PARTICULARS TO APPEAR ON THE OUTER PACKAGE

{Cartons of 3, 12, or 24 syringes}
{Buckets of 120 syringes}

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Combiclav Intramammary Suspension

2. STATEMENT OF ACTIVE SUBSTANCES

Each intramammary syringe of 3 g contains:
Amoxicillin 200 mg
Clavulanic Acid 50 mg
Prednisolone 10 mg

3. PACKAGE SIZE

3, 12, 24 or 120 syringes

4. TARGET SPECIES

Cattle (lactating cows).

5. INDICATIONS

6. ROUTES OF ADMINISTRATION

Intramammary use.

7. WITHDRAWAL PERIODS

Withdrawal period:
Meat and offal: 7 days.
Milk: 84 hours.

Combined Therapy:
When using this product and Combiclav Injection in combination:
Meat and offal: 42 days
Milk: 84 hours
From the last treatment of Combiclav Injection, following the minimum posology regime.

8. EXPIRY DATE

Exp. {mm/yyyy}

9. SPECIAL STORAGE PRECAUTIONS

Do not store above 25°C.
Store in a dry place.

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

Read the package leaflet before use.

User warnings:
Penicillins and cephalosporins may occasionally cause severe allergic reactions. See package leaflet for user warnings.

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

Norbrook Laboratories Limited

14. MARKETING AUTHORISATION NUMBERS

Vm 02000/5017
Vm 02000/3009

15. BATCH NUMBER

Lot {number}

MINIMUM PARTICULARS TO APPEAR ON SMALL IMMEDIATE PACKAGING UNITS

{3 g Syringe label}

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Combiclav

2. QUANTITATIVE PARTICULARS OF THE ACTIVE SUBSTANCES

Each intramammary syringe of 3g contains:

Amoxicillin	200 mg
Clavulanic Acid	50 mg
Prednisolone 1	0 mg

3. BATCH NUMBER

Lot {number}

4. EXPIRY DATE

Exp. {mm/yyyy}

PARTICULARS TO APPEAR ON THE PACKAGE LEAFLET

PACKAGE LEAFLET

1. Name of the veterinary medicinal product

Combiclav Intramammary Suspension for Lactating Cows

2. Composition

Each 3 g syringe contains:

Active substances:

Amoxicillin (as amoxicillin trihydrate)	200 mg
Clavulanic Acid (as potassium clavulanate)	50 mg
Prednisolone	10 mg

Cream to buff oily suspension.

3. Target species

Cattle (lactating cows).

4. Indications for use

For the treatment of clinical mastitis caused by the following bacteria susceptible to the combination of amoxicillin and clavulanic acid:
Staphylococci (including β-lactamase producing strains).
Streptococci (including *S. agalactiae, S. dysgalactiae,* and *S. uberis*).
Escherichia coli (including β-lactamase producing strains).

5. Contraindications

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

6. Special warnings

Special warnings:

Do not use in cases associated with *Psuedomonas.*

Special precautions for safe use in the target species:

Swab teat end before treatment, with cleaning towels provided.

Recommendations for prudent use:
The veterinary medicinal product should be used for treatment of clinical mastitis only.
Use of the veterinary medicinal product should be based on susceptibility testing of the bacteria isolated from the animal. If this is not possible, therapy should be based

on local (regional, farm level) epidemiological information about susceptibility of the target bacteria.
Official, national and regional antimicrobial policies should be taken into account when the veterinary medicinal product is used.
The combination of amoxicillin and clavulanic acid should be reserved for the treatment of clinical conditions which have responded poorly, or are expected to respond poorly, to other classes of antimicrobials.
Avoid use of the veterinary medicinal product in herds where no β-lactamase producing staphylococci strains have been isolated. Veterinarians should strive to use narrow spectrum antibiotics if possible. Inappropriate use of the veterinary medicinal product may increase the prevalence of bacteria resistant to β-lactam antibiotics and may decrease the effectiveness of treatment with β-lactam antibiotics, due to the potential for cross-resistance.
The feeding of waste milk containing residues of antibiotics to calves should be avoided up to the end of the milk withdrawal period, except during colostral phase, because it could select antimicrobial resistant bacteria within the intestinal microbiota of the calf and increase the faecal shedding of these bacteria.

