with the requirements of this section. The notification:
(i) Must be signed by the chief executive officer, or person in an
equivalent position, and submitted to the Farm Credit Administration as soon
as the Reporting Entity becomes aware of its inability to comply;
(ii) Must explain the reasons for its inability to prepare and submit the
report; and
(iii) May include a request that the Farm Credit Administration extend the
due date for the quarterly report of accounts and exposures.
(7) In the event there is a breach of
information, immediately provide
written notice of the breach to:
(i) The Farm Credit Administration;
and
(ii) Each bank and association
concerned so that it may determine
whether any notice of the breach to any
of its borrowers is required under
applicable laws and regulations and, if
so, each bank and association shall be
responsible for providing such
notification;
(iii) For the purposes of this section,
“breach of information” means
unauthorized acquisition of or access to
the central data repository, any
quarterly reports of accounts and
exposures or any other information
received pursuant to § 621.15(a)(1).
(8) Notify the Farm Credit
Administration in writing of any request
for data contained in the reports of
accounts and exposures that are not
explicitly allowed for in § 618.8320(b) of
this chapter.
Dated: August 5, 2013.
Dale L. Aultman,
Secretary, Farm Credit Administration Board.
[FR Doc. 2013–19231 Filed 8–8–13; 8:45 am]
BILLING CODE 6705–01–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 or we) is
extending the comment period for
the proposed rule, and for the
information collection related to the proposed rule,
“Current Good Manufacturing Practice
and Hazard Analysis and Risk-Based
Preventive Controls for Human Food,”
that appeared in the Federal Register
of January 16, 2013. We are taking this
action to allow interested persons an
opportunity to consider the
interrelationships between this proposal
and the two proposals announced in
July 2013 on the Foreign Supplier
Verification Program and on
Accreditation of Third-Party Auditors/
Certification Bodies. We also are taking
this action to keep the comment period
for the information collection provisions
associated with the rule consistent with
the comment period for the proposed
rule.

DATES: The FDA is extending the
comment period on the above proposed
rule. Submit either electronic or written
comments on the proposed rule by
November 15, 2013. Submit comments
on information collection issues under the
Paperwork Reduction Act of 1995
(the PRA) by November 15, 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 “Request for
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 Drive, P50–400T, Rockville,
MD 20850, domini.bean@fda.hhs.gov.

SUPPLEMENTARY INFORMATION:

I. Background

In the Federal Register of January 16,
2013 (78 FR 3646), we published a
proposed rule entitled “Current Good
Manufacturing Practice and Hazard
Analysis and Risk-Based Preventive
Controls for Human Food” with a 120-
day comment period on the
provisions of the proposed rule and a 30-day
comment period on the information
collection provisions that are subject to
review by OMB under the PRA (44

OMB and FDA previously received
requests for a 90-day extension of the
comment period for the information
collection provisions of the proposed
rule. We considered the requests and
extended the comment period for the
information collection for 90 days to
make the comment period for the
information collection provisions the
same as that for the proposed rule—i.e.,
until May 16, 2013 (Federal Register
of February 19, 2013, 78 FR 11611). FDA
continued to receive comments
requesting an extension of the comment
period on the proposed rule. Each
request conveyed concern that the 120-
day comment period did not allow
sufficient time to develop a meaningful
or thoughtful response to the proposed
rule. FDA considered the requests and
granted a 120-day extension of the
comment period for the proposed rule
and for the information collection provisions—i.e., until September 16, 2013 (Federal Register of April 26, 2013, 78 FR 24691). In the Federal Register of July 29, 2013 (78 FR 45729 and 78 FR 45781) we published two proposed rules entitled, “Foreign Supplier Verification Programs for Importers of Food for Humans and Animals” (Docket No. FDA–2011–N–0143) and “Accreditation of Third-Party Auditors/Certification Bodies to Conduct Food Safety Audits and to Issue Certifications” (Docket No. FDA–2011–N–0146) with a 120-day comment period. These two proposals are related to the proposed rule “Current Good Manufacturing Practice and Hazard Analysis and Risk-Based Preventive Controls for Human Food.” Therefore, FDA is granted a 60-day final extension of the comment period for the “Current Good Manufacturing Practice and Hazard Analysis and Risk-Based Preventive Controls for Human Food” proposed rule to allow interested person an opportunity to consider the interrelationships between the proposals. We also are extending the comment period for the information collection provisions for 60 days to continue 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 oira_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.”

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: August 5, 2013.

Leslie Kux,
Assistant Commissioner for Policy.

[FR Doc. 2013–19300 Filed 8–8–13; 8:45 am]
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 or we) is extending the comment period for the proposed rule, and for the information collection related to the proposed rule, “Standards for the Growing, Harvesting, Packing, and Holding of Produce for Human Consumption,” that appeared in the Federal Register of January 16, 2013. We are taking this action to allow interested persons an opportunity to consider the interrelationships between this proposal and the two proposals announced in July 2013 on the Foreign Supplier Verification Program and on Accreditation of Third-Party Auditors/Certification Bodies. We also are taking this action to keep the comment period for the information collection provisions associated with the rule consistent with the comment period for the proposed rule.

DATES: The FDA is extending the comment period on the above proposed rule. Submit either electronic or written comments on the proposed rule by November 15, 2013. Submit comments on information collection issues under the Paperwork Reduction Act of 1995 (the PRA) by November 15, 2013. (See the “Paperwork Reduction Act of 1995” section).

FOR FURTHER INFORMATION CONTACT:
With regard to the proposed rule: Samir Assar, Center for Food Safety and Applied Nutrition (HFS–317), 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 Drive, PL50–400T, Rockville, MD 20850, Domini.Bean@fda.hhs.gov.

SUPPLEMENTAL INFORMATION:

I. Background

In the Federal Register of January 16, 2013 (78 FR 3504), we published a proposed rule entitled “Standards for the Growing, Harvesting, Packing, and Holding of Produce for Human Consumption” with a 120-day comment period on the provisions of the proposed rule and a 30-day comment period on the information collection provisions that are subject to review by