





## **EC** Certificate

Production Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex V
(Devices in class I in sterile conditions, sterilised systems or procedure packs)

No. G2S 058561 0031 Rev. 00

Manufacturer IZI Medical Products, LLC

5 Easter Court, Suite J Owings Mills MD 21117

USA

Product Sterile Syringes and Imaging Markers for Category(ies): Radiology, Radiation Therapy and Surgical

**Procedures** 

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture in accordance with MDD Annex V. This quality assurance system covers those aspects of manufacture concerned with securing and maintaining sterile conditions of the respective devices / device categories and conforms to the requirements of this Directive. It is subject to periodical surveillance. See also notes overleaf.

**Report No.:** 72153904

 Valid from:
 2020-07-03

 Valid until:
 2024-05-26

Date, 2020-07-03

Christoph Dicks

Head of Certification/Notified Body