Adrenacaine Solution for Injection for Cattle

Authorised

- Procaine hydrochloride
- Adrenaline hydrogen tartrate

Product identification

Medicine name:
Adrenacaine Solution for Injection for Cattle

Active substance:
Procaine hydrochloride

Adrenaline hydrogen tartrate

Target species:
Cattle

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Procaine hydrochloride
50.00 milligram(s) / 1.00 millilitre(s)

Adrenaline hydrogen tartrate

0.04 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Subcutaneous use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN01BA52

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

Authorisation status:
Valid

Authorised in:
Ireland

Available in:
Ireland

Package description:
The product will be supplied in 100 ml amber type I glass vials with bromobutyl bungs and aluminium caps.

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:
Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:
17/07/2009

Manufacturing sites for batch release:
Norbrook Manufacturing Limited
Norbrook Laboratories Limited

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA22664/087/001

Date of authorisation status change:
17/07/2009

Reference member state:
Ireland

Procedure number:
IE/V/0597/001

Concerned member states:
United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

Combined File of all Documents