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Clavaseptin 50mg & 62.5mg Palatable Tablets for Dogs and Cats, Clavaseptin 250mg & 500mg Palatable Tablets for Dogs (NI)

Species:Cats, Dogs

Active ingredient:Amoxicillin, Clavulanic Acid

Product:Clavaseptin 50mg & 62.5mg Palatable Tablets for Dogs and Cats, Clavaseptin 250mg & 500mg Palatable Tablets for Dogs (NI)

Product index:Clavaseptin

Incorporating:Clavaseptin 50mg & 62.5mg Palatable Tablets for Dogs and Cats, Clavaseptin 250mg & 500mg Palatable Tablets for Dogs

Qualitative and quantitative composition

Each tablet contains:

	Active Substances	
	Amoxicillin (as amoxicillin trihydrate)	Clavulanic acid (as salt)
Clavaseptin 50 mg Palatable tablets for dogs and cats	40mg	10mg
Clavaseptin 62.5 mg Palatable tablets for dogs and cats	50mg	12.5mg
Clavaseptin 250 mg Palatable tablets for dogs	200mg	50mg
Clavaseptin 500 mg Palatable tablets for dogs	400mg	100mg

Excipients:

Qualitative composition of excipients and other constituents

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Crospovidone

Povidone K25

Silicon dioxide

Microcrystalline cellulose

Pig liver flavour

Dried yeast

Magnesium stearate

Hypromellose

Pharmaceutical form

Beige scored tablets that can be divided into two equal parts.

Clinical particulars

Target species

50mg & 62.5mg : Dogs and cats

250mg & 500mg: Dogs

Indications for use, specifying the target species

In dogs: treatment of infections caused by bacteria susceptible to amoxicillin in combination with clavulanic acid (including beta-lactamase producing strains), in particular:

- Skin infections (including deep and superficial pyodermas, wounds, abscesses) caused by *Staphylococcus spp*, *Streptococcus spp* and *Pasteurella spp*.
- Respiratory tract infections (sinusitis, rhino-tracheitis, bronchopneumonia) caused by *Staphylococcus spp* and *E. coli*.
- Infections of the oral cavity (mucous membranes) caused by *Streptococcus spp*, and *Pasteurella spp*.
- Urinary tract infections (nephritis, cystitis) caused by *E. coli*, *Klebsiella spp* and *Proteus mirabilis*.
- Digestive tract infections, especially gastroenteritis caused by *E. coli*.

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In cats: treatment of infections caused by bacteria susceptible to amoxicillin in combination with clavulanic acid (including beta-lactamase producing strains), in particular:

- Skin infections (including deep and superficial pyodermas, wounds, abscesses) caused by *Staphylococcus spp*, *Streptococcus spp* and *Pasteurella spp*.
- Respiratory tract infections (sinusitis, rhino-tracheitis, bronchopneumonia) caused by *Staphylococcus spp* and *E. coli*.
- Infections of the oral cavity (mucous membranes) caused by *Streptococcus spp*, and *Pasteurella spp*.
- Urinary tract infections (nephritis, cystitis) caused by *E. coli*, *Pasteurella spp*, *Klebsiella spp* and *Proteus mirabilis*.
- Digestive tract infections, especially gastroenteritis caused by *E. coli*.

Contraindications

Do not use in cases of hypersensitivity to penicillins or other substances of the β -lactam group or to any of the excipients.

Do not administer to gerbils, guinea pigs, hamsters, rabbits and chinchillas or other small herbivores.

Do not use in animals with serious dysfunction of the kidneys accompanied by anuria or oliguria.

Do not administer to horses and ruminating animals.

Special warnings

Cross-resistance has been shown between amoxicillin/clavulanic acid and β -lactam antibiotics. Use of the veterinary medicinal product should be carefully considered when susceptibility testing has shown resistance to β -lactam antibiotics because its effectiveness may be reduced.

Methicillin resistant *S. aureus* (MRSA) and methicillin resistant *S. pseudintermedius* (MRSP) have been isolated in cats and dogs with proportion of resistance that varies across EU countries.

Do not use in cases of known resistance to the combination of amoxicillin and clavulanic acid.

Do not use in cases of suspected or confirmed MRSA/MRSP infections, as isolates should be considered resistant to all β -lactam including amoxicillin/clavulanic acid combination.

High resistances (up to 100%) have been reported in *E. coli* isolates from skin and soft tissue infections in dogs.

