

# PENETHAONE Powder and solvent for suspension for injection for cattle

Authorised

- Penethamate hydriodide

## Product identification

**Medicine name:**

PENETHAONE Powder and solvent for suspension for injection for cattle

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**Active substance:**

Penethamate hydriodide

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**Target species:**

Cattle (lactating cow)

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**Route of administration:**

Intramuscular use

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## Product details

**Active substance and strength:**

Penethamate hydriodide

236.30 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Powder and solvent for suspension for injection

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**Withdrawal period by route of administration:**

**Intramuscular use:**

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**Cattle (lactating cow)**

- Meat and offal. 4 day

- Milk. no withdrawal period

Meat and offal: 4 days; Milk: 2,5 days/60 hours.

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ01CE90

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Italy

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**Available in:**

Italy

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**Package description:**

box containing 10 x 10,000,000 IU powder vial and 10 x 36 ml solvent vial

box containing 5 x 10,000,000 IU powder vial and 5 x 36 ml solvent vial

box containing 1 x 10,000,000 IU powder vial and 1 x 36 ml solvent vial

box containing 10 x 5,000,000 IU powder vial and 10 x 18 ml solvent vial

box containing 5 x 5,000,000 IU powder vial and 5 x 18 ml solvent vial

box containing 1 x 5,000,000 IU powder vial and 1 x 18 ml solvent vial

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Divasa Farmavic S.A.

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**Marketing authorisation date:**

16/06/2016

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**Manufacturing sites for batch release:**

Divasa Farmavic S.A.

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**Responsible authority:**

Ministry Of Health

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**Authorisation number:**

104774

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**Date of authorisation status change:**

16/06/2016

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**Reference member state:**

Spain

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**Procedure number:**

ES/V/0226/001

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**Concerned member states:**

Austria Belgium Bulgaria Czechia Denmark France Germany Greece  
Hungary Iceland Ireland Italy Lithuania Netherlands Norway Poland  
Portugal Romania Slovakia Sweden United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

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Summary of Product Characteristics