



KVP Pharma (Veterinär Produkte GmbH Projensdorfer Straße 324 24106 Kiel, Germany		Certificate of Analysis	Page: 1 of 2 Date: 2016-04-19
Material: 85071890 Your material:		ADVANTIX DOG >25 KG 6X4X4.0 ML ADVANTIX DOG >25 KG 6X4X4.0 ML	
Batch: KP0B4SZ Date of manufacture: 2016-01-12 Expiry date: 2021-01-31	Country: Hungary Delivery number: 107835169 Order number: 5525716		
From material: 81858748 Batch: KP0B2BJ Inspection lot: 040001559672	IMI10%/PERM50% EU PIPX4.0 ML 81322066		Insp. instruction: T.02.02 - 2 Specification: T.02.28 - 4

Inspection	Acceptance criterion	UoM	Result
Clarity (visual)	clear solution		clear solution
Colour (visual)	yellowish to brownish		slightly brownish yellow
Identity Imidacloprid (HPLC)	must comply		complies
Identity Imidacloprid (TLC)	must comply		complies
Identity Permethrin (HPLC)	must comply		complies
Identity Permethrin (TLC)	must comply		complies
Density	1.123 - 1.150	g/ml	1.136
Water	max. 1.0	g/100g	< 0.3
pH-value	3.0 - 5.0		3.5
DCVC acid	max. 1.2	%	0.4
Any unspecified degrad. product, largest	max. 1.0	%	0.2
Sum of all degradation products	max. 3.2	%	0.9
Assay Imidacloprid	9.5 - 10.5	g/hml	9.9
Assay Permethrin	47.5 - 52.5	g/hml	50.0

Diszpó: 5928021983 Vev: ALPHA-VET File név: BA00001878 Egyedi sorszám: BA000002042 13 másolat
 Szállító: Bayer Állat gyógyászati Üzletág Oldal/Lap 2 / 1 Anyaglap: 500177461
 Az eredetivel megegyez elektronikus másolat a(z):
 Dr. Gacsályi Panna min.bízt.gyógyszerész által küldve. Nyomtatás dátuma: 2016.06.09 CertEx v 3.03a



KVP Pharma+Veterinär Produkte GmbH Progersdorfer Straße 324 24105 Kiel, Germany		Certificate of Analysis		Page: 2 of 2 Date: 2016-04-19
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Batch: KPOB4SZ Date of manufacture: 2016-01-12 Expiry date: 2021-01-31	Country: Hungary Delivery number: 107835169 Order number: 6525716			
From material: 81858748 Batch: KPOB2BJ Inspection lot: 040001559672	IMI: 10%/PERM50% EU PIPX4,0 ML 81322066		Insp. instruction: T.02.02 - 2 Specification: T.02.28 - 4	
Inspection:	Acceptance criterion	UoM	Result	
Assay butylhydroxytoluene	0.09 - 0.11	g/ml	0.10	
Uniformity of content:	must comply		complies	

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.

Batch release electronically signed:

Dr. Tobias Amann (KPAM)
Qualified Person
2016-04-18 06:02:08 p.m. CET (UTC + 1 hour)
040001578040

Date/time:
Inspection lot:

This Certificate of Analysis was automatically printed.

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