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Equioxx 8.2 mg/g - Oral paste

Authorised

- Firocoxib

Product identification

Medicine name:
Equioxx 8.2 mg/g - Oral paste

Active substance:
Firocoxib

Target species:
Horse

Route of administration:
Oral use

Product details

Active substance and strength:
Firocoxib

Presentation_strength:0.06 g Reference:Hse Index:0

Pharmaceutical form:
Oral paste

Withdrawal period by route of administration:

Oral use:

- **Horse**
  - Meat and offal. 26 day 26 days

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM01AH90

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

Authorisation status:
Valid

Authorised in:
Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Package description:
Packaging:Pre-filled syringe (PP), Package_size:14 syringes, Content:7.32 g
Packaging:Pre-filled syringe (PP), Package_size:7 syringes, Content:7.32 g
Packaging:Pre-filled syringe (PP), Package_size:1 syringe, Content:7.32 g

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:
Audevard

Marketing authorisation date:
25/06/2008

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
25/06/2008

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

Documents

Combined File of all Documents

English (PDF)
Published on: 10/04/2026
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