umbilicals, and other equipment prior to disconnection. No releases or discharges of fluid from pipelines, umbilicals, and/or other equipment that have not been fully flushed prior to being disconnected or cut from the facility are authorized under this NPDES permit.

16. Calculation for WET critical dilutions and testing frequencies is based on calendar year.

17. Waiver for the minimum number of samples to be collected for WET tests, should the effluent cease discharging for produced water.

18. For Treatment, Completion, and Workover discharges, acute WET results can be derived from chronic WET test.

19. Compliance schedule for WET acute limits related to Treatment, Completion, and Workover discharges and sample holding time of 72 hours.

20. No approved Alternative Test Procedure (ATP) for WET, however they can be requested at any time following 40 CFR 136.5.

21. 72 hour hold time for WET samples for Chemically Treated Miscellaneous Discharges.

22. For Chemically Treated Miscellaneous Discharges, non-continuous discharges are discharges that occur less than or equal to once per week and last less than 24 hours. These discharges shall be monitored once per discharge.

23. State general permit or state individual permit may be required in addition to authorization under this permit.

24. Defines decommissioning and Subsea Cleaning Fluids.

25. 7-day chronic toxicity requirements for Well Treatment Fluids, Completion Fluids, and Workover Fluids has been moved from limitations to monitoring section, to provide clarity that chronic is monitoring only.

26. Free oil language has been updated to reference DMRs and twenty-four hour reporting requirements.

27. Part I.C. reflects Other Limitations, Prohibitions and Discharges not Authorized. Moved Limitations on Coverage section in Part I.A.1 to Part I.C for Prohibitions and Discharges Not Authorized.

28. Permit does not authorize radioactive materials that are under the jurisdiction of the NRC.

29. Miscellaneous Discharges of Water Which Have Been Chemically Treated includes discharges from well operations other than those covered by other sections of Part I.B of the permit.

30. Corrections to the Permit Summary Table. Table is for reference only.

31. Corrected data for Discharge Monitoring Reports (DMRs) and Other Reports must be submitted as soon as the error has been identified but no later than the following quarter. Submittal of corrected data does not excuse any permit violation.

32. If Offshore 24-Hour Reporting Application Portal is not available, an email shall be sent within 24 hours of occurrence of specified violations and electronic report shall be submitted within 14 days of the system becoming available.

33. A facility map that delineates authorized discharge locations and type must be submitted, as an attachment, when filing the eNOI.

34. Language has been updated to specify that new operators are not eligible for coverage and existing operators may not submit new NOI's during the administraive continued period.

35. Updated language to provide clarity that timely updates to ''CDX'' are required, in lieu of ''eNOI.''

36. Numeric exceedances of maximum through-screen design intake velocity and dates must also be included on DMRs, for all new facilities required to comply with intake structure monitoring requirements.

37. Definition of Mobile Offshore Drilling Unit (MODU) has been removed from the permit because it does not exist in the Code of Federal Regulations. Part I.A.2 has been updated to provide examples of MODUs.

38. Civil and administrative penalty amounts have been updated to reflect updated statutory amounts.

39. Once a month temperature monitoring for produced water.

II. Other Legal Requirements

Other statutory and regulatory requirements are discussed in the fact sheet that include: Oil Spill Requirement; Ocean Discharge Criteria Evaluation; Marine Protection, Research, and Sanctuaries Act; National Environmental Policy Act; Magnuson-Stevens Fisheries Conservation and Management Act; Endangered Species Act; State Water Quality Standards and State Certification; Coastal Zone Management Act; and Paperwork Reduction Act.

Under Executive Order 12866 (58 FR 51735, October 4, 1993), this action is a ''significant regulatory action.'' Accordingly, EPA submitted this action to the Office of Management and Budget (OMB) for review under Executive Orders 12866 and 13563 (76 FR 3821, January 21, 2011) and any changes made in response to OMB recommendations have been documented in the docket for this action.

Charles W. Maguire,

Director, Water Division, EPA Region 6.

