Somulose 400 mg/ml + 25 mg/ml solution for injection

Authorised

- Secobarbital sodium
- Cinchocaine hydrochloride

Product identification

Medicine name:
Somulose 400 mg/ml + 25 mg/ml solution for injection

Active substance:
Secobarbital sodium
Cinchocaine hydrochloride

Target species:
Cattle
Dog
Horse (non food-producing)
Cat

Route of administration:
Intravenous use

Product details

Active substance and strength:
Secobarbital sodium

400.00 milligram(s) / 1.00 millilitre(s)

Cinchocaine hydrochloride
25.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN05CB02

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

Authorisation status:
Valid

Authorised in:
Ireland

Available in:
Ireland

Package description:
50 ml in amber type I glass vial with red chlorobutyl rubber stopper and aluminium seal in cardboard box carton.
25 ml in amber type I glass vial with red chlorobutyl rubber stopper and aluminium seal in cardboard box carton.

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:
Dechra Regulatory B.V.

Marketing authorisation date:

6/10/2006

Manufacturing sites for batch release:
Eurovet Animal Health B.V.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA22622/018/001

Date of authorisation status change:
6/10/2006

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

Documents

Summary of Product Characteristics