



KVP Pharma+Veterinär Produkte GmbH Projensdorfer Straße 324 24106 Kiel, Germany		Certificate of Analysis		Page: 1 of 3 Date: 2016-04-14	
Material: 84051365 Your material:		ADVOCATE SO DOGS 3 X 4,0 ML BOX X 3 PIP X 4,0 ML ADVOCATE SO DOGS 3 X 4,0 ML			
Batch: KP0B5RK Date of manufacture: 2016-02-01 Expiry date: 2019-02-28		Country: Hungary Delivery number: 107794043 Order number: 5996519			
From material: 81245401 Batch: KP0B1DU Inspection lot: 040001554289		IMI10%/MOXI 2,5% H 4,0 ML 81094160		Insp. instruction: T.02.02 - 7 Specification: T.02.28 - 5	
Inspection	Acceptance criterion	UoM	Result		
Material (visual)	Solution		solution		
Clarity (visual)	clear		clear		
Colour (Ph.Eur.)	yellow to brownish-yellow		yellow		
Identity (HPLC)	must comply		complies		
Identity (TLC)	must comply		complies		
Assay Imidacloprid	9.5 - 10.5	g/hml	10.1		
Any unspecified degrad.prod. Imidacloprid	max. 1.0	%	0.5		
Sum of all degrad.products imidacloprid	max. 1.0	%	0.5		
Assay Moxidectin	2.38 - 2.62	g/hml	2.53		
23-Keto-F-Alpha compound	max. 0.3	%	< 0.1		
Any unspecified degrad.prod. Moxidectin	max. 1.0	%	0.6		
Sum of all degrad.products moxidectin	max. 2.0	%	0.6		
Relative density (USP)	1.093 - 1.103		1.099		

Diszpó: 5928021867 Vev: ALPHA-VET File név: BA00001873 Egyedi sorszám: BA000002033 12 másolat
 Szállító: Bayer Állat gyógyászati Üzletág Oldal/Lap 3 / 1 Anyaglap: 500176592
 Az eredetivel megegyez elektronikus másolat a(z):
 Dr. Gacsályi Panna min.bizt.gyógyszerész által küldve. Nyomtatás dátuma: 2016.06.02 CertEx v 3.03a



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Inspection	Acceptance criterion	UoM	Result
Butylhydroxytoluene	0.090 - 0.125	g/hml	0.110
Water	max. 2.0	%	< 0,3
Uniformity of content	must comply		complies
Total aerobic microbial count (TAMC)	max. 100	CFU/g	*)
Total combined yeast/mould count (TYMC)	max. 10	CFU/g	*)
Microb.purity Staphylococcus aureus/1g	absent		*)
Microbial purity Pseudomonas aerug./1g	absent		*)

*) Test is carried out on spot-check basis. However we confirm compliance with the inspection and requirements also for this batch.

This batch complies with the specification and is manufactured in accordance with EC Guide to GMP for Medicinal Products and the requirement laid down in the product license.

Release Documentation is signed by responsible Person Quality Management.

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Dr. Gacályi Panna min.bízt.gyógyszerész által küldve. Nyomtatás dátuma: 2016.06.02 CertEx v 3.03a



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Batch release electronically signed:

Dr. Carsten Bode (KPCBO)

Qualified Person

Date/time:

2016-04-13 08:30:47 p.m. CET (UTC + 1 hour)

Inspection lot:

040001577606

This Certificate of Analysis was automatically printed.

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