Page 10 – Sharon Young, Cue Health, Inc.

V. Duration of Authorization

This EUA will be effective until the declaration that circumstances exist justifying the authorization of the emergency use of in vitro diagnostics for detection and/or diagnosis of infection with the monkeypox virus, including in vitro diagnostics that detect and/or diagnose infection with non-variola *Orthopoxvirus*, is terminated under Section 564(b)(2) of the Act or the EUA is revoked under Section 564(g) of the Act.

Sincerely,

//s//

Jeffrey E. Shuren, M.D., J.D.
Director
Center for Devices and Radiological Health
Food and Drug Administration

Enclosure

Dated: April 12, 2023.

Lauren K. Roth,

Associate Commissioner for Policy.

[FR Doc. 2023–08023 Filed 4–14–23; 8:45 am]

BILLING CODE 4164–01–C

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Announcement of the Second Meeting of the 2025 Dietary Guidelines Advisory Committee

AGENCY: U.S. Department of Health and Human Services (HHS); Office of the Assistant Secretary for Health (OASH); and U.S. Department of Agriculture (USDA), Food, Nutrition, and Consumer Services (FNCS).

ACTION: Notice.

SUMMARY: The Departments of Health and Human Services and Agriculture announce the second meeting of the 2025 Dietary Guidelines Advisory Committee (Committee). This meeting will be open to the public virtually.

DATES: The second meeting of the 2025 Dietary Guidelines Advisory Committee will be held on May 10, 2023, 9 a.m. to 3:30 p.m. ET.

ADDRESSES: The meeting will be accessible online via livestream and recorded for later viewing. Registrants will receive the livestream information prior to the meeting.

SUPPLEMENTARY INFORMATION:

Authority and Purpose: Under Section 301 of Public Law 101–445 (7 U.S.C. 5341, the National Nutrition Monitoring and Related Research Act of 1990, Title III), the Secretaries of HHS and USDA are directed to publish the *Dietary Guidelines for Americans* jointly at least every five years. See 88 FR 3423, January 19, 2023, for notice of the first meeting of the 2025 Dietary Guidelines Advisory Committee, the complete Authority and Purpose, and the Committee's Task. The 2025 Dietary Guidelines Advisory Committee is formed and governed under the provisions of the Federal Advisory Committee Act (FACA), Public Law 92–463, as amended (5 U.S.C., App).

Purpose of the Meeting: The Committee will meet to discuss their prioritization of the scientific questions proposed by the Departments (see *www.DietaryGuidelines.gov*) and share draft plans for their review of the scientific evidence. In accordance with FACA, deliberations of the Committee will occur in a public forum.

Meeting Agendas: A detailed agenda will be announced in advance of the meeting at *www.DietaryGuidelines.gov.* The agenda will include presentations by each subcommittee and deliberation by the full Committee regarding the prioritization of scientific questions and initial draft protocol development and discussion of plans for future Committee work.

Public Comment: Public comments to the Committee opened on January 19, 2023 and will remain open throughout the Committee's deliberations. Comments may be submitted at *https://www.regulations.gov/document/HHS-OASH-2022-0021-0001.*

Meeting Registration: This Committee meeting is open to the public. The meeting will be accessible online via livestream and recorded for later viewing. Registration is required for the livestream. To register, go to *www.DietaryGuidelines.gov* and click on the link for ''Meeting Registration.''

Closed captioning will be available to all participants. Individuals who need accommodations should contact Kara Beckman (*Kara.Beckman@hhs.gov*). Requests should be made at least five business days in advance of the meeting.

Meeting materials for each meeting will be accessible at *www.DietaryGuidelines.gov.* Materials may be requested by email at *dietaryguidelines@hhs.gov.*

FOR FURTHER INFORMATION CONTACT: Designated Federal Officer, 2025 Dietary Guidelines Advisory Committee, Janet M. de Jesus, MS, RD; HHS/OASH/ODPHP, 1101 Wootton Parkway, Suite 420, Rockville, MD 20852; Phone: 240–453–8266; Email *DietaryGuidelines@hhs.gov.* Additional information is

available on the internet at *www.DietaryGuidelines.gov.*

Paul Reed,

Deputy Assistant Secretary for Health, Office of Disease Prevention and Health Promotion.

[FR Doc. 2023–08081 Filed 4–14–23; 8:45 am]

BILLING CODE 4150–32–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

[Document Identifier OS–0990–0323]

Agency Information Collection Request; 30-Day Public Comment Request

AGENCY: Office of the Secretary, HHS.

ACTION: Notice.

SUMMARY: In compliance with the requirement of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, is publishing the following summary of a proposed collection for public comment.

DATES: Comments on the ICR must be received on or before May 17, 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.

FOR FURTHER INFORMATION CONTACT: Sherrette Funn, *Sherrette.Funn@hhs.gov* or (202) 264–0041, or *PRA@HHS.GOV.* When submitting comments or requesting information, please include the document identifier 0990–0323–30D and project title for reference.

SUPPLEMENTARY INFORMATION: Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Title of the Collection: MedicalCountermeasures.gov.

Type of Collection: Reinstatement without chg.

OMB No.: 0990–0323.

Abstract: Department of Health and Human Services, Administration for Strategic Preparedness and Response (ASPR).

The USG seeks information from stakeholders on available medical countermeasures in development, with a particular interest in products, technologies, and capabilities that have progressed into or beyond clinical trials, have established large-scale cGMP manufacturing capability, or utilize an approved platform. Information regarding diagnostics, therapeutics, vaccines, and other products, technologies, or capabilities relevant to respond to public health emergencies are sought. The TechWatch program, run by ASPR/BARDA, provides the *Medicalcountermeasures.gov bdr.hhs.gov* portal as a single point of entry for the submission of meeting requests from interested stakeholders with promising MCM products, technologies, and capabilities.

The information collection request is seeking OMB approval for a three (3) year duration. It is expected that any given responded would submit TechWatch meeting requests no more than annually, based on program history. Developers of medical countermeasures respondents will submit a response once and never submit subsequent requests.

ESTIMATED ANNUALIZED BURDEN TABLE

Type of respondent	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours
Developers of medical countermeasures addressing naturally occurring and intentional public health threats	350	1	8/60	47
Total	350			47

Sherrette A. Funn,

Paperwork Reduction Act Reports Clearance Officer, Office of the Secretary.

[FR Doc. 2023–08075 Filed 4–14–23; 8:45 am]

BILLING CODE 4150–37–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

[Document Identifier: OS–0990–0459]

Agency Father Generic Information Collection Request; 60-Day Public Comment Request

AGENCY: Office of the Secretary, HHS.

ACTION: Notice.

SUMMARY: In compliance with the requirement of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, is publishing the following summary of a proposed collection for public comment.

DATES: Comments on the ICR must be received on or before June 16, 2023.

ADDRESSES: Submit your comments to *Sherrette.Funn@hhs.gov* or *PRA@hhs.gov* by calling (202) 264–0041.

FOR FURTHER INFORMATION CONTACT: When submitting comments or requesting information, please include the document identifier 0990–New–60D, and project title for reference, to Sherrette Funn, the Reports Clearance Officer, *Sherrette.funn@hhs.gov,* or call 202–264–0041.

SUPPLEMENTARY INFORMATION: Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Title of the Collection: Fast-Track Generic Clearance for the Collection of Routine Customer Feedback on HHS Communications.

Type of Collection: Father Generic ICR.

OMB No.: 0990–0459—ASPA.