

# Scabigard Suspension for Cutaneous Administration - product defect recall alert

2.4.2026 - | Her Majesty's Revenue and Customs

## **Product defect recall alert for Scabigard Suspension for Cutaneous Administration - Vm 42058/5173 and Vm 60021/3020 by Zoetis Belgium S.A.**

We wish to inform wholesalers, veterinarians and end users that Zoetis has initiated a Class II recall for Scabigard Suspension for Cutaneous Administration, UK(GB) Vm 42058/5173 by Zoetis UK Limited and UK(NI) Vm 60021/3020 by Zoetis Belgium S.A.

The reason for this recall is that the affected batches may not demonstrate the expected signs of take following administration.

This recall applies to the following batches:

| <b>Batch number</b> | <b>Date of manufacture</b> | <b>Expiry date</b> |
|---------------------|----------------------------|--------------------|
| 7482095A01          | June 2024                  | May 2026           |
| 7482095A02          | June 2024                  | May 2026           |
| 7482095B01          | June 2024                  | May 2026           |
| 7482095B02          | June 2024                  | May 2026           |
| 7482095D01          | June 2024                  | May 2026           |
| 7482095D02          | June 2024                  | May 2026           |
| 7482096A01          | August 2024                | July 2026          |
| 748209701           | January 2025               | December 2026      |
| 748209702           | January 2025               | December 2026      |

Zoetis is contacting wholesalers and veterinarians to immediately review inventory and quarantine products subject to this recall.

End users should return any affected product to the veterinary practice from which it was purchased.

For further information regarding this recall, please contact Zoetis on 0345 300 8034 or at [customersupportUK@zoetis.com](mailto:customersupportUK@zoetis.com).

<https://www.gov.uk/government/news/scabigard-suspension-for-cutaneous-administration-product-defect-recall-alert>