

## **PARTICULARS TO APPEAR ON THE OUTER PACKAGE**

Cardboard box of 1 vial of 100 ml

Cardboard box of 1 vial of 250 ml

Cardboard box of 12 vials of 100 ml

Cardboard box of 12 vials of 250 ml

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

Vetrimoxin L.A. 150 mg/ml suspension for injection

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

150 mg/ml amoxicillin (as trihydrate)

### **3. PACKAGE SIZE**

100 ml

250 ml

12 x 100 ml

12 x 250 ml

### **4. TARGET SPECIES**

Cattle and pigs

### **5. INDICATION(S)**

### **6. ROUTES OF ADMINISTRATION**

Intramuscular use.

Shake well before use.

Read the package leaflet before use.

### **7. WITHDRAWAL PERIODS**

Withdrawal periods:

Cattle:

Meat and offal: 18 days

Milk: 72 hours

Pigs:

Meat and offal: 16 days

### **8. EXPIRY DATE**

Exp: {mm/yyyy}

Once broached, use within 28 days.

Use by...

## **9. SPECIAL STORAGE PRECAUTIONS**

Do not refrigerate. Protect from frost. 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**

Ceva Sante Animale

## **14. MARKETING AUTHORISATION NUMBERS**

Vm 14966/5065 (GB)

Vm 14966/3064 (NI)

## **15. BATCH NUMBER**

Lot

**PARTICULARS TO APPEAR ON THE IMMEDIATE PACKAGE**

Label of 1 vial of 100 ml

Label of 1 vial of 250 ml

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

Vetrimoxin L.A. 150 mg/ml suspension for injection

**2. STATEMENT OF ACTIVE SUBSTANCES**

150 mg/ml amoxicillin (as trihydrate)

100 ml

250 ml

**3. TARGET SPECIES**

Cattle and pigs

**4. ROUTE OF ADMINISTRATION**

Intramuscular use.

Read the package leaflet before use.

**5. WITHDRAWAL PERIODS**

Withdrawal periods:

Cattle:

Meat and offal: 18 days Milk: 72 hours

Pigs:

Meat and offal: 16 days

**6. EXPIRY DATE**

Exp:{mm/yyyy}

Once broached, use within 28 days by.

Use by \_\_\_\_\_

**7. SPECIAL STORAGE PRECAUTIONS**

Do not refrigerate. Protect from frost. Keep the vial in the outer carton in order to protect from light.

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

Ceva Sante Animale

**9. BATCH NUMBER**

Lot {number}

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

### **PACKAGE LEAFLET**

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

Vetrimoxin L.A. 150 mg/ml suspension for injection for cattle and pigs

#### **2. Composition**

Each ml contains:  
150 mg amoxicillin (as trihydrate)  
Cream-beige suspension

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

Cattle and pigs

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

In cattle:

Treatment of respiratory infections caused by *Mannheimia haemolytica* and *Pasteurella multocida* susceptible to amoxicillin.

In pigs:

Treatment of respiratory infections caused by *Pasteurella multocida* susceptible to amoxicillin.

#### **5. Contraindications**

Do not use in cases of hypersensitivity to penicillins cephalosporins or to any of the excipients.

Do not use in cases of severe renal dysfunction with anuria and oliguria.

Do not use in case of infection with beta-lactamase-producing bacteria.

Do not use in rabbits, hares, hamsters, guinea pigs or other small herbivores.

Do not administer to equidae, because amoxicillin – like all aminopenicillins – may adversely affect the bacterial flora of the caecum.

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

##### Special precautions for safe use in the target species:

The choice of using amoxicillin should be based on bacterial susceptibility testing and take into account official, national and regional antimicrobial policies.

Use of the veterinary medicinal product deviating from the instructions given in the SPC may increase the prevalence of bacteria resistant to amoxicillin and may decrease the effectiveness of treatment with amoxicillin due to the potential for cross-resistance.

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

Penicillin and cephalosporin may cause an allergic reaction following accidental injection, inhalation or absorption via the skin, which may be life threatening. Hypersensitivity to penicillin may lead to cross sensitivity to cephalosporins and vice versa. Avoid direct contact of the veterinary medicinal product with the skin or the mucosae.

