

NEOSKILAB Solution for injection

Oprávněný

- NEOSTIGMINE METHYLSULFATE

Product identification

Názov lieku:

NEOSKILAB Solution for injection

NEOSKILAB 1.5 mg/ml solutie injectabila pentru bovine ovine caprine cabaline

Účinná látka:

Dostupné len v [English](#)

Cieľové druhy:

Dostupné len v [Bulgarian](#) [Spanish](#) [Czech](#) [Danish](#) [German](#) [Estonian](#) [Greek](#) [English](#) [French](#) [Italian](#) [Latvian](#) [Lithuanian](#) [Hungarian](#) [Dutch](#) [Romanian](#) [Slovenian](#) [Swedish](#) [Icelandic](#) [Norwegian](#)

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Cesta podania:

Intramuskulárne použitie

Subkutánne použitie

Product details

Účinná látka a sila:

Dostupné len v [English](#)

1.50 milligram(s) / 1.00 millilitre(s)

Lieková forma:

Injekčný roztok

Withdrawal period by route of administration:

Intramuskulárne použitie:

• 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

Subkutánne použitie:

• 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

Anatomicko-terapeuticko-chemický veterinárny kód (ATCvet):

QN07AA01

Právny stav dodávky:

Dostupné len v [Czech](#) [Estonian](#) [English](#) [French](#) [Italian](#) [Latvian](#) [Portuguese](#) [Slovenian](#) [Finnish](#) [Swedish](#) [Icelandic](#) [Norwegian](#)

Stav registrácie:

Valid

Authorised in:

Rumunsko

Opis balenia:

Dostupné len v [English](#)

Additional information

Entitlement type:

Dostupné len v [English](#) [French](#) [Croatian](#) [Italian](#) [Latvian](#) [Finnish](#) [Swedish](#) [Icelandic](#) [Norwegian](#)

Právny základ pre registráciu produktu:

Dostupné len v [English](#) [Italian](#) [Latvian](#) [Norwegian](#)

Držiteľ rozhodnutia o registrácii:

Labiana Life Sciences S.A.

Marketing authorisation date:

21/11/2021

Miesta výroby na uvoľnenie šarže:

Labiana Life Sciences S.A.

Zodpovedný orgán:

Institute For Control Of Biological Products And Veterinary Medicines

Číslo registrácie:

210194

Dátum zmeny stavu registrácie:

21/11/2021

Referenčný členský štát:

Španielsko

Číslo postupu:

ES/V/0389/001

Dotknuté členské štáty:

Belgicko Chorvátsko Cyprus Estónsko Francúzsko Grécko Maďarsko Írsko
Taliano Lotyšsko Litva Luxembursko Portugalsko Rumunsko

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

Documents

Súhrn charakteristických vlastností lieku

Tento dokument v tomto jazyku (slovenčina) neexistuje. Nájdete ho v inom jazyku nižšie.

Source URL: <https://medicines.health.europa.eu/veterinary/600000038501>