Reconciliation Act of 1990 (Pub. L. 101–508) further amended the Privacy Act regarding protections for such individuals. The Privacy Act, as amended, regulates the use of computer matching by Federal agencies when records in a system of records (SOR) are matched with other Federal, state, or local government records. It requires Federal agencies involved in computer matching programs to:

1. Negotiate written agreements with the other agencies participating in the matching programs;
2. Obtain the Data Integrity Board approval of the match agreements;
3. Furnish detailed reports about matching programs to Congress and OMB;
4. Notify applicants and beneficiaries that the records are subject to matching; and,
5. Verify match findings before reducing, suspending, terminating, or denying an individual's benefits or payments.

This matching program meets the requirements of the Privacy Act of 1974, as amended.

Dated: July 31, 2013.

Michelle Snyder,

Chief Operating Officer, Centers for Medicare & Medicaid Services.

CMS Computer Match No. 2013–12
HHS Computer Match No. 1307
SSA Computer Match No. 1097–1899

NAME:

''Computer Matching Agreement between the Department of Health and Human Services, Centers for Medicare & Medicaid Services and the Social Security Administration for Determining Enrollment or Eligibility for Insurance Affordability Programs under the Patient Protection and Affordable Care Act''.

SECURITY CLASSIFICATION:

Unclassified.

PARTICIPATING AGENCIES:

Department of Health and Human Services (HHS), Centers for Medicare & Medicaid Services (CMS), and the Social Security Administration (SSA).

AUTHORITY FOR CONDUCTING MATCHING PROGRAM:

Sections 1411 and 1413 of the Patient Protection and Affordable Care Act of 2010 (Pub. L. 111–148), as amended by the Health Care and Education Reconciliation Act of 2010 (Pub. L. 111–152) (collectively, the ACA) require the Secretary of HHS to establish a program for determining eligibility for certain Insurance Affordability Programs, certifications of Exemption, and authorize use of secure, electronic interfaces and an on-line system for the verification of eligibility.

PURPOSE(S) OF THE MATCHING PROGRAM:

The purpose of the Computer Matching Agreement (CMA) is to establish the terms, conditions, safeguards, and procedures under which SSA will disclose information to CMS in connection with the administration of Insurance Affordability Programs under the ACA and its implementing regulations. SSA will provide data to CMS and CMS will use SSA data needed to make initial Eligibility Determinations, eligibility Redeterminations and Renewal decisions, including appeal determinations, for Insurance Affordability Programs and certifications of Exemption. Insurance Affordability Programs include:

1. Qualified Health Plan through an Exchange established under the ACA,
2. Advance payments of the premium tax credit and cost sharing reductions,
3. Medicaid,
4. Children's Health Insurance Program, and
5. Basic Health Program.

As set forth in the CMA, SSA will provide CMS the following information when relevant: (1) Social Security number (SSN) verifications, (2) a death indicator, (3) an indicator of a finding of disability by SSA under title II of the Social Security Act, (4) prisoner data, (5) monthly and annual Social Security benefit information under title II of the Social Security Act, (6) quarters of coverage, and (7) confirmation that an allegation of citizenship is consistent with SSA records.

DESCRIPTION OF RECORDS TO BE USED IN THE MATCHING PROGRAM:

The matching program will be conducted with data maintained by CMS in the Health Insurance Exchanges System (HIX), CMS System No. 09–70–0560, as amended, published at 78 FR 8538 (Feb. 6, 2013) and 78 FR 32256 (May 29, 2013).

The matching program will also be conducted with data maintained by SSA in the following SORs:

- Master Files of SSN Holders and SSN Applications, SSA/OEEAS, 60–0058, 75 FR 82121 (December 29, 2010), as amended 78 FR 40542 (July 5, 2013);
- Prisoner Update Processing System (PUPS), SSA/OPB, 60–0269, 64 FR 11076 (March 8, 1999), as amended 72 FR 69723 (December 10, 2007) and 78 FR 40542 (July 5 2013);
- Master Beneficiary Record, SSA/ORSIS, 60–0090, 71 FR 1826 (January 11, 2006), as amended 72 FR 69723 (December 10, 2007) and 78 FR 40542 (July 5, 2013);
- Earnings Recording and Self-Employment Income System, SSA/OEEAS, 60–0059, 71 FR 1819 (January 11, 2006), as amended 78 FR 40542 (July 5, 2013).

INCLUSIVE DATES OF THE MATCH:

The CMP will become effective no sooner than 40 days after the report of the matching program is sent to OMB and Congress, or 30 days after publication in the **Federal Register,** whichever is later. The matching program will continue for 18 months from the effective date and may be extended for an additional 12 months thereafter, if certain conditions are met.

[FR Doc. 2013–19014 Filed 8–6–13; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects

Title: Required Data Elements for Paternity Establishment Affidavits.

OMB No.: 0970–0171.

Description: Section 466(a)(5)(C)(iv) of the Social Security Act (the Act) requires States to develop and use an affidavit for the voluntary acknowledgement of paternity. The affidavit for the voluntary acknowledgement of paternity must include the minimum requirements specified by the Secretary under section 452(a)(7) of the Act. The affidavits will be used by hospitals, birth record agencies, and other entities participating in the voluntary paternity establishment program.

Respondents: State and Tribal IV–D agencies, hospitals, birth record agencies and other entities participating in the voluntary paternity establishment program.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
None	1,113,719	1	0.17	189,332.23

Estimated Total Annual Burden Hours: 189,332.23.

In compliance with the requirements of Section 506(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. 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: ACF Reports Clearance Officer. Email address: *infocollection@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.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2013–19058 Filed 8–6–13; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2012–N–0937]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Clinical Laboratory Improvement Amendments Waiver Applications

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Clinical Laboratory Improvement Amendments Waiver Applications'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Daniel Gittleson, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, 301–796–5156, *Daniel.Gittleson@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: On April 23, 2013, the Agency submitted a proposed collection of information entitled ''Clinical Laboratory Improvement Amendments Waiver Applications'' 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 0910–0598. The approval expires on July 31, 2016. A copy of the supporting statement for this information collection is available on the Internet at *http://www.reginfo.gov/public/do/PRAMain.*

Dated: August 1, 2013.

Leslie Kux,

Assistant Commissioner for Policy.

[FR Doc. 2013–19053 Filed 8–6–13; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2012–D–1092]

Minimizing Risk for Children's Toy Laser Products; Draft Guidance for Industry and Food and Drug Administration Staff; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of the draft guidance entitled ''Minimizing Risk for Children's Toy Laser Products.'' This draft guidance is to inform manufacturers of laser products, FDA headquarters and field personnel, and the public of the Center for Devices and Radiological Health's (CDRH) proposed approach on the safety of toy laser products. This draft guidance is not final nor is it in effect at this time.

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 November 5, 2013.

ADDRESSES: Submit written requests for single copies of the draft guidance document entitled ''Minimizing Risk for Children's Toy Laser Products'' to the Division of Small Manufacturers, International and Consumer Assistance, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 66, rm. 4613, Silver Spring, MD 20993–0002. Send one self-addressed adhesive label to assist that office in processing your request, or fax your request to 301–847–8149. See the **SUPPLEMENTARY INFORMATION** section for information on electronic access to the guidance.

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. Identify comments with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Robert J. Doyle, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 66, Rm. 4672, Silver Spring, MD 20993–0002, 301–796–5863.

I. Background

This draft guidance is to inform manufacturers of laser products, FDA headquarters and field personnel, and the public of CDRH's proposed approach on the safety of children's toy laser products. Lasers with outputs above certain levels that are operated in