

05 May 2022

To Whom This May Concern

Subject: **Letter of Certification that Zoetis is Manufacturing According to FDA Regulations**

Zoetis certifies that Lincomix® 20, a Type A Medicated Article manufactured by Zoetis Suzhou Manufacturing Co. Ltd., Suzhou, Jiangsu, China facility in and imported by Zoetis Inc., is manufactured in accordance with US Food and Drug Administration (FDA) current Good Manufacturing Practices regulations as set forth in 21 CFR parts 210 ([CURRENT GOOD MANUFACTURING PRACTICE IN MANUFACTURING, PROCESSING, PACKING, OR HOLDING OF DRUGS; GENERAL](#)) and 226 ([CURRENT GOOD MANUFACTURING PRACTICE FOR TYPE A MEDICATED ARTICLES](#)).

Type A Medicated Articles are regulated as drug products (pharmaceuticals) in the United States. As such, all Zoetis manufacturing sites, and active ingredient manufacturing sites are registered with the FDA, have FDA approved New Animal Drug Applications (NADAs) or Veterinary Master Files (VMFs), and are routinely inspected by the FDA. The controls and oversight employed by Zoetis Quality Operations at Zoetis manufacturing sites, active ingredient manufacturers, and raw material suppliers assure that Zoetis products meet the requirements of the FDA Food Safety Modernization Act (FSMA) including the Foreign Supplier Verification Program (FSVP) requirements.

All Zoetis locations conform to the same Zoetis Quality Management System with Corporate Standards for compliance with the Health Authority requirements for the markets in which the products are sold. Each lot is analytically tested and verified to confirm to the registered specifications for those markets. Lot specific Certificates of Analysis (COA) provides the details of this testing. All lot records including the manufacturing and testing are reviewed and approved by the Zoetis site Quality Assurance. COA's are available upon request to customers.

Sincerely,



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FOR ANIMALS. FOR HEALTH. FOR YOU.