CEFAVEN 50MG/ML SZUSZP. INJ. AUV 100 ML
Gysz:171649/306416 Lej:19/11/30
Kisz:1X Me.:Darab
SZÉKELY KFT részére

Medimpex Gy. Zrt.
1158 Budapest
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LABORATORIOS MAYMO S.A

Via Augusta , 302
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Tel.: 932370220
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ANALYSIS CERTIFICATE

Product Name:	CEFAVEN 50 mg/ml	Manufacture date	15/11/2017
Batch Number	171649	Expiry date	30/11/2019

Characteristics	Specifications	Results
Appearance	Suspension	Conforms
Density (25ºC)	0,90 - 1,00 g/ml	0,94 g/ml
Kinetic viscosity (30ºC)	60 - 70 csk	Conforms
Resuspendability	max. 30 s	Conforms
Sedimentation time	min. 120 min	Conforms
Particle size	lower than 10 µm: not be less than 80%	Conforms
Ceftiofur (HCl) identification	Positive	Conforms
Ceftiofur (HCl) Content	47,5 - 52,5 mg/ml	48,5 mg/ml
Impurity RRT 1,8	max. 1,0%	N.D.
Any impurity	max. 1,0%	Conforms
Total impurities	max. 2,0%	1,9%
Sterility	Sterile	Meets Test

Quality Control Department
LABORATORIOS MAYMO, S.A.
This certificate was created electronically and is valid without signature.

MEDIMPEX GYÓGYSZER ZRT.
1158 Bp.,Rákospalotai határút 2.
Pantyiné Dr. Balázsi Ilona
Minőségbiztosítási Vezető
Csak a piros szín hiteles

Meoszám:180600006MBT 5.pld. Kiadva: 18/10/30

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international@maymo.es

BATCH CERTIFICATE

Product name	Cefaven 50 mg/ml szuszpenziós injekció sertések és szarvasmarhák részére		
	Cefaven 50 mg/ml zawiesina do wstrzykiwań, dla świń i bydła		
Pharmaceutical form	Injectable suspension		
M.A. Number	3409/1/13 NÉBIH ÁTI	Country	Hungria
	PL: 2386/14		Poland

Batch Number	171649	Package	100 ml
Manufacture date	November 2017	Expiry date	November 2019

Holder and manufacturer	Laboratorios Maymó, S.A.
Manufacture site	Pol. Ind. Can Pelegrí - Ferro, 9 08755 Castellbisbal (Barcelona) - Spain
Manufacturer number	4280 E
GMP Certificate number	ES / 105HV / 16
Name of the Qualified Persons for Batch Release	Nestor Maestre Eduard Esteller
Certificate of Analysis Specifications & Results	See Certificate of Analysis attached

Certification statement:

I hereby certify that this batch has been manufactured in accordance with GMP requirements in Spain, and the Certificate of Analysis results comply with the product Specifications in accordance with the Marketing Authorisation of this product.

Signature of the Qualified Person:

Name: Eduard Esteller

Date: January 24, 2018

LABORATORIOS MAYMO S.A
Via Augusta, 302
08017 BARCELONA

Meoszám:180600006MBT 5.pld. Kiadva: 18/10/30

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