

EC CERTIFICATE

According to Annex II of the Directive 93/42/EEC on Medical Devices

Full Quality Assurance System

Certificate Number: 2195-MED-2114101

Manufacturer:

Gohar Shafa Co.,

Head Office: Unit 9, First Floor, No.328, Between 25th & 27th St., Gisha Ave,
1447893539, Tehran, IRAN

Factory: 2nd Eftekhari St., Industrial Complex #1, 6814183183,
Khorramabad – Lorestan, IRAN

Product(s):

1. Sterile Thoracic Catheter (Chest Tube)
2. Sterile Ureteral Stent Set
3. Sterile Ureteral Catheter

Model(s):

1. Straight Thoracic Catheter, Right-angle Thoracic Catheter
2. Standard Double-J Ureteral Stent Set
3. Hydrophilic Coated

Reference Report No: MM0811-P001-R01, MM0811-P001-R02, MM0811-P001-R03, MM0811-P001-R04

Szutest, Notified Body 2195, declares that the aforementioned manufacturer has implemented a quality assurance system according to Annex II (excluding section 4), Section 3 of the directive 93/42/EEC on medical devices. This quality assurance system covers those aspects of manufacturing concerned with securing and maintaining safe conditions of the respective product(s) and conforms to the provisions of this Directive. The approved quality system is subject to surveillance pursuant to Annex II, Section 5 of Directive 93/42/EEC and unannounced audits.

Szutest must be informed of any significant changes in the design and/or construction of the product(s). For class I devices with sterile conditions the quality management system evaluation is restricted to the aspects of manufacture concerned with securing and maintaining sterile conditions. For class I devices with measuring function the quality management system evaluation is restricted to the aspects of manufacture concerned with the conformity of the devices with metrological requirements.

This EC certificate is valid till 2024-05-26.

Issue Date:

2021-05-21



Rukiye BALKAN
Deputy General Manager