

SUMMARY OF PRODUCT CHARACTERISTICS

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Amoxiclav-VMD 50 mg (BE, NL)
Clavucill 40 mg/10 mg (DE, DK, ES, FR, PL, PT, RO, SE)
Tablets for dogs and cats.

2. QUALITATIVE AND QUANTITATIVE COMPOSITION

Each tablet (180 mg) contains:

- Active substances: Amoxicillin (as amoxicillin trihydrate) 40 mg/tablet
Clavulanic acid (as potassium clavulanate) 10 mg/tablet
- Excipients: Erythrosine (E127) 0.05 mg.

For the full list of excipients, see section 6.1.

3. PHARMACEUTICAL FORM

Tablet.

Pale pink, rounded, one side scored, uncoated tablet with a diameter of 8 mm.
The tablets can be divided into two equal parts.

4. CLINICAL PARTICULARS

4.1 Target species

Dogs and cats.

4.2 Indications for use, specifying the target species

Dogs:

Treatment of infections caused by micro-organisms sensitive for the combination amoxicillin/clavulanic acid, especially:

- dermatitis (superficial and deep pyoderma) caused by *Staphylococcus (pseud)intermedius*.
- urinary tract infections caused by *E. coli*.
- respiratory tract infections caused by *Streptococcus* spp.
- enteritis caused by *E. coli*.

Cats:

Treatment of infections caused by micro-organisms sensitive for the combination amoxicillin/clavulanic acid, especially:

- dermatitis (superficial and deep pyoderma) caused by *Staphylococcus (pseud)intermedius*.
- urinary tract infections caused by *E. coli*
- respiratory tract infections caused by *Streptococcus* spp.
- enteritis caused by *E. coli*.

4.3 Contraindications

- Do not use in animals with known hypersensitivity to penicillin or other substances of the beta-lactam group or to any of the excipients.
- Do not use in case of serious dysfunction of the kidneys accompanied by anuria and oliguria.
- Do not use in rabbits, guinea pigs, hamsters, chinchillas or gerbils.

4.4 Special warnings for each target species

None known.

4.5 Special precautions for use

i) Special precautions for use in animals

- Do not use in case of known resistance to the combination.
- Official, national and regional antimicrobial policies with respect to the use of broad-spectrum antibiotics should be taken into account.
- Do not use in case of bacteria sensitive to narrow spectrum penicillins or to amoxicillin as single substance.
- Whenever possible, the product should only be used based on susceptibility testing. Use of the product deviating from the instructions given in the SPC may increase the prevalence of bacteria resistant to the amoxicillin/clavulanate, and may decrease the effectiveness of treatment with β -lactam antibiotics, due to the potential for cross resistance.
- In animals with hepatic and renal failure, the dosing regimen should be carefully evaluated.

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

Penicillins and cephalosporins may cause hypersensitivity reactions (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.

- People with known hypersensitivity to penicillins should avoid contact with the veterinary medicinal product. Handle this product with great care to avoid exposure, taking all recommended precautions.
- 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 or eyes or difficulty with breathing are more serious symptoms and require urgent medical attention.
- Wash hands after use.

4.6 Adverse reactions (frequency and seriousness)

- Gastro-intestinal disturbances (diarrhoea, vomiting, ...) may occur uncommonly after administration of the product. Treatment may be discontinued depending on the severity of the undesirable effects and a benefit/risk evaluation by the veterinary surgeon.
- Dose independent allergic reactions may very rarely occur, such as skin reactions or anaphylaxis. In those cases the treatment must be stopped immediately and a symptomatic treatment should be given.

The frequency of adverse reactions is defined using the following convention:

- very common (more than 1 in 10 animals displaying adverse reactions during the course of one treatment)
- common (more than 1 but less than 10 animals in 100 animals)
- uncommon (more than 1 but less than 10 animals in 1,000 animals)
- rare (more than 1 but less than 10 animals in 10,000 animals)
- very rare (less than 1 animal in 10,000 animals, including isolated reports).

4.7 Use during pregnancy or lactation

Laboratory studies in rats and mice have not produced any evidence of teratogenic or foetotoxic effects. No studies have been conducted in pregnant or lactating dogs and cats. Use only according to the benefit/risk assessment by the responsible veterinarian.

4.8 Interaction with other medicinal products and other forms of interaction

Chloramphenicol, macrolides, sulfonamides and tetracyclines may inhibit the antibacterial effects of penicillins.

The potential for allergic cross-reactivity with other penicillins should be considered.

Penicillins may increase the effect of aminoglycosides.

4.9 Amounts to be administered and administration route

Oral use.

