EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

5 December 2025
EMA/CVMP/380363/2025
Veterinary Medicines Division

MRL summary opinion[1]

Lidocaine

Porcine

On 4 December 2025, the Committee for Veterinary Medicinal Products adopted an opinion[2] recommending, by consensus, the modification of maximum residue limits for lidocaine in porcine in Table 1 ('Allowed substances') of the Annex to Commission Regulation (EU) No 37/2010, in accordance with the following table:

Pharmaco-logically active substance	Marker residue	Animal species	MRLs	Target tissues	Other provisions	Therapeutic classification
Lidocaine	NOT APPLICABLE	Equidae	No MRL required	NOT APPLICABLE	For local/regional anaesthesia only.	Local anaesthetic
		Porcine	No MRL required	NOT APPLICABLE	For cutaneous and epilesional use in piglets up to 7 days of age only. For injection into scrotum, testicles and spermatic cord in piglets up to 7 days of age only.	
	Lidocaine	Bovine	150 µg/kg 200 g/kg 1 µg/kg	Muscle Fat Liver	NOT APPLICABLE	

[1] Summaries of opinion are published without prejudice to the Commission Decision.
[2] Applicants may request re-examination of any CVMP opinion, provided they notify the European Medicines Agency in writing of their intention to request such re-examination within 15 days of receipt of the opinion

			200 µg/kg 30 µg/kg	Kidney Milk		