(7) The applicability of the insured institution's fidelity bond coverage to the person;

(8) The opinion or position of the primary Federal and/or state regulator; and

(9) Any additional factors in the specific case that appear relevant.

The foregoing criteria will also be applied by the FDIC to determine whether the interests of justice are served in seeking an exception in the appropriate court when an application is made to terminate the ten-year ban prior to its expiration date.

Some applications can be approved without an extensive review because the person will not be in a position to constitute any substantial risk to the safety and soundness of the insured institution. Persons who will occupy clerical, maintenance, service or purely administrative positions, generally fall into this category. A more detailed analysis will be performed in the case of persons who will be in a position to influence or control the management or affairs of the insured institution. Approval orders will be subject to the condition that the person shall be covered by a fidelity bond to the same extent as others in similar positions. In cases in which a waiver of the institution filing requirement has been granted to an individual, approval of the application will be conditioned upon that person disclosing the presence of the conviction to all insured institutions in the affairs of which he or she wishes to participate. When deemed appropriate, approval orders may also be subject to the condition that the prior consent of the FDIC will be required for any proposed significant changes in the person's duties and/or responsibilities. Such proposed changes may, in the discretion of the Regional Director, require a new application. In situations in which an approval has been granted for a person to participate in the affairs of a particular insured institution and subsequently seeks to participate at another insured institution, approval does not automatically follow. In such cases, another application must be submitted.

By Order of the Board of Directors.

Dated at Washington, DC, this 11th day of December, 2012.

Federal Deposit Insurance Corporation.

Robert E. Feldman,

Executive Secretary.

[FR Doc. 2012–30351 Filed 12–17–12; 8:45 am]

BILLING CODE 6714–01–P

FEDERAL DEPOSIT INSURANCE CORPORATION

Privacy Act of 1974; System of Records

AGENCY: Federal Deposit Insurance Corporation.

ACTION: Notice to Delete a System of Records.

SUMMARY: In accordance with the requirements of the Privacy Act of 1974, as amended (Privacy Act), the Federal Deposit Insurance Corporation (FDIC) deletes one system of records from its existing inventory of systems of records subject to the Privacy Act.

DATES: Effective Date is July 23, 2012.

FOR FURTHER INFORMATION CONTACT: Gary Jackson, Counsel, FDIC, 550 17th Street NW., Washington, DC 20429, (703) 562–2677.

SUPPLEMENTARY INFORMATION: The FDIC deletes its system of records for the Nationwide Mortgage Licensing System and Registry, 76 FR 15309 (March 21, 2011). Section 1100 of Title X of the Dodd-Frank Wall Street Reform and Consumer Protection Act transferred to the Bureau of Consumer Financial Protection (CFPB) authority to develop and maintain the national registration system for residential mortgage loan originators required by Section 1507 of the Secure and Fair Enforcement for Mortgage Licensing Act. The CFPB published its own notice of the establishment of a Privacy Act system of records for the Nationwide Mortgage Licensing System and Registry, 77 FR 35359 (June 13, 2012), effective as of July 23, 2012.

The deletion is not within the purview of subsection (r) of the Privacy Act, which requires submission of a report on a new or altered system of records. The FDIC's systems of records notices subject to the Privacy Act have been published in the **Federal Register** and may be viewed at *http://www.fdic.gov/regulations/laws/rules/2000-4000.html* on the FDIC's Privacy Web page.

By order of the Board of Directors.

Dated at Washington, DC, this 11th day of December 2012.

Robert E. Feldman,

Executive Secretary.

[FR Doc. 2012–30254 Filed 12–17–12; 8:45 am]

BILLING CODE P

FEDERAL ELECTION COMMISSION

Sunshine Act Meeting

AGENCY: Federal Election Commission.

DATE AND TIME: Thursday, December 20, 2012 at 10:00 a.m.

PLACE: 999 E Street NW., Washington, DC (Ninth Floor).

STATUS: This Meeting Will Be Open to the Public.

ITEMS TO BE DISCUSSED:

Correction and Approval of the Minutes for the Meeting of December 6, 2012;
Draft Advisory Opinion 2012–35: Global Transaction Services Group, Inc.;
Draft Advisory Opinion 2012–37: Yamaha Motor Corporation, U.S.A.;
Itemization of Ultimate Payee of Committee Disbursements;
Request for Comment on the Enforcement Process;
Election of Officers;
Future Meeting Dates;
Management and Administrative Matters.

