NEOSKILAB Solution for injection

Authorised

- NEOSTIGMINE METHYLSULFATE

Product identification

Medicine name:
NEOSKILAB Solution for injection
NEOSKILAB SOLUTION INJECTABLE POUR BOVINS, OVINS, CAPRINS ET CHEVAUX

Active substance:
NEOSTIGMINE METHYLSULFATE

Target species:
Cattle
Sheep
Goat
Horse

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
NEOSTIGMINE METHYLSULFATE
1.50 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. no withdrawal period withdrawal period is 0 days
  - Milk. no withdrawal period withdrawal period is 0 days
- **Sheep**
  - Meat and offal. no withdrawal period withdrawal period is 0 days
  - Milk. no withdrawal period withdrawal period is 0 days
- **Goat**
  - Meat and offal. no withdrawal period withdrawal period is 0 days
  - Milk. no withdrawal period withdrawal period is 0 days
- **Horse**
  - Meat and offal. no withdrawal period withdrawal period is 0 days
  - Milk. no withdrawal period withdrawal period is 0 days

Subcutaneous use:

- **Cattle**
  - Meat and offal. no withdrawal period withdrawal period is 0 days
  - Milk. no withdrawal period withdrawal period is 0 days
- **Sheep**
  - Meat and offal. no withdrawal period withdrawal period is 0 days
  - Milk. no withdrawal period withdrawal period is 0 days
- **Goat**
  - Meat and offal. no withdrawal period withdrawal period is 0 days
  - Milk. no withdrawal period withdrawal period is 0 days
- **Horse**

- Meat and offal. no withdrawal period — withdrawal period is 0 days
- Milk. no withdrawal period — withdrawal period is 0 days

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN07AA01

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

Authorisation status:
Valid

Authorised in:
France

Package description:
box containing 1 vial of 25 ml

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:
Labiana Life Sciences S.A.

Marketing authorisation date:
29/03/2021

Manufacturing sites for batch release:
Labiana Life Sciences S.A.

Responsible authority:
National Veterinary Medicines Agency

Authorisation number:
FR/V/9656943 6/2021

Date of authorisation status change:
29/03/2021

Reference member state:
Spain

Procedure number:
ES/V/0389/001

Concerned member states:
Belgium Croatia Cyprus Estonia France Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Portugal Romania

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

Documents

Summary of Product Characteristics

English (PDF)
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