

OXYTOCIN/PROVET 10IU/ML ΕΝΕΣΙΜΟ ΔΙΑΛΥΜΑ

Oprávněný

- Oxytocin

Product identification

Názov lieku:

OXYTOCIN/PROVET 10IU/ML ΕΝΕΣΙΜΟ ΔΙΑΛΥΜΑ

Účinná látka:

Dostupné len v English

Cieľové druhy:

Dostupné len v Bulgarian Spanish Danish German Greek English Italian Latvian
Lithuanian Hungarian Romanian Swedish

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

Dostupné len v Bulgarian Spanish Czech Danish German Estonian Greek English
French Italian Latvian Lithuanian Hungarian Dutch Romanian Swedish Icelandic
Norwegian

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Italian Latvian Lithuanian Hungarian Dutch Romanian Swedish Icelandic Norwegian

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

Cesta podania:

Intramuskulárne použitie

Subkutánne použitie

Product details

Účinná látka a sila:

Dostupné len v [English](#)

10.00 international unit(s) / 1.00 millilitre(s)

Lieková forma:

Injekčný roztok

Withdrawal period by route of administration:

Intramuskulárne použitie:

- **Equid (female)**

- Meat and offal, milk. 0 day

- **Cattle (cow)**

- Meat and offal, milk. 0 day

- **Pig (sow)**

- Meat and offal. 0 day

- **Sheep (ewe)**

- Meat and offal, milk. 0 day

- **Goat**

- Meat and offal, milk. 0 day

- **Dog (bitch)**

- Not applicable. no withdrawal period

- **Cat**

- Not applicable. no withdrawal period

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- Not applicable. no withdrawal period

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- Not applicable. no withdrawal period

Anatomicko-terapeuticko-chemický veterinárny kód (ATCvet):

QH01BB02

Právny stav dodávky:

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

Stav registrácie:

Valid

Authorised in:

Grécko

Opis balenia:

Dostupné len v [Greek](#)

Dostupné len v [Greek](#)

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](#)

Držiteľ rozhodnutia o registrácii:

Provet S.A.

Marketing authorisation date:

27/02/1985

Miesta výroby na uvoľnenie šarže:

Provet S.A.

Zodpovedný orgán:

National Organization For Medicines

Číslo registrácie:

66842/20-09-2012/K-0036301

Dátum zmeny stavu registrácie:

10/10/2021

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

Source URL: <https://medicines.health.europa.eu/veterinary/600000107585>