T-545 BROWNSVILLE, TX (BRO) TO GOOCH SPRINGS, TX (AGJ) [NEW]

Brownsville, TX (BRO)	VORTAC	(Lat. 25°55′26.66″ N, long. 097°22′30.97″ W)
Corpus Christi, TX (CRP)	VORTAC	(Lat. 27°54'13.56" N, long. 097°26'41.57" W)
SLENA, TX	FIX	(Lat. 28°32'38.31" N, long. 098°11'47.89" W)
San Antonio, TX (SAT)	VORTAC	(Lat. 29°38'38.51" N, long. 098°27'40.74" W)
Gooch Springs, TX (AGJ)	VORTAC	(Lat. 31°11′07.82″ N, long. 098°08′30.69″ W)

*

Issued in Washington, DC.

Brian Eric Konie,

Manager (A), Rules and Regulations Group. [FR Doc. 2025-08796 Filed 5-16-25; 8:45 am] BILLING CODE 4910-13-P

*

*

DELAWARE RIVER BASIN COMMISSION

18 CFR Parts 401 and 420

Regulatory Program Fees and Water Charges Rates

AGENCY: Delaware River Basin Commission. **ACTION:** Final rule.

SUMMARY: Notice is provided of the Commission's regulatory program fees and schedule of water charges for the fiscal year beginning July 1, 2025. **DATES:** This final rule is effective July 1, 2025.

FOR FURTHER INFORMATION CONTACT: Elba L. Deck, CPA, Director of Finance and Administration, (609) 477-7201. SUPPLEMENTARY INFORMATION: The

Delaware River Basin Commission ("DRBC" or "Commission") is a Federal-interstate compact agency charged with managing the water

resources of the Delaware River Basin on a regional basis without regard to political boundaries. Its members are the governors of the four basin states— Delaware, New Jersev, New York and Pennsylvania—and on behalf of the Federal Government, the North Atlantic Division Commander of the U.S. Army Corps of Engineers.

In accordance with 18 CFR 401.43(c), on July 1 of every year, the Commission's regulatory program fees as set forth in tables 1, 2 and 3 of that section are subject to an annual adjustment, commensurate with any increase in the annual April 12-month Consumer Price Index (CPI) for Philadelphia published by the U.S. Bureau of Labor Statistics during that year. Pursuant to 18 CFR 420.41(c), the same indexed adjustment applies to the Commission's schedule of water charges for consumptive and non-consumptive withdrawals of surface water within the basin. The referenced April 12-month CPI for 2025 showed an increase of 2.88%. Commensurate adjustments are thus required.

This notification is made in accordance with 18 CFR 401.43(c) and 420.41(c), which provide that a revised fee schedule will be published in the Federal Register by July 1. The revised fees also may be obtained by contacting the Commission during business hours or by checking the Commission's website.

List of Subjects

18 CFR Part 401

Administrative practice and procedure, Project review, Water pollution control, Water resources.

18 CFR Part 420

Water supply. For the reasons set forth in the preamble, the Delaware River Basin Commission amends parts 401 and 420 of title 18 of the Code of Federal Regulations as set forth below:

PART 401—RULES OF PRACTICE AND PROCEDURE

■ 1. The authority citation for part 401 continues to read as follows:

Authority: Delaware River Basin Compact (75 Stat. 688), unless otherwise noted.

Subpart C—Project Review Under Section 3.8 of the Compact

■ 2. In § 401.43, revise tables 1, 2, and 3 to read as follows:

§ 401.43 Regulatory program fees.

* * *

TABLE 1 TO §401.43—DOCKET APPLICATION FILING FEE

Project type	Docket application fee	Fee maximum
Water Allocation	\$526 per million gallons/month of allocation, ¹ not to exceed \$19,724 ¹ Fee is doubled for any portion to be exported from the basin	Greater of: \$19,724 ¹ or Alternative Review Fee.
Wastewater Discharge	Private projects: \$1,315 ¹ Public projects: \$657 ¹	Alternative Review Fee.
Other	0.4% of project cost up to \$10,000,000 plus 0.12% of project cost above \$10,000,000 (if applicable), not to exceed \$98,618.1	Greater of: \$98,618 ¹ or Alternative Review Fee.

¹ Subject to annual adjustment in accordance with paragraph (c) of this section.

