IMOXI FOR DOGS- imidacloprid and moxidectin solution Vetoquinol USA, Inc.

-----

IMOXI<sup>™</sup> Topical Solution for Dogs (imidacloprid + moxidectin)

ACPX1008

Once-a-month topical solution for the prevention of heartworm disease, the treatment of circulating microfilariae, kills adult fleas, is indicated for the treatment of flea infestations, the treatment and control of sarcoptic mange, as well as the treatment and control of intestinal parasite infections in dogs and puppies that are at least 7 weeks of age and that weigh at least 3 lbs.

#### WARNING

- DO NOT ADMINISTER THIS PRODUCT ORALLY
- For the first 30 minutes after application ensure that dogs cannot lick the product from application sites on themselves or other treated animals.
- Children should not come in contact with application sites for two (2) hours after application. (See Contraindications, Warnings, Human Warnings, and Adverse

Reactions, for more information.)

#### CAUTION:

Federal (USA) Law restricts this drug to use by or on the order of a licensed veterinarian.

#### **DESCRIPTION:**

IMOXI<sup>™</sup> Topical Solution for Dogs (10% imidacloprid + 2.5% moxidectin) is a colorless to yellow ready-to-use solution packaged in single dose applicator tubes for topical treatment of dogs. The formulation and dosage schedule are designed to provide a minimum of 4.5 mg/lb. (10 mg/kg) imidacloprid and 1.1 mg/lb. (2.5 mg/kg) moxidectin based on body weight.

Imidacloprid is a chloronicotinyl nitroguanidine insecticide. The chemical name for imidacloprid is 1- [(6-Chloro- 3-pyridinyl) methyl]-N-nitro-2-imidazolidinimine. Moxidectin is a semisynthetic macrocyclic lactone endectocide derived from the actinomycete *Streptomycetes cyaneogriseus noncyanogenus*. The chemical name for moxidectin is [6R, 23E, 25S(E)]-5-O-Demethyl-28-deoxy- 25-(1,3-dimethl-1- butenyl)-6,28-epoxy-23-(methoxyimino) milbemycin B.

# **INDICATIONS:**

IMOXI<sup>™</sup> Topical Solution for Dogs is indicated for the prevention of heartworm disease caused by *Dirofilaria immitis* and the treatment of *Dirofilaria immitis* circulating microfilariae in heartworm-positive dogs. IMOXI<sup>™</sup> Topical Solution for Dogs kills adult fleas and is indicated for the treatment of flea infestations (*Ctenocephalides felis*). IMOXI<sup>™</sup> Topical Solution for Dogs is indicated for the treatment and control of sarcoptic mange caused by *Sarcoptes scabiei* var. *canis*. IMOXI<sup>™</sup> Topical Solution for Dogs is also indicated for the treatment and control of the following intestinal parasites:

		I	ntestinal S	tage
Intestir	nal Parasite	Adult	Immature Adult	Fourth Stage Larvae
Hookworm	Ancylostoma caninum	Х	X	Х
Species	Uncinaria stenocephala	х	X	Х
Roundworm	Toxocara canis	Х		Х
Species	Toxascaris leonina	Х		
Whipworm	Trichuris vulpis	Х		

# **CONTRAINDICATIONS:**

Do not administer this product orally. (See WARNINGS)

Do not use this product (containing 2.5% moxidectin) on cats.

# WARNINGS:

For the first 30 minutes after application:

Ensure that dogs cannot lick the product from application sites on themselves or other treated dogs, and separate treated dogs from one another and from other pets to reduce the risk of accidental ingestion.

Ingestion of this product by dogs may cause serious adverse reactions including depression, salivation, dilated pupils, incoordination, panting, and generalized muscle tremors.

In avermectin sensitive dogs,<sup>1</sup> the signs may be more severe and may include coma and death.<sup>2</sup>

# HUMAN WARNINGS:

<sup>1</sup> Some dogs are more sensitive to avermectins due to a mutation in the MDR1 gene. Dogs with this mutation may develop signs of severe avermectin toxicity if they ingest this product. The most common breeds associated with this mutation include Collies and Collie crosses.

<sup>2</sup> Although there is no specific antagonist for avermectin toxicity, even severely affected dogs have completely recovered from avermectin toxicity with intensive veterinary supportive care.

# Not for human use. Keep out of the reach of children. Children should not come in contact with application sites for two (2) hours after application.

Causes eye irritation. Harmful if swallowed. Do not get in eyes or on clothing. Avoid contact with skin. Exposure to the product has been reported to cause headache; dizziness; and redness, burning, tingling, or numbness of the skin.

# Wash hands thoroughly with soap and warm water after handling.

If contact with eyes occurs, hold eyelids open and flush with copious amounts of water for 15 minutes. If eye irritation develops or persists, contact a physician. If swallowed, call poison control center or physician immediately for treatment advice. Have person sip a glass of water if able to swallow. Do not induce vomiting unless told to do so by the poison control center or physician. People with known hypersensitivity to benzyl alcohol, imidacloprid or moxidectin should administer the product with caution. In case of allergic reaction, contact a physician. If contact with skin or clothing occurs, take off contaminated clothing. Wash skin immediately with plenty of soap and water. Call a poison control center or physician for treatment advice.

The Safety Data Sheet (SDS) provides additional occupational safety information. For consumer questions call 1-800-835-9496.

For a copy of the Safety Data Sheet (SDS) or to report adverse reactions call Vetoquinol USA, Inc. at 1-800-835-9496.

# **PRECAUTIONS:**

Do not dispense dose applicator tubes without complete safety and administration information.

