emergent situations. The final rule also provided new information regarding the processes and procedures for amendments and extensions for approved waiver plans. *Form Number:* CMS–10383 (OMB Control Number 0938–1389; *Frequency:* Occasionally; *Affected Public:* State Governments; *Number of Respondents:* 19; *Total Annual Responses:* 399; *Total Annual Hours:* 5,549. (For policy questions regarding this collection contact Lina Rashid at 301–492–4193.)

3. *Type of Information Collection Request:* Revision of a currently approved collection; *Title of Information Collection:* Patient Protection and Affordable Care Act; Exchange Functions: Eligibility for Exemptions; *Use:* The data collection and reporting requirements in ''Patient Protection and Affordable Care Act; Exchange Functions: Eligibility for Exemptions; Miscellaneous Minimum Essential Coverage Provisions'' (78 FR 39494 (July 1, 2013)), address federal requirements that states must meet with regard to the Exchange minimum function of performing eligibility determinations and issuing certificates of exemption from the shared responsibility payment. In the final regulation, CMS addresses standards related to eligibility, including the verification and eligibility determination process, eligibility redeterminations, options for states to rely on HHS to make eligibility determinations for certificates of exemption, and reporting. CMS developed four appendices of application materials to illustrate the process applicants use to apply for exemptions from the shared responsibility payment. This information collection requests seeks approval for the requirements associated with the collection of information associated with these four appendices. *Form Number:* CMS–10466 (OMB Control Number 0938–1190; *Frequency:* Annually; *Affected Public:* Individuals and Households—State, Local, or Tribal Governments; *Number of Respondents:* 849; *Total Annual Responses:* 849; *Total Annual Hours:* 1,962. (For policy questions regarding this collection contact John Kenna at 301–492–4452.)

Dated: October 12, 2023.

William N. Parham, III,

Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2023–22897 Filed 10–16–23; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for Office of Management and Budget Review; Federal Case Registry (Office of Management and Budget #0970–0421)

AGENCY: Office of Child Support Enforcement, Administration for Children and Families, United States Department of Health and Human Services.

ACTION: Request for public comments.

SUMMARY: The Office of Child Support Enforcement (OCSE), Administration for Children and Families (ACF), is requesting from the Office of Management and Budget (OMB) to extend approval of the Federal Case Registry (FCR) for an additional three years. The current approval expires November 30, 2023. OCSE is proposing minor changes to punctuation, formatting, grammar, clarity, and spacing to enable easier completion of the form.

DATES: *Comments due within 30 days of publication.* OMB must make a decision about 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.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to *www.reginfo.gov/public/do/PRAMain.* Find this particular information collection by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function. You can also obtain copies of the proposed collection of information by emailing *infocollection@acf.hhs.gov.* Identify all emailed requests by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: The FCR is a national database of information pertaining to child support cases processed by state child support agencies, referred to as ''IV–D'' cases, and non-IV–D support orders privately established or modified by courts or tribunals on or after October 1, 1998. FCR information is comprised of child support orders and case information from each State Case Registry (SCR). The FCR automatically compares new SCR submissions to existing FCR information and to wage and employment information in the National Directory of New Hires. The Federal Parent Locator Service notifies state agencies if a IV–D case participant in the state matches a participant in a IV–D or non-IV–D case in another state and supplies any matched wage and employment information. Matches enable state agencies to locate parties that live in different states to establish, modify, or enforce child support obligations; to establish paternity; to enforce state law regarding parental kidnapping; and to establish or enforce child custody or visitation determinations.

The FCR instrument, Appendix G: Input Record Layout, contains minor changes in punctuation, formatting, grammar, clarity, and spacing to enable easier completion of the form.

Respondents: State child support enforcement agencies.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Annual number of responses per respondent	Average burden hours per response	Annual burden hours
Appendix G: Input Transactions Layout	54	406	0.033	730

Estimated Total Annual Burden Hours: 730.

Authority: 42 U.S.C. 653(h); 42 U.S.C. 654a(e); 42 U.S.C. 654a(f)(1).