TABLE 2 TO §401.43—ANNUAL MONITORING AND COORDINATION FEE

	Annual fee	Allocation
Water Allocation	¹ 855 ¹ 1.085	<4.99 mgm. 5.00 to 49.99 mgm. 50.00 to 499.99 mgm. 500.00 to 9,999.99 mgm. > or = to 10,000 mgm.
	Annual fee	Discharge design capacity
Wastewater Discharge	¹ 394 ¹ 802 ¹ 1,078	<0.05 mgd. 0.05 to 1 mgd. 1 to 10 mgd.

TABLE 2 TO §401.43—ANNUAL MONITORING AND COORDINATION FEE—Continued

Annual fee	Allocation
¹ 1,315	>10 mgd.

¹ Subject to annual adjustment in accordance with paragraph (c) of this section.

TABLE 3 TO §401.43—ADDITIONAL FEES

Proposed action	Fee	Fee maximum
Emergency Approval Under 18 CFR 401.40. Late Filed Renewal Surcharge	\$5,000 \$2,000.	Alternative Review Fee.
	At Executive Director's discretion, Docket Application Fee for the appropriate project type.	Alternative Review Fee.
Name change Change of Ownership		

¹ Subject to annual adjustment in accordance with paragraph (c) of this section.

PART 420—BASIN REGULATIONS— WATER SUPPLY CHARGES

■ 3. The authority citation for part 420 continues to read as follows:

Authority: Delaware River Basin Compact, 75 Stat. 688.

■ 4. In § 420.41, revise paragraphs (a) and (b) to read as follows:

§ 420.41 Schedule of water charges.

* * * * * * (a) \$105 per million gallons for consumptive use, subject to paragraph (c) of this section; and

(b) \$1.05 per million gallons for nonconsumptive use, subject to paragraph (c) of this section.

Dated: May 13, 2025.

Pamela M. Bush,

Assistant General Counsel and Commission Secretary.

[FR Doc. 2025–08900 Filed 5–16–25; 8:45 am] BILLING CODE P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 573

[Docket No. FDA-2014-F-1184]

Food Additives Permitted in Feed and Drinking Water of Animals; Selenium

AGENCY: Food and Drug Administration, Department of Health and Human Services (HHS).

ACTION: Final amendment; order.

SUMMARY: The Food and Drug Administration (FDA, we, or the Agency) is amending the regulations for food additives permitted in feed and drinking water of animals to provide for the safe use of zinc L-selenomethionine as a source of selenium in complete feed for broiler chickens. This action is in response to a food additive petition filed by Zinpro Corp.

DATES: This order is effective May 19, 2025. See section V, Objections and Hearing Requests, for further information on the filing of objections. Either electronic or written objections and requests for a hearing on the final amendment must be submitted by June 18, 2025.

ADDRESSES: You may submit objections and requests for a hearing as follows. Please note that late, untimely filed objections will not be considered. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of June 18, 2025. Objections received by mail/hand delivery/courier (for written/ paper submissions) will be considered timely if they are received on or before that date.

Electronic Submissions

Submit electronic objections in the following way:

• Federal eRulemaking Portal: https://www.regulations.gov. Follow the instructions for submitting objections. Objections submitted electronically, including attachments, to https:// www.regulations.gov will be posted to the docket unchanged. Because your objection will be made public, you are solely responsible for ensuring that your objection does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that

identifies you in the body of your objection, that information will be posted on *https://www.regulations.gov.*

• If you want to submit an objection with confidential information that you do not wish to be made available to the public, submit the objection as a written/paper submission and in the manner detailed (see "Written/Paper Submissions" and "Instructions").

Written/Paper Submissions

Submit written/paper submissions as follows:

• Mail/Hand Delivery/Courier (for written/paper submissions): Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

• For written/paper objections submitted to the Dockets Management Staff, FDA will post your objection, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in "Instructions."

Instructions: All submissions received must include the Docket No. FDA– 2014–F–1184 for "Food Additives Permitted in Feed and Drinking Water of Animals; Selenium." Received objections, those filed in a timely manner (see **ADDRESSES**), will be placed in the docket and, except for those submitted as "Confidential Submissions," publicly viewable at *https://www.regulations.gov* or at the Dockets Management Staff between 9 a.m. and 4 p.m., Monday through Friday, 240–402–7500.

• Confidential Submissions—To submit an objection with confidential information that you do not wish to be made publicly available, submit your objections only as a written/paper submission. You should submit two copies in total. One copy will include the information you claim to be