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

The veterinary medicinal product may cause skin and eye irritation. Avoid contact with the skin and eyes. In the event of skin or eye contact rinse with plenty of clean water.
The cleaning towels supplied with the veterinary medicinal product contain isopropyl alcohol, which may cause skin or eye irritation in some people.
The wearing of gloves is recommended during administration of the veterinary medicinal product and when handling the cleaning towels.
Penicillins and cephalosporins may cause hypersensitivity (allergy) following injection, inhalation, ingestion, or skin contact.
Hypersensitivity to penicillins may lead to cross reactions to cephalosporins and vice versa. Allergic reactions to these substances may occasionally be serious.
Do not handle this veterinary medicinal product if you know you are sensitised, or if you have been advised not to work with such preparations.
Handle this veterinary medicinal product with great care to avoid exposure, taking all recommended precautions.
If you develop symptoms following exposure such as a skin rash, you should seek medical advice and show the leaflet or the label to the physician.
Swelling of the face, lips or eyes or difficulty with breathing, are more serious symptoms and require urgent medical attention.
Wash hands after use.

Special precautions for the protection of the environment:

Due to the endocrine-disrupting potential of prednisolone, the veterinary medicinal product may be dangerous to fish and other aquatic organisms. Consequently treated animals should not have access to watercourses during the first 12 hours after treatment.

Pregnancy and lactation:

Can be used during pregnancy and lactation.

Interaction with other medicinal products and other forms of interaction:

None known.

Overdose:

No adverse events are to be expected from an accidental overdose.

Major incompatibilities:
Not applicable.

7. Adverse events

Cattle (lactating cows):

Very rare (<1 animal / 10,000 animals treated, including isolated reports):	Hypersensitivity reaction

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

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

Intramammary use.

The syringe must only be used once. Partly emptied syringes due to the unsuccessful use should be discarded.
The content of one syringe should be infused gently into the teat of the infected quarter every 12 hours after each of three consecutive milkings.
Milk out the infected quarters. After thoroughly cleaning and disinfecting the teat and teat orifice with the cleaning towels provided, gently infuse the contents of one syringe into each affected quarter. Disperse the product by gentle massage of the teat and udder of the affected animal.
In cases of infections caused by *Staphylococcus aureus*, a longer course of antibacterial therapy may be required. Therefore overall treatment length must be at the veterinarian's discretion but should be long enough to ensure complete resolution of intramammary infection.

Combined therapy for the treatment of bovine mastitis. In the situation where systemic treatment as well as intramammary treatment is necessary, especially in

cases of serious clinical mastitis Combiclav Injection can be administered in combination with this veterinary medicinal product.

For combined therapy the following minimum treatment regime should be followed:

Potentiated Penicillin Injection	Potentiated Penicillin Intramammary Suspension for Lactating Cows
Intramuscular injection at a dose rate of 8.75 mg/kg bodyweight (7.0 mg amoxicillin and 1.75 mg clavulanic acid) which corresponds to 1 ml of veterinary medicinal product/20 kg bodyweight daily for 3 days as follows: 8.75 mg/kg bodyweight (7.0 mg amoxicillin, 1.75 mg clavulanic acid) i.e. 1 ml/20 kg bodyweight 24 hours ↓ 8.75 mg/kg bodyweight (7.0 mg amoxicillin, 1.75 mg clavulanic acid) i.e. 1 ml/20 kg bodyweight 24 hours ↓ 8.75 mg/kg bodyweight (7.0 mg amoxicillin, 1.75 mg clavulanic acid) i.e. 1 ml/20 kg bodyweight Where necessary Potentiated Penicillin Injection may be administered for an additional two days for a total of 5 daily injections	The content of one syringe should be infused gently into the teat of the infected quarter every 12 hours after each of three consecutive milkings as follows: One syringe gently infused into the teat of the infected quarter 12 hours ↓ One syringe gently infused into the teat of the infected quarter 12 hours ↓ One syringe gently infused into the teat of the infected quarter

9. Advice on correct administration

Use each syringe only once.

Partly emptied syringes due to the unsuccessful use should be discarded.
Milk out the infected quarters. After thoroughly cleaning and disinfecting the teat and teat orifice with the cleaning towels provided, gently infuse the contents of one syringe into each affected quarter. Disperse the veterinary medicinal product by gentle massage of the teat and udder of the affected animal.

10. Withdrawal periods

Meat and offal: 7 days.
Milk: 84 hours.

Combined Therapy:
When using this product and Combiclav Injection in combination:

Meat and offal: 42 days
Milk: 84 hours
From the last treatment of Combiclav Injection, following the minimum posology regime.

11. Special storage precautions

Keep out of the sight and reach of children.

Do not store above 25 C.
Store in a dry place.
Do not use this veterinary medicinal product after the expiry date which is stated on the label and carton after Exp. The expiry date refers to the last day of that month.

