ANNUAL BURDEN ESTIMATES

| | Annual number of respondents | Number of responses per respondent | Average burden hours per response | Total burden hours |
|---|---|---|---|---|
| Field clearance instrument: | | | | |
| Discussion Guide for use with Researchers, Policy Experts, and State Level Coordinators | 10 | 1 | 1 | 10 |
| Discussion Guide for Use with Program Directors | 20 | 2 | 2 | 80 |
| Discussion Guide for Use with Program Staff | 40 | 1 | 2 | 80 |
| Focus Group Discussion Guide for use with Program Participants | 100 | 1 | 1.5 | 150 |
| Discussion Guide for Use with School Administrators | 70 | 1 | 1 | 70 |
| Short Survey with Program Directors | 70 | 1 | 0.25 | 17.5 |
| Short Survey with Program Staff | 140 | 1 | 0.25 | 35 |
| Short Survey with School Administrators | 70 | 1 | 0.25 | 17.5 |
| Estimated Annual Burden Sub-total for Field Clearance | ..... | ..... | ..... | 460 |
| In-Depth Implementation Instrument: | | | | |
| Master Topic Guide Interviews for General Staff and Community Members | 40 | 1 | 1.5 | 60 |
| Focus Group Discussion Guide with Frontline Staff | 30 | 1 | 1.5 | 45 |
| Focus Group Discussion Guide with Participating Youths | 150 | 1 | 1.5 | 225 |
| Focus Group Discussion Guide with Control Group Schools About Counterfactuals | 40 | 1 | 1 | 40 |
| Estimated Annual Burden Sub-total for In-Depth Implementation | ..... | ..... | ..... | 370 |
| Baseline Instrument: | | | | |
| Baseline Instrument for study participants | 2500 | 1 | .5 | 1250 |
| Administrative Data Collection instrument for Schools and Organizations | 100 | 1 | 4 | 400 |
| Estimated Annual Burden Sub-total for Baseline | ..... | ..... | ..... | 1650 |
| TOTAL Estimated Annual Burden | ..... | ..... | ..... | 2640 |

In compliance with the requirements of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: OPRE Reports Clearance Officer. E-mail address: *OPREinfocollection@acf.hhs.gov.* All requests should be identified by the title of the information collection.

The Department specifically requests comments on (a) whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Dated: March 9, 2011.

**Steven M. Hanmer,**

*Reports Clearance Officer.*

[FR Doc. 2011–5962 Filed 3–15–11; 8:45 am]

**BILLING CODE 4184–37–M**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

**[Docket No. FDA–2010–N–0541]**

### Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Guidance for Industry on Special Protocol Assessment

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing that a collection of information entitled "Guidance for Industry on Special Protocol Assessment" has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

**FOR FURTHER INFORMATION CONTACT:** Elizabeth Berbakos, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, 301–796–3792, *Elizabeth.Berbakos@fda.hhs.gov.*

**SUPPLEMENTARY INFORMATION:** In the **Federal Register** of October 26, 2010 (75 FR 65636), the Agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An Agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0470. The

approval expires on February 28, 2014. A copy of the supporting statement for this information collection is available on the Internet at *http://www.reginfo.gov/public/do/PRAMain.*

Dated: March 10, 2011.

**Leslie Kux,**

*Acting Assistant Commissioner for Policy.*

[FR Doc. 2011–6092 Filed 3–15–11; 8:45 am]

**BILLING CODE 4160–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

**[Docket No. FDA–2010–N–0468]**

### Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Patent Term Restoration, Due Diligence Petitions, Filing, Format, and Content of Petitions

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing that a collection of information entitled "Patent Term Restoration, Due Diligence Petitions, Filing, Format, and Content of Petitions" has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

**FOR FURTHER INFORMATION CONTACT:** Elizabeth Berbakos, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, 301–796–3792, *Elizabeth.Berbakos@fda.hhs.gov.*

**SUPPLEMENTARY INFORMATION:** In the **Federal Register** of January 12, 2011 (76 FR 2127), the Agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An Agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0233. The approval expires on February 28, 2014. A copy of the supporting statement for this information collection is available on the Internet at *http://www.reginfo.gov/public/do/PRAMain.*

Dated: March 10, 2011.

**Leslie Kux,**

*Acting Assistant Commissioner for Policy.*

[FR Doc. 2011–6091 Filed 3–15–11; 8:45 am]

**BILLING CODE 4160–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

**[Docket No. FDA–2010–N–0122]**

### Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Focus Groups About Drug Products, as Used by the Food and Drug Administration

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing that a collection of information entitled "Focus Groups About Drug Products, as Used by the Food and Drug Administration" has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

**FOR FURTHER INFORMATION CONTACT:** Elizabeth Berbakos, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, 301–796–3792, *Elizabeth.Berbakos@fda.hhs.gov.*

**SUPPLEMENTARY INFORMATION:** In the **Federal Register** of July 9, 2010 (75 FR 39541), the Agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An Agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0677. The approval expires on July 31, 2012. A copy of the supporting statement for this information collection is available on the Internet at *http://www.reginfo.gov/public/do/PRAMain.*

Dated: March 10, 2011.

**Leslie Kux,**

*Acting Assistant Commissioner for Policy.*

[FR Doc. 2011–6093 Filed 3–15–11; 8:45 am]

**BILLING CODE 4160–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

**[Docket No.FDA–2011–N–0085]**

### Agency Information Collection Activities; Proposed Collection; Comment Request; Guidance for Industry: Cooperative Manufacturing Arrangements for Licensed Biologics

**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 the proposed extension of the collection of information concerning the guidance for industry on cooperative manufacturing arrangements for licensed biologics.

**DATES:** Submit either electronic or written comments on the collection of information by May 16, 2011.

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

**FOR FURTHER INFORMATION CONTACT:** Juanmanuel Vilela, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr, PI50–400B, Rockville, MD 20850, 301–796–3794, *Juanmanuel.Vilela@fda.hhs.gov.*

**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