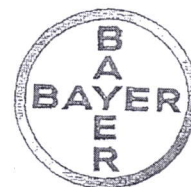




KVP Pharma+Veterinär Produkte GmbH Projensdorfer Straße 324 24106 Kiel, Germany		Certificate of Analysis		Page: 1 of 2 Date: 2018-07-19
Material: 85662805 Your material:		BAYCOX 2,5% SOLUTION 1000 ML BAYCOX 2,5% SOLUTION 1000 ML		
Batch:	KP0D83T	Country: Hungary		
Date of manufacture:	2018-06-15	Delivery number: 115293570		
Expiry date:	2023-06-30	Order number: 703201927		
From material:	05352304	BAYCOX LOESUNG 2,5% G/V		
Batch:	KP0D6DV	Insp. instruction:	T.02.02 - 10	
Inspection lot:	040001907852	Specification:	T.02.28 - 8	
Inspection	Acceptance criterion	UoM	Result	
Material (visual)	Solution		solution	
Colour (visual)	colourless to brown		colorless	
Clarity (visual)	clear		clear	
Identity Toltrazuril (retention time)	must comply		complies	
Identity Toltrazuril (UV absorption)	must comply		complies	
Relative density	1.120 - 1.140		1.133	
pH-value	8.0 - 10.0		9.1	
Appearance of solution colour	max. Y7		less intensely coloured than Y7	
Water	max. 0.5	%	0.3	
Hydrol.prod.calc.on act.subst.(CZ I/II)	max. 0.5	%	< 0.1	
Hydrol.prod.calc.on act.subst(CZ III/IV)	max. 0.5	%	< 0.1	
Sum of all impurities	max. 1.0	%	< 0.3	
Assay	2.37 - 2.63	g/hml	2.49	



KVP Pharma+Veterinär Produkte GmbH Projensdorfer Straße 324 24106 Kiel, Germany	Certificate of Analysis	Page: 2 of 2 Date: 2018-07-19	
Material: 85662805 Your material: *	BAYCOX 2,5% SOLUTION 1000 ML BAYCOX 2,5% SOLUTION 1000 ML		
Batch: KP0D83T Date of manufacture: 2018-06-15 Expiry date: 2023-06-30	Country: Hungary Delivery number: 115293570 Order number: 703201927		
From material: 05352304			
Inspection lot: 040001907852	Insp. instruction: T.02.02 - 10 Specification: T.02.28 - 8		
Inspection	Acceptance criterion	UoM	Result
Total aerobic microbial count (TAMC)	max. 1000	CFU/g	*)
Total combined yeast/mould count (TYMC)	max. 100	CFU/g	*)
Microb.purity Escherichia coli/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.

Batch release electronically signed:

Dr. Carsten Bode (KPCBO)
Qualified Person

Date/time:
Inspection lot:

2018-07-18 05:54:11 p.m. CET (UTC + 1 hour)
040001912151

This Certificate of Analysis was automatically printed.