

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:

11/10/2010

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

Documents

Combined File of all Documents

English (PDF)

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