Morbital Plus 400 mg/1 ml roztwór do wstrzykiwań

Autorizado

- Pentobarbital sodium

Identificación del medicamento

Nombre del medicamento:
Morbital Plus 400 mg/1 ml roztwór do wstrzykiwań

Principio activo:
Disponible únicamente en English

Especies de destino:
Bovino
Caballos
Porcino
Perros
Gatos

Vía de administración:
Vía intracardiaca
Vía intravenosa
Vía intraperitoneal

Datos del medicamento

Principio activo y concentración:

Disponible únicamente en English
400.00 Miligramo(s) / 1.00 Mililitro(s)

Forma farmacéutica:

Solución inyectable

Tiempo de espera por vía de administración:

Vía intracardiaca:

- **Bovino**

  - 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.

- **Caballos**

  - 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.

- **Porcino**

  - 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.

Vía intravenosa:

- **Bovino**

- 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.

- **Caballos**
  - 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.

- **Porcino**
  - 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.

Vía intraperitoneal:

- **Bovino**
  - 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.

- **Caballos**

- 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.

- **Porcino**
  - 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.

código químico anatómico-terapéutico para medicamentos veterinarios (ATCvet):
QN51AA01

Condiciones de dispensación:
Disponible únicamente en Czech Estonian English French Italian Latvian Lithuanian Portuguese Romanian Slovenian Finnish Swedish Icelandic Norwegian

Estado de la autorización:
Autorizado

Autorizado en:
Polonia

Descripción del formato:
Disponible únicamente en Polish

Información adicional

Tipo legal de la autorización:
Disponible únicamente en English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

Base legal de la autorización del medicamento:
Disponible únicamente en English Italian Latvian Norwegian

Titular de la autorización de comercialización:
Biowet Pulawy Sp. z o.o.

Fecha de autorización de comercialización:
8/12/2022

Fabricante responsable de la liberación del lote:
Biowet Pulawy Sp. z o.o.

Autoridad responsable:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Número de autorización:
3218

Fecha de modificación del estado de la autorización:
8/12/2022

Para consultar notificaciones de presuntos efectos adversos diríjase a www.adrreports.eu/vet

Documentos

Ficha técnica o resumen de las características del producto

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Prospecto

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Etiquetado

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