CLAVACILLIN- amoxicillin anhydrous and clavulanate potassium tablet Dechra Veterinary Products LLC

Clavacillin® (amoxicillin and clavulanate potassium tablets), USP

Veterinary Tablets

For use in dogs and cats

CAUTION:

Federal law restricts this drug to use by or on the order of a licensed veterinarian.

DESCRIPTION:

Clavacillin (amoxicillin and clavulanate potassium tablets), USP is an orally administered formulation comprised of the broad-spectrum antibiotic amoxicillin trihydrate and the β -lactamase inhibitor, clavulanate potassium (the potassium salt of clavulanic acid).

Amoxicillin trihydrate is a semisynthetic antibiotic with a broad spectrum of bactericidal activity against many gram-positive and gram-negative, aerobic and anaerobic microorganisms. It does not resist destruction by β -lactamases; therefore, it is not effective against β -lactamase-producing bacteria. Chemically, it is D(-)- α -amino-p-hydroxybenzyl penicillin trihydrate.

Clavulanic acid, an inhibitor of β-lactamase enzymes, is produced by the fermentation of *Streptomyces clavuligerus*. Clavulanic acid by itself has only weak antibacterial activity. Chemically, clavulanate potassium is potassium z-(3R,5R)-2-β-hydroxyethylidene clavam-3-carboxylate.

ACTIONS:

Clavacillin is stable in the presence of gastric acid and is not significantly influenced by gastric or intestinal contents. The 2 components are rapidly absorbed resulting in amoxicillin and clavulanic acid concentrations in serum, urine, and tissues similar to those produced when each is administered alone.

Amoxicillin and clavulanic acid diffuse readily into most body tissues and fluids with the exception of brain and spinal fluid, which amoxicillin penetrates adequately when meninges are inflamed. Most of the amoxicillin is excreted unchanged in the urine. Clavulanic acid's penetration into spinal fluid is unknown at this time. Approximately 15% of the administered dose of clavulanic acid is excreted in the urine within the first 6 hours.

Clavacillin combines the distinctive properties of a broad-spectrum antibiotic and a β -lactamase inhibitor to effectively extend the antibacterial spectrum of amoxicillin to include β -lactamase-producing as well as non- β -lactamase-producing aerobic and anaerobic organisms.

MICROBIOLOGY:

Amoxicillin is bactericidal in action and acts through the inhibition of biosynthesis of cell wall mucopeptide of susceptible organisms. The action of clavulanic acid extends the antimicrobial spectrum of amoxicillin to include organisms resistant to amoxicillin and

other β -lactam antibiotics. Amoxicillin/clavulanate has been shown to have a wide range of activity which includes β -lactamase-producing strains of both gram-positive and gram-negative aerobes, facultative anaerobes, and obligate anaerobes. Many strains of the following organisms, including β -lactamase-producing strains, isolated from veterinary sources, were found to be susceptible to amoxicillin/clavulanate *in vitro* but the clinical significance of this activity has not been demonstrated for some of these organisms in animals.

Aerobic bacteria, including *Staphylococcus aureus*¹, β-lactamase-producing *Staphylococcus aureus*¹ (penicillin resistant), *Staphylococcus* species¹, *Staphylococcus epidermidis*, *Staphylococcus intermedius*, *Streptococcus faecalis*, *Streptococcus species*¹, *Corynebacterium pyogenes*, *Corynebacterium species*, *Erysipelothrix rhusiopathiae*, *Bordetella bronchiseptica*, *Escherichia coli*¹, *Proteus mirabilis*, *Proteus species*, *Enterobacter species*, *Klebsiella pneumoniae*, *Salmonella dublin*, *Salmonella typhimurium*, *Pasteurella multocida*, *Pasteurella haemolytica*, *Pasteurella species*¹.

Studies have demonstrated that both aerobic and anaerobic flora are isolated from gingival cultures of dogs with clinical evidence of periodontal disease. Both gram-positive and gram-negative aerobic and anaerobic subgingival isolates indicate sensitivity to amoxicillin/clavulanic acid during antimicrobial susceptibility testing.

