making any changes in response to the objections.

The filing of the objections served to stay automatically the effectiveness of § 73.520. Section 701(e)(2) of the FD&C Act states that, until final action upon such objections is taken by the Secretary, the filing of such objections operates to stay the effectiveness of those provisions of the order to which the objections are made. Section 701(e)(3) of the FD&C Act further stipulates that, as soon as practicable, the Secretary shall by order act upon such objections and make such order public. We have completed our evaluation of the objections and conclude that a continuation of the stay of § 73.520 is not warranted.

In the absence of any other objections and requests for a hearing, we conclude that this document constitutes final action on the objections received in response to the regulation as prescribed in section 701(e)(2) of the FD&C Act. Therefore, we are ending the administrative stay of the regulation as of December 19, 2019 for the § 73.520 listing soy leghemoglobin as a color additive for use in ground beef analogue products.

VI. References

The following references marked with an asterisk (*) are on display at the Dockets Management Staff (see **ADDRESSES**) and are available for viewing by interested persons between 9 a.m. and 4 p.m., Monday through Friday; they also are available electronically at *https://www.regulations.gov.* References without asterisks are not on public display at *https://www.regulations.gov* because they have copyright restriction. Some may be available at the website address, if listed. References without asterisks are available for viewing only at the Dockets Management Staff. FDA has verified the website addresses, as of the date this document publishes in the **Federal Register**, but websites are subject to change over time.

1. Ladics, G.S., ''Current Codex Guidelines for Assessment of Potential Protein Allergenicity.'' *Food and Chemical Toxicology,* 46: S20–S23, 2008.
2. *FDA. ''Redbook 2000 Guidance for Industry and Other Stakeholders; Toxicological Principles for the Safety Assessment of Food Ingredients,'' 2007. Retrieved from *https://www.fda.gov/media/79074/download.*
3. Giknis, M.L.A. and C.B. Clifford, ''Clinical Laboratory Parameters for Crl:CD(SD) Rats,'' 2006. Retrieved from *https://www.crj.co.jp/cms/pdf/info_common/50/8250933/rm_rm_r_clinical_parameters_cd_rat_06.pdf.*
4. Giknis, M.L.A. and C.B. Clifford, ''Clinical Laboratory Parameters for Crl:WI(Han),'' 2008. Retrieved from *https://www.criver.com/sites/default/files/resources/rm_rm_r_Wistar_Han_clin_lab_parameters_08.pdf.*
5. Matsuzawa, T., M. Nomura, and T. Unno, ''Clinical Pathology Reference Ranges of Laboratory Animals. Working Group II, Nonclinical Safety Evaluation Subcommittee of the Japan Pharmaceutical Manufacturers Association.'' *Journal of Veterinary Medical Science,* 55(3): 351–362, 1993.
6. Pettersen, J.C., R.L. Morrissey, D. R. Saunders, et al., ''A 2-Year Comparison Study of Crl:CD BR and Hsd:Sprague-Dawley SD Rats.'' *Fundamental and Applied Toxicology,* 33: 196–211, 1996.
7. Petterino, C. and A. Argentino-Storino, ''Clinical Chemistry and Haematology Historical Data in Control Sprague-Dawley Rats From Pre-clinical Toxicity Studies.'' *Experimental and Toxicologic Pathology,* 57: 213–219, 2006.
8. Seibel, J., K. Bodié, S. Weber, et al., ''Comparison of Haematology, Coagulation and Clinical Chemistry Parameters in Blood Samples From the Sublingual Vein and Vena Cava in Sprague-Dawley Rats.'' *Laboratory Animals,* 44: 344–351, 2010.

List of Subjects in 21 CFR Part 73

Color additives, Cosmetics, Drugs, Foods, Medical devices.

Therefore, under the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 321, 341, 342, 343, 348, 351, 352, 355, 361, 362, 371, 379e) and under authority delegated to the Commissioner of Food and Drugs (section 1410.10 of the FDA Staff Manual Guide), notice is given that the objections and requests for hearings were filed in response to the August 1, 2019, final rule. Notice is also given that FDA is denying these objections and requests for hearings. Accordingly, the administrative stay on the effective date of the amendments is lifted as of December 19, 2019.

Dated: December 12, 2019.

Lowell J. Schiller,

Principal Associate Commissioner for Policy.

[FR Doc. 2019–27173 Filed 12–17–19; 11:15 am]

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ENVIRONMENTAL PROTECTION AGENCY

40 CFR Part 282

[EPA–R01–UST–2019–0421; FRL–10003–06–Region 1]

New Hampshire: Final Approval of State Underground Storage Tank Program Revisions, Codification, and Incorporation by Reference

AGENCY: Environmental Protection Agency (EPA).

ACTION: Direct final rule; correction.

SUMMARY: The Environmental Protection Agency (EPA) is correcting a direct final rule that appeared in the **Federal Register** on November 1, 2019. The document is taking direct final action to approve revisions to the State of New Hampshire's Underground Storage Tank (UST) program submitted by the New Hampshire Department of Environmental Services (NHDES). This action also codifies EPA's approval of New Hampshire's state program and incorporates by reference those provisions of the State regulations that meet the requirements for approval.

DATES: This rule is effective December 31, 2019, unless EPA received adverse comment by December 2, 2019. If EPA received adverse comments, it will publish a timely withdrawal in the **Federal Register** informing the public that the rule will not take effect. The incorporation by reference of certain publications listed in the regulations is approved by the Director of the Federal Register, as of December 31, 2019, in accordance with 5 U.S.C. 552(a) and 1 CFR part 51.

FOR FURTHER INFORMATION CONTACT: Susan Hanamoto, RCRA Waste Management, UST, and Pesticides Section; Land, Chemicals, and Redevelopment Division; EPA Region 1, 5 Post Office Square, Suite 100, (Mail Code 07–1), Boston, MA 02109–3912.

SUPPLEMENTARY INFORMATION: In FR Doc. 2019–23709 appearing on pages 58627 and 58631 in the **Federal Register** of Friday, November 1, 2019, the following corrections are made:

1. On page 58627, in the heading of the document, the agency heading is corrected to read ''ENVIRONMENTAL PROTECTION AGENCY'' and in the AGENCY caption, the agency is corrected to read ''Environmental Protection Agency (EPA)''.

2. On page 58627, in the first sentence of the SUMMARY, ''Environmental Services Agency'' is corrected to read ''Environmental Protection Agency''.

3. On page 58631, middle column, in the List of Subjects in 40 CFR part 282, ''Environmental Services'' is corrected to read ''Environmental Protection''.

Dated: November 5, 2019.

Nancy Barmakian,

Acting Director of Land, Chemicals, and Redevelopment Division.

[FR Doc. 2019–26690 Filed 12–18–19; 8:45 am]

BILLING CODE 6560–50–P