

Bayer HealthCare



KVP Pharma+Veterinär Produkte GmbH Kiel, Germany	Certificate of Analysis		Page: 1 of 3 Date: 2015-08-21
Material: 84051225 Your material:	ADVOCATE SO DOGS 3 X 1,0 ML BOX X 3 PIP X 1,0 ML ADVOCATE SO DOGS 3 X 1,0 ML		
Batch: <u>KP0AJGD-</u> Date of manufacture: 2015-06-08 Expiry date: 2018-06-30	Country: Hungary Delivery number: 105724122 Order number: 4891531		
From material: 81245266 Batch: KP0AFX2 Inspection lot: 040001462779	IMI10%/MOXI 2,5 % H 1,0 ML 81094136	Insp. instruction: T.02.02 - 6 Specification: T.02.28 - 4	
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	10.2
Any unspecified degrad.prod. Imidacloprid	max. 1.0	%	0.6
Sum of all degrad.products imidacloprid	max. 1.0	%	0.6
Assay Moxidectin	2.38 - 2.62	g/hml	2.52
23-Keto-F-Alpha compound	max. 0.3	%	< 0.1
Any unspecified degrad.prod. Moxidectin	max. 1.0	%	0.4
Sum of all degrad.products moxidectin	max. 2.0	%	0.4
Relative density (USP)	1.093 - 1.103		1.099

Diszpó: 5928018014 Vev: ALPHA-VET KFT. File név: BA00001724 Egyedi sorszám: BA000001820 7 másolat
 Szállító: Bayer Állat gyógyászati Üzletág Oldal/Lap 3 / 1 Anyaglap: 500151947
 Az eredetivel megegyez elektronikus másolat a(z):
 Dr. Gacsályi Panna min.bízt.gyógyszerész által küldve. Nyomtatás dátuma: 2015.09.10 **CertEx v 3.03a**



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Date of manufacture:	2015-06-08	Delivery number: 105724122		
Expiry date:	2018-06-30	Order number: 4891531		
From material:	81245266	IMI10%/MOXI 2.5 % H 1,0 ML	81094136	
Batch:	KP0AFX2			Insp. instruction: T.02.02 - 6
Inspection lot:	040001462779			Specification: T.02.28 - 4
Inspection	Acceptance criterion	UoM	Result	
Butylhydroxytoluene	0.090 - 0.126	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 Mrs. Cordula Weber, Dr. Bode or Dr. Amann

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Batch release electronically signed: Dr. Tobias Amann (KPAM)
 Qualified Person
 Date/time: 2015-08-21 12:26:53 p.m. CET (UTC + 1 hour)
 Inspection lot: 040001479620

This Certificate of Analysis was automatically printed.

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