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DEPARTMENT OF TRANSPORTATION

Federal Aviation Administration

14 CFR Part 11

[Docket No. FAA-2016-9064; Amdt. No. 11-61]

RIN 2120-AJ60

Approval of Information Collections for Operation and Certification of Small Unmanned Aircraft Systems

AGENCY: Federal Aviation Administration (FAA), Department of Transportation (DOT).

ACTION: Final rule.

SUMMARY: On June 28, 2016, the FAA published a final rule entitled Operation and Certification of Small Unmanned Aircraft Systems (81 FR 42063) which will result in new information collection requirements. This rule updates the FAA's list of OMB control numbers to display the control numbers associated with the approved information collection activities in the final rule.

DATES: Effective August 29, 2016.

FOR FURTHER INFORMATION CONTACT: Everette Rochon, Manager, Commercial Operations Branch, AFS-820, Flight Standards Service, Federal Aviation Administration, 55 M Street SE., 8th Floor, Washington, DC 20003; telephone 1-844-FLY-MYUAS; email UAShelp@faa.gov.

SUPPLEMENTARY INFORMATION:

Background

On June 28, 2016, the FAA and the Office of the Secretary of Transportation published a final rule entitled Operation and Certification of Small Unmanned Aircraft Systems. The rule adds a new part 107 to Title 14 Code of Federal Regulations (14 CFR) to allow for routine civil operation of small unmanned aircraft systems (UAS) in the

NAS and to provide safety rules for those operations. The final rule resulted in new and revised information collection requirements.

As required by the Paperwork Reduction Act of 1995 (44 U.S.C. 3507(d)), the FAA submitted these information collection amendments to OMB for its review. On August 12 and 15, 2016, OMB approved the information collection requests. The pertinent OMB control numbers are 2120-0005, 2120-0021, 2120-0027, 2120-0767, and 2120-0768.

Updating OMB Control Numbers in Part 11

The FAA lists OMB control numbers assigned to its information collection activities in 14 CFR 11.201(b). Accordingly, this final rule updates 14 CFR 11.201(b) to display OMB control numbers 2120-0005, 2120-0021, 2120-0027, 2120-0767, and 2120-0768 associated with the information collection activities in the final rule, Operation and Certification of Small Unmanned Aircraft Systems. See 81 FR 42063. This final rule also removes from 14 CFR 11.201(b) the OMB control numbers for 14 CFR parts 108 and 109, as those parts were removed by a joint FAA and Transportation Security Administration final rule, Civil Aviation Security Rules, published on February 22, 2002. See 67 FR 8340.

Because this rule concerns agency organization, procedure or practice, the FAA finds that the notice and public procedures under 5 U.S.C. 553(b)(3)(A) are unnecessary. For the same reason, the FAA finds good cause exists under 5 U.S.C. 553(d)(3) to make the amendment effective in less than 30 days.

List of Subjects in 14 CFR Part 11

Administrative practice and procedure, Reporting and recordkeeping requirements.

The Amendment

In consideration of the foregoing the Federal Aviation Administration amends Chapter I of Title 14 Code of Federal Regulations as follows:

PART 11—GENERAL RULEMAKING PROCEDURES

■ 1. The authority citation for part 11 continues to read as follows:

Authority: 49 U.S.C. 106(f), 106(g), 40101, 40103, 40105, 40109, 40113, 44110, 44502, 44701-44702, 44711, and 46102.

■ 2. In § 11.201(b), revise the entry for part 107 and remove the entries for parts 108 and 109 to read as follows:

§ 11.201 Office of Management and Budget (OMB) control numbers assigned under the Paperwork Reduction Act.

* * * * *
(b) * * *

14 CFR part or section identified and described	Current OMB control No.
Part 107	2120-0005, 2120-0021, 2120-0027, 2120-0767, 2120-0768.

Issued in Washington, DC, under the authority provided by 49 U.S.C. 106(f) and 44701(a) on August 24, 2016.

