Isoflurane 1000 mg/g Inhalation Vapour, liquid.

Authorised

- Isoflurane

Product identification

Medicine name:
Isoflurane 1000 mg/g Inhalation Vapour, liquid.
Isoflurin 1000 mg/g vloeistof voor inhalatiedamp

Active substance:
Isoflurane

Target species:
Horse
Reptile
Ornamental bird
Dog
Cat
Ferret
Rat
Hamster
Chinchilla
Guinea pig
Gerbil
Mouse

Route of administration:

Inhalation use

Product details

Active substance and strength:
Isoflurane
1000.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Inhalation vapour, liquid

Withdrawal period by route of administration:
Inhalation use:

- **Horse**
  - Meat and offal. 2 day
- **Reptile**
- **Ornamental bird**
- **Dog**
- **Cat**
- **Ferret**
- **Rat**
- **Hamster**
- **Chinchilla**
- **Guinea pig**
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Gerbil

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Mouse

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN01AB06

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

Authorisation status:
Valid

Authorised in:
Netherlands

Available in:
Netherlands

Package description:
Amber coloured glass bottle (Type III) containing 250ml isoflurane. Bottles are closed with a black polypropylene screw cap.
Amber coloured glass bottle (Type III) containing 100ml isoflurane. Bottles are closed with a black polypropylene screw cap.

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:
Vetpharma Animal Health S.L.

Marketing authorisation date:
18/12/2015

Manufacturing sites for batch release:
Chemical Iberica Productos Veterinarios S.L.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 117045

Date of authorisation status change:
28/01/2022

Reference member state:
Netherlands

Procedure number:
NL/V/0196/001

Concerned member states:
Belgium Bulgaria Croatia Cyprus Czechia Denmark Finland France Greece Hungary Ireland Latvia Lithuania Luxembourg Norway Poland Portugal Slovakia Slovenia Spain Sweden United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

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