Use with caution in sick, debilitated, or underweight animals. The safety of IMOXI<sup>™</sup> Topical Solution for Dogs has not been established in breeding, pregnant, or lactating dogs. The safe use of IMOXI<sup>™</sup> Topical Solution for Dogs has not been established in puppies and dogs less than 7 weeks of age or less than 3 lbs. body weight. Prior to administration of IMOXI<sup>™</sup> Topical Solution for Dogs, dogs should be tested for existing heartworm infection. At the discretion of the veterinarian, infected dogs should be treated with an adulticide to remove adult heartworms. The safety of IMOXI<sup>™</sup> Topical Solution for Dogs is not effective against adult *D. immitis*. Although the number of circulating microfilariae is substantially reduced in most dogs following treatment with IMOXI<sup>™</sup> Topical Solution for Dogs, the microfilaria count in some heartworm-positive dogs may increase or remain unchanged following treatment with IMOXI<sup>™</sup> Topical Solution for Dogs, alone or in a dosing regimen with melarsomine dihydrochloride.

# (See ADVERSE REACTIONS and ANIMAL SAFETY - Safety Study in Heartworm-Positive Dogs.)

IMOXI<sup>™</sup> Topical Solution for Dogs has not been evaluated in heartworm-positive dogs with Class 4 heartworm disease.

# ADVERSE REACTIONS:

# Heartworm-Negative Dogs

**Field Studies:** Following treatment with imidacloprid and moxidectin topical solution or an active control, dog owners reported the following post-treatment reactions:

OBSERVATION	imidacloprid and moxidectin topical solution n=128	Active Control n=68
Pruritus	19 dogs (14.8%)	7 dogs (10.3%)
Residue	9 dogs (7.0%)	5 dogs (7.4%)
Medicinal Odor	5 dogs (3.9%)	None observed
Lethargy	1 dog (0.8%)	1 dog (1.5%)
Inappetence	1 dog (0.8%)	1 dog (1.5%)
Hyperactivity	1 dog (0.8%)	None observed

During a field study using 61 dogs with pre-existing flea allergy dermatitis, one (1.6 %) dog experienced localized pruritus immediately after imidacloprid application, and one investigator noted hyperkeratosis at the application site of one dog (1.6 %).

In a field safety and effectiveness study, imidacloprid and moxidectin topical solution was administered to 92 client-owned dogs with sarcoptic mange. The dogs ranged in age from 2 months to 12.5 years and ranged in weight from 3 to 231.5 pounds. Adverse reactions in dogs treated with imidacloprid and moxidectin topical solution included hematochezia, diarrhea, vomiting, lethargy, inappetence, and pyoderma.

**Laboratory Effectiveness Studies:** One dog in a laboratory effectiveness study experienced weakness, depression, and unsteadiness between 6 and 9 days after application with imidacloprid and moxidectin topical solution. The signs resolved without intervention by day 10 post-application. The signs in this dog may have been related to peak serum levels of moxidectin, which vary between dogs, and occur between 1 and 21 days after application of imidacloprid and moxidectin topical solution.

The following clinical observations also occurred in laboratory effectiveness studies following application with imidacloprid and moxidectin topical solution and may be directly attributed to the drug or may be secondary to the intestinal parasite burden or other underlying conditions in the dogs: diarrhea, bloody stools, vomiting, anorexia, lethargy, coughing, ocular discharge and nasal discharge. Observations at the application sites included damp, stiff or greasy hair, the appearance of a white deposit on the hair, and mild erythema, which resolved without treatment within 2 to 48 hours.

# <u>Heartworm-Positive Dogs</u>

**Field Study:** A 56-day field safety study was conducted in 214 *D. immitis* heartworm and microfilaria positive dogs with Class 1, 2 or 3 heartworm disease. All dogs received imidacloprid and moxidectin topical solution on Study Days 0 and 28; 108 dogs also received melarsomine dihydrochloride on Study Days – 14, 14, and 15. All dogs were hospitalized for a minimum of 12 hours following each treatment. Effectiveness against circulating *D. immitis* microfilariae was > 90 % at five of six sites; however, one site had an effectiveness of 73.3 %. The microfilaria count in some heartworm-positive dogs increased or remained unchanged following treatment with imidacloprid and moxidectin topical solution alone or in a dosing regimen with melarsomine dihydrochloride. Following treatment with imidacloprid and moxidectin topical solution alone or in a dosing regimen with melarsomine dihydrochloride, the following adverse reactions were observed:

Adverse Reaction	Dogs Treated with imidacloprid and moxidectin topical solution n=106	Dogs Treated with imidacloprid and moxidectin topical solution + Melarsomine n=108
Cough	24 (22.6%)	25 (23.1%)
Lethargy	14 (13.2%)	42 (38.9%)
Vomiting	11 (10.4%)	18 (16.7%)
Diarrhea, including hemorrhagic	10 (9.4%)	22 (20.4%)
Inappetence	7 (6.6%)	19 (17.6%)
Dyspnea	6 (5.7%)	10 (9.3%)
Tachypnea	1 (<1%)	7 (6.5%)
Pulmonary Hemorrhage	0	1 (<1%)
Death	0	3 (2.8%)

Three dogs treated with imidacloprid and moxidectin topical solution in a dosing regimen with melarsomine dihydrochloride died of pulmonary embolism from dead and dying heartworms. One dog, treated with imidacloprid and moxidectin topical solution and melarsomine dihydrochloride, experienced pulmonary hemorrhage and responded to supportive medical treatment. Following the first treatment with imidacloprid and moxidectin topical solution alone, two dogs experienced adverse reactions (coughing, vomiting, and dyspnea) that required hospitalization. In both groups, there were more adverse reactions to imidacloprid and moxidectin topical solution following the first treatment than the second treatment.

