

**PARTICULARS TO APPEAR ON THE OUTER PACKAGE Cardboard Box**

**1. NAME OF THE VETERINARY MEDICINAL PRODUCT**

Depocillin 300 mg/ml Suspension for injection

**2. STATEMENT OF ACTIVE SUBSTANCES**

Procaine benzylpenicillin 300 mg/ml

**3. PACKAGE SIZE**

100 ml







**4. TARGET SPECIES**

Cattle, horses, sheep, pigs, dogs, cats.

**5. INDICATIONS**

**6. ROUTES OF ADMINISTRATION**

Cattle, horses, sheep, pigs: Intramuscular use.  
Dogs, cats: Subcutaneous use.

 <b>Cattle</b> 500 kg 20 ml	 <b>Horse</b> 500 kg 20 ml	 <b>Sheep</b> 50 kg 2.5 ml
 <b>Pigs</b> 50 kg 2.5 ml	 <b>Dogs</b> 10 kg 1 ml	 <b>Cats</b> 5 kg 0.5 ml

**7. WITHDRAWAL PERIODS**

Meat and offal:

Cattle, pigs, sheep: 5 days for the treatment duration 3-5 days  
7 days for the treatment duration 6-7 days

Horses: Not to be used in horses intended for human consumption. Treated horses may never be slaughtered for human consumption. The horse must have been declared as not intended for human consumption under national horse passport legislation.

Milk:

Cattle: 264 hours (11 days)

Sheep, horses: Not authorised for use in animals producing milk for human consumption.

## **8. EXPIRY DATE**

Exp. {mm/yyyy}

Once opened use within 28 days.

## **9. SPECIAL STORAGE PRECAUTIONS**

Store in a refrigerator. Do not freeze.

Keep the vial in the outer carton in order to protect from light.

## **10. THE WORDS “READ THE PACKAGE LEAFLET BEFORE USE”**

Read the package leaflet before use.

## **11. THE WORDS “FOR ANIMAL TREATMENT ONLY”**

For animal treatment only.

## **12. THE WORDS “KEEP OUT OF THE SIGHT AND REACH OF CHILDREN”**

Keep out of the sight and reach of children.

## **13. NAME OF THE MARKETING AUTHORISATION HOLDER**

Intervet International B.V.

## **14. MARKETING AUTHORISATION NUMBERS**

Vm 06376/4081

## **15. BATCH NUMBER**

Lot {number}

## **PARTICULARS TO APPEAR ON THE IMMEDIATE PACKAGE VIAL LABEL**

### **1. NAME OF THE VETERINARY MEDICINAL PRODUCT**

Depocillin 300 mg/ml Suspension for injection

### **2. STATEMENT OF ACTIVE SUBSTANCES**

Procaine benzylpenicillin 300 mg/ml

### **3. TARGET SPECIES**

Cattle, horses, sheep, pigs, dogs, cats.

### **4. ROUTES OF ADMINISTRATION**

Read the package leaflet before use.

### **5. WITHDRAWAL PERIODS**

Meat and offal:

Cattle, pigs, sheep: 5 days for the treatment duration 3-5 days  
7 days for the treatment duration 6-7 days

Horses: Not to be used in horses intended for human consumption. Treated horses may never be slaughtered for human consumption. The horse must have been declared as not intended for human consumption under national horse passport legislation.

Milk:

Cattle: 164 hours (11 days)  
Sheep, horses: Not authorised for use in animals producing milk for human consumption.

### **6. EXPIRY DATE**

Exp. {mm/yyyy}

Once opened use within 28 days.

Once broached use by...

### **7. SPECIAL STORAGE PRECAUTIONS**

Store in a refrigerator.

Do not freeze.

Keep the vial in the outer carton in order to protect from light.

### **8. NAME OF THE MARKETING AUTHORISATION HOLDER**

Intervet International B.V.

## 9. BATCH NUMBER

Lot {number}

## **PARTICULARS TO APPEAR ON THE PACKAGE LEAFLET:**

### **PACKAGE LEAFLET**

#### **1. Name of the veterinary medicinal product**

Depocillin 300 mg/ml Suspension for injection

#### **2. Composition**

Each ml contains:

##### **Active substance:**

Procaine benzylpenicillin 300 mg

##### **Excipients:**

Methyl parahydroxybenzoate 1.10 mg

White to off-white suspension.

