Bayer HealthCare

KVP Pharma+Veterinär Produkte GmbH Kiel, Germany	**Certificate of Analysis**	Page: 1 of 2 Date: 2013-11-15

Material: 82258354 Your Material:	ADVOCATE S.O. FOR SMALL CATS <4KG3X0,4ML BOX X 3 PIP X 0,4 CC ADVOCATE S.O. FOR SMALL CATS < 4 KG

Batch:	KP0945H	Country: Hungary
Date of manufacture:	2013-08-05	Delivery number: 101391366
Expiry date:	2016-08-31	Order number: 1880620

From Material:	83196816	IMI 10%/MOXI 1% K 0,4 ML 80142277	
Batch:	KP090XZ	Insp. instruction:	T.02.02 - 7
Inspection lot:	040001185989	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	0.90 - 1.05	g/hml	0.99
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		1.097
Butylhydroxytoluene	0.050 - 0.116	g/hml	0.103
Water	max. 6.0	%	< 0.3

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Material: 82256354 Your Material:	ADVOCATE S.O. FOR SMALL CATS <4KG3X0,4ML BOX X 3 PIP X 0,4 CC ADVOCATE S.O. FOR SMALL CATS < 4 KG

Batch: Date of manufacture: Expiry date:	KP0945H 2013-08-05 2016-08-31	Country: Hungary Delivery number: 101391366 Order number: 1880620

From Material: Batch: Inspection lot:	83196816 IMI 10%/MOXI 1% K 0,4 ML 80142277 KP090XZ 040001185989	Insp. instruction: Specification:	T.02.02 - 7 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.