To report a suspected adverse reaction, call 1-800-835-9496.

# Post-Approval Experience:

The following adverse events are based on postapproval adverse drug experience reporting. Not all adverse reactions 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 this data. The following adverse events in dogs are listed in decreasing order of reporting frequency: depression/lethargy, vomiting, pruritus, diarrhea, anorexia, hyperactivity, ataxia, trembling, hypersalivation, application site reactions (alopecia, pruritus, lesions, and erythema), seizures, and anaphylaxis/anaphylactic reactions (hives, urticaria, facial swelling, edema of the head).

# Serious reactions, including neurologic signs and death have been reported when cats have been exposed (orally and topically) to this product.

In humans, nausea, numbness or tingling of the mouth/lips and throat, ocular and dermal irritation, pruritus, headache, vomiting, diarrhea, depression and dyspnea have been reported following exposure to this product.

To report suspected adverse drug events, for technical assistance or to obtain a copy of the SDS, contact Vetoquinol USA, Inc. at 1-800-835-9496.

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:**

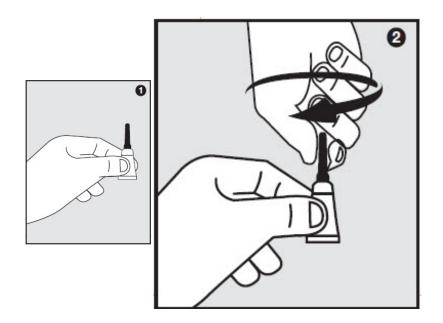
The recommended minimum dose is 4.5 mg/lb. (10 mg/kg) imidacloprid and 1.1 mg/lb. (2.5 mg/kg) moxidectin, once a month, by topical administration.

Do not apply to irritated skin.

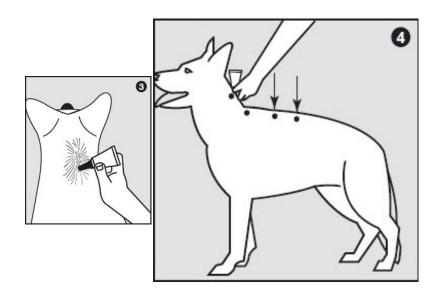
**1.** Remove one dose applicator tube from the package. As specified in the table, administer the entire contents of the IMOXI<sup>™</sup> Topical Solution for Dogs tube that correctly corresponds with the body weight of the dog.

Dog (lbs.)	Volume (mL)	Imidacloprid (mg)	Moxidectin (mg)
3-9	0.4	40	10
9.1-20	1.0	100	25
20.1-55	2.5	250	62.5
55.1-88	4.0	400	100
88.1-110 *	5.0	500	125

\* Dogs over 110 lbs. should be treated with the appropriate combination of IMOXI<sup>™</sup> Topical Solution for Dogs tubes.



2. While holding the Twist•N•Go<sup>™</sup> tube in an upright position, twist the dispensing cap clockwise about ½ turn to break the tube's seal. Remove the cap from the tube.



**3.** The dog should be standing for application. Position the dispensing tip on the dog's back between the shoulder blades. The dispensing tip of the tube can be used to part the dog's hair until the skin is visible.

**4.** For dogs weighing 20 lbs. or less, place the tip of the tube on the skin and apply the entire contents directly on the exposed skin at one spot between the shoulder blades. For dogs weighing more than 20 lbs., place the tip of the tube on the skin and apply the entire contents directly on the exposed skin at 3 or 4 spots on the top of the backline from the base of the neck to the upper back in an area inaccessible to licking. Do not apply an amount of solution at any one location that could run off the side of the dog.

Do not let this product get in your dog's mouth or eyes. **Do not allow the dog to lick any of the application sites for 30 minutes.** In households with multiple pets, keep each treated dog separated from other treated dogs and other pets for 30 minutes after application to prevent licking the application sites. **(See WARNINGS)** 

Stiff hair, a damp appearance of the hair, pink skin, or a slight powdery residue may be observed at the application site on some animals. This is temporary and does not affect the safety and effectiveness of the product.

Shampooing 90 minutes after treatment does not reduce the effectiveness of IMOXI<sup>™</sup> Topical Solution for Dogs in the prevention of heartworm disease. Shampooing or water immersion 4 days after treatment will not reduce the effectiveness of IMOXI<sup>™</sup> Topical Solution for Dogs in the treatment of flea infestations. However, shampooing as often as once weekly may reduce the effectiveness of the product against fleas.

Heartworm Prevention: For prevention of heartworm disease, IMOXI<sup>™</sup> Topical Solution for Dogs should be administered at one-month intervals. IMOXI<sup>™</sup> Topical Solution for Dogs may be administered year-round or at a minimum should start one month before the first expected exposure to mosquitoes and should continue at monthly intervals until one month after the last exposure to mosquitoes. If a dose is missed and a 30-day interval between doses is exceeded, administer IMOXI<sup>™</sup> Topical Solution for Dogs immediately and resume the monthly dosing schedule. When replacing another heartworm preventative product in a heartworm prevention program, the first treatment with IMOXI<sup>™</sup> Topical Solution for Dogs should be given within one month of the last dose of the former medication.

