UBAC Emulsion for injection

Authorised

- Streptococcus uberis, strain 5616, lipoteichoic acid from Biofilm Adhesion Component

Product identification

Medicine name:
UBAC Emulsion for injection

Active substance:
Streptococcus uberis, strain 5616, lipoteichoic acid from Biofilm Adhesion Component

Target species:
Cattle

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Streptococcus uberis, strain 5616, lipoteichoic acid from Biofilm Adhesion Component
Presentation_strength:≥ 1 RP Reference:Hse Index:0

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Not applicable. 0 day Zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI02AB

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

Authorisation status:
Valid

Authorised in:
Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Package description:
Packaging:Vial (PET), Package_size:1 vial, Content:25 doses (50 ml)
Packaging:Vial (PET), Package_size:1 vial, Content:5 doses (10 ml)
Packaging:Vial (glass), Package_size:20 vials, Content:1 dose (2 ml)
Packaging:Vial (PET), Package_size:1 vial, Content:50 doses (100 ml)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Laboratorios Hipra, S.A.

Marketing authorisation date:

26/07/2018

Manufacturing sites for batch release:
Laboratorios Hipra S.A.

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
26/07/2018

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

Documents

Combined File of all Documents

English (PDF)
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