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2023–22809 Filed 10–16–23; 8:45 am]

BILLING CODE 4184–41–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2020–D–2016]

Policy for Testing of Alcohol (Ethanol) and Isopropyl Alcohol for Methanol; Guidance for Industry; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of availability.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing the availability of a final guidance for industry entitled ''Policy for Testing of Alcohol (Ethanol) and Isopropyl Alcohol for Methanol.'' This guidance is intended to alert pharmaceutical manufacturers and pharmacists in State-licensed pharmacies or Federal facilities who engage in drug compounding to the potential public health hazard of alcohol (ethyl alcohol or ethanol) or isopropyl alcohol contaminated with or substituted with methanol. During the Coronavirus Disease 2019 (COVID–19) public health emergency (PHE), FDA became aware of reports of fatal methanol poisoning of consumers who ingested alcohol-based hand sanitizer products that were manufactured with methanol or methanol-contaminated ethanol. FDA is concerned that other drug products containing ethanol or isopropyl alcohol (pharmaceutical alcohol), which are widely used active ingredients in a variety of drug products, could be similarly vulnerable to methanol contamination. This guidance replaces the guidance for industry entitled ''Policy for Testing Alcohol (Ethanol) and Isopropyl Alcohol for Methanol, Including During the Public Health Emergency (COVID–19)'' published in January 2021.

DATES: The announcement of the guidance is published in the **Federal Register** on October 17, 2023.

ADDRESSES: You may submit either electronic or written comments on Agency guidances at any time as follows:

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *https://www.regulations.gov* will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on *https://www.regulations.gov.*
- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see ''Written/Paper Submissions'' and ''Instructions'').

Written/Paper Submissions

Submit written/paper submissions as follows:

- *Mail/Hand Delivery/Courier (for written/paper submissions):* Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Dockets Management Staff, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in ''Instructions.''

Instructions: All submissions received must include the Docket No. FDA–2020–D–2016 for ''Policy for Testing of Alcohol (Ethanol) and Isopropyl Alcohol for Methanol.'' Received comments will be placed in the docket and, except for those submitted as ''Confidential Submissions,'' publicly viewable at *https://www.regulations.gov* or at the Dockets Management Staff between 9 a.m. and 4 p.m., Monday through Friday, 240–402–7500.

- Confidential Submissions—To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states ''THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION.'' The Agency will review this copy, including the claimed confidential information, in its consideration of comments. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on *https://www.regulations.gov.* Submit both copies to the Dockets Management Staff. If you do not wish your name and contact information to be made publicly available, you can provide this information on the cover sheet and not in the body of your comments and you must identify this information as ''confidential.'' Any information marked as ''confidential'' will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA's posting of comments to public dockets, see 80 FR 56469, September 18, 2015, or access the information at: *https://www.govinfo.gov/content/pkg/FR-2015-09-18/pdf/2015-23389.pdf.*

Docket: For access to the docket to read background documents or the electronic and written/paper comments received, go to *https://www.regulations.gov* and insert the docket number, found in brackets in the heading of this document, into the ''Search'' box and follow the prompts and/or go to the Dockets Management Staff, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852, 240–402–7500.

You may submit comments on any guidance at any time (see 21 CFR 10.115(g)(5)).

Submit written requests for single copies of the guidance to the Division of Drug Information, Center for Drug Evaluation and Research, Food and Drug Administration, 10001 New Hampshire Ave., Hillandale Building, 4th Floor, Silver Spring, MD 20993–0002; or to the Office of Communication, Outreach and Development, Center for Biologics Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 71, Rm. 3128, Silver Spring, MD 20993–0002; or to Policy and Regulations Staff, HFV–6, Center for Veterinary Medicine, Food and Drug Administration, 7500 Standish Place, 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 guidance document.

FOR FURTHER INFORMATION CONTACT: Francis Godwin, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 51, Rm. 4342, Silver Spring, MD 20993–0002, 301–796–5362; or Anne Taylor, Center for Biologics Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 71, Rm. 7301, Silver Spring, MD 20993–0002,