Treatment of Circulating Microfilaria: For the treatment of circulating D. immitis

microfilaria in heartworm-positive dogs, IMOXI<sup>™</sup> Topical Solution for Dogs should be administered at one-month intervals. Treatment with an approved adulticide therapy is recommended because IMOXI<sup>™</sup> Topical Solution for Dogs is not effective for the treatment of adult *D. immitis*. **(See PRECAUTIONS)** 

**Flea Treatment:** For the treatment of flea infestations, IMOXI<sup>™</sup> Topical Solution for Dogs should be administered at one-month intervals. If the dog is already infested with fleas when the first dose of IMOXI<sup>™</sup> Topical Solution for Dogs is administered, adult fleas on the dog will be killed. However, reinfestation from the emergence of preexisting pupae in the environment may continue to occur for six weeks or longer after treatment is initiated. Dogs treated with imidacloprid, including those with pre-existing flea allergy dermatitis have shown clinical improvement as a direct result of elimination of fleas from the dog.

**Treatment and Control of Intestinal Nematode Infections:** For the treatment and control of intestinal hookworm infections caused by *Ancylostoma caninum* and *Uncinaria stenocephala* (adults, immature adults and fourth stage larvae) and roundworm infections caused by *Toxocara canis* (adults and fourth stage larvae), and *Toxascaris leonina* (adults), and whipworm infections caused by *Trichuris vulpis* (adults), IMOXI<sup>™</sup> Topical Solution for Dogs should be administered once as a single topical dose.

**Treatment and Control of Sarcoptic Mange:** For the treatment and control of sarcoptic mange caused by *Sarcoptes scabiei* var. *canis*, IMOXI<sup>™</sup> Topical Solution for Dogs should be administered as a single topical dose. A second monthly dose may be administered if necessary.

# ANIMAL SAFETY:

# Heartworm-Negative Dogs

**Field Study:** In a controlled, double-masked, field safety study, imidacloprid and moxidectin topical solution was administered to 128 dogs of various breeds, 3 months to 15 years of age, weighing 4 to 157 pounds. Imidacloprid and moxidectin topical solution was used safely in dogs concomitantly receiving ACE inhibitors, anticonvulsants, antihistamines, antimicrobials, chondroprotectants, corticosteroids,

imrnunotherapeutics, MAO inhibitors, NSAIDs, ophthalmic medications,

sympathomimetics, synthetic estrogens, thyroid hormones, and urinary acidifiers. Owners reported the following signs in their dogs after application of imidacloprid and moxidectin topical solution: pruritus, flaky/greasy residue at the treatment site, medicinal odor, lethargy, inappetence, and hyperactivity.

# (See ADVERSE REACTIONS)

**Safety Study in Puppies:** Imidacloprid and moxidectin topical solution was applied topically at 1, 3 and 5X the recommended dose to 7-week-old Beagle puppies once every 2 weeks for 6 treatments on days 0, 14, 28, 42, 56, and 70. Loose stools and diarrhea were observed in all groups, including the controls, throughout the study. Vomiting was seen in one puppy from the 1X treatment group (day 57), in two puppies from the 3X treatment group (days 1 and 79), and in one puppy from the 5X treatment group (day 1). Two puppies each in the 1X, 3X, and 5X groups had decreased appetites within 24 hours post-dosing. One puppy from the 1X treatment group had pruritus for one hour following the fifth treatment. A puppy from the 5X treatment group displayed

rapid, difficult breathing from 4 to 8 hours following the second treatment.

**Dermal Dose Tolerance Study:** Imidacloprid and moxidectin topical solution was administered topically to 8-month-old Beagle dogs at 10X the recommended dose once. One dog showed signs of treatment site irritation after application. Two dogs vomited, one at 6 hours and one at 6 days post-treatment. Increased RBC, hemoglobin, activated partial thromboplastin, and direct bilirubin were observed in the treated group. Dogs in the treated group did not gain as much weight as the control group.

**Oral Safety Study in Beagles:** Imidacloprid and moxidectin topical solution was administered once orally at the recommended topical dose to 12 dogs. Six dogs vomited within 1 hour of receiving the test article, 2 of these dogs vomited again at 2 hours, and 1 dog vomited again up to 18 hours post-dosing. One dog exhibited shaking (nervousness) 1 hour post-dosing. Another dog exhibited abnormal neurological signs (circling, ataxia, generalized muscle tremors, and dilated pupils with a slow pupillary light response) starting at 4 hours post-dosing through 18 hours post-dosing. Without treatment, this dog was neurologically normal at 24 hours and had a normal appetite by 48 hours post-dosing.

# (See CONTRAINDICATIONS)

**Dermal Safety Study in Ivermectin-Sensitive Collies:** Imidacloprid and moxidectin topical solution was administered topically at 3 and 5X the recommended dose every 28 days for 3 treatments to Collies which had been prescreened for avermectin sensitivity. No clinical abnormalities were observed.

**Oral Safety Study in Ivermectin-Sensitive Collies:** Imidacloprid and moxidectin topical solution was administered orally to 5 pre-screened ivermectin-sensitive Collies. The Collies were asymptomatic after ingesting 10% of the minimum labeled dose. At 40% of the minimum recommended topical dose, 4 of the dogs experienced neurological signs indicative of avermectin toxicity including depression, ataxia, mydriasis, salivation, muscle fasciculation, and coma, and were euthanized.

# (See CONTRAINDICATIONS)

# <u>Heartworm-Positive Dogs</u>

**Laboratory Safety Study in Heartworm-Positive Dogs:** Imidacloprid and moxidectin topical solution was administered topically at 1 and 5X the recommended dose every 14 days for 3 treatments to dogs with adult heartworm infections and circulating microfilaria. At 5X, one dog was observed vomiting three hours after the second treatment. Hypersensitivity reactions were not seen in the 5X treatment group. Microfilaria counts decreased with treatment.