Handle the product with great care to avoid exposure.

Wear gloves and wash hands after use of the veterinary medicinal product.

In case of contact with the skin or eyes, wash immediately with water.

Do not smoke, eat or drink during use of the product.

If you develop symptoms following exposure, such as a skin rash, seek medical advice immediately and show the package leaflet or the label to the physician. Swelling of the face, lips and eyes or difficulty with breathing are more serious symptoms and require urgent medical attention.

Pregnancy and lactation:

Laboratory studies in rats and rabbits have not produced any evidence of harmful or toxic effect to the foetus or the mother (sow or cow). However, the tolerance of the medicinal product in cattle and pigs during pregnancy and lactation was not investigated. In these cases, use only according to the benefit/risk assessment by the responsible veterinarian.

Interaction with other medicinal products and other forms of interaction:

Do not use with antibiotics, which inhibit bacterial protein synthesis and can produce a contrasting action to the bactericidal effect of penicillins.

Overdose:

Amoxicillin has a wide safety margin.

Major incompatibilities:

Due to the lack of compatibility studies, this veterinary medicinal product must not be mixed with other veterinary medicinal products.

**7. Adverse events**

Cattle, pigs:

Rare (1 to 10 animals / 10,000 animals treated):	Injection site irritation <sup>1</sup>
Undetermined frequency (cannot be estimated from the available data)	Allergic reaction <sup>2</sup> (e.g. urticaria, anaphylactic shock)

<sup>1</sup> Of low intensity and recedes spontaneously and quickly, frequency may be decreased by reducing the volume of injection per injection site (see section “Advice on correct administration”).

<sup>2</sup> Varying in severity. In case of allergic reaction treatment should be discontinued and a symptomatic treatment should be initiated.

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 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**

### **Intramuscular use only.**

Intramuscular use.

Shake well before use.

15 mg amoxicillin per kg bodyweight corresponding to 1 ml of the veterinary medicinal product per 10 kg.

Administration should be repeated after 48 hours.

To ensure a correct dosage and to avoid underdosing, body weight should be determined as accurately as possible.

In cattle, do not administer more than 20 ml of the veterinary medicinal product per injection site.

In pigs, do not administer more than 6 ml of the veterinary medicinal product per injection site.

A separate injection site should be used for each administration.

As with other injectable preparations normal aseptic precautions should be observed.

If no distinct clinical response is seen after the second treatment, a check of the diagnosis and eventually a change of treatment are required.

Do not broach the vial more than 10 times: if necessary, use automatic syringes.

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

## **10. Withdrawal periods**

### Cattle:

Meat and offal: 18 days

Milk: 72 hours

### Pigs:

Meat and offal: 16 days

## **11. Special storage precautions**

Keep out of the sight and reach of children.

Do not refrigerate.

Protect from frost.

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 after first opening the immediate packaging: 28 days.

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

Medicines should not be disposed of via wastewater or household waste.

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 14966/5065 (GB)

Vm 14966/3064 (NI)

### Pack sizes

- 100 ml
- 12 x 100 ml
- 250 ml
- 12 x 250 ml

Not all pack sizes may be marketed.

## **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

Ceva Sante Animale  
8 rue de Logrono  
33500 Libourne  
France

### Contact details to report suspected adverse reactions:

Ceva Animal Health Ltd  
Explorer House  
Mercury Park  
Wycombe Lane  
Wooburn Green  
High Wycombe  
Buckinghamshire  
HP10 0HH  
United Kingdom  
Tel: 00800 35 22 11 51  
Email for the reporting of adverse events: [pharmacovigilance@ceva.com](mailto:pharmacovigilance@ceva.com)

### Manufacturer responsible for batch release:

Ceva Santé Animale, 10 av. de La Ballastière, 33500 Libourne, France

## **17. Other information**

POM-V

Veterinary Medicinal product subject to prescription  
For animal treatment only

*Gavin Hall*

Approved: 07 April 2026