

NEOSKILAB Solution for injection

Geautoriseerd

- NEOSTIGMINE METHYLSULFATE

Product identification

Naam van het geneesmiddel:

NEOSKILAB Solution for injection

NEOSKILAB 1,5 mg/ml ενέσιμο διάλυμα για βοοειδή, πρόβατα, αίγες και άλογα

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:

Cyprus

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:

23/05/2021

Productielocaties partijvrijgifte:

Labiana Life Sciences S.A.

Verantwoordelijke instantie:

Ministry Of Agriculture Rural Development And Environment

Toelatingsnummer:

CY00833V

Wijzigingsdatum status toelating:

23/05/2021

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.

Source URL: <https://medicines.health.europa.eu/veterinary/600000038483>