Lirio Liu,

Director, Office of Rulemaking.

[FR Doc. 2016-20687 Filed 8-26-16; 8:45 am]

BILLING CODE 4910-13-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 1, 100, 101, and 104

[Docket No. FDA-2016-N-0011]

Food Labeling; Technical Amendments

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; technical amendments.

SUMMARY: The Food and Drug Administration (FDA or we) is amending the food labeling regulations by redesignating a provision, updating any references to that provision to reflect the redesignation, and revising the section heading. The rule does not alter the content or application of the redesignated provision in any substantive manner. This action is editorial in nature and is intended to provide clarity and consistency to our regulations.

DATES: This rule is effective August 29, 2016.

FOR FURTHER INFORMATION CONTACT:

Carole Adler, Center for Food Safety and Applied Nutrition (HFS-820), Food and Drug Administration, 5001 Campus Dr., College Park, MD 20740, 240-402-2371.

SUPPLEMENTARY INFORMATION:

I. Background

The Fair Packaging and Labeling Act (FPLA) (15 U.S.C. 1451 *et seq.*) requires certain consumer commodities sold in interstate commerce to be honestly and informatively labeled to facilitate value comparisons and enable consumers to make informed choices. FDA administers and enforces the FPLA with respect to drugs, cosmetics, medical devices, and certain foods. Pursuant to our authority under FPLA, FDA issued certain food labeling regulations, including specifications regarding the statement of identity and the net quantity of contents statement, which must be present on the labels of most packaged foods. Our regulations affecting the declaration of a food product's net quantity of contents are currently located in § 101.105 (21 CFR 101.105). Section 101.105 specifies how the net quantity of contents must be expressed on the package, including the required units of measurement, wording, typeface, and size to be used in the declaration, as well as the location of the declaration on the label or package. Currently, § 101.105 requires that the units of measurement be expressed using the most appropriate units of the customary inch/pound (avoirdupois) system carried out to not more than two decimal places, as applicable.

In the *Federal Register* of May 21, 1993 (58 FR 29716), we proposed to amend our food labeling regulations to require that the net quantity of contents declaration be expressed using the most appropriate units of both imperial units (inches/pounds) and the metric system (International System of Units (SI)). The rule also proposed to provide examples of the quantity of contents declaration, include the SI equivalents to the avoirdupois terms used in the regulation, provide a specific conversion chart for use in calculating the conversion between the two systems of measurement, include SI terminology, provide exemptions from SI labeling requirements, permit the expression of the net quantity of contents to be carried out up to three decimal places, and make the SI declaration of the net quantity of contents on random weight packages optional. Additionally, the rule

proposed certain technical amendments, including redesignating § 101.105 as new § 101.7, revising the section heading to correct the title of the section, and making other editorial changes to the net quantity of contents regulations. In the *Federal Register* of April 22, 2003 (68 FR 19766), we announced our intent to withdraw the proposed rule, along with several other unrelated proposed actions that had been published more than 5 years before the withdrawal, but never finalized. The withdrawal was part of an overall regulatory reform initiative to reduce our regulatory backlog and focus our resources on higher priority regulations.

Through this final rule, we are making some of the same technical amendments to the declaration of net contents provision that were proposed in 1993. However, we are not proposing at this time to reinstitute rulemaking proceedings concerning the remaining amendments proposed in 1993, such as those pertaining to the declaration of net quantity in SI units or those modifying the expression of net quantity in decimal fractions.

II. Provisions of Technical Amendments

We are making technical amendments in our regulations at parts 1, 100, 101, and 104 (21 CFR parts 1, 100, 101, and 104) to redesignate § 101.105 as new § 101.7, update references in other provisions to reflect this redesignation, and revise the section heading of the redesignated provision. Nothing in these technical amendments is to be construed as modifying the applicability of the current regulations affecting the declaration of net quantity of contents.

