



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

18 June 2026
EMA/CVMP/PhVWP/10418/2009-Rev.17
Committee for Veterinary Medicinal Products

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

Revision agreed by Pharmacovigilance Working Party (PhVWP-V)	27 May 2026
Adoption by CVMP	16-18 June 2026
Date for coming into effect	1 October 2026

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



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



VeDDRA_Rev17_EVV
etv22.xlsx