#### **3. Target species**

Cattle, horses, sheep, pigs, dogs, cats.

#### **4. Indications for use**

The veterinary medicinal product is indicated for the treatment of infections caused by bacteria sensitive to penicillin.

#### **5. Contraindications**

Do not use in cases of hypersensitivity to the active substance or to any of the excipients.

Do not use in rabbits, guinea pigs, hamsters or gerbils. Caution is advised in its use in any other small herbivores.

Not effective against beta-lactamase producing organisms.

#### **6. Special warnings**

##### Special warnings:

Occasionally in sucking and fattening pigs, administration of products containing procaine penicillin may cause a transient pyrexia, vomiting, shivering, listlessness and incoordination. Additionally in pregnant sows and gilts, a vulval discharge which could be associated with abortion has been reported.

After absorption, benzylpenicillin poorly penetrates biological membranes (e.g., blood-brain barrier) since it is ionised and poorly lipid soluble. Use of the product for treatment of meningitis or CNS infections due to e.g., *Streptococcus suis* or *Listeria monocytogenes* may not be efficacious. Furthermore, benzylpenicillin penetrates mammalian cells poorly and hence this product may have little effect in treating intracellular pathogens e.g., *Listeria monocytogenes*.

Elevated MIC values or bi-modal distribution profiles suggesting acquired resistance have been reported for the following bacteria:

- *Glaesserella parasuis*, *Staphylococcus* spp. causing MMA/PPDS, *Streptococcus* spp. and *S. suis* in pigs;
- *Fusobacterium necrophorum* causing metritis and *Mannheimia haemolytica* (only in some member states), as well as *Bacteroides* spp., *Staphylococcus chromogenes*, *Actinobacillus lignieresii* and *Trueperella pyogenes* in cattle;
- *S. aureus*, coagulase negative Staphylococci and *Enterococcus* spp. in dogs;
- *Staphylococcus aureus* and *Staphylococcus felis* in cats.

Use of the veterinary medicinal product may result in a lack of clinical efficacy when treating infections caused by these bacteria.

Special precautions for safe use in the target species:

Not recommended for intravenous or intrathecal administration.

Special precautions to be taken by the person administering the veterinary medicinal product to animals:

Penicillin and cephalosporins may cause hypersensitivity (allergy) following injection, inhalation, ingestion or skin contact. Hypersensitivity to penicillin may lead to cross reactions to cephalosporins and vice versa. Allergic reactions to these substances are occasionally serious.

1. Do not handle this product if you know you are sensitised, or if you have been advised not to work with such preparations.
2. Handle this product with great care to avoid exposure, taking all recommended precautions.
3. If you develop symptoms following exposure, such as a skin rash, you should seek medical advice and show the doctor this warning. Swelling of the face, lips or eyes or difficulty with breathing are more serious symptoms and require urgent medical attention.

Wash hands after use.

Pregnancy and lactation:

No special precautions necessary.

Interaction with other medicinal products and other forms of interaction:

Antagonism between the product and bacteriostatic preparations may occur. Resistant bacteria, particularly gram-negative, that show a cross-resistance with other beta-lactam antibiotics might occur.

Overdose:

Penicillin is a compound with a very high therapeutic ratio. It is very unlikely that an overdose of the product will have adverse effects on the treated animal.

## 7. Adverse events

Target species:

Cattle, horses, sheep, dogs, cats:

Very rare (<1 animal / 10,000 animals treated, including isolated reports):
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Hypersensitivity reaction, Anaphylaxis <sup>1</sup> ; Injection site reaction (e.g. swelling, pain)
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Pigs

Very rare (<1 animal / 10,000 animals treated, including isolated reports):
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Hypersensitivity reaction, Anaphylaxis <sup>1</sup> ; Vomiting <sup>2</sup> ; Injection site reaction (e.g. swelling, pain)
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<sup>1</sup> May be life-threatening. Potentially fatal reactions associated with the administration of procaine penicillin in horses have been observed. If such reactions occur appropriate treatment is recommended.

<sup>2</sup> In sucking and fattening pigs.

Systemic toxic effects have been observed in young piglets, which are transient but can be potentially lethal, especially at higher doses.

