administrative, provider, and customer services process.

- Facilitate plans for IT Integration of data resources and data services.
- Coordinate policy analysis, development and execution for CMS.
- Build and maintain agency capacity to perform analysis of regional variation in the quality and cost of care.
- Conduct and manage surveys to capture information about beneficiary populations that our programs serve that is not available in the administrative data. This includes the Medicare Current Beneficiary Survey (MCBS) and the Medicare Health Outcomes Survey (HOS).
- Conduct and manage the Research Data Assistance Center (RESDAC), Research Data Distribution Center (RDDC) and Chronic Condition Warehouse (CCW) activities.
- Operationalize research-usable files for Medicare, Medicaid, and CHIP administrative data.

Dated: March 24, 2011.

Marilyn Tavenner,

Principal Deputy Administrator and Chief Operating Officer, Centers for Medicare & Medicaid Services.

[FR Doc. 2011-7903 Filed 4-1-11; 8:45 am]

BILLING CODE 4120-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Reunification Procedures for Unaccompanied Alien Children. *OMB No.*: 0970–0278.

Description: Following the passage of the 2002 Homeland Security Act (Pub. L. 107–296), the Administration for Children and Families (ACF), Office of Refugee Resettlement (ORR), is charged with the care and placement of

unaccompanied alien children in Federal custody, and implementing a policy for the release of these children, when appropriate, upon the request of suitable sponsors while awaiting immigration proceedings. In order for ORR to make determinations regarding the release of these children, the potential sponsors must meet certain conditions pursuant to section 462 of the Homeland Security Act and the Flores v. Reno Settlement Agreement No. CV85 4544-RJK (C.D. Cal. 1997). The proposed information collection requests information to be utilized by ORR for determining the suitability of a sponsor/respondent for the release of a minor from ORR custody. The proposed instruments are the Sponsors Agreement to Conditions of Release, Verification of Release, Family Reunification Packet, and the Authorization for Release of Information.

Respondents: Sponsors requesting release of unaccompanied alien.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Verification of Release (UAC)	4,595 4,595 4,595 4,595 4,595 4,595 4,595 4,595	1 1 1 1 1 1 1	0.25 0.25 1 0.25 0.25 0.25 1 0.25	1,148.75 1,148.75 4,595 1,148.75 1,148.75 4,595 1,148.75

Estimated Total Annual Burden Hours: 16,082.50.

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, Fax: 202–395–7285, E-mail:

OIRA_SUBMISSION@OMB.EOP.GOV, Attn: Desk Officer for the Administration for Children and Families.

Dated: March 29, 2011.

Robert Sargis,

Reports Clearance Officer. [FR Doc. 2011–7823 Filed 4–1–11; 8:45 am]

BILLING CODE 4184-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration [Docket No. FDA-2010-N-0474]

Maja S. Ruetschi: Debarment Order

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is issuing an

order under the Federal Food, Drug, and Cosmetic Act (the FD&C Act) debarring Maja S. Ruetschi, MD for 5 years from providing services in any capacity to a person that has an approved or pending drug product application. FDA bases this order on findings that Dr. Ruetschi was convicted of a misdemeanor under Federal law for conduct relating to the regulation of a drug product under the FD&C Act and that the type of conduct underlying the conviction undermines the process for the regulation of drugs. Dr. Ruetschi was given notice of the proposed debarment and an opportunity to request a hearing within the timeframe prescribed by regulation. Dr. Ruetschi failed to respond. Dr. Ruetschi's failure to respond constitutes a waiver of her right to a hearing concerning this action.

DATES: This order is effective April 4,

ADDRESSES: Submit applications for termination of debarment to the Division of Dockets Management (HFA—