NEOSKILAB Solution for injection

Viðurkennt

- NEOSTIGMINE METHYLSULFATE

Product identification

Heiti lyfs:
NEOSKILAB Solution for injection

Virkt efni:
Aðeins í boði í English

Dýrategundir:
Nautgripir
Sauðkind
Geit
Hestur

Íkomuleið:
Til notkunar í vöðva
Til notkunar undir húð

Product details

Virkt efni / Styrkur:
Aðeins í boði í English
1.50 milligram(s) / 1.00 millilitre(s)

Lyfjaform:
Stungulyf, lausn

Withdrawal period by route of administration:

Til notkunar í vöðva:

- **Nautgripir**
  - Kjöt og innmatur. no withdrawal period withdrawal period is 0 days
  - Mjólk. no withdrawal period withdrawal period is 0 days
- **Sauðkind**
  - Kjöt og innmatur. no withdrawal period withdrawal period is 0 days
  - Mjólk. no withdrawal period withdrawal period is 0 days
- **Geit**
  - Kjöt og innmatur. no withdrawal period withdrawal period is 0 days
  - Mjólk. no withdrawal period withdrawal period is 0 days
- **Hestur**
  - Kjöt og innmatur. no withdrawal period withdrawal period is 0 days
  - Mjólk. no withdrawal period withdrawal period is 0 days

Til notkunar undir húð:

- **Nautgripir**
  - Kjöt og innmatur. no withdrawal period withdrawal period is 0 days
  - Mjólk. no withdrawal period withdrawal period is 0 days
- **Sauðkind**
  - Kjöt og innmatur. no withdrawal period withdrawal period is 0 days
  - Mjólk. no withdrawal period withdrawal period is 0 days
- **Geit**
  - Kjöt og innmatur. no withdrawal period withdrawal period is 0 days
  - Mjólk. no withdrawal period withdrawal period is 0 days
- **Hestur**

- Kjöt og innmatur. no withdrawal period	withdrawal period is 0 days
- Mjólk. no withdrawal period	withdrawal period is 0 days

ATC flokkun (dýralyf):
QN07AA01

Lögformleg staða:
Ávísunarskylt dýralyf

Staða markaðsleyfis:
Gilt

Authorised in:
Lúxemborg

Áletrun:
Aðeins í boði í English

Additional information

Entitlement type:
Marketing Authorisation

Lagagrundvöllur vöruleyfis:
Aðeins í boði í English Italian Latvian Norwegian

Markaðsleyfishafi:
Labiana Life Sciences S.A.

Marketing authorisation date:
7/12/2021

Framleiðandi sem ber ábyrgð á lokasamþykkt:
Labiana Life Sciences S.A.

Ábyrgt yfirvald:
Ministere De La Sante Division De La Pharmacie Et Des Medicaments

Markaðsleyfisnúmer:
V 006/21/12/2171

Dagsetning leyfisbreytingar:
7/12/2021

Umsjónarland (RMS):
Spánn

Númer verkferlis:
ES/V/0389/001

Þátttökulönd (CMS):
Belgía Króatía Kýpur Eistland Frakkland Grikkland Ungverjaland Írland Ítalía Lettland Litáen Lúxemborg Portúgal Rúmenía

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

Documents

Samantekt á eiginleikum lyfs

Þetta skjal er ekki til á þessu tungumáli (íslenzkan). Þú getur fundið það á öðru tungumáli hér að neðan.