Reporting adverse events is important. It allows continuous safety monitoring of a product. If you notice any side effects, even those not already listed in this package leaflet, or you think that the medicine has not worked, please contact, in the first instance, your veterinarian. You can also report any adverse events to the marketing authorisation holder or the local representative of the marketing authorisation holder using the contact details at the end of this leaflet, or via your national reporting system at:

Website: <https://www.gov.uk/report-veterinary-medicine-problem/animal-reacts-medicine>

e-mail: [adverse.events@vmd.gov.uk](mailto:adverse.events@vmd.gov.uk)

## 8. Dosage for each species, routes and method of administration

Horses and cattle 12 mg/kg body weight, sheep and pigs 15 mg /kg. Intramuscular use. Dogs and cats 30 mg/kg (1 ml per 10 kg body weight). Subcutaneous use.

Suggested doses are:

Cattle	500 kg	20	ml
Horses	500 kg	20	ml
Sheep	50 kg	2.5	ml
Pigs	50 kg	2.5	ml
Dogs	10 kg	1	ml
Cats	5 kg	0.5	ml

Clean the area of the injection site and swab with spirit.

The treatment duration is 3 to 7 days and should be repeated at 24 hour intervals.

Do not use the same injection site more than once during a course of treatment.

Do not inject more than 20 ml per injection site in cattle.

Do not inject more than 5 ml per injection site in pigs and sheep.

## **9. Advice on correct administration**

Shake the container well before use.

## **10. Withdrawal periods**

Meat and offal:

Cattle, pigs, sheep: 5 days for the treatment duration 3-5 days

7 days for the treatment duration 6-7 days

Horses: Not to be used in horses intended for human consumption. Treated horses may never be slaughtered for human consumption. The horse must have been declared as not intended for human consumption under national horse passport legislation.

Milk:

Cattle: 264 hours (11 days)

Sheep, horses: Not authorised for use in animals producing milk for human consumption.

## **11. Special storage precautions**

Keep out of the sight and reach of children.

Store in a refrigerator (2 °C – 8 °C).

Do not freeze.

Keep the vial in the outer carton in order to protect from light.

Do not use this veterinary medicinal product after the expiry date which is stated on the label after Exp. The expiry date refers to the last day of that month.

Shelf life of the veterinary medicinal product as packaged for sale: 3 years.

Shelf life after first opening the immediate packaging: 28 days.

## **12. Special precautions for disposal**

Medicines should not be disposed of via wastewater.

Use take-back schemes for the disposal of any unused veterinary medicinal product or waste materials derived thereof in accordance with local requirements and with any applicable national collection systems. These measures should help to protect the environment.

Ask your veterinary surgeon or pharmacist how to dispose of medicines no longer required.

### **13. Classification of veterinary medicinal products**

Veterinary medicinal product subject to prescription.

### **14. MARKETING AUTHORISATION NUMBERS AND PACK SIZES**

Vm 06376/4081

Pack size:

Multidose glass or PET vials of 100 ml.

### **15. PID link (Do not print heading)**

*[The following statement must be included where reference to the European Union Product Database is included on the product information. This statement is relevant to both UK(GB) and UK(NI) products:]*

Find more product information by searching for the 'Product Information Database' on [www.gov.uk](http://www.gov.uk).

### **16. Contact details**

Marketing authorisation holder:

Intervet International B.V.  
Wim de Körverstraat 35  
5831 AN Boxmeer  
Netherlands

Manufacturer responsible for batch release:

Intervet Productions S.r.l.  
Via Nettunense Km 20, 300  
04011 Aprilia  
Italy

Local representative:

MSD Animal Health UK Limited  
Walton Manor, Walton  
Milton Keynes  
MK7 7AJ, United Kingdom

Contact details to report suspected adverse reactions:

**UK(GB)**

MSD Animal Health UK Ltd.  
Tel.: +44 (0)1908 685685

**UK(NI)**

Intervet Ireland Ltd.  
Tel.: +353 (0)1 2970220

Distributor in Northern Ireland:

Intervet Ireland Ltd.  
Magna Drive, Magna Business Park, Citywest Road  
Dublin 24, Ireland

For any information about this veterinary medicinal product, please contact the local representative of the marketing authorisation holder.

**17. Other information**

POM-V

*Gavin Hall*  
Approved: 28 October 2025