Valid from: 28-01-2022 Replaces: 19-02-2021

Batch Release Certificate

Product name: Dextran T40

Specification No.: 40029

Batch No.: ABxxx

Manufacturing sites: Pharmacosmos A/S, Roervangsvej 30, DK-4300 Holbaek, Denmark

254629 DKMA* No.: N/A GMP certificate No.: FDA establishment No.: N/A FDA facility classification: N/A EDQM* certificate No.: N/A

Method:	Parameter:	Results of analysis:	_	Limits:
_	Description:	Complies		White or almost white powder
DF034	Average molecular mass, Mw:	xx,000		35,000 - 45,000
Pharmacosmos	Loss on drying (105°C, 5h), % w/w:	x	**	≤7
Pharmacosmos	Color of solution (Absorbance at 375 nm, 10% sol., 1 cm):	x.xx		≤0.05
Pharmacosmos	Specific rotation (+/-) °:	+x		+195 – +201
Pharmacosmos	Nitrogen containing impurities, ppm N:	x		≤100

We confirm that no solvents are used in the production of this product.

We confirm that no metal catalysts or metal reagents are used in the manufacturing of this product.

I hereby certify that the above information is authentic and accurate. This batch of technical quality dextran has been manufactured, including packaging and quality control in full compliance with the above mentioned specifications.

Date (dd.mm.yyyy):

Heidi Skjødt Andersen, M.Sc. Pharm., Quality Control

PHARMACOSMOS A/S ROERVANGSVEJ 30 DK-4300 HOLBAEK DENMARK

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^{*)} EDQM refers to 'European Directorate for the Quality of Medicines and Healthcare'. DKMA refers to 'Danish Medicines Agency'
**) Due to the hygroscopic nature of the powder, loss on drying may change during storage. We recommend keeping small container well closed and protected against moisture.