REPORTING ADVERSE EFFECTS FROM SOLENSIA/LIBRELA

Paws Over Profits

LFPCSA
LOST AND FOUND PETS IN CAMBRIDGE AND SURROUNDING AREAS

Please make sure any side effects are reported not only to Zoetis, but also, to the Regulator, as this will significantly help with the class action.

Region	Zoetis	Regulator
UK	• Zoetis UK **To report adverse reactions, you can either:** - Use this form, - call 0345 300 8034, - email customersupportuk@zoetis.com	⚠ **VMD – Temporary changes to reporting:** You cannot currently report adverse reactions directly to VMD, but are required to do so to Zoetis, who must then notify VMD within 30 days (more info can be found here). We will notify you as soon as the form has been re-opened.
EU/EEA	• Zoetis website *(please scroll down to 'International Contact Information' and choose your region/country)*	• HMA National Contacts
Australia	Zoetis AU **E-mail:** productsupport.au@zoetis.com **Phone:** 1800 814 883	• APVMA Adverse Experience Reporting Program • APVMA Adverse Experience Reporting Form **E-mail:** enquiries@apvma.gov.au **Phone:** +61 2 6770 2300
Canada	• Zoetis Canada • Online Adverse Event Reporting Form **E-mail:** productsupportservice@zoetis.com **Phone:** 1-800-461-0917	• Health Canada: Report a side effect to a veterinary drug - form **E-mail:** pv-vet@hc-sc.gc.ca **Phone:** 1-877-838-7322
US	• Zoetis US • Product Support Online Form **Phone:** 1-888-963-8471	• How Do I Report Adverse Events for Animal Drugs and Devices? Download Form 1932a, (instructions under the link above), fill in and send to CVM1932a@fda.hhs.gov **E-mail:** AskCVM@fda.hhs.gov **Phone:** 1-888-FDA-VETS (1-888-332-8387)
Other	• Zoetis website *(please scroll down to 'International Contact Information' and choose your region/country)*	• WHO - Regulators' list

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