HY-50 Vet. 17 mg/ml, solution for injection

Authorised

- SODIUM HYALURONATE

Product identification

Medicine name:
HY-50 Vet. 17 mg/ml, solution for injection

Active substance:
SODIUM HYALURONATE

Target species:
Horse

Route of administration:
Intravenous use
Intraarticular use

Product details

Active substance and strength:
SODIUM HYALURONATE
17.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intravenous use:

- **Horse**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM09AX01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Denmark

Available in:
Denmark

Package description:
Pre-filled syringe, 12 x 3 ml
Pre-filled syringe, 1 x 3 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Dechra Regulatory B.V.

Marketing authorisation date:
21/07/1999

Manufacturing sites for batch release:
Eurovet Animal Health B.V.
Dales Pharmaceuticals Limited

Responsible authority:
Danish Medicines Agency

Authorisation number:
30718

Date of authorisation status change:
21/07/1999

Reference member state:
Sweden

Procedure number:
SE/V/0106/001

Concerned member states:
Belgium Denmark Germany Spain United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 19/06/2025
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Package Leaflet