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ESTRUMATE FLACON UNIDOSE

Authorised

- Cloprostenol sodium

Product identification

Medicine name:

ESTRUMATE FLACON UNIDOSE

Active substance:

Cloprostenol sodium

Target species:

Cattle (cow)

Horse (mare)

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Cloprostenol sodium

0.26 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Cattle (cow)

- Meat and offal. 1 day
- Milk. 0 day

-

Horse (mare)

- Meat and offal. 2 day
- Milk. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QG02AD90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Available in:

France

Package description:

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Intervet

Marketing authorisation date:

19/04/1989

Manufacturing sites for batch release:

Vet Pharma Friesoythe GmbH

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/0030299 9/1989

Date of authorisation status change:

19/04/2009

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

Documents

Package Leaflet and Labelling

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.