A. Redesignation

FDA is amending the food labeling provisions in § 101.105 by redesignating § 101.105 as new § 101.7. Section 101.105 is currently located in part 101, subpart G, which is entitled "Exemptions From Food Labeling Requirements." However, § 101.105 contains no information pertaining to when a food is exempt from a declaration of the net quantity of contents. Instead, § 101.105 establishes general provisions for the declaration of the statement of net quantity of contents on the labels of most packaged foods. By redesignating § 101.105 as new § 101.7, we are moving the provision to subpart A, entitled "General Provisions." Subpart A provides general food labeling regulations and is a more appropriate location for a provision regulating the declaration of net quantity of contents.

B. Revising Section Heading

We also are revising the section heading of new § 101.7 to read: "Declaration of net quantity of contents" instead of "Declaration of net quantity of contents when exempt." The revised heading, by removing any reference to exemptions, is more reflective of the section's general provisions for the declaration of the statement of net quantity of contents on all food labels. The revised heading does not alter the substance of the provision.

C. Revising References

Several existing regulations refer to § 101.105. Therefore, because we are redesignating § 101.105 as a new § 101.7, we are making corresponding editorial changes to reflect the redesignation in parts 1, 100, 101, and 104. These corresponding changes replace any mention of § 101.105 with § 101.7.

III. Notice and Public Comment

Publication of this document constitutes final action of these changes under the Administrative Procedure Act (5 U.S.C. 553). FDA has determined that notice and public comment are unnecessary because these amendments are nonsubstantive and provide only technical changes to redesignate an existing regulation, make corresponding changes to other regulations to reflect the redesignated section number, and make an editorial change to the section heading. These technical amendments are being made to improve the accuracy of our regulations.

List of Subjects

21 CFR Part 1

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

21 CFR Part 100

Administrative practice and procedure, Food labeling, Food packaging, Foods, Intergovernmental relations.

21 CFR Part 101

Food labeling, Nutrition, Reporting and recordkeeping requirements.

21 CFR Part 104

Food grades and standards, Frozen foods, Nutrition.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs, the Food and Drug Administration amends 21 CFR parts 1, 100, 101, and 104 as follows:

PART 1—GENERAL ENFORCEMENT REGULATIONS

■ 1. The authority citation for part 1 continues 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, 342i, 343, 350c, 350d, 350e, 352, 355, 360b, 360ccc, 360ccc-1, 360ccc-2, 362, 371, 373, 374, 381, 382, 387, 387a, 387c, 393; 42 U.S.C. 216, 241, 243, 262, 264.

§ 1.1 [Amended]

■ 2. Amend § 1.1(c) as follows:

- a. Remove “101.105(f)” and add in its place “101.7(f)”.
- b. Remove “101.105(i)” and add in its place “101.7(i)”.
- c. Remove “101.105(j)” and add in its place “101.7(j)”.
- d. Remove “101.105(o)” and add in its place “101.7(o)”.

§ 1.20 [Amended]

■ 3. In § 1.20, by removing “§ 101.105(f)” and adding in its place “§ 101.7(f)”.

§ 1.24 [Amended]

■ 4. Amend § 1.24 as follows:

- a. Remove “§ 101.105” in paragraph (a)(2) and add in its place “§ 101.7”.
- b. Remove “§ 101.105(b)(2)” wherever it appears and add in its place “§ 101.7(b)(2)”.
- c. Remove “§ 101.105(f)” wherever it appears and add in its place “§ 101.7(f)”.
- d. Remove “§ 101.105(j)” wherever it appears and add in its place “§ 101.7(j)”.
- e. Remove “§ 101.105(j)(1)” wherever it appears and add in its place “§ 101.7(j)(1)”.

PART 100—GENERAL

■ 5. The authority citation for part 100 continues to read as follows:

Authority: 21 U.S.C. 321, 331, 337, 342, 343, 348, 371.

§ 100.155 [Amended]

■ 6. Amend § 100.155 in paragraphs (a) and (b) by removing “§ 101.105” and adding in its place “§ 101.7”.