Amounts to be administered: The recommended dose rate is 10 mg amoxicillin / 2.5 mg clavulanic acid per kg bodyweight (= 12.5 mg of combined active substances) twice a day by the oral route in dogs and cats, i.e. 1 tablet per 4 kg body weight every 12 h.

| Body weight (kg) | Number of tablets (twice daily) |
|------------------|---------------------------------|
| (1.0 – 2.0) | ½ |
| (2.1 – 4.0) | 1 |
| (4.1 – 6.0) | 1 ½ |
| (6.1 – 8.0) | 2 |
| > 8 | Use 250 or 500 mg tablets |

In case of complicated infections, especially respiratory infections, a better cure rate is obtained with a double dose, up to 25 mg of the combination of the active substances per kg weight, twice daily.

Treatment duration:

In the majority of cases, a treatment of 5 to 7 days is sufficient.

For chronic and refractory infections, longer courses of antibacterial therapy may be required.

Treatment length should be adapted by the veterinarian, and should be long enough to ensure complete bacteriological cure.

To ensure a correct dosage, body weight should be determined as accurately as possible to avoid under-dosing.

4.10 Overdose (symptoms, emergency procedures, antidotes), if necessary

Mild gastrointestinal symptoms (diarrhoea, vomiting) may occur more frequently after overdose of the product.

4.11 Withdrawal period

Not applicable.

5. PHARMACOLOGICAL PROPERTIES

Pharmacotherapeutic group: Antibacterials for systemic use, amoxicillin and enzyme inhibitor
ATCvet code: QJ01CR02.

4.1 Pharmacodynamic properties

Amoxicillin is a beta-lactam antibiotic, and as such interferes with the synthesis of cell wall peptidoglycan; it has a bactericidal activity on growing bacteria. It is considered as a broad-spectrum penicillin; it is active in vitro against many aerobic and anaerobic, Gram+ and Gram- bacteria, However it is inactivated in bacteria producing beta-lactamases. Susceptible bacterial species include: *Staphylococcus intermedius*, β -haemolytic streptococci and *Escherichia coli*.

Clavulanic acid is a potent inhibitor of many β -lactamases produced by Gram positive and Gram negative bacteria, of plasmid or chromosomal origin. Inhibition is allowed by structural similarity with beta-lactams, and occurs through the formation of a stable molecule-enzyme complex. During this process, clavulanic acid is destroyed leading to the protection of amoxicillin against inactivation by these enzymes.

Acquired resistance may be high in *E. coli*. Resistance notably develops through the production of inhibitor-resistant beta-lactamases or the hyperproduction of beta-lactamases.

In some strains of *Staphylococcus aureus* (methicillin-resistant *S. aureus*, MRSA), and of *Staphylococcus pseudintermedius*, resistance to all beta-lactams is conferred by the alteration of the cell wall target proteins (Penicillin-Binding Proteins). This is often associated to resistance to multiple other antimicrobial compounds.

Pseudomonas aeruginosa and *Enterobacter spp.* can be regarded as intrinsically resistant to the combination.

4.2 Pharmacokinetic particulars

Amoxicillin is well absorbed after oral intake. The mean bioavailability associated with the tablets is of approximately 53% in dogs and 69% in cats. Following absorption the highest concentrations are found in the kidneys (urine) and the bile, then the liver, the lungs, the heart and the spleen. The distribution of amoxicillin in the cerebrospinal fluid is limited, unless the meninges are inflamed.

Clavulanic acid is also well absorbed after oral administration. The mean bioavailability associated with the tablets is of approximately 43% in dogs and 69% in cats. The distribution to the cerebrospinal fluid is limited, unless the meninges are inflamed. Clavulanic acid is excreted mainly through the kidneys (unchanged in the urine).

The main pharmacokinetic parameters after a single dose of 25 mg of the combination of active substances per kg body weight were summarized in the following tables:

Cat:

| <i>Parameter</i> | <i>Mean value</i> | |
|----------------------------|-------------------|-----------------|
| | Amoxicillin | Clavulanic acid |
| C _{max} (µg/mL) | 12,1 | 8,09 |
| T _{max} (hr) | 1,87 | 0,97 |
| t _{1/2} (hr) | 1,31 | 0,84 |
| AUC _∞ (µg.h/ml) | 41,9 | 11,8 |

Dog:

| <i>Parameter</i> | <i>Mean value</i> | |
|----------------------------|-------------------|-----------------|
| | Amoxicillin | Clavulanic acid |
| C _{max} (µg/mL) | 12,49 | 4,23 |
| T _{max} (hr) | 1,18 | 0,97 |
| t _{1/2} (hr) | 1,57 | 0,63 |
| AUC _∞ (µg.h/ml) | 31,1 | 5,54 |

6. PHARMACEUTICAL PARTICULARS

6.1 List of excipients

- Colloidal silica, anhydrous
- Sodium starch glycolate, type A
- Microcrystalline cellulose (E460)
- Erythrosine (E127)
- Magnesium stearate (E572)

6.2 Incompatibilities

Not applicable.

