Bayer HealthCare

KVP Pharma+Veterinär Produkte GmbH Kiel, Germany	**Certificate of Analysis**	Page: 1 of 2 Date: 2013-06-10

Material: 81899909 Your Material:	BAYCOX BOVIS 5 % 1000 ML PLASTIC BOTTLE X 1000 CC BAYCOX BOVIS VET 5 % 1000 ML

Batch:	KP08R0V	Country: Hungary	
Date of manufacture:	2013-03-18	Delivery number: 100708944	
Expiry date:	2018-03-31	Order number: 1493163	
From Material:	05352290 BAYCOX SUSPENSION 5% G/V N		
Batch:	KP08LH1	Insp. instruction:	T.02.02 - 7
Inspection lot:	040001109135	Specification:	T.02.01 - 11

Inspection	Acceptance criterion	UoM.	Result
Material (visual)	suspension		suspension
Colour (visual)	white to yellowish		white
Identity	must comply		complies
Relative density	1.020 - 1.040		1.032
pH-value	4.0 - 5.0		4.3
Particle size D90	<10 µm		<10
Particle size D99	<30 µm		<30
Sodium benzoate	0.18 - 0.23	g/hml	0.21
Sodium propionate	0.18 - 0.23	g/hml	0.21
Any unspecified impurity	max. 0.5	%	< 0.3
Sum of all impurities	max. 1.0	%	< 0.3
Viscosity	20 - 100	mPa.s	44
Assay	4.75 - 5.25	g/hml	4.95
Total aerobic microbial count (TAMC)	max. 100	CFU/g	*)
Total combined yeast/mould count (TYMC)	max. 10	CFU/g	*)
Escherichia coli	Absence in 1 g		*)

Bayer HealthCare

KVP Pharma+Veterinär Produkte GmbH Kiel, Germany	**Certificate of Analysis**	Page: 2 of 2 Date: 2013-06-10
Material: 81899909 Your Material:	BAYCOX BOVIS 5 % 1000 ML PLASTIC BOTTLE X 1000 CC BAYCOX BOVIS VET 5 % 1000 ML	
Batch: KP08R0V Date of manufacture: 2013-03-18 Expiry date: 2018-03-31		Country: Hungary Delivery number: 100708944 Order number: 1493163
From Material: 05352290 BAYCOX SUSPENSION 5% G/V N Batch: KP08LH1 Inspection lot: 040001109135		Insp. instruction: T.02.02 - 7 Specification: T.02.01 - 11

Inspection	Acceptance criterion	UoM.	Result

*) 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.