12. Special precautions for disposal

Medicines should not be disposed of via wastewater.

The veterinary medicinal product should not enter water courses as it is extremely dangerous for fish and aquatic life. Do not contaminate ponds, waterways or ditches with the veterinary medicinal product or used container.

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 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 02000/5017
Vm 02000/3009

Single dose 3g white LDPE syringes with a white LDPE dual push-fit cap.
Cartons of 3, 12, and 24 syringes, including 3, 12, or 24 individually wrapped teat cleaning towels containing isopropyl alcohol, or buckets of 120 syringes including 120 individually wrapped teat cleaning towels containing isopropyl alcohol.
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.

16. Contact details

Marketing authorisation holder, manufacturer responsible for batch release and contact details to report suspected adverse reactions:
Norbrook Laboratories Limited
Station Works
Newry
Co. Down
BT35 6JP
United Kingdom
Tel: +44 (0)28 3026 4435
E-mail: phvdept@norbrook.co.uk

Manufacturer responsible for batch release:
Norbrook Laboratories Limited
Station Works
Newry
Co. Down
BT35 6JP
United Kingdom

17. Other information

Amoxicillin is a broad spectrum bactericidal β-lactam antibiotic. Clavulanic acid inactivates β-lactamases. This combination is effective against β-lactamase producing organisms.

Prednisolone is an anti-inflammatory corticosteroid.

In vitro, clavulanic acid and amoxicillin in combination are active against a wide range of clinically important bacteria including the following organisms which are commonly associated with bovine mastitis:
Staphylococci (including β-lactamase producing strains).
Streptococci (including *S. agalactiae, S. dysgalactiae* and *S. uberis*).
Escherichia coli (including β-lactamase producing strains).

The Minimum Inhibitory Concentrations (MICs) of these target organisms determined from samples collected in nine EU countries (namely Belgium, Czech Republic, Denmark, France, Germany, Italy, Netherlands, Spain, and the UK)[1], show susceptibility to amoxicillin and clavulanic acid used in combination in accordance with the Clinical and Laboratory Standards Institute (CLSI) guidelines[2] on breakpoints (Table 1 and 2).

Table 1: Minimum Inhibitory Concentrations (mg/L) of Amoxicillin/Clavulanic Acid against strains from mastitis in dairy cattle in nine EU countries

	E. coli	*S. aureus*	*CNS*	*S. uberis*	*S. dysgalactiae*
Amoxicillin/Clavulanic Acid	8	1	0.5	0.5	≤0.03

Table 2: Clinical Laboratory Standards Institute (CLSI) resistance breakpoints (mg/L) for target bacteria

	E. coli	*S. aureus*	*CNS*[3]	*S. uberis*	*S. agalactiae*	*S. dysgalactiae*
Amoxicillin/Clavulanic Acid	≥32	≥8	≥8	≥32	≥8	≥32

[1]Antimicrobial susceptibility of mastitis pathogens isolated from diseased dairy cows across Europe: VetPath monitoring results, European society of clinical microbiology and infectious diseases (ECCMID), 2015.
[2]Clinical and Laboratory Standards Institute (2013). Approved standards- fourth edition, CLSI document VETO01-A4, Wayne, PA, USA.
[3]CNS – Coagulase Negative Staphylococci

The mechanisms underlying antimicrobial resistance in *Streptococcus* can be acquired through the mutation of intrinsic genes or horizontal exchange of genetic material encoding resistance determinants. Mastitic strains of *E. coli* and *Staphylococcus*, are known to acquire resistance through horizontal gene transfer and bacteriophages/plasmid transfer, and also through their ability to form a biofilm.

Acquired resistance prevalence in particular to be high in *E. coli*. In some strains of *Staphylococcus aureus* (methicillin-resistant *S. aureus*, MRSA), and of *Staphylococcus pseudintermedius*, resistance to all β-lactams is conferred by the alteration of the cell wall target proteins (penicillin-binding proteins). This is often associated with resistance to multiple other antimicrobial compounds with cross resistance.

Mastitic strains of *E. coli* and *Staphylococcus* are known to acquire resistance through horizontal gene transfer and bacteriophages/plasmid transfer, and also through their own ability to form a biofilm.

It has been documented that the pharmacokinetic characteristics of penicillins (including amoxicillin) after intramammary administration indicate rapid elimination of the drug from milk. The mean residence time has a several-fold lower value than the designated elimination half-life and amounts to only 3.4 hours. The concentration of the drug in the milk drops relatively quickly and the process is very dynamic.

POM-V

Gavin Hall

Approved 02 April 2025