Individuals who plan to attend and require special assistance, such as sign language interpretation or other reasonable accommodations, should contact Shawn Woodhead Werth, Secretary and Clerk, at (202) 694–1040, at least 72 hours prior to the meeting date.

PERSON TO CONTACT FOR INFORMATION: Judith Ingram, Press Officer, Telephone: (202) 694–1220.

Shawn Woodhead Werth,

Secretary and Clerk of the Commission.

[FR Doc. 2012–30492 Filed 12–14–12; 11:15 am]

BILLING CODE 6715–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Unaccompanied Refugee Minor Placement and Outcomes Reports; ORR–3 and ORR–4.

OMB No.: 0970–0034.

Description: The two reports collect information necessary to administer the Unaccompanied Refugee Minor (URM) program. The ORR–3 (Placement Report) is submitted to the Office of Refugee Resettlement (ORR) by the State agency at initial placement within 30 days of the placement, and whenever there is a change in the child's status, including termination from the program, within 60 days of the change or closure of the case. The ORR–4 (Outcomes Report) is submitted within approximately 12 months of the initial placement and each subsequent 12 months to record outcomes of the

child's progress toward the goals listed in the child's case plan and particularly for youth 17 years of age and above related to independent living and/or educational plans. ORR–4 is also submitted as a baseline report along with the initial ORR–3 report for 17 years old and above youth, and as a follow-up annual report for cases that have terminated and are 17 to 21 years old. ORR regulations at 45 CFR 400.120 describes specific URM program reporting requirements.

Respondents: State governments.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
ORR–3	15	75	0.25 (15 Minutes).	281.25
ORR–4	15	119	1.5 (1 Hour and 30 Minutes).	2,677.5

Estimated Total Annual Burden Hours: 2,958.75.

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 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. Email 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, Email: *OIRA_SUBMISSION@OMB.EOP.GOV.* Attn: Desk Officer for the Administration for Children and Families.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2012–30390 Filed 12–17–12; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2012–D–1197]

Draft Guidance for Industry on Certification of Designated Medical Gases; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of a draft guidance for industry entitled ''Certification Process for Designated Medical Gases.'' This draft guidance describes the new certification process created by the Food and Drug Administration Safety and Innovation Act (FDASIA) for certain medical gases and explains how FDA plans to implement that process.

DATES: Although you can comment on any guidance at any time (see 21 CFR 10.115(g)(5)), to ensure that the Agency considers your comment on this draft guidance before it begins work on the final version of the guidance, submit either electronic or written comments on the draft guidance by February 19, 2013. Submit either electronic or written comments concerning the collection of information proposed in the draft guidance and attached Form 3864 by February 19, 2013.

ADDRESSES: Submit written requests for single copies of this draft guidance to the Division of Drug Information, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 51, Rm. 2201, Silver Spring, MD 20993–0002; or the Communications Staff (HFV–12), Center for Veterinary Medicine, 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 draft guidance document.

Submit electronic comments on the draft guidance to *http://www.regulations.gov.* Submit written comments to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT:

Michael Folkendt, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Silver Spring, MD 20993–0002, 301–796–1900; or Germaine Connolly, Center for Veterinary Medicine (HFV–116), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 240–276–8331.

SUPPLEMENTARY INFORMATION:

I. Background

FDA is announcing the availability of a draft guidance for industry entitled ''Certification Process for Designated Medical Gases.'' This guidance is intended to help persons or entities interested in requesting a certification for a designated medical gas under the new approval process for designated medical gases created by FDASIA (Pub. L. 112–144, 126 Stat. 993).

Title XI, subtitle B, of FDASIA added sections 575 and 576 to the Federal Food, Drug, and Cosmetic Act (the FD&C Act), which created a certification process for designated medical gases. Specifically, section 575 provides that oxygen, nitrogen, nitrous oxide, carbon dioxide, helium, carbon monoxide, and medical air are designated medical gases. Section 576 permits any person, beginning on January 5, 2013, to request a certification of a medical gas for certain indications and describes when FDA will grant or deny these requests.

This draft guidance explains how FDA plans to implement this new certification process. Specifically, the draft guidance describes the medical gases that are eligible for certification, who should submit a certification request, what information should be submitted, and how FDA will evaluate and act on the request. The draft guidance also describes how the new certification requirement will be enforced and describes FDA's intent to exercise enforcement discretion in certain instances.

FDA has also developed a form to help requestors submit their certification requests. FDA recommends that requestors use this form. The form