFIELD, MO	1/19/2005
RENICK, LISA BOISE, ID	1/19/2005
RENKEN, CAREY DENVER, CO	1/19/2005
RIVERA, ANGEL HONDO, TX	1/19/2005
ROBERTS, FAITH MIAMI, FL	1/19/2005
SACCONE, MARGUERITE FT LAUDERDALE, FL	1/19/2005
SALTER, DEBORAH COLUMBIA, AL	1/19/2005
SANFORD, CYNTHIA ENGLEWOOD, CO	1/19/2005
SANTIAGO, JAMIE CLEVELAND, OH	1/19/2005
SCOTT, CLINTON LONG BEACH, CA	1/19/2005
SHAH, KAMLESH ONTARIO, CA	1/19/2005
SIMMONS, CAROLYN BRISTOL, VA	1/19/2005
SMITH, KAREN SALT LAKE CITY, UT	1/19/2005
SMITH, RAMONA CAROLINA BEACH, NC	1/19/2005
SOARES, KIM SWANSEA, RI	1/19/2005
SOWLES, CHARLENE PHOENIX, AZ	1/19/2005
STROBLE, LAURA WEST DOVER, VT	1/19/2005
STUDLEY, KATHLEEN RUMFORD, RI	1/19/2005
SWAIN, JEREMY COVENTRY, RI	1/19/2005
TAYLOR, DONALD HAVRE, MT	1/19/2005
TAYLOR, MATTHEW CLEARFIELD, UT	1/19/2005
TAYLOR, MEL SAN LEANDRO, CA	1/19/2005
THAN-RIDDLE, KHIN GAMBRILLS, MD	1/19/2005
TILLEY-BISBEE, COURTNEY SCOTTSDALE, AZ	1/19/2005
TOPPING, GARY DERBY, KS	1/19/2005
VALDIVIA, CARLOS OXNARD, CA	1/19/2005
VARGOVICH, KAREN OROFINO, ID	1/19/2005
VARNUM, ROLAND LAKE CHARLES, LA	1/19/2005
VINLUAN, ELEANOR CHEYENNE, WY	1/19/2005
WALLACE, FREDRICK BESSEMER, AL	1/19/2005
WALLACE, SELENA LAKE CITY, FL	1/19/2005
WASHINGTON, CARNELLA OAKLAND, CA	1/19/2005
WEBER, SUSAN COLORADO SPRINGS, CO	1/19/2005
WILKERSON, NINA PHOENIX, AZ	1/19/2005
WILLIAMS, APRIL PARACHUTE, CO	1/19/2005
WILLIAMS, REBECCA AURORA, CO	1/19/2005
WOLAN, NANCY TAMPA, FL	1/19/2005
YOUNG, KISHA LONGVIEW, WA	1/19/2005
FEDERAL/STATE EXCLUSION/ SUSPENSION	
RYDER, JON EUGENE, OR	1/19/2005
FRAUD/KICKBACKS/PROHIBITED ACTS/ SETTLEMENT AGREEMENTS	
GARCIA, GULLERMO MIAMI, FL	11/4/2004
GONSALVES, WALLACE MINERSVILLE, PA	9/23/2004
PATE, JIVA MYRTLE BEACH, SC	11/1/2004
OWNED/CONTROLLED BY CONVICTED ENTITY	
ALPHA HERBS & VITAMINS HOUSTON, TX	1/19/2005
ALPHA SENIOR HEALTH GROUP HOUSTON, TX	1/19/2005
AVIONIX MEDICAL DEVICES, INC MIDLAND, TX	1/19/2005
HEALTH RESOURCES & REHAB HOUSTON, TX	1/19/2005
HEALTH RESOURCES & REHAB CENTER HOUSTON, TX	1/19/2005
JORGENSON DRUG, INC ROUNDUP, MT	1/19/2005
MARITIME HEALTH SERVICES OF TAMPA, INC TAMPA, FL	1/19/2005
SEELIN MEDICAL, INC MIDLAND, TX	1/19/2005
DEFAULT ON HEAL LOAN	
BEAMS, JEFFREY TAMPA, FL	1/19/2005
BULEN, JERRY BRANDON, FL	1/19/2005
EAGLE, DONALD GRAND ISLAND, FL	1/19/2005
GREEN, MICHAEL MAITLAND, FL	1/19/2005
HASLEY, STEVEN MELBOURNE, FL	1/19/2005
HEISLER, HOPE PUNTA GORDA, FL	1/19/2005
JOSEPH, BRAD PITTSBURGH, PA	1/19/2005
LIGHT, DAVID WINTER GARDEN, FL	1/19/2005
MARIN, MELODY VAN NUYS, CA	1/19/2005
MARQUEZ, EVELYN LOS ANGELES, CA	1/19/2005
NEALY, JOY ATLANTA, GA	1/19/2005
ROGEL-ELLIOTT, VALERIE SEMINOLE, FL	1/19/2005
SMITH, MICHAEL TORRANCE, CA	1/19/2005
STANBRIDGE, GARY WHITTIER, CA	1/19/2005
VAFAEE, MOHAMMAD SANTA MONICA, CA	1/19/2005
YANG, SHENG FOUNTAIN VALLEY, CA	1/19/2005
OWNERS OF EXCLUDED ENTITIES	
RATHOD, BABUBHAI MT PLEASANT, MI	1/19/2005

Dated: January 24, 2005.

