Equiparin 5.000 IU/100 g gel for horses

Authorised

- Levomenthol
- Hydroxyethyl salicylate
- Heparin sodium

Product identification

Medicine name:
Equiparin 5.000 IU/100 g gel for horses

Active substance:
Levomenthol

Hydroxyethyl salicylate

Heparin sodium

Target species:
Horse

Route of administration:
Cutaneous use

Product details

Active substance and strength:
Levomenthol

0.50 gram(s) / 100.00 gram(s)

Hydroxyethyl salicylate
5.00 gram(s) / 100.00 gram(s)

Heparin sodium
5000.00 international unit(s) / 100.00 gram(s)

Pharmaceutical form:
Gel

Withdrawal period by route of administration:
Cutaneous use:

- **Horse**
  - Meat and offal. 0 day
  - Milk. no withdrawal period

Not permitted for use in mares producing milk for human consumption

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Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM02AC99

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

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Authorisation status:
Valid

Authorised in:
Iceland

Package description:
(ID1) 285 gram(s): unspecified outer container with 1 Bottle (Polyethylen) with 285 gram(s)
(ID2) 1710 gram(s): unspecified outer container with 6 Bottle (Polyethylen) each with 285 gram(s)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
aniMedica GmbH

Marketing authorisation date:
30/12/2009

Manufacturing sites for batch release:
aniMedica GmbH

Responsible authority:
Icelandic Medicines Agency

Authorisation number:
IS/2/09/018/01

Date of authorisation status change:
15/01/2015

Reference member state:
Germany

Procedure number:
DE/V/0128/001

Concerned member states:
Austria Hungary Iceland Ireland Poland 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