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).