

# Morbital Plus 400 mg/1 ml roztwór do wstrzykiwań

Registrováno

- Pentobarbital sodium

## Identifikace přípravku

### Název léčiva:

Morbital Plus 400 mg/1 ml roztwór do wstrzykiwań

### Léčivá látka:

Dostupné pouze v [English](#)

### Cílové druhy:

Skot

Kůň

Prase

Pes

Kočka

### Cesta podání:

Intrakardiální podání

Intravenózní podání

Intraperitoneální podání

## Údaje o přípravku

### Léčivá látka a síla:

Dostupné pouze v English  
400.00 milligram(s) / 1.00 millilitre(s)

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**Léková forma:**

Injekční roztok

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**Ochranná lhůta podle cesty podání:**

**Intrakardiální podání:**

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**Skot**

- 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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**Kůň**

- 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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**Prase**

- 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.

**Intravenózní podání:**

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**Skot**

- 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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#### **Kůň**

- 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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#### **Prase**

- 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.

### **Intraperitoneální podání:**

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#### **Skot**

- 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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#### **Kůň**

- 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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**Prase**

- 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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**Anatomicko-terapeuticko-chemická klasifikace veterinárních léčiv (ATCvet):**  
QN51AA01

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**Právní status výdeje:**

Na předpis

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**Stav registrace:**

Valid

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**Registrováno v:**

Polsko

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**Popis balení:**

Dostupné pouze v Polish

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## Doplňující informace

**Typ oprávnění:**

Dostupné pouze v English French Croatian Italian Latvian Finnish Swedish Icelandic  
Norwegian

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**Právní základ registrace přípravku:**

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

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**Držitel rozhodnutí o registraci:**

Biowet Pulawy Sp. z o.o.

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**Datum registrace:**

8/12/2022

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**Výrobní místa s propouštěním šarží:**

Biowet Pulawy Sp. z o.o.

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**Příslušný orgán:**

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

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**Registrační číslo:**

3218

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**Datum změny stavu registrace:**

8/12/2022

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Informace o nežádoucích účincích veterinárních léčivých přípravků naleznete na adrese [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Dokumenty

Souhrn údajů o přípravku

Tento dokument v tomto jazyce neexistuje (čeština). Níže jej najdete v jiném jazyce.

Příbalová informace

Tento dokument v tomto jazyce neexistuje (čeština). Níže jej najdete v jiném jazyce.

Označení na obalu

Tento dokument v tomto jazyce neexistuje (čeština). Níže jej najdete v jiném jazyce.