information line to learn about possible modifications before coming to the meeting.

**Agenda:** On March 7, 2013, the committee will discuss the new drug application (NDA) 204275, for fluticasone furoate and vilanterol dry powder inhaler (proposed tradename BREO ELLIPTA), sponsored by GlaxoSmithKline, for the long-term maintenance treatment of airflow obstruction and for reducing exacerbations in patients with chronic obstructive pulmonary disease.

FDA intends to make background material available to the public no later than 2 business days before the meeting. If FDA is unable to post the background material on its Web site prior to the meeting, the background material will be made publicly available at the location of the advisory committee meeting, and the background material will be posted on FDA’s Web site after the meeting. Background material is available at http://www.fda.gov/AdvisoryCommittees/ default.htm. Scroll down to the appropriate advisory committee meeting link.

**Procedure:** Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person on or before February 21, 2013. Oral presentations from the public will be scheduled between approximately 1 p.m. and 2 p.m. Those individuals interested in making formal oral presentations should notify the contact person and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the time requested to make their presentation on or before February 12, 2013. Time allotted for each presentation may be limited. If the number of registrants requesting to speak is greater than can be reasonably accommodated during the scheduled open public hearing session, FDA may conduct a lottery to determine the speakers for the scheduled open public hearing session. The contact person will notify interested persons regarding their request to speak by February 13, 2013.

Persons attending FDA’s advisory committee meetings are advised that the Agency is not responsible for providing access to electrical outlets.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with physical disabilities or special needs. If you require special accommodations due to a disability, please contact Cindy Hong at least 7 days in advance of the meeting.

FDA is committed to the orderly conduct of its advisory committee meetings. Please visit our Web site at http://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm111462.htm for procedures on public conduct during advisory committee meetings.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

**Dated:** December 12, 2012.

**Leslie Kux,**

Assistant Commissioner for Policy.

Note: The address listed in this notice should only be used to submit comments concerning this information collection. Please do not submit requests for individual case status inquiries to this address. If you are seeking information about the status of your individual case, please check “My Case Status” online at https://egov.uscis.gov/cris/Dashboard.do, or call the USCIS National Customer Service Center at 1–800–375–5283.

Written comments and suggestions from the public and affected agencies should address one or more of the following four points:

1. Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;
2. Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;
3. Enhance the quality, utility, and clarity of the information to be collected; and
4. Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses.
Overview of This Information Collection

(1) Type of Information Collection Request: Revision of a currently approved information collection.
(2) Title of the Form/Collection: Petition for Alien Fiancé(e).
(3) Agency form number, if any, and the applicable component of the DHS sponsoring the collection: USCIS Form I–129F; USCIS.
(4) Affected public who will be asked or required to respond, as well as a brief abstract: Primary: Individuals or households. Form I–129F must be filed with USCIS by a citizen of the United States in order to petition for an alien fiancé(e), spouse, or his/her children.
(5) An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond: 46,936 responses at 1 hour and 35 minutes (1.58 hours) per response.
(6) An estimate of the total public burden (in hours) associated with the collection: 74,158 annual burden hours.

If you need a copy of the information collection instrument with supplementary documents, or need additional information, please visit http://www.regulations.gov. We may also be contacted at: USCIS, Office of Policy and Strategy, Regulatory Coordination Division, 20 Massachusetts Avenue NW., Washington, DC 20529–2140; Telephone 202–272–8377.

Dated: December 11, 2012
Laura Dawkins,
Chief, Regulatory Coordination Division,

[FR Doc. 2012–30215 Filed 12–13–12; 8:45 am]
BILLING CODE 9111–97–P

DEPARTMENT OF HOMELAND SECURITY

U.S. Citizenship and Immigration Services

[OMB Control Number 1615–0124]

Agency Information Collection Activities: Consideration of Deferred Action for Childhood Arrivals, Form I–821D, Revision of a Currently Approved Collection

ACTION: 60-day notice.

SUMMARY: In accordance with the Paperwork Reduction Act of 1995 (Pub. L. 104–13, 44 U.S.C. 35), on August 15, 2012, the Department of Homeland Security (DHS), U.S. Citizenship and Immigration Services (USCIS), submitted an information collection request, utilizing emergency review procedures, to the Office of Management and Budget (OMB) for review and clearance. OMB approved the information collection request. DHS is now requesting OMB approval of a revision and extension of the approved information collection.

DATES: Comments are encouraged and will be accepted for sixty days until February 12, 2013.

ADDRESSES: Written comments and suggestions regarding items contained in this notice, and especially with regard to the estimated public burden and associated response time should be directed to: DHS, USCIS, Office of Policy and Strategy, Chief, Regulatory Coordination Division, 20 Massachusetts Avenue NW., Washington, DC 20529–2140. Comments may be submitted to DHS via email at uscisfrcomment@uscis.dhs.gov and must include OMB Control Number 1615–0038 in the subject box. Comments may also be submitted via the Federal eRulemaking Portal at www.Regulations.gov under e-Docket ID number USCIS–2012–0012.

All submissions received must include the agency name and Docket ID. Regardless of the method used for submitting comments or material, all submissions will be posted, without change, to the Federal eRulemaking Portal at www.Regulations.gov, and will include any personal information you provide. Therefore, submitting this information makes it public. You may wish to consider limiting the amount of personal information that you provide in any voluntary submission you make to DHS. DHS may withhold information provided in comments from public viewing that it determines may impact the privacy of an individual or that is offensive. For additional information, please read the Privacy Act notice that is available via the link in the footer of www.Regulations.gov.

Issues for Comment Focus

DHS, USCIS invites the general public and other Federal agencies to comment upon this proposed revision of a currently approved collection of information. In accordance with the Paperwork Reduction Act of 1995, the information collection notice is published in the Federal Register to obtain comments regarding the nature of the information collection, the categories of respondents, the estimated burden (i.e., the time, effort, and resources used by the respondents to respond).

For Form I–821D, USCIS is especially interested in the public’s experience, input, and estimates on the burden in terms of time and money incurred by applicants for the following aspects of this information collection:
• The time burden incurred by preparers (persons who assist the respondent with the preparation of the form) who are not paid.
• The amount that paid preparers charge for their services.
• The time required to obtain supporting documents for Form I–821D.
• The monetary costs incurred to obtain supporting documents from sources such as a landlord, church, utility, public agency (housing, social services, law enforcement), school, medical care provider, advocacy group, law firm, or military service.
• The average time required and money expended to secure secondary evidence such as an affidavit.
• The percentage of total applicants who require English translations of their supporting documents.
• The percentage of supporting documents for each individual applicant that require translation into English.
• The time required to find, hire, or otherwise obtain translations of supporting documents for immigration benefit requests.

In addition, in order to truly be helpful to the improvement of this form and program written comments and suggestions concerning the collection of information are requested to provide clear and specific suggestions on the data elements on the form and the evidence required to be submitted with a focus on one or more of the following four points:
(1) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;
(2) The accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;
(3) How to enhance the quality, utility, and clarity of the information to be collected; and
(4) How to reduce or minimize the burden of the collection of information on those who are to respond, including through the use of appropriate