

CERTIFICATE OF ANALYSIS

Customer

METAPHARMACEUTICAL IND., S.L.

Customer PO

2017 1 / 293 / 05-JUL-2017

Order Number

81700228

Product Name

(6S)-CALCIUM FOLINATE / CALCIUM LEVOFOLINATE EP

Description

Lot

 $\ensuremath{\mathsf{PA/PE}}$ bag in $\ensuremath{\mathsf{PE/Alu}}$ bag, both vacuum sealed, in cardboard box

Product Code

610003-03-02

Manufacturing Date

C00270-B00051 25-APR-2017

Release Date

20 / 11 / 11 20 / /

ricicase Date

20-JUL-2017

Retest Date

15-APR-2019

Storage Conditions

Store at 2-8 °C under vacuum, protected from light.

Tests	Specifications			Unit of	
	Target	Min	Max	Measure	Results
Packaging	Packaging conform		Conform		
Labeling	Label conform		Conform		
Appearance	White to yellowish p		Conform		
IR spectrum	Comparable to refer		Comparable		
Test for calcium	Positive for calcium			Positive	
Specific optical rotation (1.0% in THAM buffer, pH 8.1, 25 °C)		-15	-10	-	-13
10-formylfolic acid			0.8	%	0.11
p-aminobenzoylglutamic acid			0.8	%	< 0.05
5,10-diformyltetrahydrofolic acid			0.8	%	0.06
Folic acid			0.8	%	< 0.05
5-formyltetrahydropteroic acid			0.8	%	< 0.05
10-formyldihydrofolic acid			8.0	%	0.08
7,8-dihydrofolic acid			8.0	%	< 0.05
Any unspecified impurity			0.10	%	0.08
(major) Sum of related substances (excluding 10-formylfolic acid) (6R)-folinic acid (impurity H)			2.0	%	0.26
Residual ethanol			3.0	%	0.4
Residual acetone			0.5	%	< 0.10
рН		7.5	8.5	-	7.9
(0.8% in water) Clarity (0.8% in water)	Clear solution				Clear
Absorbance (0.8% in water, 420 nm)			0.25	Au	0.03
Heavy metals			50	ppm	10
Water content (by Karl Fischer)		10.0	17.0	%	12.5
Chlorides	Not more than 0.5%				Conform

CoA No. 6075159

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Tests	Specifications			Unit of	
	Target	Min	Max	Measure	Results
Total aerobic microbial count			500	cfu/g	0
Escherichia coli	Absent/g				Absent/g
Salmonella spp.	Absent/10 g				Absent/10 g
Staphylococcus aureus	Absent/g				Absent/g
Pseudomonas aeruginosa	Absent/g				Absent/g
Total molds and yeasts			100	cfu/g	0
Bacterial endotoxins (LAL test)			0.500	EU/mg	< 0.100
(6S)-calcium folinate (anhydrous and solvent free substance)		97.0	102.0	%	99.6
Calcium		7.54	8.14	%	7.89

Conformity Statement:

We herewith certify that this lot was manufactured, packaged and analysed by Cerbios-Pharma SA, CH-6917 Barbengo/Lugano, Switzerland, according to cGMP and to the current registration dossier by trained personnel; results were found to comply with the registered specifications.

No quality relevant deviations occurred during the lot manufacturing and quality control.

The manufacturing and analytical records have been reviewed and approved.

Based on the above, this lot is conform and released.

Note: With regard to platinum to be controlled according to the EP monograph for this drug substance, we state that the drug substance is manufactured without using platinum catalyst, and it is not tested for routinely, but that the drug substance would meet specification if it were tested for.

Barbengo, July 20, 2017

Cerbios-Pharma SA

Alessandro Perenna

Quality Assurance

Unit

Web