Ultrapen LA 300 mg/ml suspensija injekcijām liellopiem un cūkām

Awtorizzat

- Benzylpenicillin procaine

Product identification

Isem tal-mediċina:
Ultrapen LA 300 mg/ml suspensija injekcijām liellopiem un cūkām

Sustanza attiva:
Disponibbli biss fi English

Speċi li fuqhom ser jintużа l-prodott:
Disponibbli biss fi Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Slovenian Finnish Swedish Icelandic Norwegian
Disponibbli biss fi Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian

Metodu ta' amministrazzjoni:
Użu għal taħt il-ġilda
Użu għal ġol-muskoli

Product details

Sustanza attiva / Qawwa:
Disponibbli biss fi English
300.00 milligram(s) / 1.00 millilitre(s)

Forma farmaċewtika:
Suspensjoni għall-injezzjoni

Withdrawal period by route of administration:
Użu għal taħt il-ġilda:

- **Cattle**
  - Meat and offal. 13 day

Nav reģistrēts subkutānai lietošanai dzīvniekiem, kuru pienu paredzēts izmantot cilvēku uzturā.

Użu għal ġol-muskoli:

- **Cattle**
  - Meat and offal. 23 day
  - Milk. 132 hour
- **Pig**
  - Meat and offal. 10 day

Kodiċi veterinarju anatomiku terapewtiku kimiku (ATCvet):
QJ01CE09

Status legali tal-provvista:
Disponibbli biss fi Czech Estonian English French Italian Latvian Lithuanian Portuguese Romanian Slovenian Finnish Swedish Icelandic Norwegian

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Status tal-awtorizzazzjoni:
Valid

Authorised in:
Disponibbli biss fi Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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Deskrizzjoni tal-pakkett:
Disponibbli biss fi Latvian

Disponibbli biss fi Latvian

Additional information

Entitlement type:
Disponibbli biss fi English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

Bażi ġuridika tal-awtorizzazzjoni tal-prodott:
Disponibbli biss fi English French Italian Latvian Lithuanian Norwegian

Id-detentur tal-awtorizzazzjoni għat-tqegħid fis-suq:
Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:
3/12/1998

Siti ta' manifattura b'rilaxx tal-lott:
Norbrook Manufacturing Limited
Norbrook Laboratories Limited

Awtorità responsabbli:
Food And Veterinary Service

Numru tal-awtorizzazzjoni:
V/NRP/98/0939

Data tal-bidla fl-istatus tal-awtorizzazzjoni:
3/12/1998

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

Documents

Sommarju tal-karatteristiċi tal-prodott

Dan id-dokument ma jeżistix f'dan il-lingwa (Malti). Tista 'ssibha f'lingwa oħra hawn taħt.

Tikkettar

Dan id-dokument ma jeżistix f'dan il-lingwa (Malti). Tista 'ssibha f'lingwa oħra hawn taħt.

Fuljett ta’ tagħrif

Dan id-dokument ma jeżistix f'dan il-lingwa (Malti). Tista 'ssibha f'lingwa oħra hawn taħt.

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