Special precautions for use

Special precautions for safe use in the target species:

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In animals with impaired liver and kidney function, the use of the veterinary medicinal product should be subject to a benefit/risk evaluation by the veterinary surgeon and the posology evaluated carefully.

Use of the veterinary medicinal product should be based on identification and susceptibility testing of the target pathogen(s). If this is not possible, therapy should be based on epidemiological information and knowledge of susceptibility of the target pathogens at local/regional level.

Use of the veterinary medicinal product should be in accordance with official, national and regional antimicrobial policies. Aminopenicillins in combination with beta-lactamase inhibitors are in AMEG category „C”. An antibiotic with a lower risk of antimicrobial resistance selection (lower AMEG category) should be used for first line treatment where susceptibility testing suggests the likely efficacy of this approach. Narrow spectrum antibiotic therapy with a lower risk of antimicrobial resistance selection should be used for first line treatment where susceptibility testing suggests the likely efficacy of this approach.

The tablets are flavoured. In order to avoid any accidental ingestion, store tablets out of reach of the animals.

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

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

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 veterinary medicinal preparations. Handle this veterinary medicinal product with great care to avoid exposure, taking all recommended precautions.

If you develop symptoms following exposure, such as skin rash, you should seek medical advice and show the doctor this warning. Swelling of the face, lips or eyes or difficulty breathing are more serious symptoms and require urgent medical attention.

Wash hands after handling the tablets.

Accidental ingestion of the veterinary medicinal product by a child may be harmful. To avoid accidental ingestion, particularly by a child, unused part-tablets should be returned to the open blister space and inserted back into the carton. In case of accidental ingestion, seek medical advice immediately and show the package leaflet or the label to the physician.

Special precautions for the protection of the environment:

Not applicable.

Adverse events

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Dogs and cats.

Very rare (<1 animal / 10,000 animals

Vomiting¹, Diarrhoea.¹

treated, including isolated reports):

Hypersensitivity reaction (Allergic reactions²), anaphylaxis²

1) Treatment may be discontinued depending on the severity of the undesirable effects and a benefit/risk evaluation by the veterinary surgeon

2) In these cases, administration should be discontinued and a symptomatic treatment given

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 its local representative or the national competent authority via the national reporting system.

Use during pregnancy, lactation or lay

Pregnancy and lactation:

The safety of the product has not been established during pregnancy and lactation. Laboratory studies in rats have not produced any evidence of teratogenic, foetotoxic or maternotoxic effects. Use only in accordance with the the benefit/risk assessment by the responsible veterinarian.

Interaction with other medicinal products and other forms of interaction

The bactericidal activity of amoxicillin may be reduced by the simultaneous use of bacteriostatic substances such as macrolides, tetracyclines, sulfonamides and chloramphenicol. Penicillins may increase the effect of aminoglycosides.

Administration routes and dosage

Oral use.

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

The recommended dose of the product is 10 mg amoxicillin /2.5 mg clavulanic acid per kg body weight twice a day by the oral route in dogs and cats, every 12 h, according to the following tables:

Clavaseptin 50 mg Palatable tablets for dogs and cats:

Bodyweight (kg)	Number of tablets twice daily
[1.0- 2.0]	½
[2.1- 4.0]	1
[4.1- 6.0]	1 ½

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[6.1- 8.0] 2

Clavaseptin 62.5 mg Palatable tablets for dogs and cats:

Bodyweight (kg)	Number of tablets twice daily
[1.0- 2.5]	½
[2.6- 5.0]	1
[5.1- 7.5]	1 ½
[7.6 - 10.0]	2

Clavaseptin 250 mg Palatable tablets for dogs:

Bodyweight (kg)	Number of tablets twice daily
[8.1 - 10]	½
[10.1- 20]	1
[20.1- 30]	1 ½
[30.1- 40]	2

Clavaseptin 500 mg Palatable tablets for dogs:

Bodyweight (kg)	Number of tablets twice daily
[30.1 - 40]	1
[40.1 - 60]	1 ½
[60.1 - 80]	2

In severe cases, the dose can be doubled at the discretion of the responsible veterinarian.

Duration of the treatment:

For all indications, a treatment of 5 to 7 days is sufficient in the majority of cases.

For chronic or recurrent cases, it may be necessary to continue treatment for 2 to 4 weeks.

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

At three times the recommended dose for a period of 28 days, a decrease in cholesterol values and episodes of vomiting were observed in cats and diarrhoea was observed in dogs. In the event of an overdose symptomatic treatment is advised.

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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.