6.3 Shelf life

Shelf-life of the veterinary medicinal product as packaged for sale: 2 years.
Any divided tablet portion remaining after 24 hours should be discarded.

6.4. Special precautions for storage

Do not store above 25°C.
Divided tablets should be stored in the blister pack.

6.5 Nature and composition of immediate packaging

Alu-Alu-blister packs consisting of aluminium foil (Polyester / Aluminium foil / LD Polyethylene), heat sealed, in strips of 10 tablets. Cartons containing 10, 100 or 500 tablets.

Not all pack sizes may be marketed.

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

Any unused veterinary medicinal product or waste materials derived from such veterinary medicinal products should be disposed of in accordance with local requirements.

7. MARKETING AUTHORISATION HOLDER

V.M.D. n.v.
Hoge Mauw 900
2370 Arendonk.
Belgium
Tel.: +32 14.67.20.51
Fax: +32 14.67.21.52
e-mail: vmd@vmdvet.be

8. MARKETING AUTHORISATION NUMBER**9. DATE OF FIRST AUTHORISATION/RENEWAL OF THE AUTHORISATION**

Date of first authorisation:
Date of last renewal:

10 DATE OF REVISION OF THE TEXT**PROHIBITION OF SALE, SUPPLY AND/OR USE**

Delivery: veterinary prescription only.

LABELLING**PARTICULARS TO APPEAR ON THE OUTER PACKAGE AND THE IMMEDIATE PACKAGE****{Carton}****1. NAME OF THE VETERINARY MEDICINAL PRODUCT**

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Clavucill 40 mg/10 mg (DE, DK, ES, FR, PL, PT, RO, SE)
Tablets for dogs and cats.
Amoxicillin + clavulanic acid

2. STATEMENT OF ACTIVE AND OTHER SUBSTANCES

Active substances:
Amoxicillin (as amoxicillin trihydrate) 40 mg/tablet
Clavulanic acid (as potassium clavulanate) 10 mg/tablet
Excipients:
Erythrosine (E127) 0.05 mg.

3. PHARMACEUTICAL FORM

Tablet.

4. PACKAGE SIZE

10, 100 or 500 tablets.

5. TARGET SPECIES

Dogs and cats.

6. INDICATION(S)**7. METHOD AND ROUTE(S) OF ADMINISTRATION**

Oral use. Read the package leaflet before use.

8. WITHDRAWAL PERIOD

Not applicable.

9. SPECIAL WARNING(S), IF NECESSARY**10. EXPIRY DATE**

EXP {month/year}
Any divided tablet portion remaining after 24 hours should be discarded.

11. SPECIAL STORAGE CONDITIONS

Do not store above 25°C.
Divided tablets should be stored in the blister pack.

12. SPECIAL PRECAUTIONS FOR THE DISPOSAL OF UNUSED PRODUCTS OR WASTE MATERIALS, IF ANY

Disposal: read package leaflet.

13. THE WORDS “FOR ANIMAL TREATMENT ONLY” AND CONDITIONS OR RESTRICTIONS REGARDING SUPPLY AND USE, IF APPLICABLE

For animal treatment only.
To be supplied only on veterinary prescription.
Administration by a veterinary surgeon or under his responsibility.

14. THE WORDS “KEEP OUT OF THE SIGHT AND REACH OF CHILDREN”

Keep out of the sight and reach of children.

15. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER

V.M.D. n.v.
Hoge Mauw 900
2370 Arendonk.
Belgium

16. MARKETING AUTHORISATION NUMBER(S)**17. MANUFACTURER’S BATCH NUMBER**

Lot {number}

MINIMUM PARTICULARS TO APPEAR ON BLISTERS OR STRIPS

{Blister}

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2. NAME OF THE MARKETING AUTHORISATION HOLDER

V.M.D. n.v.

3. EXPIRY DATE

EXP {month/year}

4. BATCH NUMBER

Lot {number}

5. THE WORDS “FOR ANIMAL TREATMENT ONLY”

For animal treatment only.

PACKAGE LEAFLET FOR:
Amoxiclav-VMD 50 mg (BE, NL)
Clavucill 40 mg/10 mg (DE, DK, ES, FR, PL, PT, RO, SE)
Tablets for dogs and cats.

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V.M.D. n.v.
Hoge Mauw 900
2370 Arendonk.
Belgium

2. NAME OF THE VETERINARY MEDICINAL PRODUCT

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- enteritis caused by *E. coli*.