# **STORAGE INFORMATION:**

Store at temperatures between  $20^{\circ}$  C (68 °F) and  $25^{\circ}$  C (77 °F), avoiding excess heat or cold.

# HOW SUPPLIED:

	tubes	6 × 4.0 mL
Applications Per	6 × 1.0 mL	tubes
Package:	tubes	6 × 5.0 mL
	6 × 2.5 mL	tubes
	tubes	

Approved by FDA under ANADA # 200-615 Made in U.S.A. © 2019 Vetoquinol USA, Inc.

Vetoquinol, the Vetoquinol logo, and IMOXI<sup>™</sup> Topical Solution for Dogs are registered trademarks of Vetoquinol USA, Inc.

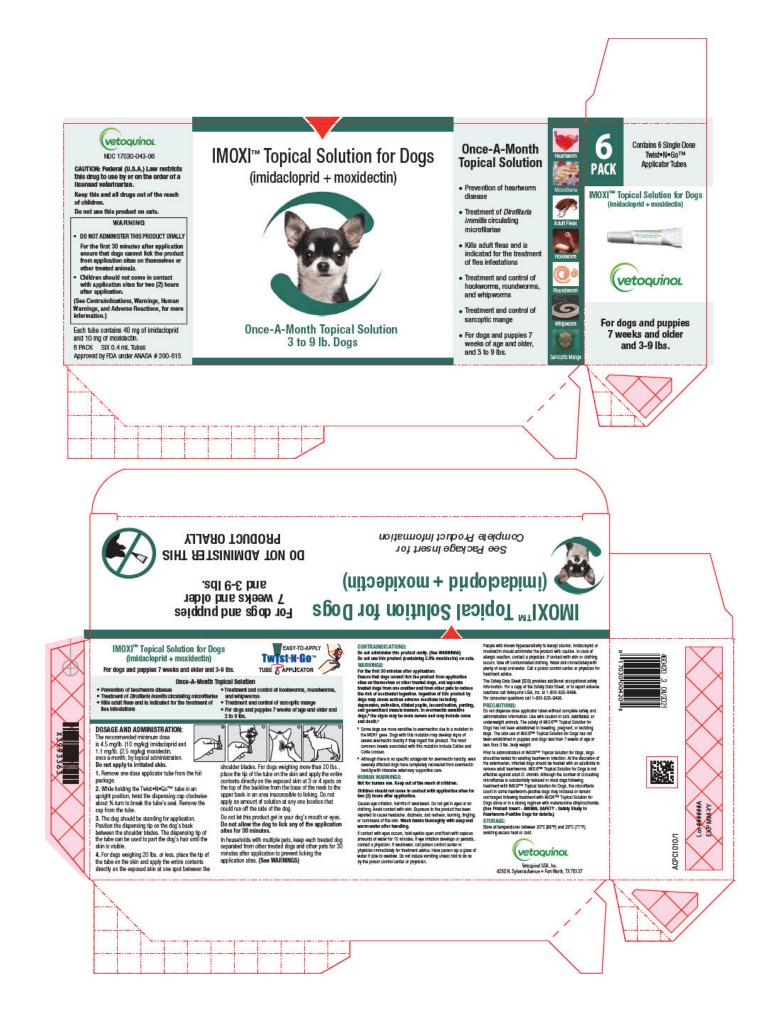
Vetoquinol USA, Inc. 4250 N. Sylvania Avenue Fort Worth, TX 76137

R1 - 05/2020

# **PRINCIPAL DISPLAY PANEL - 0.4 mL Tube Blister Pack Carton**

IMOXI<sup>™</sup> Topical Solution for Dogs (imidacloprid + moxidectin)

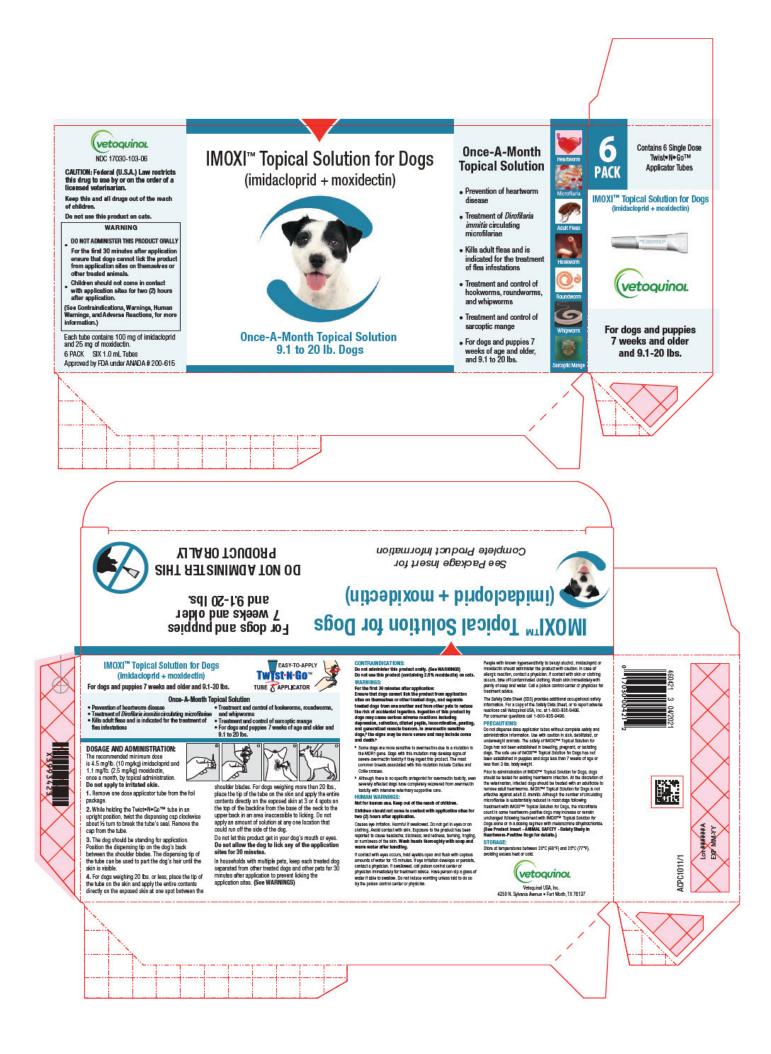
Once-A-Month Topical Solution 3 to 9 lb. Dogs



