## PART 1273—UNIFORM ADMINISTRATIVE REQUIREMENTS FOR GRANTS AND COOPERATIVE AGREEMENTS TO STATE AND LOCAL GOVERNMENTS

■ 3. The authority citation for 14 CFR part 1273 is revised to read as follows:

**Authority:** 51 U.S.C. 20113(e), Pub. L. 97–258, 96 Stat. 1003 (31 U.S.C. 6301, *et seq.*), and 2 CFR Part 200.

**§§ 1273.50 and 1273.51 [Removed and Reserved]**

■ 4. Sections 1273.50 and 1273.51 are removed and reserved.

## PART 1274—COOPERATIVE AGREEMENTS WITH COMMERCIAL FIRMS

■ 5. The authority citation for 14 CFR part 1274 is revised to read as follows:

**Authority:** 51 U.S.C. 20113(e), Pub. L. 97–258, 96 Stat. 1003 (31 U.S.C. 6301, *et seq.*).

**§§ 1274.803 and 1274.804 [Removed and Reserved]**

■ 6. Sections 1274.803 and 1274.804 are removed and reserved.

[FR Doc. 2014–08372 Filed 4–14–14; 8:45 am]

**BILLING CODE 7510–01–P**

## DEPARTMENT OF ENERGY

### Federal Energy Regulatory Commission

### 18 CFR Part 341

**[Docket Nos. RM12–15–000 and RM01–5–000]**

### Filing, Indexing and Service Requirements for Oil Pipelines

**AGENCY:** Federal Energy Regulatory Commission, DOE.

**ACTION:** Notice of extension of compliance date.

**SUMMARY:** This document revises the date to comply with the terms of the Final Rule (RM12–15–000) which was published in the **Federal Register** of Wednesday, May 29, 2013. The rule amended regulations under the Interstate Commerce Act to update requirements governing the form, composition and filing of rates and charges by interstate oil pipelines for transportation in interstate commerce.

**DATES:** Effective May 15, 2014.

**FOR FURTHER INFORMATION CONTACT:** Aaron Kahn (Technical Issues), 888 First Street, NE., Washington, DC 20426, (202) 502–8339, *aaron.kahn@ferc.gov.*

**SUPPLEMENTARY INFORMATION:**

#### Notice Regarding Compliance Date

On June 14, 2013, the Commission granted an indefinite extension of time for compliance with the Final Rule in Docket No. RM12–15–000 (May 16, 2013 Order) [1] pending final clearance from the Office of Management and Budget (OMB) and further notice from the Commission. The Commission received clearance from OMB on September 30, 2013. Beginning May 15, 2014, covered entities are required to comply with the terms of the Final Rule published May 29, 2013 at 78 FR 32090.

Dated: April 9, 2014.

**Kimberly D. Bose,**

*Secretary.*

[FR Doc. 2014–08510 Filed 4–14–14; 8:45 am]

**BILLING CODE 6717–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

### 21 CFR Part 522

**[Docket No. FDA–2014–N–0002]**

### New Animal Drugs; Ceftiofur Sodium; Gentamicin; Xylazine

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

**ACTION:** Final rule; technical amendment.

**SUMMARY:** The Food and Drug Administration (FDA) is amending the animal drug regulations to reflect approval actions for new animal drug applications (NADAs) and abbreviated new animal drug applications (ANADAs) during March 2014. 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 a change of sponsorship for an ANADA.

**DATES:** This rule is effective April 15, 2014.

**FOR FURTHER INFORMATION CONTACT:** George K. Haibel, Center for Veterinary Medicine (HFV–6), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240–276–9019, *george.haibel@fda.hhs.gov.*

**SUPPLEMENTARY INFORMATION:** FDA is amending the animal drug regulations to reflect approval actions for NADAs and ANADAs during March 2014, 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 Center for Veterinary Medicine 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.*

Also, the regulations are being amended to reflect the previous approval of revised food safety warnings for ceftiofur sodium powder for injection. This amendment is being made to improve the accuracy of the regulations.

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.

[1] *Filing, Indexing and Service Requirements for Oil Pipelines,* Order No. 780, 78 FR 32090 (May 29, 2013), FERC Stats. & Regs. ¶ 31,347 (2013).

TABLE 1—ORIGINAL AND SUPPLEMENTAL NADAS AND ANADAS APPROVED DURING MARCH 2014

| NADA/ ANADA | Sponsor | New animal drug product name | Action | 21 CFR Section | FOIA Summary | NEPA Review |
|---|---|---|---|---|---|---|
| 200–468 ... | Cross Vetpharm Group Ltd., Broomhill Rd., Tallaght, Dublin 24, Ireland. | GENTAMED–P for Poultry (gentamicin sulfate) Injection. | Original approval as a generic copy of NADA 101–862. | 522.1044 | yes ..... | CE.[1] [2] |
| 200–529 ... | Cross Vetpharm Group Ltd., Broomhill Rd., Tallaght, Dublin 24, Ireland. | XYLAMED (xylazine) Injection. | Original approval as a generic copy of NADA 047–956. | 522.2662 | yes ..... | CE.[1] [2] |

[1] The Agency has determined under 21 CFR 25.33 that this action is categorically excluded (CE) from the requirement to submit an environmental assessment or an environmental impact statement because it is of a type that does not individually or cumulatively have a significant effect on the human environment.