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- Do not use in animals with known hypersensitivity to penicillin or other substances of the beta-lactam group or to any of the excipients.
- Do not use in case of serious dysfunction of the kidneys accompanied by anuria and oliguria.
- Do not use in rabbits, guinea pigs, hamsters, chinchillas or gerbils.

6. ADVERSE REACTIONS

- Gastro-intestinal disturbances (diarrhoea, vomiting, ...) may occur uncommonly after administration of the product. Treatment may be discontinued depending on the severity of the undesirable effects and a benefit/risk evaluation by the veterinary surgeon.
- Dose independent allergic reactions may very rarely occur, such as skin reactions or anaphylaxis. In those cases the treatment must be stopped immediately and a symptomatic treatment should be given.

The frequency of adverse reactions is defined using the following convention:

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- uncommon (more than 1 but less than 10 animals in 1,000 animals)
- rare (more than 1 but less than 10 animals in 10,000 animals)
- very rare (less than 1 animal in 10,000 animals, including isolated reports).

If you notice any serious effects or other effects not mentioned in this package leaflet, please inform your veterinary surgeon.

7. TARGET SPECIES

Dogs and cats.

8. DOSAGE FOR EACH SPECIES, ROUTE(S) AND METHOD OF ADMINISTRATION

Oral use.

Amounts to be administered: The recommended dose rate is 10 mg amoxicillin / 2.5 mg clavulanic acid per kg bodyweight (= 12.5 mg of combined active substances) twice a day by the oral route in dogs and cats, i.e. 1 tablet per 4 kg body weight every 12 h.

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Treatment duration:

In the majority of cases, a treatment of 5 to 7 days is sufficient.

For chronic and refractory infections, longer courses of antibacterial therapy may be required.

Treatment length should be adapted by the veterinarian, and should be long enough to ensure complete bacteriological cure.

To ensure a correct dosage, body weight should be determined as accurately as possible to avoid under-dosing.

9. ADVICE ON CORRECT ADMINISTRATION

- Do not use in case of known resistance to the combination.
- Official, national and regional antimicrobial policies with respect to the use of broad-spectrum antibiotics should be taken into account.
- Do not use in case of bacteria sensitive to narrow spectrum penicillins or to amoxicillin as single substance.
- Whenever possible, the product should only be used based on susceptibility testing. Use of the product deviating from the instructions given in the leaflet may increase the prevalence of bacteria resistant to the amoxicillin/clavulanate, and may decrease the effectiveness of treatment with β -lactam antibiotics, due to the potential for cross resistance.
- In animals with hepatic and renal failure, the dosing regimen should be carefully evaluated.

10. WITHDRAWAL PERIOD

Not applicable.

11. SPECIAL STORAGE PRECAUTIONS

Keep out of the sight and reach of children.

Do not store above 25°C.

Divided tablets should be stored in the blister pack.

Any divided tablet portion remaining after 24 hours should be discarded.

Do not use this veterinary medicinal product after the expiry date which is stated on the blister and the carton.

12. SPECIAL WARNING(S)

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

Penicillins and cephalosporins may cause hypersensitivity reactions (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.

- People with known hypersensitivity to penicillins should avoid contact with the veterinary medicinal product. Handle this product with great care to avoid exposure, taking all recommended precautions.
- 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 or eyes or difficulty with breathing are more serious symptoms and require urgent medical attention.
- Wash hands after use.

Pregnancy or lactation:

Laboratory studies in rats and mice have not produced any evidence of teratogenic or foetotoxic effects. No studies have been conducted in pregnant or lactating dogs and cats. Use only according to the benefit/risk assessment by the responsible veterinarian.

Interaction with other medicinal products and other forms of interaction:

Chloramphenicol, macrolides, sulfonamides, and tetracyclines may inhibit the antibacterial effects of penicillins.

The potential for allergic cross-reactivity with other penicillins should be considered.

Penicillins may increase the effect of aminoglycosides.

Overdose (symptoms, emergency procedures, antidotes):

Mild gastrointestinal symptoms (diarrhoea, vomiting) may occur more frequently after overdose of the product.

Incompatibilities

Not applicable.

13. SPECIAL PRECAUTIONS FOR THE DISPOSAL OF UNUSED PRODUCT OR WASTE MATERIALS, IF ANY

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

Ask your veterinary surgeon how to dispose of medicines no longer required. These measures should help to protect the environment.

14. DATE ON WHICH THE PACKAGE LEAFLET WAS LAST APPROVED**15. OTHER INFORMATION**

Delivery: veterinary prescription only.

Pack size: 10, 100 or 500 tablets.

Not all pack sizes may be marketed.