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: *http://www.fda.gov/regulatoryinformation/dockets/default.htm.*

Docket: For access to the docket to read background documents or the electronic and written/paper comments received, go to *http://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 Division of Dockets Management, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Philip L. Chao, Center for Food Safety and Applied Nutrition (HFS–24), Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740, 240–402–2378.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of April 6, 2016 (81 FR 19976), FDA published a notice announcing the availability of a draft guidance for industry entitled ''Inorganic Arsenic in Rice Cereals for Infants: Action Level,'' a supporting document entitled ''Supporting Document for Action Level for Inorganic Arsenic in Rice Cereals for Infants,'' and a risk assessment report entitled ''Arsenic in Rice and Rice Products Risk Assessment: Report.'' Although you can comment on any guidance at any time, to ensure that we consider comments on this draft guidance before we begin work on the final version, interested persons were originally given until July 5, 2016, to comment on the draft guidance, the supporting document, or the risk assessment report. In early July 2016, the Federal eRulemaking Portal, *http://www.regulations.gov,* is undergoing maintenance. We are, therefore, extending the comment period for the draft guidance, the supporting document, and the risk assessment report. The extended comment period will close on July 19, 2016.

Dated: June 24, 2016.

Leslie Kux,

Associate Commissioner for Policy.

[FR Doc. 2016–15478 Filed 6–29–16; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2016–N–0321]

Risk Assessment of Foodborne Illness Associated With Pathogens From Produce Grown in Fields Amended With Untreated Biological Soil Amendments of Animal Origin; Request for Scientific Data, Information, and Comments; Extension of Comment Period

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; extension of comment period.

SUMMARY: The Food and Drug Administration (FDA or we) is extending the comment period for the notice entitled ''Risk Assessment of Foodborne Illness Associated With Pathogens From Produce Grown in Fields Amended With Untreated Biological Soil Amendments of Animal Origin; Request for Scientific Data, Information, and Comments'' that appeared in the **Federal Register** of March 4, 2016. The notice requested scientific data, information, and comments that would assist in the development of a risk assessment for produce grown in fields or other growing areas amended with untreated biological soil amendments of animal origin (including raw manure). In the **Federal Register** notice of April 22, 2016, the comment period for this request was initially extended to July 5, 2016. We are taking this action due to maintenance on the Federal eRulemaking portal in early July 2016.

DATES: FDA is extending the comment period on the notice published March 4, 2016 (81 FR 11572). Submit either electronic or written comments by July 19, 2016.

ADDRESSES: You may submit comments as follows:

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal: http://www.regulations.gov.* Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *http://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 *http://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):* Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Division of Dockets Management, 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–2016–N–0321 for ''Risk Assessment of Foodborne Illness Associated With Pathogens From Produce Grown in Fields Amended With Untreated Biological Soil Amendments of Animal Origin; Request for Scientific Data, Information, and Comments.'' Received comments will be placed in the docket and, except for those submitted as ''Confidential Submissions,'' publicly viewable at *http://www.regulations.gov* or at the Division of Dockets Management 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 *http://www.regulations.gov.* Submit both copies to the Division of Dockets Management. 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: *http://www.fda.gov/regulatoryinformation/dockets/default.htm.*

Docket: For access to the docket to read background documents or the electronic and written/paper comments received, go to *http://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 Division of Dockets Management, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Jane Van Doren, Center for Food Safety and Applied Nutrition (HFS–005), Food and Drug Administration, 5100 Campus Dr., College Park, MD 20740, 240–402–2927.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of April 22, 2016 (81 FR 23733), FDA published a notice giving interested persons until July 5, 2016, to comment on our request for scientific data, information, and comments that would assist us in our plan to develop a risk assessment for produce grown in fields or other growing areas amended with untreated biological soil amendments of animal origin (including raw manure).

In early July 2016, the Federal eRulemaking Portal, *http://www.regulations.gov,* is undergoing maintenance. We are, therefore, extending the comment period for our request for scientific data, information, and comments. The extended comment period will close on July 19, 2016.