1 The susceptibility of these organisms has also been demonstrated in *in vivo* studies.

SUSCEPTIBILITY TEST:

The recommended quantitative disc susceptibility method (FEDERAL REGISTER 37:20527-29; Bauer AW, Kirby WMM, Sherris JC, *et al:* Antibiotic susceptibility testing by standardized single disc method. *Am J Clin Path* 45:493, 1966) utilized 30 mcg Augmentin[®] (AMC) discs for estimating the susceptibility of bacteria to amoxicillin and clavulanate potassium tablets.

INDICATIONS:

Clavacillin Tablets are indicated in the treatment of:

Dogs: Skin and soft tissue infections such as wounds, abscesses, cellulitis, superficial/juvenile and deep pyoderma due to susceptible strains of the following organisms: β-lactamase-producing *Staphylococcus aureus*, non-β-lactamase-producing *Staphylococcus aureus*, non-β-lactamase-producing *Staphylococcus* spp., and *E. coli*.

Periodontal infections due to susceptible strains of both aerobic and anaerobic bacteria. Clavacillin has been shown to be clinically effective for treating cases of canine periodontal disease.

Cats: Skin and soft tissue infections such as wounds, abscesses, and cellulitis/dermatitis due to susceptible strains of the following organisms: β-lactamase-producing *Staphylococcus aureus*, non-β-lactamase-producing *Staphylococcus aureus*, *Staphylococcus* spp., *Streptococcus* spp., *E. coli*, and *Pasteurella* spp. Urinary tract infections (cystitis) due to susceptible strains of *E. coli*.

Therapy may be initiated with Clavacillin prior to obtaining results from bacteriological and susceptibility studies. A culture should be obtained prior to treatment to determine susceptibility of the organisms to Clavacillin. Following determination of susceptibility results and clinical response to medication, therapy may be reevaluated.

CONTRAINDICATIONS:

The use of this drug is contraindicated in animals with a history of an allergic reaction to any of the penicillins or cephalosporins.

WARNINGS:

Keep Clavacillin Tablets in a secure location out of reach of dogs, cats, and other animals to prevent accidental ingestion or overdose.

Safety of use in pregnant or breeding animals has not been determined. Store at controlled room temperature, 68-77°F (20-25°C).

Do not remove from foil strip until ready to use.

ADVERSE REACTIONS:

Clavacillin contains a semisynthetic penicillin (amoxicillin) and has the potential for producing allergic reactions. If an allergic reaction occurs, administer epinephrine and/or steroids.

Post-Approval Experience (July, 2017)

The following adverse events are based on post-approval adverse drug experience reporting. Not all adverse events are reported to FDA/CVM. It is not always possible to reliably estimate the adverse event frequency or establish a causal relationship to product exposure using these data.

The following adverse events reported for dogs and cats are listed in decreasing order of reporting frequency for amoxicillin and clavulanate potassium tablets: Anorexia, lethargy, vomiting and diarrhea.

To report suspected adverse events, for technical assistance or to obtain a copy of the Safety Data Sheet (SDS), contact Dechra at (866) 933-2472.

For additional information about adverse drug experience reporting for animal drugs, contact FDA at 1-888-FDA-VETS, or online at http://www.fda.gov/reportanimalae

DOSAGE AND ADMINISTRATION:

Dogs: The recommended dosage is 6.25 mg/lb of body weight twice a day.

Skin and soft tissue infections such as abscesses, cellulitis, wounds, superficial/juvenile pyoderma, and periodontal infections should be treated for 5-7 days or for 48 hours after all symptoms have subsided. If no response is seen after 5 days of treatment, therapy should be discontinued and the case reevaluated. Deep pyoderma may require treatment for 21 days; the maximum duration of treatment should not exceed 30 days.

Cats: The recommended dosage is 62.5 mg twice a day.

