

# PRODUCT INFORMATION

## Elution medium for itG Ge-68/Ga-68 Generator

Product No. D010

### Product Description

This product is manufactured according to GMP requirements. It is tested for sterility and bacterial endotoxins and is intended for the elution of the itG Ge-68/Ga-68 Generator. To be used by qualified personnel only.

Content: 5 vials containing 4.15 ml 0.05 M HCl each  
> 5 ml PP bottle, crimp top  
> Chlorobutyl rubber stopper, 20 mm, with FluroTec™ Barrier Film center tear-off crimp cap, 20 mm, red

### Application

This product is only to be used for the elution of the itG Ge-68/Ga-68 Generator. It is not intended for direct use in patients.

### Packaging

This product is packed ready to use. It is labeled according to ITG manufacturing documents. Each package, packed in white cardboard boxes, contains one to five vials.

### Storage

Store this product in a dry place, protected from light at temperatures up to 30 °C. Keep this product out of reach of children.

### Product Release

This product has been released according to the product specification. Each shipment includes a Certificate of Analysis (CoA) which contains all batch specific information. For further information please contact us directly (contact details see below).

### Toxicology

Handle, transport and store with care, avoid inhalation, ingestion, eye or skin contact. Wear suitable protective clothing<sup>[1]</sup>. For detailed information we refer to the cited reference.

### References

<sup>[1]</sup> GESTIS -Database on hazardous substances  
(<http://www.dguv.de/ifa/Gefahrstoffdatenbanken/GESTIS-Stoffdatenbank/index.jsp>)

### Notification

**This document does not exempt you from performing the standard control upon receipt of incoming goods!**

This product has been manufactured according to the regulations applicable at the site of manufacture. It is a solution with defined specifications as declared in the certificate of analysis – suitable for the elution of the itG Ge-68/Ga-68 Generator.

The quality of a potential final pharmaceutical product has to be checked by the producer, the quality of the product is only partially determined by the quality of the ingredients.

The product is not intended and not suitable to be used directly and/or unprocessed in humans. The customer has to ensure himself that he is in compliance with all applicable legal requirements from all competent authorities for the site of use.