



September 11, 2026

Zoetis Inc.
Attention: John Hallberg, DVM, PhD
Senior Director of Regulatory Affairs
333 Portage St.
Kalamazoo, MI 49007

Re: Emergency Use Authorization 006700 Revised Fact Sheets

Dear Dr. Hallberg:

Please refer to your Emergency Use Authorization (EUA) for the emergency use of DECTOMAX/DECTOMAX-CA1 (doramectin injection) for the prevention and treatment of infestations caused by *Cochliomyia hominivorax* larvae (myiasis) in dairy cattle (lactating dairy cows, dry dairy cows, and replacement dairy heifers 20 months of age and older), except for calves that will be processed for veal, and, for the prevention of infestations caused by *Cochliomyia hominivorax* larvae (myiasis) in horses one year and older, swine, sheep except for lactating sheep, and deer, issued on May 19, 2026, pursuant to Section 564 of the Federal Food, Drug, and Cosmetic Act (FD&C Act) (21 U.S.C. § 360bbb-3).

We have updated the authorized “Fact Sheet: Emergency Use Authorization of Dectomax/Dectomax-CA1 (doramectin injection) for New World Screwworm (NWS) Dairy Cattle, Swine, Sheep, and Deer” and “Fact Sheet: Emergency Use Authorization of Dectomax/Dectomax-CA1 (doramectin injection) for New World Screwworm (NWS) Horses” to clarify the Directions sections and clarify the cattle subclasses which are subject to this EUA (i.e., lactating dairy cows, dry dairy cows, and replacement dairy heifers 20 months of age and older), compared to the conditional and original approvals. The updated authorized “Fact Sheet: Emergency Use Authorization of Dectomax/Dectomax-CA1 (doramectin injection) for New World Screwworm (NWS) Dairy Cattle, Swine, Sheep, and Deer” and “Fact Sheet: Emergency Use Authorization of Dectomax/Dectomax-CA1 (doramectin injection) for New World Screwworm (NWS) Horses” are attached to this correspondence for your reference with the new revision date of 09/10/2026. Please update your website with the current authorized Fact Sheets.

If you submit correspondence relating to this letter, your correspondence should reference the date and EUA submission number identified in this letter. For correspondence relating to this letter, or if you have questions or need assistance with the product development process or project updates, contact your project manager. If you do not know who your project manager is, send an email to AskONAPEPM@fda.hhs.gov.

Sincerely,

{see appended electronic signature page}

Matthew A. Lucia, DVM

Director

Office of New Animal Product Evaluation

Center for Veterinary Medicine

Enclosure:
Revised Fact Sheets (2)