

# Morbital Plus 400 mg/1 ml roztwór do wstrzykiwań

Zugelassen

- Pentobarbital sodium

## Produktidentifikation

**Arzneimittel:**

Morbital Plus 400 mg/1 ml roztwór do wstrzykiwań

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**Wirkstoff:**

Verfügbar nur in [englisch](#)

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**Zieltierarten:**

Rind

Pferd

Schwein

Hund

Katze

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**Art der Anwendung:**

intracardiale Anwendung

intravenöse Anwendung

intraperitoneale Anwendung

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## Produktdetails

**Wirkstoff und Stärke:**

Verfügbar nur in englisch  
400.00 milligram(s) / 1.00 millilitre(s)

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**Darreichungsform:**

Injektionslösung

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**Wartezeit(en) per Anwendungsart:**

**intracardiale Anwendung:**

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**Rind**

- Alle Zielgewebe. 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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**Pferd**

- Alle Zielgewebe. 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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**Schwein**

- Alle Zielgewebe. 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.

**intravenöse Anwendung:**

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**Rind**

- Alle Zielgewebe. 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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### **Pferd**

- Alle Zielgewebe. 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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### **Schwein**

- Alle Zielgewebe. 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.

## **intraperitoneale Anwendung:**

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### **Rind**

- Alle Zielgewebe. 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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### **Pferd**

- Alle Zielgewebe. 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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### **Schwein**

- Alle Zielgewebe. 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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### **Anatomisch-therapeutisch-chemischer Veterinär-code (ATCvet-Code):**

QN51AA01

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### **Abgaberegung:**

Verfügbar nur in tschechisch estnisch englisch französisch italienisch lettisch litauisch portugiesisch rumänisch slowenisch finnisch schwedisch isländisch Norwegian

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### **Zulassungsstatus:**

Zulassung gültig

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### **Zugelassen in:**

Polen

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### **Packungsbeschreibung:**

Verfügbar nur in polieren

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## Zusätzliche Informationen

### **Anspruchstyp:**

Verfügbar nur in englisch französisch kroatisch italienisch lettisch finnisch schwedisch isländisch Norwegian

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**Rechtsgrundlage der Produktzulassung:**

Verfügbar nur in [englisch](#) [italienisch](#) [lettisch](#) [Norwegian](#)

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**Zulassungsinhaber:**

Biowet Pulawy Sp. z o.o.

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**Zulassungsdatum:**

8/12/2022

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**Für die Chargenfreigabe zuständige Produktionsstätten:**

Biowet Pulawy Sp. z o.o.

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**Zuständige Behörde:**

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

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**Zulassungsnummer:**

3218

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**Tag der Änderung des Zulassungsstatus:**

8/12/2022

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Dokumente

Zusammenfassung der Merkmale des Arzneimittels

Dieses Dokument existiert in dieser Sprache nicht (deutsch). Sie finden es in einer anderen Sprache unten.

## Packungsbeilage

Dieses Dokument existiert in dieser Sprache nicht (deutsch). Sie finden es in einer anderen Sprache unten.

## Etikettierung

Dieses Dokument existiert in dieser Sprache nicht (deutsch). Sie finden es in einer anderen Sprache unten.