

CALFOSET injekčný roztok pre kone, hovädzí dobytok, ovce, kozy, ošípané a odstavné a odstavné

Pooblaščeno

- Calcium gluconate monohydrate
- CALCIUM GLYCEROPHOSPHATE
- Magnesium chloride hexahydrate

Product identification

Ime zdravila:

CALFOSET injekčný roztok pre kone, hovädzí dobytok, ovce, kozy, ošípané a odstavné a odstavné

Učinkovina:

Na voljo samo v [English](#)

Na voljo samo v [English](#)

Na voljo samo v [English](#)

Ciljne živalske vrste:

govedo

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

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

Na voljo samo v [Bulgarian](#) [Spanish](#) [Czech](#) [Danish](#) [German](#) [Estonian](#) [Greek](#) [English](#) [French](#) [Italian](#) [Latvian](#) [Lithuanian](#) [Hungarian](#) [Dutch](#) [Romanian](#) [Finnish](#) [Swedish](#)

Icelandic Norwegian

Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish

Icelandic Norwegian

Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish

Icelandic Norwegian

Pot uporabe:

Subkutana uporaba

Intravenska uporaba

intramuskularna uporaba

Product details

Učinkovina / Jakost:

Na voljo samo v English

328.20 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English

81.30 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English

41.80 milligram(s) / 1.00 millilitre(s)

Farmaceutska oblika:

Raztopina za injiciranje

Withdrawal period by route of administration:

Subkutana uporaba:

- **govedo**

- All relevant tissues. 0 day

- **Sheep**

- All relevant tissues. 0 day

- **Goat**

- All relevant tissues. 0 day

- **Pig**

- All relevant tissues. 0 day

- **Pig (weaned piglet)**

- All relevant tissues. 0 day

Intravenska uporaba:

- **Horse**

- All relevant tissues. no withdrawal period

All relevant tissues: Zero days

- **govedo**

- All relevant tissues. no withdrawal period

All relevant tissues: Zero days

- **Sheep**

- All relevant tissues. no withdrawal period

All relevant tissues: Zero days

- **Goat**

- All relevant tissues. no withdrawal period

All relevant tissues: Zero days

- **Pig**

- All relevant tissues. no withdrawal period

All relevant tissues: Zero days

intramuskularna uporaba:

- **govedo**

- All relevant tissues. no withdrawal period

All relevant tissues: Zero days

- **Sheep**

- All relevant tissues. no withdrawal period

All relevant tissues: Zero days

- **Goat**

- All relevant tissues. no withdrawal period

All relevant tissues: Zero days

- **Pig**

- All relevant tissues. no withdrawal period

All relevant tissues: Zero days

- **Pig (weaned piglet)**

- All relevant tissues. no withdrawal period

All relevant tissues: Zero days

Anatomsko-terapevtsko-kemična veterinarska oznaka (ATCvet):

QA12AX

Pravni status za dobavo / izdajo zdravila:

Zdravilo, ki se izdaja na veterinarski recept

Status dovoljenja:

Valid

Authorised in:

Na voljo samo v [Spanish](#) [Czech](#) [German](#) [Estonian](#) [English](#) [French](#) [Italian](#) [Dutch](#) [Portuguese](#) [Slovak](#) [Swedish](#) [Icelandic](#) [Norwegian](#)

Opis ovojnine:

Na voljo samo v [Slovak](#)

Na voljo samo v [Slovak](#)

Additional information

Entitlement type:

Na voljo samo v [English](#) [French](#) [Croatian](#) [Italian](#) [Latvian](#) [Finnish](#) [Swedish](#) [Icelandic](#) [Norwegian](#)

Pravna podlaga za izdajo dovoljenja za promet z zdravilom:

Na voljo samo v [English](#) [French](#) [Italian](#) [Latvian](#) [Norwegian](#)

Imetnik dovoljenja za promet z zdravilom:

KRKA tovarna zdravil d.d. Novo mesto

Marketing authorisation date:

28/03/1995

Proizvodna mesta, odgovorna za sproščanje serij:

Krka d.d. Novo Mesto

Pristojni organ:

Institute For State Control Of Veterinary Biologicals And Medicaments

Številka dovoljenja :

96/0014/95-S

Datum spremembe statusa dovoljenja:

28/03/1995

Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinarski medicini obiščite stran: www.adrreports.eu/vet

Documents

Combined File of all Documents

Ta dokument ne obstaja v tem jeziku (slovenščina). Spodaj ga najdete v drugem jeziku.

Source URL: <https://medicines.health.europa.eu/veterinary/600000023188>