[FR Doc. 2023–11770 Filed 6–1–23; 8:45 am]

BILLING CODE 6560–50–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Agency for Healthcare Research and Quality, HHS.

ACTION: Notice

SUMMARY: This notice announces the intention of the Agency for Healthcare Research and Quality (AHRQ) to request that the Office of Management and Budget (OMB) approve the proposed information collection project ''AHRQ Research Reporting System (ARRS).'' This proposed information collection was previously published in the **Federal Register** on March 29th, 2023 and allowed 60 days for public comment. AHRQ received no substantive comments from members of the public. The purpose of this notice is to allow an additional 30 days for public comment.

DATES: Comments on this notice must be received by July 3, 2023.

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.

Copies of the proposed collection plans, data collection instruments, and specific details on the estimated burden can be obtained from the AHRQ Reports Clearance Officer.

FOR FURTHER INFORMATION CONTACT: Doris Lefkowitz, AHRQ Reports Clearance Officer, (301) 427–1477, or by email at *doris.lefkowitz@AHRQ.hhs.gov.*

SUPPLEMENTARY INFORMATION:

Proposed Project

AHRQ Research Reporting System (ARRS)

AHRQ has developed a systematic method for its grantees to report project progress and important preliminary findings for grants funded by the

Agency. This system, the AHRQ Research Reporting System (ARRS), previously known as the Grants Reporting System (GRS), was last approved by OMB on August 31, 2020. The system addressed the shortfalls in the previous reporting process and established a consistent and comprehensive reporting solution for grants in AHRQ. The ARRS provides a centralized repository of grants research progress and additional information that can be used to support initiatives within the Agency. This includes future research planning and support for administrative activities such as performance monitoring, budgeting, and knowledge transfer, as well as for strategic planning.

This Project has the following goals:

(1) To promote the transfer of critical information more frequently and efficiently and enhance the Agency's ability to support research designed to improve the outcomes and quality of health care, reduce its costs, and broaden access to effective services

(2) To increase the efficiency of the Agency in responding to ad-hoc information requests

(3) To support Executive Branch requirements for increased transparency and public reporting

(4) To establish a consistent approach throughout the Agency for information collection regarding grant progress and a systematic basis for oversight and for facilitating potential collaborations among grantees

(5) To decrease the inconvenience and burden on grantees of unanticipated ad-hoc requests for information by the Agency in response to particular (one-time) internal and external requests for information

This project is being conducted by AHRQ through its contractor, Science Applications International Corporation, Inc, pursuant to AHRQ's statutory authority to conduct and support research on health care and on systems for the delivery of such care, including activities with respect to the quality, effectiveness, efficiency, appropriateness, and value of healthcare services and with respect to quality measurement and improvement. 42 U.S.C 299a(a)(1) and (2).

Method of Collection

Grantees use the ARRS system to report project progress and important preliminary findings for grants funded by the Agency. Grantees submit progress reports on a monthly or quarterly basis, which are reviewed by AHRQ personnel. All users access the ARRS system through a secure online interface which requires a user id and password entered through the ARRS Login screen. When status reports are due AHRQ notifies Principal Investigators (PI) via email.

The ARRS is an automated user-friendly resource that is utilized by AHRQ staff for preparing, distributing, and reviewing reporting requests to grantees for the purpose of information sharing. AHRQ personnel are able to systematically search the information collected and stored in the ARRS database. Personnel will also use the information to address internal and/or external requests for information regarding grant progress, preliminary findings, and other requests, such as Freedom of Information Act requests, and producing responses related to federally mandated programs and regulations.

Estimated Annual Respondent Burden

Exhibit 1 shows the estimated annualized burden hours for the respondents. It will take grantees an estimated 15 minutes to enter the necessary data into the ARRS System. Frequency of reporting varies from monthly to once a year. The total number of responses submitted for the past year is considered for this estimation. Based on that, the total annualized burden hours are estimated to be 125 hours.

Exhibit 2 shows the estimated annualized cost burden for the respondents. The total estimated cost burden for respondents is $5,475.