Skin and soft tissue infections such as abscesses and cellulitis/dermatitis should be treated for 5-7 days or for 48 hours after all symptoms have subsided, not to exceed 30 days. If no response is seen after 3 days of treatment, therapy should be discontinued and the case reevaluated.

Urinary tract infections may require treatment for 10-14 days or longer. The maximum duration of treatment should not exceed 30 days.

HOW SUPPLIED:

Clavacillin Tablets in the following strengths are supplied in strip packs. Each carton can hold 5 strips with 14 tablets (70 tablets per carton) or 15 strips with 14 tablets (210 tablets per carton).

Each 62.5-mg tablet contains amoxicillin trihydrate equivalent to 50 mg of amoxicillin activity and 12.5 mg of clavulanic acid as the potassium salt. For use in dogs and cats.

Each 125-mg tablet contains amoxicillin trihydrate equivalent to 100 mg of amoxicillin activity and 25 mg of clavulanic acid as the potassium salt. For use in dogs only.

Each 250-mg tablet contains amoxicillin trihydrate equivalent to 200 mg of amoxicillin activity and 50 mg of clavulanic acid as the potassium salt. For use in dogs only.

Each 375-mg tablet contains amoxicillin trihydrate equivalent to 300 mg of amoxicillin activity and 75 mg of clavulanic acid as the potassium salt. For use in dogs only.

Dispense according to recommendations outlined in Dosage and Administration section.

Approved by FDA under ANADA # 200-592

Augmentin is a trademark owned by GlaxoSmithKline.

Manufactured for:

Dechra Veterinary Products 7015 College Boulevard, Suite 525 Overland Park, KS 66211 USA

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Rev. April 2022



PRINCIPAL DISPLAY PANEL - 62.5 mg Tablet Blister Pack Carton

Clavacillin[®] (amoxicillin and clavulanate potassium tablets), USP 62.5 mg

Veterinary Tablets

For use in dogs and cats

15 strips, 14 tablets each 210 tablets

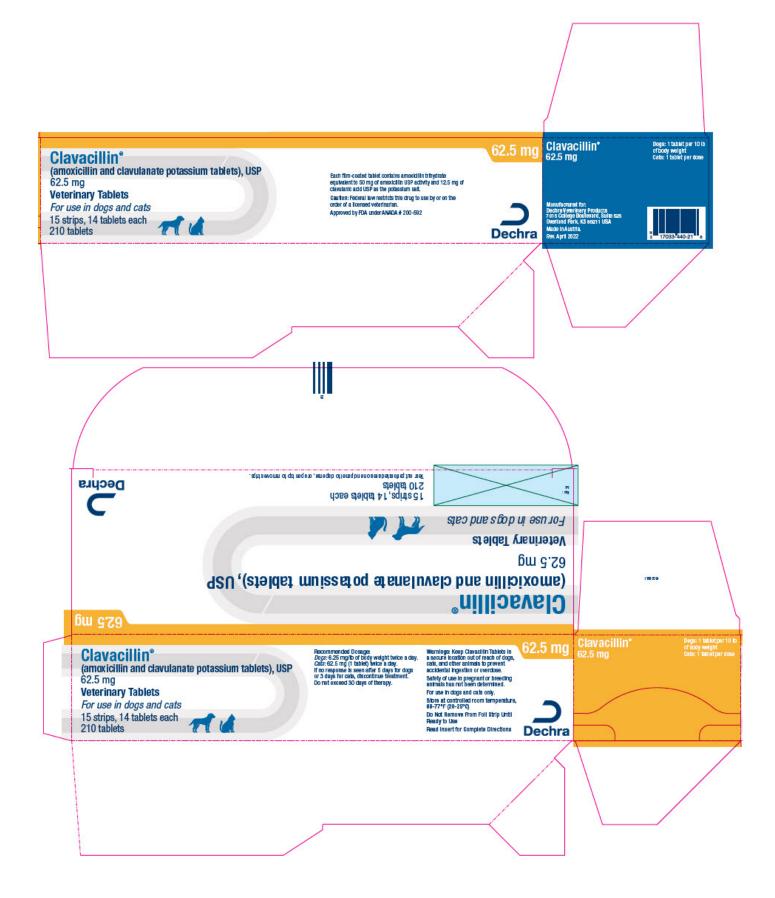
62.5 mg

Each film-coated tablet contains amoxicillin trihydrate

equivalent to 50 mg of amoxicillin USP activity and 12.5 mg of clavulanic acid USP as the potassium salt.

