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DUPHALYTE

Ni odobreno

- DL-Tryptophan
- L-THREONINE
- PHENYLALANINE DL
- L-METHIONINE
- L-lysine hydrochloride
- L-LEUCINE
- L-ISOLEUCINE
- L-HISTIDINE MONOHYDROCHLORIDE MONOHYDRATE
- SODIUM L-GLUTAMATE
- L-CYSTEINE HYDROCHLORIDE MONOHYDRATE
- L-arginine hydrochloride
- GLUCOSE, ANHYDROUS
- Potassium chloride
- Magnesium sulfate
- Calcium chloride hexahydrate
- Dexpanthenol
- Nicotinamide
- Cyanocobalamin
- Pyridoxine hydrochloride
- Riboflavin sodium phosphate hydrate
- Thiamine hydrochloride

Informacije o zdravilu

Ime zdravila:
DUPHALYTE

Učinkovina:

Na voljo samo v English
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Ciljne živalske vrste:

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

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Pot uporabe:
Intravenska uporaba
Subkutana uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
0.01 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
0.02 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
0.03 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
0.01 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
0.03 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
0.04 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
0.01 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
0.01 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
0.04 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
0.01 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
0.03 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
45.46 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
0.20 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
0.29 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
0.23 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
0.05 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
1.50 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
0.05 microgram(s) / 1.00 millilitre(s)

Na voljo samo v English
0.10 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
0.04 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
0.10 milligram(s) / 1.00 millilitre(s)

Farmacevtska oblika:

Raztopina za injiciranje

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):

QA11A

Pravni status za dobavo/izdajo zdravila:

Zdravilo, ki se izdaja na veterinarski recept

Status dovoljenja za promet z zdravilom:

Surrendered

Ima dovoljenje za promet v:

Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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Opis ovojnine zdravila:

Na voljo samo v Bulgarian

Dodatne informacije

Vrsta dovoljenja:
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 Italian Latvian Lithuanian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Zoetis Belgium

Datum dovoljenja za promet z zdravilom:
9/11/2008

Proizvodna mesta, odgovorna za sproščanje serij:
Zoetis Manufacturing & Research Spain S.L.

Pristojni organ:
Bulgarian Food Safety Authority

Številka dovoljenja za promet z zdravilom:
0022-2050-12.06.2013

Datum spremembe statusa dovoljenja za promet:
14/05/2026

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

Dokumenti

Povzetek glavnih značilnosti zdravila

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