



27 August 2026

(26-5948)

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Committee on Sanitary and Phytosanitary Measures

Original: English

NOTIFICATION

1.	Notifying Member: UNITED STATES OF AMERICA If applicable, name of local government involved:
2.	Agency responsible: Food and Drug Administration (FDA)
3.	Products covered (provide tariff item number(s) as specified in national schedules deposited with the WTO; ICS numbers should be provided in addition, where applicable): Preparations of a kind used in animal feeding (HS code(s): 2309); Animal feeding stuffs (ICS code(s): 65.120)
4.	Regions or countries likely to be affected, to the extent relevant or practicable: ☒ All trading partners ☐ Specific regions or countries:
5.	Title of the notified document: Food Additives Permitted in Feed and Drinking Water of Animals; Chromium DL-methionine Chelate; Final Amendment; Order. Language(s): English. Number of pages: 3 https://www.federalregister.gov/d/2026-16942 https://members.wto.org/crnattachments/2026/SPS/USA/26_04407_00_e.pdf
6.	Description of content: 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 chromium DLMethionine chelate as a nutritional source of chromium in cattle feed. This action is in response to a food additive petition filed by Zinpro Corp. This order is effective 19 August 2026.
7.	Objective and rationale: ☒ food safety, ☒ animal health, ☐ plant protection, ☐ protect humans from animal/plant pest or disease, ☐ protect territory from other damage from pests.
8.	Is there a relevant international standard? If so, identify the standard: ☐ Codex Alimentarius Commission (e.g. title or serial number of Codex standard or related text): ☐ World Organization for Animal Health (OIE) (e.g. Terrestrial or Aquatic Animal Health Code, chapter number): ☐ International Plant Protection Convention (e.g. ISPM number): ☒ None Does this proposed regulation conform to the relevant international standard? ☐ Yes ☐ No If no, describe, whenever possible, how and why it deviates from the international standard:
9.	Other relevant documents and language(s) in which these are available:

10.	Proposed date of adoption (dd/mm/yy): 19 August 2026 Proposed date of publication (dd/mm/yy): 19 August 2026
11.	Proposed date of entry into force: ☐ Six months from date of publication, and/or (dd/mm/yy): 19 August 2026 ☒ Trade facilitating measure
12.	Final date for comments: ☐ Sixty days from the date of circulation of the notification and/or (dd/mm/yy): Either electronic or written objections and requests for a hearing on the final amendment must be submitted by 18 September 2026. Agency or authority designated to handle comments: ☐ National Notification Authority, ☐ National Enquiry Point. Address, fax number and e-mail address (if available) of other body: 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 18 September 2026. 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-2017-F-4399 for "Food Additives Permitted in Feed and Drinking Water of Animals; Chromium DL-methionine Chelate". 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.
13.	Text(s) available from: ☐ National Notification Authority, ☐ National Enquiry Point. Address, fax number and e-mail address (if available) of other body: Text can be found in the Federal Register, Vol. 91, No. 159, Page 53524 or on the Internet at: https://www.federalregister.gov/d/2026-16942