Dated: June 24, 2016.

Leslie Kux,

Associate Commissioner for Policy.

[FR Doc. 2016–15480 Filed 6–29–16; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

Administration for Children and Families

Delegation of Authority

AGENCY: Office of the Secretary, Administration for Children and Families, HHS.

ACTION: Delegation of authority.

SUMMARY: Notice is hereby given that I delegate to the Assistant Secretary for the Administration for Children and Families (ACF) the following authorities vested in the Secretary of Health and Human Services under the Trafficking Victims Protection Act of 2000 (TVPA), Public Law 106–386, as amended.

Authority under section 107(b)(1)(B)(i) of the TVPA (22 U.S.C. 7105(b)(1)(B)(i)) to expand benefits and services to victims of severe forms of trafficking in persons in the United States, without regard to immigration status. In the case of non-entitlement programs funded by the Secretary of Health and Human Services, such benefits and services may include services to assist potential victims of trafficking in achieving certification and to assist minor dependent children of victims of severe forms of trafficking in persons or potential victims of trafficking.

Authority under section 107(b)(1)(B)(ii) of the TVPA (22 U.S.C. 7105(b)(1)(B)(ii)) to make grants for a national communication system to assist victims of severe forms of trafficking in persons in communicating with service providers.

Authority under section 107(f) of the TVPA (22 U.S.C. 7105(f)) to establish a program to assist United States citizens and aliens lawfully admitted for permanent residence who are victims of severe forms of trafficking. In addition to the authority to provide such victims with specialized services, the program also has the authority to identify current providers and provide a means to make referrals to programs for which such victims are already eligible. In the course of exercising the authority to conduct activities, personnel in the Administration for Children and Families will consult with the Attorney General, the Secretary of Labor, and non-governmental organizations that provide services to victims of severe forms of trafficking in the United States.

These authorities may be redelegated.

These authorities shall be exercised under the Department's policy on regulations and the existing delegation of authority to approve and issue regulations.

These delegations shall be exercised under financial and administrative requirements applicable to the Administration for Children and Families authorities.

I hereby affirm and ratify any actions taken by the Assistant Secretary for Children and Families, or your subordinates, which involved the exercise of these authorities delegated herein prior to the effective date of this delegation.

This delegation supersedes all existing delegations of these authorities.

DATES: This delegation is effective upon signature.

Dated: June 21, 2016.

Sylvia M. Burwell,

Secretary.

[FR Doc. 2016–15470 Filed 6–29–16; 8:45 am]

BILLING CODE 4184–34–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Mandatory Guidelines for Federal Workplace Drug Testing Programs

AGENCY: Substance Abuse and Mental Health Services Administration (SAMHSA), Department of Health and Human Services (HHS).

ACTION: HHS approval of entities that certify Medical Review Officers (MRO).

SUMMARY: The current version of the Department of Health and Human Services (HHS) Mandatory Guidelines for Federal Workplace Drug Testing Programs (Mandatory Guidelines), effective on October 1, 2010, addresses the role and qualifications of Medical Review Officers (MROs) and HHS approval of entities that certify MROs.

DATES: HHS approval is effective June 30, 2016.

FOR FURTHER INFORMATION CONTACT: Jennifer Fan, Pharm.D., J.D., Division of Workplace Programs (DWP), Center for Substance Abuse Prevention (CSAP), Substance Abuse and Mental Health Services Administration (SAMHSA), 5600 Fishers Lane, Room 16N02B, Rockville, MD 20857; Telephone: (240) 276–1759; Email: *jennifer.fan@samhsa.hhs.gov.*

SUPPLEMENTARY INFORMATION: Subpart M-Medical Review Officer (MRO), section 13.1(b) of the Mandatory Guidelines, ''Who may serve as an MRO?'' states as follows: ''Nationally recognized entities that certify MROs or subspecialty boards for physicians performing a review of Federal