EXHIBIT 1—ESTIMATED ANNUALIZED BURDEN HOURS

Form name	Number of responses	Hours per response	Total burden hours
Data entry into ARRS	500	15/60	125
Total	500	N/A	125

EXHIBIT 2—ESTIMATED ANNUALIZED COST BURDEN

Form name	Number of responses	Total burden hours	Average hourly wage rate *	Total cost burden
Data entry into ARRS	500	125	$43.80	$5,475
Total	500	125	N/A	$5,475

* Based upon the average wages for Healthcare Practitioner and Technical Occupations (29–0000), ''National Compensation Survey: Occupational Wages in the United States, May 2021,'' U.S. Department of Labor, Bureau of Labor Statistics, *http://www.bls.gov/oes/current/oes_nat.htm#29-0000.*

Request for Comments

In accordance with the Paperwork Reduction Act, 44 U.S.C. 3501–3520, comments on AHRQ's information collection are requested with regard to any of the following: (a) whether the proposed collection of information is necessary for the proper performance of AHRQ's health care research and health care information dissemination functions, including whether the information will have practical utility; (b) the accuracy of AHRQ's estimate of burden (including hours and costs) of the proposed collection(s) of information; (c) ways to enhance the quality, utility and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information upon the respondents, including the use of automated collection techniques or other forms of information technology.

Comments submitted in response to this notice will be summarized and included in the Agency's subsequent request for OMB approval of the

proposed information collection. All comments will become a matter of public record.

Dated: May 26, 2023.

Marquita Cullom,

Associate Director.

[FR Doc. 2023–11723 Filed 6–1–23; 8:45 am]

BILLING CODE 4160–90–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2012–D–0322]

Action Level for Inorganic Arsenic in Apple Juice: Guidance for Industry; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of availability.

SUMMARY: The Food and Drug Administration (FDA or we) is announcing the availability of a final guidance for industry entitled ''Action Level for Inorganic Arsenic in Apple Juice.'' The guidance identifies for industry an action level for inorganic arsenic in apple juice that is intended to help protect public health by reducing exposure to inorganic arsenic and is achievable with the use of current good manufacturing practices. It also describes our intended sampling and enforcement approach. Thus, the guidance finalizes the approach presented in the draft guidance issued in 2013.

DATES: The announcement of the guidance is published in the **Federal Register** on June 2, 2023.

ADDRESSES: You may submit either electronic or written comments on FDA 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–2012–D–0322 for ''Action Level for Inorganic Arsenic in Apple Juice: Guidance for Industry.'' 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.'' We will review this copy, including the claimed confidential information, in our 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 Plant Products and Beverages, Office of Food Safety, Center for Food Safety and Applied Nutrition (HFS–317), Food and Drug Administration, 5001 Campus Dr., College Park, MD 20740. Send two self-addressed adhesive labels to assist that office in processing your request. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the guidance.

FOR FURTHER INFORMATION CONTACT: Eileen Abt, Center for Food Safety and Applied Nutrition (HFS–317), Food and Drug Administration, 5001 Campus Dr., College Park, MD 20740, 240–402–1529; or Denise See, Center for Food Safety and Applied Nutrition, Office of Regulations and Policy (HFS–024), Food and Drug Administration, 5001 Campus Dr., College Park, MD 20740, 240–402–2378.

SUPPLEMENTARY INFORMATION:

I. Background

We are announcing the availability of a guidance for industry entitled ''Action Level for Inorganic Arsenic in Apple Juice.'' We are issuing this guidance consistent with our good guidance practices regulation (21 CFR 10.115). The guidance represents the current thinking of FDA on this topic. It does not establish any rights for any person and is not binding on FDA or the public. You can use an alternative approach if it satisfies the requirements of the applicable statutes and regulations.

In the **Federal Register** of July 15, 2013 (78 FR 42086), we announced the availability of a draft guidance for industry entitled ''Arsenic in Apple Juice: Action Level.'' We also announced the availability of two related scientific documents: a document entitled ''Supporting Document for Action Level for Arsenic in Apple Juice'' (supporting document),