

SynVet-50; 50 mg solution for injection for horses

Authorised

- SODIUM HYALURONATE

Product identification

Medicine name:

SynVet-50; 50 mg solution for injection for horses

Active substance:

SODIUM HYALURONATE

Target species:

Horse

Route of administration:

Intraarticular use

Product details

Active substance and strength:

SODIUM HYALURONATE

50.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:

Solution for injection

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

-

Horse

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM09AX01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Package description:

Single-dose glass syringe barrel with luer tip and rigid tip capType 1 glass syringe, lubricated with dimethicone. Styrene-butadiene rubber cap.Bromobutyl rubber plunger.Available as multipack with six single dose cartons aggregated and overwrapped with plastic foil and label.

Single-dose glass syringe barrel with luer tip and rigid tip capType 1 glass syringe, lubricated with dimethicone. Styrene-butadiene rubber cap.Bromobutyl rubber plunger.Available as single dose syringe in a sealed translucent plastic blister packed in one carton folding box.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:

Equi Pharma Limited

Marketing authorisation date:

18/07/2014

Manufacturing sites for batch release:

CROMA-PHARMA GmbH

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA22962/001/001

Date of authorisation status change:

18/07/2014

Reference member state:

Ireland

Procedure number:

IE/V/0580/001

Concerned member states:

Austria Belgium Denmark France Germany Netherlands Norway Sweden
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: 7/06/2026

Download

Package Leaflet

Combined File of all Documents