# **PRINCIPAL DISPLAY PANEL - 1.0 mL Tube Blister Pack Carton**

IMOXI<sup>™</sup> Topical Solution for Dogs (imidacloprid + moxidectin)

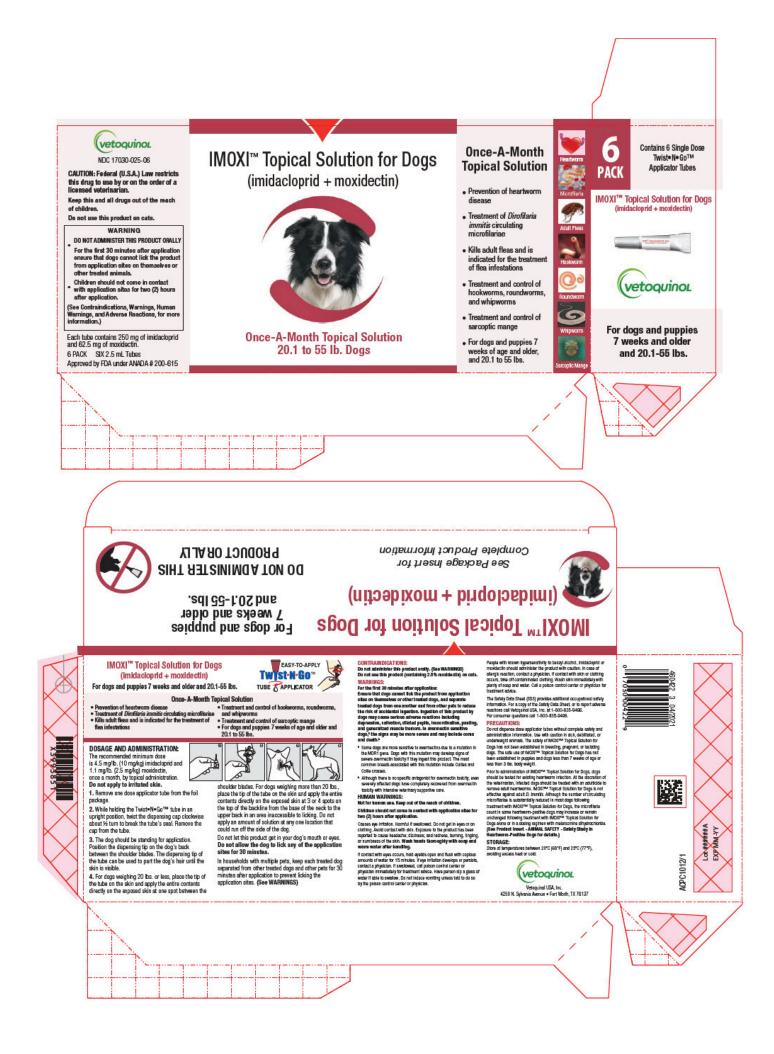
Once-A-Month Topical Solution 9.1 to 20 lb. Dogs



# PRINCIPAL DISPLAY PANEL - 2.5 mL Tube Blister Pack Carton

IMOXI<sup>™</sup> Topical Solution for Dogs (imidacloprid + moxidectin)

Once-A-Month Topical Solution 20.1 to 55 lb. Dogs



# PRINCIPAL DISPLAY PANEL - 4.0 mL Tube Blister Pack Carton

IMOXI<sup>™</sup> Topical Solution for Dogs (imidacloprid + moxidectin)

Once-A-Month Topical Solution 55.1 to 88 lb. Dogs



# **PRINCIPAL DISPLAY PANEL - 5.0 mL Tube Blister Pack Carton**

IMOXI<sup>™</sup> Topical Solution for Dogs (imidacloprid + moxidectin)

Once-A-Month Topical Solution 88.1 to 110 lb. Dogs



	000						
IMOXI FOR D		- hatie -					
midacloprid and mo	sidectin s	olution					
Product Informa	stion						
	ation						
Product Type		PRESCRIPTION ANIM	al drug	ltem	Code (Sour	ce)	NDC:17030-043
Route of Administr	ation	TOPICAL					
Active Ingredien	t/Active	Moietv					
jj		lient Name			Basis of St	trenath	Strength
IMIDACLOPRID (UNII: 3	-		III:3BN7M937V	(8)	IMIDACLOPRID	-	100 mg in 1 mL
MOXIDECTIN (UNII: NO	GU5H31YO9)	(MOXIDECTIN - UNII:N	IGU5H31YO9)		MOXIDECTIN		25 mg in 1 mL
	-						
Inactive Ingredie							
		gredient Name				S	trength
<b>GRAPEFRUIT</b> (UNII: 08	2C39RR8C)						
Packaging							
# Item Code	Packa	ge Description	Marketin	g Sta	art Date	Market	ing End Date
<b>1</b> NDC:17030-043-06	1 in 1 CAF						
<b>1</b> NDC:17030-043-01		STER PACK					
1	0.4 mL in	IIUBE					
Marketing In	format	ion					
Marketing	Applica	tion Number or M	onograph	м	arketing Sta	art N	larketing End
Category		Citation		12/0	Date		Date
ANADA	ANADA2006	15		12/0	7/2020		
	NUCS						
midacloprid and mo		olution					
	Maccun 5						
Product Informa	ation						
Product Type		PRESCRIPTION ANIM	AL DRUG	ltem	Code (Sour	ce)	NDC:17030-103
					5040 (50di	,	