[2] CE granted under 21 CFR 25.33(a)(1).

**List of Subjects in 21 CFR Part 522**

Animal drugs.

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 522 is amended as follows:

## PART 522—IMPLANTATION OR INJECTABLE DOSAGE FORM NEW ANIMAL DRUGS

■ 1. The authority citation for 21 CFR part 522 continues to read as follows:

**Authority:** 21 U.S.C. 360b.

■ 2. In 522.313c, revise paragraph (d) to read as follows:

**§ 522.313c Ceftiofur sodium.**

* * * * *

(d) *Special considerations.* Federal law restricts this drug to use by or on the order of a licensed veterinarian. Federal law prohibits extra-label use of this drug in cattle, swine, chickens, and turkeys for disease prevention purposes; at unapproved doses, frequencies, durations, or routes of administration; and in unapproved major food-producing species/production classes.

* * * * *

**§ 522.1044 [Amended]**

■ 3. In § 522.1044, in paragraph (b)(4), remove ''No. 000859'' and in its place add ''Nos. 000859 and 061623''.

**§ 522.2662 [Amended]**

■ 4. In § 522.2662, in paragraph (b)(2), remove ''No. 000010'' and in its place add '' Nos. 000010 and 061623''.

Dated: April 9, 2014.

**Bernadette Dunham,**

*Director, Center for Veterinary Medicine.*

[FR Doc. 2014–08445 Filed 4–14–14; 8:45 am]

BILLING CODE 4160–01–P

## PENSION BENEFIT GUARANTY CORPORATION

### 29 CFR Part 4022

### Benefits Payable in Terminated Single-Employer Plans; Interest Assumptions for Paying Benefits

**AGENCY:** Pension Benefit Guaranty Corporation.

**ACTION:** Final rule.

**SUMMARY:** This final rule amends the Pension Benefit Guaranty Corporation's regulation on Benefits Payable in Terminated Single-Employer Plans to prescribe interest assumptions under the regulation for valuation dates in May 2014. The interest assumptions are used for paying benefits under terminating single-employer plans covered by the pension insurance system administered by PBGC.

**DATES:** Effective May 1, 2014.

**FOR FURTHER INFORMATION CONTACT:** Catherine B. Klion (*Klion.Catherine@pbgc.gov*), Assistant General Counsel for Regulatory Affairs, Pension Benefit Guaranty Corporation, 1200 K Street NW., Washington, DC 20005, 202–326–4024. (TTY/TDD users may call the Federal relay service toll-free at 1–800–877–8339 and ask to be connected to 202–326–4024.)

**SUPPLEMENTARY INFORMATION:** PBGC's regulation on Benefits Payable in Terminated Single-Employer Plans (29 CFR Part 4022) prescribes actuarial assumptions—including interest assumptions—for paying plan benefits under terminating single-employer plans covered by title IV of the Employee Retirement Income Security Act of 1974. The interest assumptions in the regulation are also published on PBGC's Web site (*http://www.pbgc.gov*).

PBGC uses the interest assumptions in Appendix B to Part 4022 to determine whether a benefit is payable as a lump sum and to determine the amount to pay. Appendix C to Part 4022 contains interest assumptions for private-sector pension practitioners to refer to if they wish to use lump-sum interest rates determined using PBGC's historical methodology. Currently, the rates in Appendices B and C of the benefit payment regulation are the same.

The interest assumptions are intended to reflect current conditions in the financial and annuity markets. Assumptions under the benefit payments regulation are updated monthly. This final rule updates the benefit payments interest assumptions for May 2014.[1]

The May 2014 interest assumptions under the benefit payments regulation will be 1.50 percent for the period during which a benefit is in pay status and 4.00 percent during any years preceding the benefit's placement in pay status. In comparison with the interest assumptions in effect for April 2014, these interest assumptions are unchanged.

PBGC has determined that notice and public comment on this amendment are impracticable and contrary to the public interest. This finding is based on the need to determine and issue new interest assumptions promptly so that the assumptions can reflect current market conditions as accurately as possible.

Because of the need to provide immediate guidance for the payment of benefits under plans with valuation dates during May 2014, PBGC finds that good cause exists for making the assumptions set forth in this amendment effective less than 30 days after publication.

PBGC has determined that this action is not a ''significant regulatory action'' under the criteria set forth in Executive Order 12866.

[1] Appendix B to PBGC's regulation on Allocation of Assets in Single-Employer Plans (29 CFR Part 4044) prescribes interest assumptions for valuing benefits under terminating covered single-employer plans for purposes of allocation of assets under ERISA section 4044. Those assumptions are updated quarterly.