



KVP Pharma+Veterinär Produkte GmbH Kiel, Germany	Certificate of Analysis		Page: 1 of 2 Date: 2014-05-19
Material: 81594783 Your material:	ADVOCATE SO CAT 3 X 0,8 ML BOX X 3 PIP X 0,8 ML ADVOCATE SO CAT 3 X 0,8 ML		
Batch: KP09J0N Date of manufacture: 2013-12-02 Expiry date: 2016-12-31	Country: Hungary Delivery number: 102399751 Order number: 2467094		
From material: 81591067 Batch: KP09BKD Inspection lot: 040001228819	IMI 10%/MOXI 1% K 0,8 ML 81266107		Insp. instruction: T.02.02 - 7 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	10.2
Any unspecified degrad.prod.Imidacloprid	max. 1.0	%	0.3
Sum of all degrad.products imidacloprid	max. 1.0	%	0.3
Assay Moxidectin	0.90 - 1.05	g/hml	1.01
23-Keto-F-Alpha compound	max. 2.0	%	< 0.1
Any unspecified degrad.prod.Moxidectin	max. 1.0	%	0.5
Sum of all degrad.products moxidectin	max. 4.0	%	1.0
Relative density (USP)	1.093 - 1.103		1.097



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Inspection	Acceptance criterion	UoM	Result
Butylhydroxytoluene	0.050 - 0.116	g/hml	0.103
Water	max. 6.0	%	< 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 - Dr. Amann -QP