Active Ingredient/Active Moiety		
Ingredient Name	<b>Basis of Strength</b>	Strength
IMIDACLOPRID (UNII: 3BN7M937V8) (IMIDACLOPRID - UNII:3BN7M937V8)	IMIDACLOPRID	100 mg in 1 mL
MOXIDECTIN (UNII: NGU5H31YO9) (MOXIDECTIN - UNII:NGU5H31YO9)	MOXIDECTIN	25 mg in 1 mL

TOPICAL

Route of Administration

Inactive Ingr	edients					
		Ingredient Name				Strength
GRAPEFRUIT (UN	I: 082C39RR80	C)				
Packaging						
# Item Cod	e Pack	kage Description	Marketing	Start Date	Marke	eting End Date
<b>1</b> NDC:17030-103	-06 1 in 1 C	CARTON				
<b>1</b> NDC:17030-103	-01 6 in 1 B	BLISTER PACK				
1	1.0 mL	in 1 TUBE				
Marketing	Informa	ation				
<b>_</b>				No. of Latin 2		
Marketing Marketing Category		ation cation Number or M Citation	onograph	Marketing S Date	tart	Marketing End Date
Marketing Category		cation Number or M Citation	onograph		tart	
Marketing Category	Applic	cation Number or M Citation	onograph	Date	tart	
Marketing Category	Applic	cation Number or M Citation	onograph	Date	tart	
<b>Category</b> ANADA	Applic	cation Number or M Citation	onograph	Date	tart	
Marketing Category ANADA	Applid ANADA20	cation Number or M Citation 0615	onograph	Date	tart	
Marketing Category ANADA MOXI FO	Applid ANADA20	cation Number or M Citation 0615	onograph	Date	tart	
Marketing Category ANADA MOXI FO midacloprid and	Applid ANADA200 R DOGS	cation Number or M Citation 0615	onograph	Date	tart	
Marketing Category ANADA	Applid ANADA200 R DOGS	cation Number or M Citation 0615		Date		

Active Ingredient/Active Moiety		
Ingredient Name	<b>Basis of Strength</b>	Strength
IMIDACLOPRID (UNII: 3BN7M937V8) (IMIDACLOPRID - UNII:3BN7M937V8)	IMIDACLOPRID	100 mg in 1 mL
MOXIDECTIN (UNII: NGU5H31YO9) (MOXIDECTIN - UNII:NGU5H31YO9)	MOXIDECTIN	25 mg in 1 mL

Ir	nactive Ingredie	ents		
		Ingredient Name		Strength
GI	RAPEFRUIT (UNII: 08	2C39RR8C)		
P	ackaging			
#	ltem Code	<b>Package Description</b>	Marketing Start Date	Marketing End Date
1	NDC:17030-025-06	1 in 1 CARTON		
1	NDC:17030-025-01	6 in 1 BLISTER PACK		
1		2.5 mL in 1 TUBE		