Withdrawal periods

Not applicable

Pharmacological particulars

ATCvet code: QJ01CR02

Pharmacodynamics

Amoxicillin is an aminobenzylpenicillin from the β -lactam penicillin family which prevents the bacterial cell wall formation by interfering with the final step of peptidoglycan synthesis.

Clavulanic acid is an irreversible inhibitor of intracellular and extracellular β -lactamases which protects amoxicillin from inactivation by many β -lactamases.

Amoxicillin/clavulanate has a wide range of activity which includes β -lactamase producing strains of both Gram-positive and Gram-negative aerobes, facultative anaerobes and obligate anaerobes. The antimicrobial spectrum relevant for dog's and cat's indications is summarized in the two tables below.

Summary of susceptibility for dog target bacteria:

Target bacteria in each indication	n	Range of MIC ($\mu\text{g/mL}$)	MIC50 ($\mu\text{g/mL}$)	MIC90 (
Skin and soft tissues				
<i>Staphylococcus</i> spp	431*	0.03-32	0.12	1
<i>S. aureus</i>	38*	0.12-16	0.5	2
<i>S. intermedius</i> group	343*	0.03-8	0.12	0.5
Coagulase-negative <i>Staphylococcus</i> spp	49*	0.03-32	0.12	2
<i>Streptococcus</i> spp	142*	0.015-0.06	≤ 0.015	≤ 0.015
<i>Streptococcus canis</i>	127*	0.015-0.06	≤ 0.015	≤ 0.015
<i>Streptococcus</i> <i>dysgalactiae</i>	12*	0.015	≤ 0.015	≤ 0.015
<i>Pasteurella</i> spp	22*	0.03-0.25	0.12	0.25

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Respiratory

<i>Staphylococcus</i> spp	112*	0.06-8	0.12	0.5
<i>S. intermedius</i> group	90*	0.06-8	0.12	0.25
<i>S. aureus</i>	22*	0.12-8	0.25	1

Dental

<i>Streptococcus</i> spp	16**	0.008 - 1	0.014	0.4
<i>Pasteurella</i> spp	68**	0.03 - 64	0.124	0.4

Urinary

<i>Escherichia coli</i>	236*	1-32	4	16
<i>Klebsiella</i> spp	33*	0.5-32	2	32
<i>Proteus</i> spp	66*	0.5-16	1	8

Digestive

<i>Escherichia coli</i>	- *	1-32	4	8
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Clavaseptin: Dog

Table

Breakpoints are from CLSI VET01-S7.

* MIC values determined from bacteria collected in Europe in 2021-2022 (ComPath-IV survey).
Susceptibility of digestive isolates is assumed similar to that of the same bacteria in other types of infection.

** MIC values determined from bacteria collected from dog dental infections in Europe in 2002.

- Missing information.

Summary of susceptibility for cat target bacteria:

Target bacteria in each indication	n	Range of MIC (µg/mL)	MIC50 (µg/mL)	MIC90 (µg/mL)
Skin and soft tissues				
<i>Staphylococcus</i> spp	150*	0.03-32	0.12	1
<i>S. aureus</i>	50*	0.03-32	0.25	1
<i>S. intermedius</i> group	32*	0.03-32	0.12	8
Coagulase-negative	66*	0.03-8	0.06	0.25

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Staphylococcus spp

<i>S. felis</i>	50*	0.03-0.12	0.06	0.12
<i>Streptococcus</i> spp	30*	0.015-0.06	≤0.015	≤0.015
<i>Streptococcus canis</i>	27*	0.015-0.03	≤0.015	≤0.015
<i>Pasteurella</i> spp	58*	0.015-2	0.25	0.25

Respiratory

<i>Staphylococcus</i> spp	89*	0.03-8	0.12	1
Coagulase-negative	77*	0.03-8	0.12	1
<i>Staphylococcus</i> spp				
<i>S. intermedius</i> group	12*	0.03-2	0.12	1
<i>S. aureus</i>	30*	0.12-8	0.5	4
<i>S. felis</i>	40*	0.03-0.12	0.06	0.12

Dental

<i>Streptococcus</i> spp	–	–	–	–
<i>Pasteurella</i> spp	–	–	–	–

Urinary

<i>Escherichia coli</i>	132*	1-32	4	8
<i>Pasteurella multocida</i>	–	–	–	–
<i>Klebsiella</i> spp	19*	1-32	2	32
<i>Proteus</i> spp	17*	0.5-32	1	4

Digestive

<i>Escherichia coli</i>	- *	1-32	4	8
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Clavaseptin: Cat

Table

Breakpoints are from CLSI VET01-S7.

