

identified the application of SBA's size standards as well as other size standards used by Federal agencies (60 FR 57988 (November 24, 1995)). SBA is not aware of any Federal rule that would duplicate or conflict with establishing size standards.

However, the Small Business Act and SBA's regulations allow Federal agencies to develop different size standards if they believe that SBA's size standards are not appropriate for their programs, with the approval of SBA's Administrator (13 CFR 121.903). The SBA's regulations (13 CFR 121.903(c)) authorize an agency to establish an alternative small business definition for the sole purpose of performing a regulatory flexibility analysis pursuant to the Regulatory Flexibility Act (5 U.S.C. 601(3)), after consultation with the Office of Advocacy of the U.S. Small Business Administration.

*5. What alternatives will allow the Agency to accomplish its regulatory objectives while minimizing the impact on small entities?*

By law, SBA is required to develop numerical size standards for establishing eligibility for Federal small business assistance programs. Other than varying size standards by industry and changing the size measures, no practical alternative exists to the systems of numerical size standards.

SBA's only other consideration was whether to adopt the size standards presented in the interim final rule with no further increase for the inflation. However, SBA believes that the inflation that has occurred since the publication of the June 12, 2014 interim final rule is not sufficient to warrant an additional increase at this time.

#### List of Subjects in 13 CFR Part 121

Administrative practice and procedure, Government procurement, Government property, Grant programs—business, Individuals with disabilities, Loan programs—business, Reporting and recordkeeping requirements, Small businesses.

#### PART 121—SMALL BUSINESS SIZE REGULATIONS

■ For the reasons set forth in the preamble, the interim rule amending 13 CFR part 121, which was published at 79 FR 33647 on June 12, 2014, is adopted as a final rule without change.

Dated: January 12, 2016.

**Maria Contreras-Sweet,**  
Administrator.

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

**21 CFR Parts 1, 11, 16, 106, 110, 114, 117, 120, 123, 129, 179, and 211**

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

RIN 0910-AG36

#### Current Good Manufacturing Practice, Hazard Analysis, and Risk-Based Preventive Controls for Human Food; Correction

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

**ACTION:** Final rule; correction.

**SUMMARY:** The Food and Drug Administration (FDA or we) is correcting a final rule that published in the **Federal Register** of September 17, 2015. That final rule amended our regulation for current good manufacturing practice in manufacturing, packing, or holding human food to modernize it, and to add requirements for domestic and foreign facilities that are required to register under the Federal Food, Drug, and Cosmetic Act (the FD&C Act) to establish and implement hazard analysis and risk-based preventive controls for human food. That final rule also revised certain definitions in our current regulation for registration of food facilities to clarify the scope of the exemption from registration requirements provided by the FD&C Act for “farms.” The final rule published with some editorial and inadvertent errors. This document corrects those errors.

**DATES:** *Effective:* January 26, 2016.

#### FOR FURTHER INFORMATION CONTACT:

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.

**SUPPLEMENTARY INFORMATION:** In the **Federal Register** of Thursday, September 17, 2015 (80 FR 55908), FDA published the final rule “Current Good Manufacturing Practice, Hazard Analysis, and Risk-Based Preventive Controls for Human Food” with some editorial and inadvertent errors. This action is being taken to correct inadvertent errors in the preamble and codified.

In FR Doc. 2015-21920, appearing on page 55908 in the **Federal Register** of Thursday, September 17, 2015, the following corrections are made:

1. On page 55908, in the first column, the headings section of the document,

under the line containing “[Docket No. FDA-2011-N-0920],” is corrected by adding “RIN 0910-AG36”.

2. On page 55938, in the second column, in the first paragraph under “VII. Comments on Proposed General Revisions to Current Part 110 (Final Part 117),” “revising provisions directed to preventing contamination of food and food-contact substances” is corrected to read “revising provisions directed to preventing contamination of food and food-contact surfaces.”

■ 3. On page 56151, beginning in the second column, revise § 117.8 to read as follows:

**“§ 117.8 Applicability of subpart B of this part to the off-farm packing and holding of raw agricultural commodities.**

Except as provided by § 117.5(k)(1), subpart B of this part applies to the off-farm packaging, packing, and holding of raw agricultural commodities. Compliance with this requirement for raw agricultural commodities that are produce as defined in part 112 of this chapter may be achieved by complying with subpart B of this part or with the applicable requirements for packing and holding in part 112 of this chapter.”

#### § 117.405 [Corrected]

■ 4. On page 56164, in the first column, in § 117.405 *Requirements to establish and implement a supply chain program*, paragraph (c) introductory text is corrected to read as follows:

“(c) When a supply-chain-applied control is applied by an entity other than the receiving facility's supplier (e.g., when a non-supplier applies controls to certain produce (i.e., produce covered by part 112 of this chapter), because growing, harvesting, and packing activities are under different management), the receiving facility must:”

Dated: January 14, 2016.

**Leslie Kux,**

*Associate Commissioner for Policy.*

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