

Data Sheet

CellGenix® GMP Recombinant Human Transforming Growth Factor-beta 1 (rh TGF-β1)

Order No.: 1026-050 (50 µg)

Product Characteristics

Source Human amniocyte cell line (CAP®1)

Description Human Transforming Growth Factor-beta 1, accession # P01137, Ala279-Ser390

Molecular mass 25.6 kDa per homodimer

Formulation Lyophilized from a 0.2 µm-filtered solution containing 1 % Mannitol

Intended use For further manufacturing use.

Quality Parameters

Identity Confirmed by Immunoblotting and N-terminal sequencing of the final product.

Activity 9 - 36 x 10⁶ IU/mg calibrated against NIBSC #89/514

Measured in a cell proliferation assay using a TGF-β1-dependent cell line,

HT2 clone A5E

Purity \geq 97 %, as determined by SDS-PAGE

(under reducing and non-reducing conditions, visualized by Coomassie staining)

Product-related proteins ≤ 5 %, as determined by SDS-PAGE

(under non-reducing conditions, visualized by Coomassie staining)

Host-cell DNA \leq 20 ng/mg, as determined by gPCR

Endotoxin ≤ 10 EU/mg, as determined by LAL gel clot test according to Ph. Eur. and USP

Mycoplasma No mycoplasma detected, as determined by gPCR according to Ph. Eur.

Sterility Sterility test according to Ph. Eur. and USP of the vialed product

Mass per vial $\geq 43 \mu g$, as determined by spectrophotometrical measurement

Animal-derived ADCF Level 1: The final product contains neither animal- nor human-derived

component-free materials.



 $^{^{}f 1}$ CAP $^{f 8}$ is a registered trademark of CEVEC Pharmaceuticals GmbH, Germany.



Shipment & Storage

Transport Ambient temperature. Please refer to Technote (www.cellgenix.com)

Shelf life 1 year from date of final release

Storage & Stability Store lyophilized cytokine at -20 °C to -80 °C.

Avoid repeated freeze/thaw cycles.

Handling Instructions

Reconstitution Recommended in sterile water to a final concentration of 250 μg/ml (for 50 μg vials).

Dilution Recommended in CellGenix® serum-free media. For dilution with protein free

medium, a carrier protein (0.1–1 % albumin or 1–10 % appropriate serum) has to be included. Failure to dilute product according to these instructions may result in loss

of activity.

Quality Statement

This product is manufactured, tested and released in compliance with the relevant GMP-guidelines. No animal or human-derived components are present in the final product, and with the exception of the production cell line, no animal or human-derived materials were used in production (ADCF Level 1). USP chapter <1043> has been considered in the design of this product. This product is compliant with the Ph. Eur. general chapter 5.2.12 and ISO Technical Standard 20399.

The production cell line was derived from a human amniocyte cell line (CAP®) and was characterized following ICH Guidelines Q5A and Q5D. For the originating CAP® cell bank a Biologics Master File (BB-MF) was submitted to the U.S. Food and Drug Administration (FDA).