PART 101—FOOD LABELING

■ 7. The authority citation for part 101 continues to read as follows:

Authority: 15 U.S.C. 1453, 1454, 1455; 21 U.S.C. 321, 331, 342, 343, 348, 371; 42 U.S.C. 243, 264, 271.

§ 101.2 [Amended]

■ 8. Amend § 101.2 in paragraph (c) introductory text by removing “§ 101.105(h)(1)” and adding in its place “§ 101.7(h)(1)”.

§ 101.105 [Redesignated as § 101.7]

■ 9. Redesignate § 101.105 as § 101.7.
 ■ 10. Revise newly designated § 101.7 section heading to read as follows:

§ 101.7 Declaration of net quantity of contents.

* * * * *

§ 101.13 [Amended]

■ 11. Amend paragraphs (d)(2), (h)(4)(i), and (i)(2) by removing “§ 101.105(i)” and adding in its place “§ 101.7(i)”.

§ 101.30 [Amended]

■ 12. Amend § 101.30(g) by removing “§ 101.105(i)” and adding in its place “§ 101.7(i)”.

PART 104—NUTRITIONAL QUALITY GUIDELINES FOR FOODS

■ 13. The authority citation for part 104 continues to read as follows:

Authority: 21 U.S.C. 321, 343, 371(a).

§ 104.5 [Amended]

■ 14. Amend § 104.5(b) by removing “§ 101.105” and adding in its place “§ 101.7”.

Dated: August 16, 2016.

Jeremy Sharp,

Deputy Commissioner for Policy, Planning, Legislation, and Analysis.

[FR Doc. 2016-19925 Filed 8-26-16; 8:45 am]

BILLING CODE 4164-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES**Food and Drug Administration****21 CFR Parts 510, 520, 522, 524, and 558**

[Docket No. FDA-2016-N-0002]

New Animal Drugs; Approval of New Animal Drug Applications; Withdrawal of Approval of New Animal Drug Applications; Changes of Sponsorship; Change of Sponsor's Name and Address; Change of Sponsor's Address

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; technical amendment.

SUMMARY: The Food and Drug Administration (FDA or we) is amending the animal drug regulations to reflect application-related actions for new animal drug applications (NADAs) and abbreviated new animal drug applications (ANADAs) during May and June 2016. FDA is also informing the public of the availability of summaries of the basis of approval and of environmental review documents, where applicable. The animal drug regulations are also being amended to reflect changes of sponsorship of applications, changes of sponsors' names and addresses, and the voluntary withdrawals of approval of applications.

DATES: This rule is effective August 29, 2016, except for the amendments to 21 CFR 558.274, 58.355, 58.363, 58.550, 558.625, and 558.630, which are effective September 8, 2016.

FOR FURTHER INFORMATION CONTACT:

George K. Haibel, Center for Veterinary Medicine (HFV-6), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240-402-5689, george.haibel@fda.hhs.gov.

SUPPLEMENTARY INFORMATION:**I. Approval Actions**

FDA is amending the animal drug regulations to reflect approval actions for NADAs and ANADAs during May and June 2016, as listed in table 1. In addition, FDA is informing the public of the availability, where applicable, of documentation of environmental review required under the National Environmental Policy Act (NEPA) and, for actions requiring review of safety or effectiveness data, summaries of the basis of approval (FOI Summaries) under the Freedom of Information Act (FOIA). These public documents may be seen in the Division of Dockets Management (HFA-305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852, between 9 a.m. and 4 p.m., Monday through Friday. Persons with access to the Internet may obtain these documents at the CVM FOIA Electronic Reading Room: <http://www.fda.gov/AboutFDA/CentersOffices/OfficeofFoods/CVM/CVMFOIAElectronicReadingRoom/default.htm>. Marketing exclusivity and patent information may be accessed in FDA's publication, Approved Animal Drug Products Online (Green Book) at: <http://www.fda.gov/AnimalVeterinary/Products/ApprovedAnimalDrugProducts/default.htm>.