





Test Certificate No.: 2237/2024

Testing laboratory

Eurofins BioPharma Product Testing Slovakia s.r.o.

Radlinského 9, 811 07 Bratislava

IČO: 31 329 209 Place of work:

Testing laboratory Bratislava

Kollárovo nám. 9, 811 07 Bratislava

Tel.: 0911 810 533

cspharmask@bpt.eurofinseu.com, www.eurofins.sk

Customer

Alchimica s.r.o.

Žirovnického 694

25263 Roztoky, Praha-západ Česká republika

Product information No.:

Sample description:

2237

Chondroitin sulfát Bovine směsný vzorek - batch.: DCS2312505, DCS2312506

Sample reception date:

26.02.2024

Date of Testing:

26.02.2024 - 08.03.2024

Certificate issued on: 25.04.2024

Physical and chemical tests

Parameter	Unit	Allowed Value	Measured Value	Uncertainty*	Method	Testing method	Е	SL	TT
Loss on drying	%	-	9,07	1,9%	G	ŠPP 004-F (Ph. Eur.)	-	BA	Α
Content of chondroitin sulphate calculated on	%	(E)	90,472	20%	HPLC-DAD	ŠPP 027-F	-	ВА	Α
sulphate calculated on USP standard									

Result is calculated on dry basis.

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Eurofins BioPharma Product Testing Slovakia s.r o Radlinského 9, 811 07 Bratislava IČO: 31329209 DIČ: 2020297697

IČ DPH: SK2020297697

Method

HPLC-DAD

high performance liquid chromatography with diode array detector

Gravimetry

Notes:

- evaluation E

 satisfied NS - not satisfied

ŠPP, LS-PP-CH - Standard operation procedure

ND - not detected by given method

CFU - Colony forming unit

NM - necessary quantity

TT - type of test

(A) - accredited sampling

- accredited test executed at the own test laboratory

N - non accredited test executed at the own test laboratory

SA - accredited test executed under the subcontract

SN - unaccredited test executed under the subcontract

TM - testing outside the laboratory at the customer

m - the highest allowed value at the case of one sample

M, c - "M" highest allowed value for the number "c" at the case of 5 sample's evaluation

- uncertainty determined by extension coefficient k=2 (with probability of 95%) does not include the uncertainty of sampling.

- uncertainty given in units of analysed parameter reflects the uncertainty to the result of measurement.

- uncertainty given in % reflects the uncertainty from the result of measurement.

SL - analysing laboratory: BA-Bratislava, PN-Piešťany

Disclaimer:

The laboratory is not responsible for the information provided by the customer, which can affect the validity of the results.

If the sample has been provided by the customer, the results refer to the sample as it was received.

Gauges and measuring equipment used for testing were calibrated or attested in accordance with the valid metrological instructions.

The above mentioned test results refer to the tested sample only!

The result given in this Test Certificate and marked as non accredited test shall not be a subject of accreditation.

The result given in this Test Certificate and marked as sub-delivery is the result of a Subcontractor's gauging made under the terms and conditions of a contract concluded eith him.

It's not possible reproduce or incorporate the test certificate into promotional materials without laboratory written authorization!

SNAS is a Signatory to the Multilateral Agreement MRA ILAC.

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Test results have been electronically validated by:

Ing. Terézia Lopatová

Worked out by: Bc. Martin Tóth Document No.: 3052/2024



Eurofins BioPharma Product Testing Slovakia s.r.o. Radlinského 9, 811 07 Bratislava IČO: 31329209 DIČ: 2020297697 IČ DPH: SK2020297697 Test Certificate approved by: Ing. Terézia Lopatová Specialist

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Tel.: 0911 810 533

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Customer

Alchimica s.r.o.

Žirovnického 694

25263 Roztoky, Praha-západ Česká republika

Product information No.:

Sample description:

Chondroitin sulfát Porcine směsný vzorek - batch.: DCS2311519, DCS2311520

Sample reception date:

26.02.2024

Date of Testing:

26.02.2024 - 08.03.2024

Certificate issued on: 25.04.2024

Physical and chemical tests

Parameter	Unit	Allowed Value	Measured Value	Uncertainty*	Method	Testing method	Е	SL	TI
Loss on drying	%		9,69	1,9%	G	ŠPP 004-F (Ph. Eur.)	-	ВА	Α
Content of chondroitin sulphate calculated on USP standard	%	-	90,484	20%	HPLC-DAD	ŠPP 027-F	=	ВА	Α

Result is calculated on dry basis.

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Eurofins BioPharma Product Testing Slovakia sind Radlinského 9, 811 07 Bratislava IČO: 31329209 DIČ: 2020297697

IČ DPH: SK2020297697

Method

HPLC-DAD G

high performance liquid chromatography with diode array detector

Gravimetry

Notes:

- evaluation

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CFU - Colony forming unit

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- accredited test executed at the own test laboratory - non accredited test executed at the own test laboratory N

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Test results have been electronically validated by:

Ing. Terézia Lopatová

Worked out by: Bc. Martin Tóth Document No.: 3053/2024

eurofins BioPharma Product Testing

Eurofins BioPharma Product Testing Slovakia s.r.o. Radlinského 9, 811 07 Bratislava IČO: 31329209 DIČ: 2020297697 IČ DPH: SK2020297697 Test Certificate approved by: Ing. Terézia Lopatová Specialist

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