application process. The employer is expected to continue to pay at least the prevailing wage as promised in the employer's labor certification (ETA Form 9142) for any work performed before November 30, 2011. However, employers who received a supplemental H–2B prevailing wage determination must pay at least that wage to any H–2B worker and any U.S. worker recruited in connection with the labor certification for work performed on or after November 30, 2011.

Signed at Washington, DC, this 27th of September 2011.

Jane Oates,

Assistant Secretary for Employment and Training.

Nancy J. Leppink,

Deputy Administrator, Wage and Hour Division.

[FR Doc. 2011–25302 Filed 9–28–11; 11:15 am]

BILLING CODE 4510–FP–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 558

[Docket No. FDA–2011–N–0003]

New Animal Drugs for Use in Animal Feeds; Melengestrol; Monensin

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations to reflect approval of a supplemental abbreviated new animal drug application (ANADA) filed by Ivy Laboratories, Division of Ivy Animal Health, Inc. The supplemental ANADA provides for use of increased dose levels of melengestrol acetate and monensin in two-way, combination drug Type C medicated feeds for heifers fed in confinement for slaughter.

DATES: This rule is effective September 30, 2011.

FOR FURTHER INFORMATION CONTACT: John K. Harshman, Center for Veterinary Medicine (HFV–170), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 240–276–8197, e-mail: *john.harshman@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: Ivy Laboratories, Division of Ivy Animal Health, Inc., 8857 Bond St., Overland Park, KS 66214, filed a supplement to ANADA 200–422 for use of HEIFERMAX 500 (melengestrol acetate) and RUMENSIN (monensin, USP) single-ingredient Type A medicated articles to make two-way, combination drug Type C medicated feeds for heifers fed in confinement for slaughter. The supplemental ANADA provides for use of increased dose levels of melengestrol acetate and monensin. The supplemental application is approved as of July 1, 2011, and the regulations in 21 CFR 558.342 are amended to reflect the approval and minor revisions.

In accordance with the freedom of information provisions of 21 CFR part 20 and 21 CFR 514.11(e)(2)(ii), a summary of safety and effectiveness data and information submitted to support approval of this application 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.

The Agency has determined under 21 CFR 25.33 that this action is of a type that does not individually or cumulatively have a significant effect on the human environment. Therefore, neither an environmental assessment nor an environmental impact statement is required.

This rule does not meet the definition of ''rule'' in 5 U.S.C. 804(3)(A) because it is a rule of ''particular applicability.'' Therefore, it is not subject to the congressional review requirements in 5 U.S.C. 801–808.

List of Subjects in 21 CFR Part 558

Animal drugs, Animal feeds.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs and redelegated to the Center for Veterinary Medicine, 21 CFR part 558 is amended as follows:

PART 558—NEW ANIMAL DRUGS FOR USE IN ANIMAL FEEDS

■ 1. The authority citation for 21 CFR part 558 continues to read as follows:

Authority: 21 U.S.C. 360b, 371.

■ 2. In § 558.342, in the table in paragraph (e)(1), remove and reserve paragraphs (e)(1)(v) and (e)(1)(vi); in paragraph (e)(1)(x), in the ''Sponsor'' column, add ''021641''; and revise paragraph (d)(2) to read as follows:

§ 558.342 Melengestrol.

* * * * *

(d) * * *

(2) A physically stable melengestrol acetate liquid Type B or C feed will not be subject to the requirements for mixing directions prescribed in paragraph (d)(1) of this section provided it has a pH of 4.0 to 8.0 and contains a suspending agent(s) sufficient to maintain a viscosity of not less than 300 centipoises per second for 3 months.

* * * * *

Dated: September 20, 2011.

Steven D. Vaughn,

Director, Office of New Animal Drug Evaluation, Center for Veterinary Medicine.

[FR Doc. 2011–25220 Filed 9–29–11; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF THE TREASURY

Internal Revenue Service

26 CFR Parts 1 and 602

[TD 9551]

RIN 1545–BF94

Deduction for Qualified Film and Television Production Costs

AGENCY: Internal Revenue Service (IRS), Treasury.

ACTION: Final regulations and removal of temporary regulations.

SUMMARY: This document contains final regulations relating to deductions for the costs of producing qualified film and television productions. These final regulations reflect changes to the law made by the American Jobs Creation Act of 2004 and the Gulf Opportunity Zone Act of 2005, and affect persons that produce film and television productions within the United States.

DATES: *Effective Date:* These regulations are effective on September 29, 2011.

Applicability Dates: For dates of applicability, see § 1.181–6.

FOR FURTHER INFORMATION CONTACT: Bernard P. Harvey, (202) 622–4930 (not a toll-free number).

SUPPLEMENTARY INFORMATION:

Paperwork Reduction Act

The collection of information contained in these final regulations has been reviewed and approved by the Office of Management and Budget in accordance with the Paperwork Reduction Act of 1995 (44 U.S.C. 3507(d)) under control number 1545–2059. The collection of information in these final regulations is in §§ 1.181–1, 1.181–2, and 1.181–3. This information is required to enable the IRS to verify that a taxpayer is entitled to the deduction.

An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a valid control number.

Books and records relating to a collection of information must be