Pub. L. 101–649, 104 Stat. 4978, 5027 (8 U.S.C. 1184 note); sec. 303(a)(8), Pub. L. 102–232, 105 Stat. 1733, 1748 (8 U.S.C. 1101 note); sec. 323(c), Pub. L. 103–206, 107 Stat. 2428; sec. 412(e), Pub. L. 105–277, 112 Stat. 2681 (8 U.S.C. 1182 note); sec. 2(d), Pub. L. 106–95, 113 Stat. 1312, 1316 (8 U.S.C. 1182 note); 29 U.S.C. 49k; Pub. L. 109–423, 120 Stat. 2900; 8 CFR 214.2(h)(4)(i); and 8 CFR 214.2(h)(6)(iii).

Subparts A and C issued under 8 U.S.C. 1101(a)(15)(H)(ii)(b) and 1184; 29 U.S.C. 49 *et seq.;* and 8 CFR 214.2(h)(4)(i).

Subpart C—[Removed and Reserved]

■ 2. Remove and reserve subpart C, consisting of §§ 655.200 through 655.215.

Signed in Washington, DC, this 17th day of October 2013.

Eric M. Seleznow,

Acting Assistant Secretary, Employment and Training Administration.

[FR Doc. 2013–27693 Filed 11–19–13; 8:45 am]

BILLING CODE 4510–FP–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 1

[Docket No. FDA–2010–N–0560]

Amendments to General Regulations of the Food and Drug Administration; Technical Amendments

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; technical amendments.

SUMMARY: The Food and Drug Administration (FDA or we) published a final rule in the **Federal Register** on November 30, 2010, amending certain regulations to include tobacco products, where appropriate, in light of FDA's authority to regulate these products under the Family Smoking Prevention and Tobacco Control Act (Tobacco Control Act). The final rule inadvertently deleted an authority citation and language related to the definition of ''package.'' We are restoring the inadvertent deletions and making a corresponding technical change.

DATES: This rule is effective November 20, 2013.

FOR FURTHER INFORMATION CONTACT: Felicia Billingslea, Center for Food Safety and Applied Nutrition (HFS–820), Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740–3835, 240–402–2371.

SUPPLEMENTARY INFORMATION: We are making technical amendments to our regulations under 21 CFR part 1.

In the **Federal Register** of November 30, 2010 (75 FR 73951), we amended certain regulations in part 1 (21 CFR part 1), ''General Enforcement Regulations,'' in light of our authority under the Tobacco Control Act. The final rule, among other things:

- Revised the authority citation for part 1 by removing a reference to section 302 of the Federal Food, Drug, and Cosmetic Act (FD&C Act) (21 U.S.C. 332);
- Revised § 1.1(c), ''General,'' by removing the terms ''package in § 1.20 and of'', and
- Revised § 1.20, ''Presence of mandatory label information,'' by removing the terms ''package in § 1.20 and of''.

The preamble to the final rule explained that the revisions to part 1 reflected our authority over tobacco products under the Tobacco Control Act (75 FR 73951 at 73952). However, the revisions inadvertently created one inconsistency (in that other provisions in part 1 did, in fact, rely on section 302 of the FD&C Act as part of their legal authority) or created confusion over whether the definition of ''package'' was limited to the regulations in part 1 or whether it also applied to other FDA regulations.

Therefore, through this rule, we are amending part 1 as follows:

- We are restoring section 302 of the FD&C Act to the authority citation for part 1. Because the authority citation is expressed in terms of the U.S. Code, the amendment is to insert ''332'' in the list of U.S. Code sections.
- We are revising § 1.1(c) to restore the terms ''package in § 1.20 and of''.
- We are revising § 1.20 to add a cross-reference to § 1.1(c).

Publication of this document constitutes final action of these changes under the Administrative Procedure Act (5 U.S.C. 553). These amendments are merely correcting inadvertent deletions. FDA, therefore, for good cause, finds under 5 U.S.C. 553(b)(3)(B) and (d)(3) that notice and public comment are unnecessary.

List of Subjects in 21 CFR Part 1

Cosmetics, Drugs, Exports, Food labeling, Imports, Labeling, Reporting and recordkeeping requirements.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs, 21 CFR part 1 is amended as follows:

PART 1—GENERAL ENFORCEMENT REGULATIONS

■ 1. The authority citation for 21 CFR part 1 is revised to read as follows:

Authority: 15 U.S.C. 1333, 1453, 1454, 1455, 4402; 19 U.S.C. 1490, 1491; 21 U.S.C. 321, 331, 332, 333, 334, 335a, 343, 350c, 350d, 352, 355, 360b, 362, 371, 374, 381, 382, 387, 387a, 387c, 393; 42 U.S.C. 216, 241, 243, 262, 264.

§ 1.1 [Amended]

■ 2. Amend § 1.1 by adding the phrase ''of *package* in § 1.20 and'' after the word ''definition'' in the first sentence of paragraph (c).

■ 3. In § 1.20, revise the introductory text to read as follows:

§ 1.20 Presence of mandatory label information.

In the regulations specified in § 1.1(c) of this chapter, the term *package* means any container or wrapping in which any food, drug, device, or cosmetic is enclosed for use in the delivery or display of such commodities to retail purchasers, but does not include:

* * * * *

Dated: November 14, 2013.

Leslie Kux,

Assistant Commissioner for Policy.

[FR Doc. 2013–27773 Filed 11–19–13; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF LABOR

Occupational Safety and Health Administration

29 CFR Part 1910

[Docket No. OSHA–2013–0010]

RIN 1218–AC80

Record Requirements in the Mechanical Power Presses Standard

AGENCY: Occupational Safety and Health Administration (OSHA), Labor.

ACTION: Direct final rule; request for comments.

SUMMARY: OSHA is making two main revisions to its Mechanical Power Presses Standard. First, OSHA is revising a provision that requires employers to develop and maintain certification records of periodic inspections performed on the presses by adding a requirement that they develop and maintain certification records of any maintenance and repairs they perform on the presses during the periodic inspections. Second, OSHA is removing the requirement from another provision that employers develop and