

Drug Administration, 1350 Piccard Dr., PI50-400T, Rockville, MD 20850, [Domini.Bean@fda.hhs.gov](mailto:Domini.Bean@fda.hhs.gov).

**SUPPLEMENTARY INFORMATION:**

**I. Background**

In the **Federal Register** of July 29, 2013 (78 FR 45782), we published a proposed rule entitled “Accreditation of Third-Party Auditors/Certification Bodies to Conduct Food Safety Audits and to Issue Certifications” with a 120-day comment period on the provisions of the proposed rule and on the information collection provisions that are subject to review by OMB under the PRA (44 U.S.C. 3501-3520).

FDA has received requests for an extension of the comment period on the proposed rule to allow interested persons an opportunity to consider the interrelationship between this proposed rule and the proposed rule entitled “Current Good Manufacturing Practice and Hazard Analysis and Risk-Based Preventive Controls for Food for Animals” (78 FR 64736, October 29, 2013). FDA has considered the requests and is granting a 60-day extension of the comment period for the “Accreditation of Third-Party Auditors/Certification Bodies to Conduct Food Safety Audits and to Issue Certifications” proposed rule to allow interested persons an opportunity to consider the interrelationships between the proposed rules. We also are extending the comment period for the information collection provisions for 60 days to make the comment period for the information collection provisions the same as the comment period for the provisions of the proposed rule. To clarify, FDA is requesting comment on all issues raised by the proposed rule.

**II. Paperwork Reduction Act of 1995**

Interested persons may either submit electronic comments regarding the information collection to [aira\\_submission@omb.eop.gov](mailto:aira_submission@omb.eop.gov) or fax written comments to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, FAX: 202-395-7285. All comments should be identified with the title “Accreditation of Third-Party Auditors/Certification Bodies to Conduct Food Safety Audits and to Issue Certifications.”

**III. Request for Comments**

Interested persons may submit either electronic comments regarding the proposed rule to <http://www.regulations.gov> or written comments to the Division of Dockets Management (see **ADDRESSES**). It is only necessary to send one set of comments. Identify comments with the docket

number found in brackets in the heading of this document. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday, and will be posted to the docket at <http://www.regulations.gov>.

Dated: November 13, 2013.

**Leslie Kux,**

*Assistant Commissioner for Policy.*

[FR Doc. 2013-27644 Filed 11-19-13; 8:45 am]

**BILLING CODE 4160-01-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

**21 CFR Parts 1, 16, 106, 110, 114, 117, 120, 123, 129, 179, and 211**

**[Docket No. FDA-2011-N-0920]**

**RIN 0910-AG36**

**Current Good Manufacturing Practice and Hazard Analysis and Risk-Based Preventive Controls for Human Food; Extension of Comment Periods**

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Proposed rule; extension of comment period for the proposed rule and for its information collection provisions.

**SUMMARY:** The Food and Drug Administration (FDA) is extending the comment period for the notice of proposed rulemaking that appeared in the **Federal Register** of January 16, 2013 (78 FR 3646), entitled “Current Good Manufacturing Practice and Hazard Analysis and Risk-Based Preventive Controls for Human Food” and its information collection provisions.

**DATES:** The FDA is extending the comment period for the proposed rule referenced in the Summary. Submit either electronic or written comments on the notice of proposed rulemaking by November 22, 2013. Submit comments on information collection issues under the Paperwork Reduction Act of 1995 (the PRA) by November 22, 2013 (see the “Paperwork Reduction Act of 1995” section).

**ADDRESSES:** You may submit comments, identified by Docket No. FDA-2011-N-0920 and/or Regulatory Information Number (RIN) 0910-AG36, by any of the following methods, except that comments on information collection issues under the PRA must be submitted to the Office of Information and Regulatory Affairs, Office of Management and Budget (OMB) (see the

“Paperwork Reduction Act of 1995” section).

**Electronic Submissions**

Submit electronic comments in the following way:

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

**Written Submissions**

Submit written submissions in the following ways:

- *Mail/Hand delivery/Courier (for paper or CD-ROM submissions):* Division of Dockets Management (HFA-305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

*Instructions:* All submissions received must include the Agency name and Docket No. FDA-2011-N-0920, and RIN 0910-AG36 for this rulemaking. All comments received may be posted without change to <http://www.regulations.gov>, including any personal information provided. For additional information on submitting comments, see the “How to Submit Comments” heading of the **SUPPLEMENTARY INFORMATION** section of this document.

*Docket:* For access to the docket to read background documents or 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:**

*With regard to the proposed rule:* Jenny Scott, Center for Food Safety and Applied Nutrition (HFS-300), Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740, 240-402-2166. *With regard to the information collection:* Domini Bean, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50-400T, Rockville, MD 20850, [domini.bean@fda.hhs.gov](mailto:domini.bean@fda.hhs.gov).

**SUPPLEMENTARY INFORMATION:**

**I. Background**

In the **Federal Register** of January 16, 2013 (78 FR 3646), FDA published a proposed rule entitled “Current Good Manufacturing Practice and Hazard Analysis and Risk-Based Preventive Controls for Human Food.” The original comment period of 120 days was extended several times and interested persons were most recently given until November 15, 2013 (**Federal Register** of August 9, 2013, 78 FR 48636), to

comment on the proposed rule and its information collection provisions.