Marketing Category	Applica	tion Number or M Citation	onograph	M	arketing S Date	tart	Marketin Dat	-
NADA	ANADA2006	15		12/0	7/2020			
MOXI FOR	DOCS							
midacloprid and n		olution						
Product Inforn	nation							
Product Type		PRESCRIPTION ANIM	AL DRUG	ltem	Code (So	urce)	NDC:170	30-400
Route of Adminis	tration	TOPICAL						
Active Ingredie	nt/Active	Moiety						
Active migreate		-			Basis of	Strongt	.h. C+ra	ength
	Ingred	lient Name			Dasis Ul	Scienge	.n stre	FIIGUI
MIDACLOPRID (UNI	-	l <b>ient Name</b> 8) (IMIDACLOPRID - UN	III:3BN7M937V8	3)	IMIDACLOPR	-		in 1 m
MOXIDECTIN (UNII: I	: 3BN7M937V NGU5H31YO9)			3)		ID		in 1 m
MOXIDECTIN (UNII: I	: 3BN7M937V3 NGU5H31YO9) <b>lients</b>	3) (IMIDACLOPRID - UN		3)	IMIDACLOPR	ID I	100 mg	in 1 m in 1 mL
MOXIDECTIN (UNII: I	: 38N7M937V4 NGU5H31YO9) <b>lients</b> In	3) (IMIDACLOPRID - UN (MOXIDECTIN - UNII:N		3)	IMIDACLOPR	ID I	100 mg 25 mg	in 1 m in 1 mL
MOXIDECTIN (UNII: I Inactive Ingred GRAPEFRUIT (UNII: (	: 38N7M937V4 NGU5H31YO9) <b>lients</b> In	3) (IMIDACLOPRID - UN (MOXIDECTIN - UNII:N		3)	IMIDACLOPR	ID I	100 mg 25 mg	in 1 m in 1 mL
MOXIDECTIN (UNII: I Inactive Ingred GRAPEFRUIT (UNII: ( Packaging	: 3BN7M937V NGU5H31YO9) <b>lients</b> In D82C39RR8C)	B) (IMIDACLOPRID - UN (MOXIDECTIN - UNII:N gredient Name	NGU5H31YO9)		IMIDACLOPR MOXIDECTIN	ID I	100 mg 25 mg Strength	in 1 m in 1 mL
MOXIDECTIN (UNII: I Inactive Ingred GRAPEFRUIT (UNII: ( Packaging # Item Code	: 3BN7M937V NGU5H31YO9) lients In D82C39RR8C) Packa	B) (IMIDACLOPRID - UN (MOXIDECTIN - UNII:N gredient Name ge Description			IMIDACLOPR MOXIDECTIN	ID I	100 mg 25 mg	in 1 m in 1 mL
MOXIDECTIN (UNII: I Inactive Ingred GRAPEFRUIT (UNII: ( Packaging # Item Code 1 NDC:17030-400-06	: 3BN7M937V NGU5H31YO9) lients In D82C39RR8C) Packa 5 1 in 1 CAF	B) (IMIDACLOPRID - UN (MOXIDECTIN - UNII:N gredient Name ge Description	NGU5H31YO9)		IMIDACLOPR MOXIDECTIN	ID I	100 mg 25 mg Strength	in 1 m in 1 mL
MOXIDECTIN (UNII: I Inactive Ingred GRAPEFRUIT (UNII: C Packaging # Item Code 1 NDC:17030-400-01 1 NDC:17030-400-01	: 3BN7M937V NGU5H31YO9) lients In D82C39RR8C) Packa 5 1 in 1 CAF	B) (IMIDACLOPRID - UN (MOXIDECTIN - UNII:N gredient Name ge Description RTON STER PACK	NGU5H31YO9)		IMIDACLOPR MOXIDECTIN	ID I	100 mg 25 mg Strength	in 1 m in 1 mL
MOXIDECTIN (UNII: I Inactive Ingred GRAPEFRUIT (UNII: C Packaging # Item Code 1 NDC:17030-400-00 1 NDC:17030-400-01	: 3BN7M937V3 NGU5H31YO9) Ilients In D82C39RR8C) Packa 5 1 in 1 CAF 6 in 1 BL!	B) (IMIDACLOPRID - UN (MOXIDECTIN - UNII:N gredient Name ge Description RTON STER PACK	NGU5H31YO9)		IMIDACLOPR MOXIDECTIN	ID I	100 mg 25 mg Strength	in 1 m in 1 mL
MOXIDECTIN (UNII: I Inactive Ingred GRAPEFRUIT (UNII: C Packaging # Item Code 1 NDC:17030-400-00 1 NDC:17030-400-00 1	: 3BN7M937V3 NGU5H31YO9) lients In D82C39RR8C) 5 1 in 1 CAF 6 in 1 BL1 4.0 mL in	B) (IMIDACLOPRID - UN (MOXIDECTIN - UNII:N agredient Name ge Description RTON STER PACK 1 TUBE	NGU5H31YO9)		IMIDACLOPR MOXIDECTIN	ID I	100 mg 25 mg Strength	in 1 m in 1 mL
MOXIDECTIN (UNII: I Inactive Ingred GRAPEFRUIT (UNII: ( Packaging	: 3BN7M937V NGU5H31YO9) lients In D82C39RR8C) 282C39RR8C) 4.0 mL in 4.0 mL in	B) (IMIDACLOPRID - UN (MOXIDECTIN - UNII:N agredient Name ge Description RTON STER PACK 1 TUBE	Marketing	g Sta	IMIDACLOPR MOXIDECTIN	ID I Marke	100 mg 25 mg Strength	in 1 m in 1 mL

# IMOXI FOR DOGS inidacloprid and moxidectin solution Product Information Product Type PRESCRIPTION ANIMAL DRUG Route of Administration TOPICAL

4	ctive Ingredien	t/Active Moiety				
		Ingredient Name		Basis of	Strengt	h Strength
IM	IDACLOPRID (UNII: 3	3BN7M937V8) (IMIDACLOPRID - UN	NII:3BN7M937V8)	IMIDACLOPF	RID	100 mg in 1 m
M	OXIDECTIN (UNII: NO	GU5H31YO9) (MOXIDECTIN - UNII:M	NGU5H31YO9)	MOXIDECTIN	N	25 mg in 1 mL
In	active Ingredie	ents				
		Ingredient Name			9	Strength
GF	RAPEFRUIT (UNII: 08	•				
Pa	ackaging					
	ackaging Item Code	Package Description	Marketing	Start Date	Marke	ting End Date
#		Package Description 1 in 1 CARTON	Marketing	Start Date	Marke	ting End Date
# 1	ltem Code		Marketing	Start Date	Marke	ting End Date
# 1 1	Item Code NDC:17030-500-06	1 in 1 CARTON	Marketing	Start Date	Marke	ting End Date
# 1	Item Code NDC:17030-500-06	1 in 1 CARTON 6 in 1 BLISTER PACK	Marketing	Start Date	Marke	ting End Date
# 1 1	Item Code NDC:17030-500-06	1 in 1 CARTON 6 in 1 BLISTER PACK 5.0 mL in 1 TUBE	Marketing	Start Date	Marke	ting End Date
# 1 1	Item Code NDC:17030-500-06 NDC:17030-500-01	1 in 1 CARTON 6 in 1 BLISTER PACK 5.0 mL in 1 TUBE		Start Date		ting End Date Marketing End Date

# Labeler - Vetoquinol USA, Inc. (106824209)

Revised: 12/2022

Vetoquinol USA, Inc.