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Committee for Veterinary Medicinal Products

# Combined VeDDRA list of clinical terms for reporting suspected adverse events in animals and humans to veterinary medicinal products

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| Revision agreed by Pharmacovigilance Working Party (PhVWP-V) | 20-21 May 2025 |
| Adoption by CVMP | 10-12 June 2025 |
| Date for coming into effect | 1 October 2025 |

## Glossary of abbreviations used in VeDDRA:

**SOC:** System organ class

**HLT:** High level term

**PT:** Preferred term

**LLT:** Low level term

**NOS:** Not otherwise specified

**LLT term types:**

**A:** animal

**C:** common

**H:** human

The combined VeDDRA list of clinical terms for reporting suspected adverse events in animals and humans to veterinary medicinal products is available at the following link.

https://www.ema.europa.eu/en/documents/other/combined-veddra-list-clinical-terms-reporting-suspected-adverse-events-animals-humans-veterinary-medicinal-products_en.xlsx

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