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PRINCIPAL DISPLAY PANEL - 125 mg Tablet Blister Pack Carton

Clavacillin[®] (amoxicillin and clavulanate potassium tablets), USP 125 mg Veterinary Tablets

For use in dogs

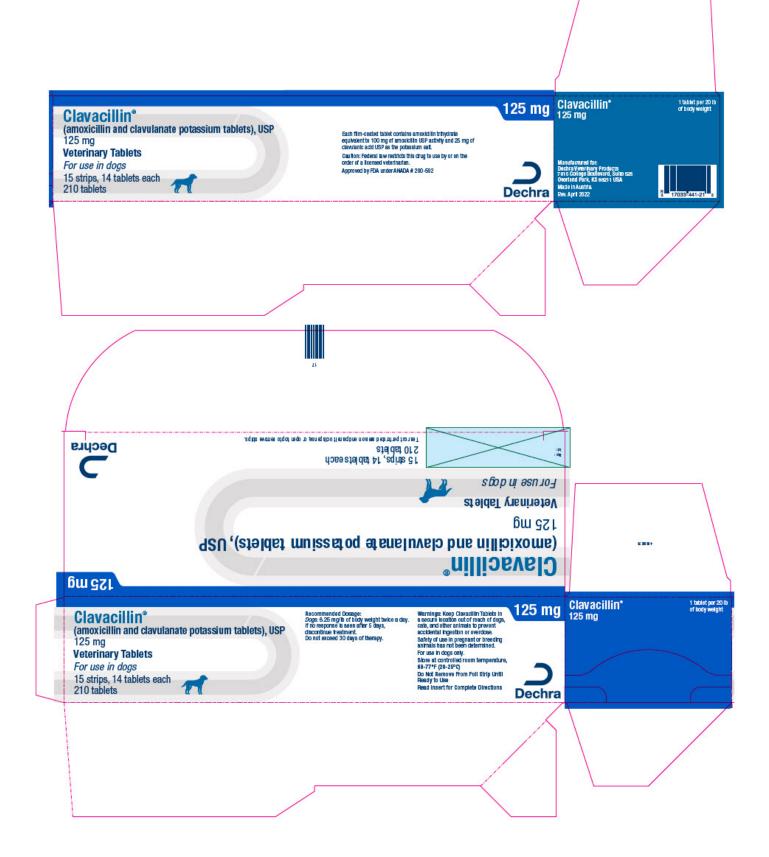
15 strips, 14 tablets each 210 tablets

125 mg

Each film-coated tablet contains amoxicillin trihydrate equivalent to 100 mg of amoxicillin USP activity and 25 mg of clavulanic acid USP as the potassium salt.

Caution: Federal law restricts this drug to use by or on the order of a licensed veterinarian.

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PRINCIPAL DISPLAY PANEL - 250 mg Tablet Blister Pack Carton

Clavacillin[®] (amoxicillin and clavulanate potassium tablets), USP 250 mg Veterinary Tablets

For use in dogs

15 strips, 14 tablets each 210 tablets

250 mg

Each film-coated tablet contains amoxicillin trihydrate equivalent to 200 mg of amoxicillin USP activity and 50 mg of clavulanic acid USP as the potassium salt.

Caution: Federal law restricts this drug to use by or on the order of a licensed veterinarian.

