

Vetalgin vet. 500 mg/ml Injektionsvätska, lösning

Authorised

- Metamizole sodium monohydrate

Product identification

Medicine name:

Vetalgin vet. 500 mg/ml Injektionsvätska, lösning

Active substance:

Metamizole sodium monohydrate

Target species:

Dog

Horse

Cattle

Pig

Route of administration:

Intramuscular use

Intravenous use

Product details

Active substance and strength:

Metamizole sodium monohydrate

500.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:**Intramuscular use:**

-

Horse

- Meat and offal. 21 day

Intravenös behandling 2-3 ggr med 8 timmars intervall med en dos av högst 20-50 mg/kg kroppsvikt:

- Meat and offal. 9 day

Intravenös behandling som engångsdos med 20-50 mg/kg kroppsvikt

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Cattle

- Milk. 4 day

Intravenös eller intramuskulär behandling som engångsdos med 20-50 mg/kg kroppsvikt

- Meat and offal. 21 day

Intravenös eller intramuskulär behandling som engångsdos med 20-50 mg/kg kroppsvikt

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Pig

- Meat and offal. 15 day

Intravenös eller intramuskulär som engångsdos med 20-50 mg/kg kroppsvikt

Intravenous use:

-

Horse

- Meat and offal. 9 day

Intravenös behandling som engångsdos med 20-50 mg/kg kroppsvikt

- Meat and offal. 21 day

Intravenös behandling 2-3 ggr med 8 timmars intervall med en dos av högst 20-50 mg/kg kroppsvikt:

-

Cattle

- Milk. 4 day

Intravenös eller intramuskulär behandling som engångsdos med 20-50 mg/kg kroppsvikt

- Meat and offal. 21 day

Intravenös eller intramuskulär behandling som engångsdos med 20-50 mg/kg kroppsvikt

-

Pig

- Meat and offal. 15 day

Intravenös eller intramuskulär som engångsdos med 20-50 mg/kg kroppsvikt

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN02BB02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Sweden

Available in:

Sweden

Package description:

Available only in Swedish

Available only in Swedish

Available only in Swedish

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Intervet International B.V.

Marketing authorisation date:

3/02/1939

Manufacturing sites for batch release:

Intervet International B.V.

Oriola Sweden AB

Responsible authority:

Swedish Medical Products Agency

Authorisation number:

1678

Date of authorisation status change:

3/02/1939

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet

Documents

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.