



SZABO SCANDIC

Part of Europa Biosite

Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

Weitere Information auf den folgenden Seiten!
See the following pages for more information!



Lieferung & Zahlungsart

siehe unsere [Liefer- und Versandbedingungen](#)

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

SZABO-SCANDIC HandelsgmbH

Quellenstraße 110, A-1100 Wien

T. +43(0)1 489 3961-0

F. +43(0)1 489 3961-7

mail@szabo-scandic.com

www.szabo-scandic.com

[linkedin.com/company/szaboscandic](https://www.linkedin.com/company/szaboscandic) 

Catalog Number: 11503

Product Description: Corning® TRANSFER CAP, 5L ERL, STERILE, MPC

Component Materials:

Cap - Virgin High Density Polyethylene, meets USP Class VI requirements for plastic containers and closures. Heavy metal free (meets CONEG requirements) color concentrate.

Accessories:

- Tubing - C-Flex, Thermoplastic Elastomer, meets USP, Class VI requirements for containers and closures.
- Filter - 50mm/0.0um Polytetrafluoroethylen (PTFE), meets USP, Class VI requirements for Biological Plastics.
- Diptube - High Density Polyethylene, meets USP, Class VI requirements for plastic containers and closures.

Product Dimensions:

Length of Diptube:	- 12.75 in.	Length of Tubing:	- 13.25 in.
Tolerances	- +/-0.05 in.	Diameter of Cap	- 4.16 in.

Sterilization - Product has been sterilized and dosimetrically released per the requirements of ANSI/AAMI/ISO 11137 "Sterilization of health care products- Radiation". Products meet a minimum Sterility Assurance Level (SAL) of 10^{-6} .

BSE/TSE - Product complies with the latest revision of EMA/410/01 "Note for Guidance on minimizing the risk of transmitting animal spongiform encephalopathy agents via human and veterinary medicinal products" by virtue of all bovine derived material having been processed per specific conditions of section 6.4 of EMA/410/01.

Non-Pyrogenic - Tested and has met the criteria established in the current version of ANSI/AAMI ST 72, Bacterial Endotoxins - Test methodologies, routine monitoring, and alternative to batch testing". The acceptance level for product is ≤ 0.1 EU/mL or ≤ 4 EU/device.

Quality Control Testing - Representative production samples are collected and inspected in accordance with current applicable product specifications. Inspection records are reviewed and approved by qualified personnel for product release. Key inspections and inline tests are listed below:

Visual Inspection – Pass
Packaging Inspection - Pass
Integrity Test – Pass

Lot Number Designation -

8 Digit Lot Number: First 3 digits – Julian date, start of manufacturing; Next 2 digits – Year of manufacture; Last 3 digits – Batch identification.

Rev 1