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PRINCIPAL DISPLAY PANEL - 375 mg Tablet Blister Pack Carton

Clavacillin[®] (amoxicillin and clavulanate potassium tablets), USP 375 mg Veterinary Tablets

For use in dogs

15 strips, 14 tablets each 210 tablets

375 mg

Each film-coated tablet contains amoxicillin trihydrate equivalent to 300 mg of amoxicillin USP activity and 75 mg of clavulanic acid USP as the potassium salt.

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CLAVACILLIN

amoxicillin anhydrous and clavulanate potassium tablet

Product Information

Product Type	PRESCRIPTION ANIMAL DRUG	Item Code (Source)	NDC:17033-440
Route of Administration	ORAL		

A	ctive Ingredien	t/Active Moi	ety					
		Ingredi	ent Name				is of ngth	Strength
					amoxicillin anhydrous		50 mg	
cla	clavulanate potassium (UNII: Q420MW3AT8) (clavulanic acid - UNII:23521W1S24) clavulanic					clavulanic	acid	12.5 mg
Ρ	roduct Charact	eristics						
С	olor	YELLOW	Sco	ore			no score	
SI	hape	ROUND	Siz	e			7mm	
FI	avor		Im	print Code		1	P1	
Co	ontains							
P	ackaging							
P #		Package D	Description	Marketing	y Start Dat	e Mai	rketing	End Date
#		Package D 5 in 1 CARTON	Description	Marketing	g Start Dat	e Mai	rketing	End Date
#	ltem Code		•	Marketing) Start Dat	e Mai	rketing	End Date
# 1	Item Code NDC:17033-440-07	5 in 1 CARTON	- R PACK	Marketing	9 Start Dat	e Mai	rketing	End Date
# 1 1	Item Code NDC:17033-440-07	5 in 1 CARTON 14 in 1 BLISTEF	PACK	Marketing	g Start Dat	e Mai	rketing	End Date
# 1 1 2	Item Code NDC:17033-440-07	5 in 1 CARTON 14 in 1 BLISTEF 15 in 1 CARTON	PACK	Marketing	9 Start Dat	e Mai	rketing	End Date
# 1 1 2	Item Code NDC:17033-440-07	5 in 1 CARTON 14 in 1 BLISTEF 15 in 1 CARTON	PACK	Marketing	g Start Dat	e Mai	rketing	End Date
# 1 1 2 2	Item Code NDC:17033-440-07	5 in 1 CARTON 14 in 1 BLISTEF 15 in 1 CARTON 14 in 1 BLISTEF	R PACK N R PACK	Marketing	9 Start Dat	e Mai	rketing	End Date
# 1 2 2	Item Code NDC:17033-440-07 NDC:17033-440-21	5 in 1 CARTON 14 in 1 BLISTEF 15 in 1 CARTON 14 in 1 BLISTEF	R PACK N R PACK		9 Start Dat Marketir Da	ng Start	Mark	End Date

CLAVACILLIN amoxicillin anhydrous and clav	ulanate potassium tablet					
Product Information						
Product Type	PRESCRIPTION ANIMAL DRUG	ltem Code	(Source)	NDC:1	7033-441	
Route of Administration	ORAL					
Active Ingredient/Active	Moiety					
Ing	redient Name		Basis o Strengt		Strength	
amoxicillin anhydrous (UNII: 9EM UNII:9EM05410Q9)	105410Q9) (amoxicillin anhydrous -		amoxicillin anhydrous		100 mg	
clavulanate potassium (UNII: Q4	20MW3AT8) (clavulanic acid - UNII:	23521W1S24)	clavulanic acio	I	25 mg	

Product Characteristics							
Co	olor	YELLOW	Sco	ore		no score	
Sł	nape	ROUND	Siz	e		9mm	
Fla	avor		Imp	orint Code		P2	
Co	ontains						
Pa	ackaging						
		Package Descr	ription	Marketing Start Date	Ma	arketing End Dat	
#		Package Descr 5 in 1 CARTON	iption	Marketing Start Date	Ma	arketing End Dat	
# 1	ltem Code		-	Marketing Start Date	Ma	arketing End Dat	
# 1 1	ltem Code	5 in 1 CARTON	-	Marketing Start Date	Ma	arketing End Dat	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
ANADA	ANADA200592	01/10/2019		