Katherine B. Petrowski,

Director, Exclusions Staff, Office of Inspector General.

[FR Doc. 05–1788 Filed 1–31–05; 8:45 am]

BILLING CODE 4150–04–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Reporting of Pregnancy Success Rates From Assisted Reproductive Technology Programs

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (DHHS).

ACTION: Notice.

SUMMARY: The CDC is tasked with implementing the Fertility Clinic Success Rate and Certification Act of 1992 (FCSRCA), Public Law 102–493. As mandated by this law CDC publishes annual reports of pregnancy success rates from ART clinics and embryo laboratory certification status of these clinics. Section 2(a) of Public Law 102–493 (42 U.S.C. 263a–1) requires that each assisted reproductive technology (ART) program shall annually report to the Secretary, through the Centers for Disease Control and Prevention, (a) pregnancy success rates achieved by such ART programs, and (b) the identity of each embryo laboratory used by such ART programs, and whether the laboratory is certified or has applied for such certification under this act. Section (6) states that the Secretary, through the CDC, shall annually publish and distribute to the States and the public, pregnancy success rates reported to the Secretary under section 2(a)(1) and, in the case of an assisted reproductive technology program which failed to report one or more success rates as required under such section, the name

of each such program and each pregnancy success rate which the program failed to report.

This Announcement includes information on the change in the data collection contractor and the change in the approved data reporting system for the 2004, 2005, 2006, 2007, and 2008 ART data reporting years in accordance with the FCSRCA. This Announcement supplements the September 1, 2000 and the February 5, 2004, notices.

SUPPLEMENTARY INFORMATION: CDC has contracted with Westat to develop a data reporting system and to collect annual clinic-specific and cycle-specific data from all practicing assisted reproductive technology clinics in the U.S. and its territories for the 2004, 2005, 2006, 2007, and 2008 ART data reporting years. The contract covers clinic tracking, data collection and quality assurance, and validation activities. As such, Westat is the new contractor for ART data collection for the 2004 through 2008 ART data reporting years.

The new Web-based data reporting system (developed by Westat) for the 2004, 2005, 2006, 2007, and 2008 ART data reporting years will be called the National ART Surveillance System (NASS). As such, NASS will be the only approved data reporting system for 2004 through 2008 ART data submissions. ART programs should be aware that Westat will develop and provide all necessary instruction materials for extracting and importing data from other electronic medical record systems into NASS and for checking imported data to ensure that it retains the accuracy and compatibility of the data entry system from which it was extracted.

The anticipated deadline for reporting is December 15 of the year 1 year subsequent to the reporting year in question. (For example, the anticipated deadline to report data on cycles initiated in 2004 is December 15, 2005.) An ART program will not be considered to be in compliance with the federal reporting requirements of FCSRCA if the ART program was in operation in the full year that is being reported, *i.e.*, the clinic was in operation after January 1 of the reporting year, and fails to submit a dataset to Westat in the required data reporting system (NASS) by the reporting deadline. ART programs considered to not be compliant with the federal reporting requirements of FCSRCA will be listed as non-reporters in the published report.

The data reporting activities and the amount and type of data collected will be similar to the current system requirements outlined in the September 1, 2000 **Federal Register** notice (Volume 65, No. 171, pages 53310–53316). CDC has completely funded the data reporting activities for the 2004 through 2008 reporting years. Thus, ART programs will not be charged fees to obtain the new reporting system or to submit data using the new reporting system.

Validation activities for the 2004 through 2008 data reporting years will be similar to those described in the September 1, 2000 **Federal Register** notice (Volume 65, No. 171, pages 53310–53316). Westat will provide the necessary personnel to perform the validation site visits.

Each ART program should be aware that the Paperwork Reduction Act is applicable to this data collection. Under the Paperwork Reduction Act of 1995, a Federal agency shall not conduct or sponsor a collection of information from ten or more persons other than Federal employees, unless the agency has submitted a Standard Form 83, Clearance Request, to the Director of the Office of Management and Budget (OMB), and OMB has approved the collection of information. A person is not required to respond to a collection of information unless it displays a currently valid OMB control number. CDC has obtained OMB approval to collect this data under OMB control No. 0920–0556.

CDC will continue to provide information to all ART programs regarding data collection activities as information becomes available.

FOR FURTHER INFORMATION CONTACT: Victoria Wright, Assisted Reproductive Technology Epidemiology Unit at (770) 488–6384.

Dated: January 25, 2005.

James D. Seligman,

Associate Director for Program Services, Centers for Disease Control and Prevention.

[FR Doc. 05–1787 Filed 1–31–05; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2005N–0029]

Agency Information Collection Activities; Proposed Collection; Comment Request; Infant Formula Recall Regulations

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on requirements related to the recall of infant formula.

DATES: Submit written or electronic comments on the collection of information by April 4, 2005.

ADDRESSES: Submit written comments to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments to *http://www.fda.gov/dockets/ecomments*. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Peggy Robbins, Office of Management Programs (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1223.

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites comments on these topics: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA's estimate of the burden of the proposed collection of information, including the