NEOSKILAB Solution for injection

Geautoriseerd

- NEOSTIGMINE METHYLSULFATE

Product identification

Naam van het geneesmiddel:
NEOSKILAB Solution for injection

Werkzame stof:
Alleen beschikbaar in English

Doeldiersoort(en):
Rund
Schaap
Geit
Paard

Toedieningsweg:
Intramusculair gebruik
Subcutaan gebruik

Product details

Werkzame stof / Sterkte:
Alleen beschikbaar in English
1.50 milligram(s) / 1.00 millilitre(s)

Farmaceutische vorm:

Oplossing voor injectie

Withdrawal period by route of administration:

Intramusculair gebruik:

- **Rund**
  - Meat and offal. no withdrawal period withdrawal period is 0 days
  - Milk. no withdrawal period withdrawal period is 0 days
- **Schaap**
  - Meat and offal. no withdrawal period withdrawal period is 0 days
  - Milk. no withdrawal period withdrawal period is 0 days
- **Geit**
  - Meat and offal. no withdrawal period withdrawal period is 0 days
  - Milk. no withdrawal period withdrawal period is 0 days
- **Paard**
  - Meat and offal. no withdrawal period withdrawal period is 0 days
  - Milk. no withdrawal period withdrawal period is 0 days

Subcutaan gebruik:

- **Rund**
  - Meat and offal. no withdrawal period withdrawal period is 0 days
  - Milk. no withdrawal period withdrawal period is 0 days
- **Schaap**
  - Meat and offal. no withdrawal period withdrawal period is 0 days
  - Milk. no withdrawal period withdrawal period is 0 days
- **Geit**
  - Meat and offal. no withdrawal period withdrawal period is 0 days
  - Milk. no withdrawal period withdrawal period is 0 days

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

Anatomisch therapeutisch chemische veterinaire classificatie (ATCvet code) :
QN07AA01

Afleverstatus:
Alleen beschikbaar in Czech Estonian English French Italian Latvian Portuguese Slovenian Finnish Swedish Icelandic Norwegian

Status toelating:
Valid

Authorised in:
Luxemburg

Package description:
Alleen beschikbaar in English

Additional information

Entitlement type:
Alleen beschikbaar in English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

Rechtsgrondslag productvergunning:
Alleen beschikbaar in English Italian Latvian Norwegian

Handelsvergunninghouder:
Labiana Life Sciences S.A.

Marketing authorisation date:
7/12/2021

Productielocaties partijvrijgifte:

Labiana Life Sciences S.A.

Verantwoordelijke instantie:
Ministere De La Sante Division De La Pharmacie Et Des Medicaments

Toelatingsnummer:
V 006/21/12/2171

Wijzigingsdatum status toelating:
7/12/2021

Rapporterende lidstaat:
Spanje

Procedurenummer:
ES/V/0389/001

Betrokken lidstaten:
België Kroatië Cyprus Estland Frankrijk Griekenland Hongarije Ierland Italië Letland Litouwen Luxemburg Portugal Roemenië

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

Documents

Samenvatting van de productkenmerken

Dit document bestaat niet in deze taal (Nederlands). Je kunt het hieronder vinden in een andere taal.
