

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