* MIC values determined from bacteria collected in Europe in 2021-2022 (ComPath-IV survey).
Susceptibility of digestive isolates is assumed similar to that of the same bacteria in other types of infection.

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- Missing information.

The two main mechanisms of resistance to amoxicillin/clavulanic acid are inactivation by β -lactamases that are not inhibited by clavulanic acid, and alteration of penicillin binding proteins, which lead to co-resistance to other β -lactam antibiotics. Impermeability of bacteria or efflux pump mechanisms may also contribute to bacterial resistance, including co- and cross-resistance. Susceptibility and resistance patterns can vary with geographical area and bacterial strain, and may change over time.

Pseudomonas spp are naturally resistant to the amoxicillin – clavulanic acid combination. Methicillin resistant *S. aureus* (MRSA) and methicillin resistant *S. pseudintermedius* (MRSP) isolates have been identified in cats and dogs and should be considered resistant to all β -lactam including amoxicillin/clavulanic acid combination.

High resistances (up to 100%) have been reported in *E. coli* isolates from skin and soft tissue infections in dogs.

Pharmacokinetic particulars

After oral administration at the recommended dose in dogs and cats, the absorption of amoxicillin and clavulanic acid is fast. In dogs, the maximum plasma concentration of amoxicillin of 8.5 $\mu\text{g/ml}$ is reached in 1.4 hours and the maximum plasma concentration of clavulanic acid of 0.9 $\mu\text{g/ml}$ is reached in 0.9 hours. Half-life is 1 hour in dogs for both substances.

In cats, the maximum plasma concentration of amoxicillin of 6.6 $\mu\text{g/ml}$ is reached in 1.8 hours and the maximum plasma concentration of clavulanic acid of 3.7 $\mu\text{g/ml}$ is reached in 0.75 hours. Half-life is 1 to 2 hours in cats for both substances.

Elimination is also fast. 12 % of the amoxicillin and 17 % of clavulanic acid is excreted in the urine. The remainder is excreted as inactive metabolites. After repeated oral administration of the recommended dose in dogs and cats, there is no accumulation of amoxicillin or clavulanic acid and the steady state is reached rapidly after first administration.

Pharmaceutical particulars

Major incompatibilities

None known.

Shelf life

50mg and 62.5mg:

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

Shelf life after first opening the immediate packaging: 16 hours.

250mg and 500mg:

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

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Shelf life after first opening the immediate packaging: 16 hours.

Special precautions for storage

This veterinary medicinal product does not require any special storage conditions.

Return any halved tablet to the opened blister-pack and use within 16h.

Nature and composition of immediate packaging

50mg, 250mg and 500mg:

Aluminium/aluminium (oPA/Alu/PE) blister pack with 10 tablets/blister

Cardboard box: Pack sizes of 10, 20, 50, 100, 120, 150, 200, 250, 300, 400, 500, 600, 750 and 1000 tablets.

Not all pack sizes may be marketed.

62.5mg

Aluminium/aluminium (oPA/Alu/PE) blister pack with 10 tablets/blister

Cardboard box: Pack sizes of 10, 100, 250 and 500 tablets.

Not all pack sizes may be marketed.

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

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 national collection systems applicable to the veterinary medicinal product concerned.

Marketing Authorisation Holder (if different from distributor)

Vetoquinol SA

Marketing Authorisation Number

50mg: Vm 06462/3001

62.5mg: Vm 06462/3003

250mg: Vm 06462/3000

500mg: Vm 06462/3002

Significant changes

Date of the first authorisation or date of renewal

62.5mg: 17 April 2013

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50mg, 250mg, 500mg: 18 June 2004

Date of revision of the text

July 2025

Any other information

Legal category

Legal category:POM-V

GTIN

GTIN description:CLAVASEPTIN 50MG 100

GTIN:03605874367821

GTIN description:CLAVASEPTIN 50MG 500

GTIN:03605874367838

GTIN description:CLAVASEPTIN 62.5MG 100

GTIN:03605874367845

GTIN description:CLAVASEPTIN 62.5MG 500

GTIN:03605874367852

GTIN description:CLAVASEPTIN 250MG 100

GTIN:03605874367791

GTIN description:CLAVASEPTIN 250MG 250

GTIN:03605874367807

GTIN description:CLAVASEPTIN 500MG 100

GTIN:03605874367814