



KVP Pharma+Veterinär-Produkte GmbH Projensdorfer Straße 324 24106 Kiel, Germany		Certificate of Analysis	Page: 1 of 2 Date: 2016-04-20
Material: 83934379 Your material:		CATOSAL 10% 100ML GLASS BOTTLE X 100 ML CATOSAL 10% 100ML	
Batch: KP0B5ZD Date of manufacture: 2016-01-20 Expiry date: 2021-01-31	Country: Hungary Delivery number: 107848586 Order number: 6134716		
From material: 80816189 Batch: AH01CHA Inspection lot: 010002751021	CATOSAL C 100ML PRI PKG KVP (ELBEUF)		Insp. instruction: T.02.02 - 2 Specification: T.02.28 - 3

Inspection	Acceptance criterion	UoM	Result
Formulation	Solution		solution
Colour (visual)	pink-coloured		pink
Clarity	clear		clear
Identity (Butaphosphane)	must comply		complies
Identity vitamin B 12	must comply		complies
Identity (n-Butanol)	must comply		complies
Extractable volume	min. 100.0	ml	102.7
Purity (visual)	must comply		complies
Density	1.007 - 1.013	g/ml	1.011
pH-value	5.0 - 6.0		5.5
n-Butanol	2.7 - 3.3	g/hml	3.0
Assay butaphosphane	9.5 - 10.5	g/hml	10.0
Assay vitamin B 12	5.2 - 5.8	mg/hml	5.2
Oxa-Butaphosphane	max. 0.5	%	0.1
Any unspecified degradation product	max. 1.0	%	< 0.3

Diszpó: 5928022108 Vev: ALPHA-VET File név: BA00001880 Egyedi sorszám: BA000002050 6 másolat  
 Szállító: Bayer Állat gyógyászati Üzletág Oldal/Lap 2 / 1 Anyaglap: 500177465  
 Az eredetivel megegyez elektronikus másolat a(z):  
 Dr. Gacsályi Panna min.bizt.gyógyszerész által küldve. Nyomtatás dátuma: 2016.06.20 CertEx v 3.03a



KVP Pharma+Veterinär Produkte GmbH Projensdorfer Straße 324 24106 Kiel, Germany		Certificate of Analysis		Page: 2 of 2 Date: 2016-04-20	
Material: 83934379 Your material:		CATOSAL 10% 100ML GLASS BOTTLE X 100 ML CATOSAL 10% 100ML			
Batch: KP0B52D Date of manufacture: 2016-01-20 Expiry date: 2021-01-31		Country: Hungary Delivery number: 107848586 Order number: 6134716			
From material: 80816189 Batch: AH01CHA Inspection lot: 010002751021		CATOSAL C 100ML PRI PKG KVP (ELBEUF) Insp. instruction: T.02.02 - 2 Specification: T.02.28 - 3			
Inspection	Acceptance criterion	UoM	Result		
Sum of all degradation products	max. 1.2	%	0,3		
Sterility	sterile		sterile		
Endotoxins	<89.29 E.U./ml		<3.84		

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

2016-04-19 08:44:41 p.m. CET (UTC + 1 hour)

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