

OMB control number: 7100–NEW.
Frequency: Quantitative Survey, annually; Qualitative Survey, occasionally.

Respondents: Consumers.

Estimated number of respondents: Quantitative Survey, 17,000 respondents; Qualitative Survey, 30 respondents.

Estimated average hours per response: Quantitative Survey, 0.47 hours; Qualitative Survey, 2 hours.

Estimated annual burden hours: Quantitative Survey, 7,990 hours; Qualitative Survey, 180 hours.

General description of report: The information collected could be used for the Board's *Report on the Economic Well-Being of U.S. Households*, for Board studies or working papers, professional journals, the *Federal Reserve Bulletin*, testimony and reports to the Congress, or other vehicles. Such event-driven consumer data collections could also be used to inform Board policy, regulatory, supervisory and operational decisions.

The Board anticipates that the SHED would include such topics as individuals' overall financial well-being, employment experiences, income and savings behaviors, economic preparedness, access to banking and credit, housing and living arrangement decisions, education and human capital, student loans, and retirement planning. The overall content of the SHED instrument would depend on changing economic, regulatory, or legislative developments as well as changes in the financial services industry.

Legal authorization and confidentiality: The Board's Legal Division has determined that Section 2A of the Federal Reserve Act (FRA) requires that the Federal Reserve Board and the Federal Open Market Committee maintain long run growth of the monetary and credit aggregates commensurate with the economy's long run potential to increase production, so as to promote effectively the goals of maximum employment, stable prices, and moderate long-term interest rates (12 U.S.C. 225a). Under section 12A of the FRA, the Federal Open Market Committee is required to implement regulations relating to the open market operations conducted by Federal Reserve Banks with a view to accommodating commerce and business and with regard to their bearing upon the general credit situation of the country (12 U.S.C. 263). Because the Board and the Federal Open Market Committee use the information obtained on the FR 3077 to fulfill these obligations, these statutory provisions provide the legal authorization for the

collection of information on the FR 3077. The FR 3077 is a voluntary survey.

The ability of the Board to maintain the confidentiality of information provided by respondents to the FR 3077 will have to be determined on a case-by-case basis depending on the type of information provided for a particular survey. To the extent that a respondent's answers reveal information "the disclosure of which would constitute a clearly unwarranted invasion of personal privacy," such information would likely be exempt from disclosure under exemption 6 of the Freedom of Information Act, (5 U.S.C. 552(b)(6)).

Consultation outside the agency: The Board will consult with outside subject matter experts for specific questions, as needed.

Board of Governors of the Federal Reserve System, August 4, 2017.

Margaret McCloskey Shanks,

Deputy Secretary of the Board.

[FR Doc. 2017–16773 Filed 8–8–17; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Docket No. CDC–2017–0068, NIOSH–299]

Draft National Occupational Research Agenda for Cancer, Reproductive, Cardiovascular and Other Chronic Disease Prevention

AGENCY: National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Request for comments.

SUMMARY: As steward of the National Occupational Research Agenda (NORA), the National Institute for Occupational Safety and Health of the Centers for Disease Control and Prevention announces the availability of the draft National Occupational Research Agenda for Cancer, Reproductive, Cardiovascular and Other Chronic Disease Prevention Agenda for public comment. Written by the NORA Cancer, Reproductive, Cardiovascular and Other Chronic Disease Prevention Cross-Sector Council, the Agenda identifies the most important occupational safety and health research needs for the next decade, 2016–2026. A copy of the draft Agenda is available at <http://www.regulations.gov> (search Docket Number CDC–2017–0068).

DATES: Electronic or written comments must be received by October 10, 2017.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2017–0068 and docket number NIOSH–299, by any of the following methods:

- *Federal eRulemaking Portal:* <http://www.regulations.gov>. Follow the instructions for submitting comments.

- *Mail:* National Institute for Occupational Safety and Health, NIOSH Docket Office, 1090 Tusculum Avenue, MS C–34, Cincinnati, Ohio 45226–1998.

Instructions: All submissions received must include the agency name and Docket Number [CDC–2017–0068; NIOSH–299]. All relevant comments received will be posted without change to <http://www.regulations.gov>, including any personal information provided. For access to the docket to read background documents or comments received, go to <http://www.regulations.gov>.

FOR FURTHER INFORMATION CONTACT:

Emily Novicki (*NORACoordinator@cdc.gov*), National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention, Mailstop E–20, 1600 Clifton Road NE., Atlanta, GA 30329.

SUPPLEMENTARY INFORMATION: The National Occupational Research Agenda (NORA) is a partnership program created to stimulate innovative research and improved workplace practices. The national agenda is developed and implemented through the NORA sector and cross-sector councils. Each council develops and maintains an agenda for its sector or cross-sector. The National Occupational Research Agenda for Cancer, Reproductive, Cardiovascular and other Chronic Disease Prevention (CRC) is intended to identify the research, information, and actions most urgently needed to prevent occupational cancer, adverse reproductive outcomes and cardiovascular disease. The National Occupational Research Agenda for CRC provides a vehicle for industry stakeholders to describe the most relevant issues, gaps, and safety and health needs for the sector. It is meant to be broader than any one agency or organization. It is a strategic plan for the entire country and all of its research and development entities, whether government, higher education, or industry.

This is the first CRC Agenda, developed for the third decade of NORA (2016–2026). The agenda was developed considering new information about injuries and illnesses, the state of the science, and the probability that new information and approaches will make a difference.

As the steward of the NORA process, NIOSH invites comments on the draft National Occupational Research Agenda for CRC. A copy of the draft Agenda is available at <http://www.regulations.gov> (see Docket Number CDC-2017-0068, NIOSH-299).

John Howard,

Director, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention.

[FR Doc. 2017-16801 Filed 8-8-17; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2017-D-0829]

Expiration Dating of Unit-Dose Repackaged Solid Oral Dosage Form Drug Products; Revised Draft 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 revised draft guidance for industry entitled “Expiration Dating of Unit-Dose Repackaged Solid Oral Dosage Form Drug Products.” The last few decades have seen an increasing demand in various health care settings for solid oral dosage form drug products repackaged into unit-dose containers, which hold a quantity of drug for administration as a single dose. The increase in unit-dose repackaging has led to questions regarding stability studies and appropriate expiration dates for these repackaged products. This revised draft guidance describes the conditions under which FDA does not intend to take action regarding required stability studies for these repackaged products and the expiration date to assign under those conditions. Through this notice, FDA is hoping to decrease the regulatory burdens of drug regulations on manufacturers of these products, while at the same time ensuring patient safety. Since FDA’s guidance documents do not bind the public or FDA to any requirements, they have not been considered to be subject to Executive Order 12866.

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 revised draft guidance before it begins work on the final version of the guidance, submit

either electronic or written comments on the draft guidance by October 10, 2017.

ADDRESSES: You may submit comments 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-2017-D-0829 for “Expiration Dating of Unit-Dose Repackaged Solid Oral Dosage Form Drug Products.” 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.

- **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.gpo.gov/fdsys/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.

Submit written requests for single copies of the revised draft 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. Send one self-addressed adhesive label to assist that office in processing your requests. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the revised draft guidance document.

FOR FURTHER INFORMATION CONTACT: Bill Harvey, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 51, Rm. 4214, Silver Spring, MD 20993-0002, 240-402-4180.

SUPPLEMENTARY INFORMATION:

I. Background

FDA is announcing the availability of a revised draft guidance for industry