

Morbital Plus 400 mg/1 ml roztwór do wstrzykiwań

Registruotas

- Pentobarbital sodium

Vaisto identifikaciniai duomenys

Vaisto pavadinimas:

Morbital Plus 400 mg/1 ml roztwór do wstrzykiwań

Veiklioji medžiaga:

Pateikiama tik [Anglų](#)

Paskirties gyvūnų rūšis (-ys):

Galvijai
Žirgas
Kiaulė
Šuo
Katė

Naudojimo būdas:

Leisti į širdį
Leisti į veną
Leisti į pilvaplėvės ertmę

Vaisto duomenys

Veiklioji medžiaga ir stiprumas:

Pateikiama tik Anglų
400.00 miligramai / 1.00 mililitrai

Vaisto forma:

Injekcinis tirpalas

Pasitraukimo laikotarpis administravimo būdu:

Leisti į širdį:

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Galvijai

- All relevant tissues. no withdrawal period

Not applicable. Appropriate measures must be taken to ensure that tissues of animals that have received this product, as well as animal by-products derived from these animals, do not enter the food chain and are not used for consumption by humans or other animals.

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Žirgas

- All relevant tissues. no withdrawal period

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Kiaulė

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Leisti į veną:

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Galvijai

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Leisti į pilvaplėvės ertmę:

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Galvijai

- All relevant tissues. no withdrawal period

Not applicable. Appropriate measures must be taken to ensure that tissues of animals that have received this product, as well as animal by-products derived from these animals, do not enter the food chain and are not used for consumption by humans or other animals.

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Anatominės, terapinės ir cheminės veterinarinių vaistų klasifikacijos (ATCvet) kodas:

QN51AA01

Tiekimo teisinis statusas:

Parduodama tik su veterinariniu receptu

Registracijos statusas:

Valid

Registruota:

Pateikiama tik [Ispanų](#) [Čekų](#) [Vokiečių](#) [Estų](#) [Anglų](#) [Prancūzų](#) [Italų](#) [Nyderlandų](#) [Portugalų](#) [Slovakų](#) [Švedų](#) [Islandų](#) [Norwegian](#)

Pakuotės aprašymas:

Pateikiama tik [Lenkų](#)

Papildoma informacija

Teisių tipas:

Pateikiama tik [Anglų](#) [Prancūzų](#) [Kroatų](#) [Italų](#) [Latvių](#) [Suomių](#) [Švedų](#) [Islandų](#) [Norwegian](#)

Vaisto registracijos teisinis pagrindas:

Pateikiama tik Anglų Italų Latvių Norwegian

Registruotojas:

Biowet Pulawy Sp. z o.o.

Rinkodaros leidimo data:

8/12/2022

Serijos išleidimo gamybos vietos:

Biowet Pulawy Sp. z o.o.

Atsakinga institucija:

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Registracijos numeris:

3218

Registracijos statuso pasikeitimo data:

8/12/2022

Pranešimai apie įtariamus nepageidaujamus reiškinius: www.adrreports.eu/vet

Dokumentai

Veterinarinio vaisto aprašas

Šis dokumentas neegzistuoja šia kalba (@Language). Jį galite rasti kita kalba žemiau.

Pakuotės lapelis

Šis dokumentas neegzistuoja šia kalba (@Language). Jį galite rasti kita kalba žemiau.

Ženkinimas

Šis dokumentas neegzistuoja šia kalba (@Language). Jį galite rasti kita kalba žemiau.