SUMMARY OF PRODUCT CHARACTERISTICS

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Combiclav Intramammary Suspension for Lactating Cows

2. QUALITATIVE AND QUANTITATIVE COMPOSITION

Each intramammary syringe of 3 g contains:

Active substances:

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

Excipients:

Qualitative composition of excipients and other constituents
Aluminium Sodium Silicate
Cetostearyl Alcohol (Type B), emulsifying
Paraffin, White Soft
Paraffin, Light Liquid

Cream to buff oily suspension.

3. CLINICAL INFORMATION

3.1 Target species

Cattle (lactating cows).

3.2 Indications for use for each target species

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).

3.3 Contraindications

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

3.4 Special warnings

Do not use in cases associated with *Pseudomonas.*

3.5 Special precautions for use

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 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 product in herds where no β-lactamase producing staphylococci strains have been isolated. *E.coli* mastitis with mild to moderate clinical signs, a non-antimicrobial approach should be the first option. Veterinarians should strive to use narrow spectrum antibiotics if possible. Inappropriate use of the 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.
Most of the ESBLs and AmpC β-lactamases producing *E.coli* strains may not be inhibited by the combination of amoxicillin/clavulanic acid.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:

This 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 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.

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.

3.6 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 veterinary medicinal product. Reports should be sent, preferably via a veterinarian, to either the marketing authorisation holder or the national competent authority via the national reporting system. See the package leaflet for respective contact details.

3.7 Use during pregnancy, lactation or lay

Pregnancy and lactation:
Can be used during pregnancy and lactation.

3.8 Interaction with other medicinal products and other forms of interaction

None known.

3.9 Administration routes and dosage

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 veterinary medicinal 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 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 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

3.10 Symptoms of overdose (and where applicable, emergency procedures and antidotes)

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

3.11 Special restrictions for use and special conditions for use, including restrictions on the use of antimicrobial and antiparasitic veterinary medicinal products in order to limit the risk of development of resistance

Not applicable.

3.12 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.

4. PHARMACOLOGICAL INFORMATION

4.1 ATCvet code:

QJ51RV01

4.2 Pharmacodynamics

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.

4.3 Pharmacokinetics

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.

5. PHARMACEUTICAL PARTICULARS

5.1 Major incompatibilities

Not applicable.

5.2 Shelf life

Shelf life of the veterinary medicinal product as packaged for sale: 2 years.

5.3 Special precautions for storage

Do not store above 25^{o}C.
Store in a dry place.

5.4 Nature and composition of immediate packaging

Single dose 3g white LDPE syringes with a white LDPE dual push-fit cap.
Pack sizes:
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.

5.5 Special precautions for the disposal of unused veterinary medicinal products or waste materials derived from the use of such products

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 national collection systems applicable to the veterinary medicinal product concerned.

6. NAME OF THE MARKETING AUTHORISATION HOLDER

Norbrook Laboratories Limited

7. MARKETING AUTHORISATION NUMBERS

Vm 02000/5017
Vm 02000/3009

8. DATE OF FIRST AUTHORISATION

03 October 2018

9. DATE OF THE LAST REVISION OF THE SUMMARY OF THE PRODUCT CHARACTERISTICS

December 2024

10. CLASSIFICATION OF VETERINARY MEDICINAL PRODUCTS

Veterinary medicinal product subject to prescription.

Find more product information by searching for the 'Product Information Database' on www.gov.uk.

Gavin Hall

Approved 02 April 2025