CLAVACILLIN						
amoxicillin anhydrous and	clavulanate potas	sium tablet				
Product Information						
Product Type	PRESCRIPTION A	ANIMAL DRUG	Item Code	(Source)	NDC:	17033-442
Route of Administration	ORAL					
Active Ingredient/Acti	ive Moiety					
	Ingredient Nam	е			is of ngth	Strength
amoxicillin anhydrous (UNII: UNII:9EM05410Q9)	9EM05410Q9) (amo>	kicillin anhydrous -		amoxicillin anhydrous		200 mg
clavulanate potassium (UNI	I: Q42OMW3AT8) (clav	ulanic acid - UNII:2	23521W1S24)	clavulanic	acid	50 mg
Product Characteristi	cs					
Color	YELLOW	Score		1	no score	
Shape	ROUND	Size			11mm	
Flavor		Imprint Code			Р3	
Contains						
Packaging						

# Item Code	Packa	ge Description	Marketing	Start Dat	e Marl	keting	End Date
1 NDC:17033-442-07	5 in 1 CAF	RTON					
1	14 in 1 BL	ISTER PACK					
2 NDC:17033-442-21	15 in 1 CA	ARTON					
2	14 in 1 BL	ISTER PACK					
Marketing In	nformat	ion					
Marketing Category	Applica	tion Number or M Citation	onograph	Marketin Dat	-	Mark	ceting End Date
ANADA	ANADA2005	92		01/10/2019			
LAVACILLIN							
		vulanate potassiu	m tablet				
amoxicillin anhydro	us and clav	vulanate potassiu	m tablet				
nmoxicillin anhydro Product Inform	us and clav	vulanate potassiu PRESCRIPTION ANIM	_	tem Code ((Source)	NDC	:17033-443
moxicillin anhydro Product Inform Product Type	us and clav		_	tem Code ((Source)	NDC	:17033-443
amoxicillin anhydro Product Inform Product Type	us and clav	PRESCRIPTION ANIM	_	tem Code ((Source)	NDC	:17033-443
CLAVACILLIN amoxicillin anhydro Product Inform Product Type Route of Administ Active Ingredie	us and clav	PRESCRIPTION ANIM	_	tem Code ((Source)	NDC	:17033-443
moxicillin anhydro Product Inform Product Type Route of Administ	us and clav nation tration nt/Active	PRESCRIPTION ANIM	_	tem Code ((Source) Basis Strer	s of	
moxicillin anhydro Product Inform Product Type Route of Administ Active Ingredie	us and clav nation tration nt/Active Ing	PRESCRIPTION ANIM ORAL Moiety redient Name	AL DRUG	tem Code (Basi	s of	:17033-443 Strength 300 mg
moxicillin anhydro Product Inform Product Type Route of Administ	us and clav nation tration nt/Active Ing	PRESCRIPTION ANIM ORAL Moiety redient Name	AL DRUG		Basis Strer amoxicillin	s of ngth	Strengt
Product Inform Product Type Route of Administ Active Ingredie	us and clav nation tration nt/Active Ing ous (UNII: 9EM	PRESCRIPTION ANIM ORAL Moiety redient Name	AL DRUG		Basis Strer amoxicillin anhydrous	s of ngth	Strengt 300 mg

Color	YELLOW	Score	no score
Shape	ROUND	Size	13mm
Flavor		Imprint Code	P4
Contains			

Packaging					
#	ltem Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:17033-443-07	5 in 1 CARTON			
1		14 in 1 BLISTER PACK			
2	NDC:17033-443-21	15 in 1 CARTON			
2		14 in 1 BLISTER PACK			

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	Date

Labeler - Dechra Veterinary Products LLC (362142734)

Revised: 3/2024

Dechra Veterinary Products LLC