## II. Request for Comments

FDA is extending the comment period due to the inability of some commenters to submit comments through the <http://www.regulations.gov> Web site from November 4, 2013, through November 14, 2013, because of technical difficulties at that Web site.

## III. Paperwork Reduction Act of 1995

Interested persons may either submit electronic comments regarding the information collection to [aira\\_submission@omb.eop.gov](mailto:aira_submission@omb.eop.gov) or fax written comments to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, FAX: 202-395-7285. All comments should be identified with the title "Current Good Manufacturing Practice and Hazard Analysis and Risk-Based Preventive Controls for Human Food."

## IV. How To Submit Comments

Interested persons may submit either electronic comments regarding this document to <http://www.regulations.gov> or written comments to the Division of Dockets Management (see **ADDRESSES**). It is only necessary to send one set of comments. Identify comments with the docket number found in brackets in the heading of this document. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday, and will be posted to the docket at <http://www.regulations.gov>.

Dated: November 15, 2013.

**Leslie Kux,**

*Assistant Commissioner for Policy.*

[FR Doc. 2013-27783 Filed 11-15-13; 4:15 pm]

**BILLING CODE 4160-01-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

#### 21 CFR Parts 16 and 112

[Docket No. FDA-2011-N-0921]

RIN 0910-AG35

### Standards for the Growing, Harvesting, Packing, and Holding of Produce for Human Consumption; Extension of Comment Periods

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Proposed rule; extension of comment period for the proposed rule and for its information collection provisions.

**SUMMARY:** The Food and Drug Administration (FDA) is extending the comment period for the notice of proposed rulemaking that appeared in the **Federal Register** of January 16, 2013 (78 FR 3504), entitled "Standards for the Growing, Harvesting, Packing and Holding of Produce for Human Consumption" and for its information collection provisions.

**DATES:** The FDA is extending the comment period for the proposed rule referenced in the Summary. Submit either electronic or written comments on the notice of proposed rulemaking by November 22, 2013. Submit comments on information collection issues under the Paperwork Reduction Act of 1995 (the PRA) by November 22, 2013 (see the "Paperwork Reduction Act of 1995" section).

**ADDRESSES:** You may submit comments, identified by Docket No. FDA-2011-N-0921 and/or Regulatory Information Number (RIN) 0910-AG35, by any of the following methods, except that comments on information collection issues under the PRA must be submitted to the Office of Information and Regulatory Affairs, Office of Management and Budget (OMB) (see the "Paperwork Reduction Act of 1995" section).

#### Electronic Submissions

Submit electronic comments in the following way:

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

#### Written Submissions

Submit written submissions in the following ways:

- *Mail/Hand delivery/Courier (for paper or CD-ROM submissions):* Division of Dockets Management (HFA-305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

*Instructions:* All submissions received must include the Agency name and Docket No. FDA-2011-N-0921, and RIN 0910-AG35 for this rulemaking. All comments received may be posted without change to <http://www.regulations.gov>, including any personal information provided. For additional information on submitting comments, see the "How to Submit Comments" heading of the **SUPPLEMENTARY INFORMATION** section of this document.

*Docket:* For access to the docket to read background documents or 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:

*With regard to the proposed rule:* Samir Assar, Center for Food Safety and Applied Nutrition (HFS-300), Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740, 240-402-1636. *With regard to the information collection:* Domini Bean, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50-400T, Rockville, MD 20850, [domini.bean@fda.hhs.gov](mailto:domini.bean@fda.hhs.gov).

#### SUPPLEMENTARY INFORMATION:

#### I. Background

In the **Federal Register** of January 16, 2013 (78 FR 3504), FDA published a proposed rule entitled "Standards for the Growing, Harvesting, Packing, and Holding of Produce for Human Consumption." The original comment period of 120 days was extended several times and interested persons were most recently given until November 15, 2013 (**Federal Register** of August 9, 2013, 78 FR 48637), to comment on the proposed rule and its information collection provisions.

#### II. Request for Comments

FDA is extending the comment period due to the inability of some commenters to submit comments through the <http://www.regulations.gov> Web site from November 4, 2013, through November 14, 2013, because of technical difficulties at that Web site.

#### III. Paperwork Reduction Act of 1995

Interested persons may either submit electronic comments regarding the information collection to [aira\\_submission@omb.eop.gov](mailto:aira_submission@omb.eop.gov) or fax written comments to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, FAX: 202-395-7285. All comments should be identified with the title "Standards for the Growing, Harvesting, Packing, and Holding of Produce for Human Consumption."

#### IV. How To Submit Comments

Interested persons may submit either electronic comments regarding the proposed rule to <http://www.regulations.gov> or written comments to the Division of Dockets Management (see **ADDRESSES**). It is only necessary to send one set of comments. Identify comments with the docket number found in brackets in the heading of this document. Received comments may be seen in the Division of Dockets Management between 9 a.m.