Technical Quality Dextran

Technical Quality Dextran is a high purity dextran fraction with selected average molecular weight and molecular weight distribution.

Easily soluble Pharmacosmos Technical Quality Dextran fractions are used as starting or intermediate reagents in several different processes in biotech, photographic and chemical manufacturing industries.

- Dextran T1
- Dextran T1.5
- Dextran T3.5
- Dextran T5
- Dextran T6
- Dextran T10
- Dextran T20
- Dextran T25
- Dextran T40
- Dextran T60
- Dextran T70
- Dextran T110
- Dextran T150
- Dextran T250
- Dextran T500
- Dextran T750
- Dextran T2000

Suggested reading

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Dextran T1 Technical Quality

Dextran T1

Technical Quality Dextran T1 is a high purity dextran fraction with selected average molecular weight and molecular weight distribution.

- Documentation
- Item no. & Packaging
- Use of Dextran

If you require a dextran product manufactured according to Good Manufacturing Practice (GMP), we recommend Pharmacosmos' Pharmaceutical Quality Dextrans.

Download Section

BRC Dextran T1

Batch Release Certificate | Dextran 1

GMP Active Substances Human | April 2014

Statement on Good Manufacturing Practices for Dextran API

MSDS on Dextran Powder

Safety Data Sheet

Technical Quality Dextrans from Pharmacosmos are packed and stored in safe and clean conditions.

Technical Quality Dextran T1 is available in the following pack sizes:

Item no	Weight	Packaging
5510 0001 9006	100 g.	Polyethylene container, FDA registered, DMF type III.
5510 0001 9007	500 g.	Polyethylene container, FDA registered, DMF type III.
5510 0001 9001	5 kg.	Polypropylene container. Dextran powder sealed in double polyethylene bags
5510 0001 9005	50 kg.	Fiber drum with the dextran powder sealed in double polyethylene bags.

Pharmacosmos supplies Technical Quality Dextrans to all countries. The product packing meets all national and international regulations.

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Suggested reading

Valid from: HSA/LC 10-12-2012
Replaces: HSA/LC 01-10-2012

Batch Release Certificate

Product name: **Dextran T1**

Specification No.: **40036**

Batch No.: ABxxx

Manufacturing sites: Pharmacosmos A/S, Roervangsvej 30, DK-4300 Holbaek, Denmark
DKMA* No.: 254629
GMP certificate No.: N/A
FDA establishment No.: N/A
FDA facility classification: N/A
EDQM* certificate No.: N/A

Method:	Parameter:	Results of analysis:	Limits:
–	Description:	Complies	White Powder
LI030-1	Average molecular mass, Mw:	xx,100	800 – 1,200
DF001	Loss on drying (105°C, 5h), % w/w:	x.x	≤7
DF030	Color of solution (Absorbance at 375 nm, 6% sol., 1 cm):	x.xx	≤0.12
DF005	Acidity or Alkalinity:	Complies	Complies
DF009	Specific rotation (+/-) °:	+x.x	+140 – +170
DF012	Heavy metals, ppm lead:	x	≤10
DF019	Nitrogen containing impurities, ppm N:	x	≤100

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

I hereby certify that the above information is authentic and accurate. This batch has been manufactured, including packaging and quality control in full compliance with the above mentioned specifications. NOTE: This product is not manufactured according to Good Manufacturing Practice (GMP).

Date (dd.mm.yyyy):

Hans Juhl, M.Sc., Quality Control

PHARMACOSMOS A/S
ROERVANGSVEJ 30
DK-4300 HOLBAEK
DENMARK

TELEPHONE: +45 59 48 59 59
TELEFAX: +45 59 48 59 60
E-MAIL: info@pharmacosmos.com
WEB-SITE: www.pharmacosmos.com

CVR NO.: DK15517085
VAT NO. EXPORT: DK29127204

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