

## Batch Release Certificate

**Product name:** Dextran 40 JP

**Specification No.:** 40010

**Batch No.:** xxxxxx

**Manufacturing date:** mmmm\_yyyy

**Retest date (3 years):** mmmm\_yyyy

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

**DKMA\* No.:** 254629

**GMP certificate No.:** DK API-H 00083617, DK API-V 00083617

**FDA establishment No.:** FEI 3002807874

**FDA facility classification:** Acceptable

**EDQM\* certificate No.:** Certificates for Dextran 1, 40, 60 & 70 Ph.Eur. available

Method:	Parameter:	Results of analysis:	Limits:
JP	Appearance of solution:	Complies	Clear and colorless
JP	Identification:	Complies	** Solution does not change in color
JP	Assay (dextran after drying, % w/w):	x.x	98.0 – 102.0
JP	Average molecular mass, Mw:	x,xxx	app. 40,000
JP	Viscosity, intrinsic, dL/g:	x.xx	0.16 – 0.19
JP	Viscosity, intrinsic, high fraction, dL/g:	x.xx	≤0.27
JP	Viscosity, intrinsic low fraction, dL/g:	x.xx	≥0.09
JP	pH (10% solution):	x.x	5.0 – 7.0
JP	Nitrogen containing impurities, % w/w N:	x.xxx	≤0.010
JP	Chloride, % w/w:	Complies	≤0.018
JP	Reducing substances per g:	Complies	≤15 (mg glucose)
JP	Heavy metals, ppm lead:	x	≤20
JP	Arsenic, ppm:	Complies	≤1.3
JP	Loss on drying (105°C, 6h), % w/w:	x.x	≤5.0
JP	Residue on ignition, % w/w:	x.x	≤0.1
JP	Bacterial endotoxins, IU/g:	Complies	<2.5
JP	Antigenicity:	Not tested	** Complies

References to official monographs are to be considered as current editions.

\*) EDQM refers to 'European Directorate for the Quality of Medicines and Healthcare'. DKMA refers to 'Danish Medicines Agency'

\*\*\*) Test is not carried out routinely.

We hereby confirm that no class 1, class 2 and class 3 solvent, cf. ICH Q3C and VICH GL 18, is used in the manufacturing of this product.

We hereby confirm that no metal catalysts or metal reagents, cf. EMEA/CHMP/4446/2000 and Ph. Eur. 5.20 are used in the manufacturing of this product

## CERTIFICATE OF CONFORMITY

I hereby certify that the above information is authentic and accurate. This batch of Active Pharmaceutical Ingredient has been manufactured, including packaging and quality control at the above mentioned site in full compliance with the GMP requirements for active starting materials and with the above mentioned specifications. The batch processing, packaging and analysis records were reviewed and found to be in compliance with GMP.

Date (dd.mm.yyyy):

Qualified Person, Heidi Skjødt Andersen, M.Sc. Pharm