SYNULOX Bolus 400 mg/100 mg filmom obalené tablety

Authorised

- Potassium clavulanate
- Amoxicillin trihydrate

Product identification

Medicine name:
SYNULOX Bolus 400 mg/100 mg filmom obalené tablety

Active substance:
Potassium clavulanate

Amoxicillin trihydrate

Target species:
Cattle (calf)

Route of administration:
Oral use

Product details

Active substance and strength:
Potassium clavulanate
120.00 milligram(s) / 1.00 Tablet

Amoxicillin trihydrate

460.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Film-coated tablet

Withdrawal period by route of administration:
Oral use:

- **Cattle (calf)**
  - Meat and offal. 9 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CR02

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

Authorisation status:
Valid

Authorised in:
Slovakia

Available in:
Slovakia

Package description:
Available only in Slovak

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Zoetis Ceska Republika s.r.o.

Marketing authorisation date:
28/04/1994

Manufacturing sites for batch release:
Haupt Pharma Latina S.r.l.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/226/94-S

Date of authorisation status change:
28/04/1994

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

Documents

Combined File of all Documents

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