

## BATCH CERTIFICATE / CERTIFICATE OF ANALYSIS RF-781.5

Dopharma Holding B.V. Zalmweg 24 4941 VX Raamsdonksveer The Netherlands Tel: + 31 (0) 162 58 20 00

Productname:

PENSTREP-JECT 100 ML

Lotno:

17F20-01C4

Production date:

06-2017

Expiry date: Storage conditions: 12-2019 2-8 °C Reg.number(incl. country): RO:150515

Ordemumber:

12473

Remark:

PACKING SLIP: 892872 / CUSTOMER NO: 10182

TEST	RESULT	UNIT	SPEC
CHARACTERS			
Appearance	CONFORM	24	CONFORM
RESUSPENDABILITY			
Resuspendability	CONFORM	is.	CONFORM
PH	1474		15
рН	6.4	<b>%</b>	>=5
			<b>&lt;=</b> 8
RELATIVE DENSITY	5.009		17.20
Relative density	1.1481	8	>=1.134
			<=1.149
VISCOSITY	Land Control Control		
Viscosity	212.63	mPa.s	>=50
			<=150
PARTICLE SIZE	44-	120	
Particles <=10µm	95	.%	>=80
Particles >50µm	0	%	<=0
DENTIFICATION SULPHATES	5250027141		120278129.0340
Identification	CONFORM		CONFORM
DENTIFICATION OF PROC BENZYLPEN	CONFORM	*	CONFORM
dentification of proc.benz.pen. by LC	CONFORM	*	CONFORM
IDENTIFICATION OF DHS Identification of DHS by LC	CONFORM	4	CONFORM
DATE TO THE STATE OF STATE OF THE STATE OF T	CONFORM		CONFORM
IDENTIFICATION OF MOB Identification of MOB by LC	CONFORM		CONFORM
ASSAY-LC	2011 01111		W 307 11 W 1111
Benzy Ipenicillin	111.56	mg/ml	>=107.7
			<=119
Benzy Ipenicillin	198573.24	(U/ml	
St. 1-350460			>=152
Dihy drostreptomy cin	159 63	mg/ml	
			<=168
Dihy drostreptomy cin sulphate	199.89	mg/ml	
Procaine	96.07	mg/ml	>=87.8
			<=107.4
ASSAY-LC			
Methy I parahy droxy benzoate	1.50	mg/ml	>=1.35
			<=1.65
BIOBURDEN		(A = 4 4 5 5 4 )	1274
Total viable bacteria	.0	CFU/ml	c-1
STERILITY	CONFORM	A Resident Services	CONFORM
Sterility	CONFORM	Microbial growth	CONFORM
FILLING VOLUME	100	mi	>=100
Filling volume	TOO	1986	
			<=105



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Dopharma Holding B.V. Zalmweg 24 4941 VX Raamsdonksveer The Netherlands Tel: + 31 (0) 162 58 20 00

Release Date:

14-JUL-2017

Disposition Code:

FULL RELEASE

Released By:

Job Title:

E. van Kuppeveld Q.P.

I hereby certify that the above information is authentic and accurate. This batch of product has been manufactured, including packaging/labelling and quality control in full compliance with the GMP requirements of the local Regulatory Authority and within the specifications in the Marketing Authorisation in the Importing country. The batch processing, packaging and analysis records were reviewed and found to be in compliance with GMP. The Batch certificate/certificate of analysis has been produced by a validated Laboratory Information Management System and therefore bears no handwritten signature. GMP Cert.no.: NL/V 14/0005; GMP+: GMP010348.