



KVP Pharma+Veterinär Produkte GmbH Kiel, Germany	Certificate of Analysis		Page: 1 of 2 Date: 2013-06-06
Material: 80993316 Your Material:	ADVOCATE S.O. FOR SMALL DOGS <4KG3X0,4ML BOX X 3 PIP X 0,4 CC ADVOCATE S.O. FOR SMALL DOGS < 4 KG		
Batch: KP08R1K Date of manufacture: 2013-03-25 Expiry date: 2016-03-31	Country: Hungary Delivery number: 100895856 Order number: 1323071		
From Material: 83196700 Batch: KP08KPS Inspection lot: 040001102176	IMI10%/MOXI 2,5 % H 0,4 ML 80142331	Insp. instruction: T.02.02 - 6 Specification: T.02.01 - 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	10.0
Any unspecified degrad. prod. Imidacloprid	max. 1.0	%	0.3
Sum of all degrad. products imidacloprid	max. 1.0	%	0.3
Assay Moxidectin	2.25 - 2.62	g/hml	2.49
23-Keto-F-Alpha compound	max. 2.0	%	< 0.1
Any unspecified degrad. prod. Moxidectin	max. 1.0	%	0.4
Sum of all degrad. products moxidectin	max. 4.0	%	0.4
Relative density (USP)	1.093 - 1.103		1.098
Butylhydroxytoluene	0.050 - 0.125	g/hml	0.110
Water	max. 6.0	%	< 0.3



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Material: 80993315 Your Material:		ADVOCATE S.O. FOR SMALL DOGS <4KG3X0,4ML BOX X 3 PIP X 0,4 CC ADVOCATE S.O. FOR SMALL DOGS < 4 KG		
Batch:	KP08R1K	Country: Hungary		
Date of manufacture:	2013-03-25	Delivery number: 100695856		
Expiry date:	2016-03-31	Order number: 1323071		
From Material:	83195700	IMI10%/MOXI 2,5 % H 0,4 ML	80142331	
Batch:	KP08KP9	Insp. Instruction: T.02.02 - 6		
Inspection lot:	040001102176	Specification: T.02.01 - 4		
Inspection	Acceptance criterion	UoM.	Result	
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 - Dr. Amann - QP

This Certificate of Analysis was automatically printed.