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Ubropen 600 mg intramammary suspension for lactating cows

Authorised

- Benzylpenicillin (procaine) monohydrate

Product identification

Medicine name:
Ubropen 600 mg intramammary suspension for lactating cows

Active substance:
Benzylpenicillin (procaine) monohydrate

Target species:
Cattle (lactating cow)

Route of administration:
Intramammary use

Product details

Active substance and strength:
Benzylpenicillin (procaine) monohydrate
600.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:
Intramammary suspension

Withdrawal period by route of administration:

Intramammary use:

- **Cattle (lactating cow)**
  - Meat and offal. 3 day
  - Milk. 6 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51CE09

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

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Authorisation status:
Valid

Authorised in:
Ireland

Package description:
Pack size: 100 x 10 g with 100 cleaning towels. White intramammary syringe (LDPE) with a double tip (LDPE) packed in a cardboard container.
Pack size: 40 x 10 g with 40 cleaning towels. White intramammary syringe (LDPE) with a double tip (LDPE) packed in a cardboard container.
Pack size: 20 x 10 g with 20 cleaning towels. White intramammary syringe (LDPE) with a double tip (LDPE) packed in a cardboard container.
Pack size: 5 x 10 g with 5 cleaning towels. White intramammary syringe (LDPE) with a double tip (LDPE) packed in a cardboard container.
Pack size: 3 x 10 g with 3 cleaning towels. White intramammary syringe (LDPE) with a double tip (LDPE) packed in a cardboard container.

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:
Vetcare Oy

Marketing authorisation date:
22/04/2016

Manufacturing sites for batch release:
KELA Kempisch Laboratorium Kela Laboratoria

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10832/002/001

Date of authorisation status change:
22/04/2016

Reference member state:
Finland

Procedure number:
FI/V/0110/001

Concerned member states:
Austria Belgium Denmark Estonia France Germany Iceland Ireland Italy Latvia Lithuania Netherlands Norway Poland Spain Sweden

United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

Package Leaflet

Labelling