Notified Body 1023
INSTITUTE FOR TESTING AND CERTIFICATION, Inc.,
třída Tomáše Bati 299, Louky, 763 02 Zlín, Czech Republic

EC Certificate - Full Quality Assurance System
No. 17 0030 QS/NB

The quality system of manufacturer

Auro Technology Co.,Ltd

No.508,Taishan Street Shijiazhuang Gaoxin Technological Zone,Hebei,China Zip 050083

has been certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex II excluding (4)

for the following product category(ies):

Cross-linked Sodium Hyaluronate Gel for plastic surgery

The Notified Body No. 1023 declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subjected to periodical surveillance. For placing on the market of Class III devices covered by this certificate, an EC Design-Examination Certificate according to Annex II (Section 4) is required.

Valid from: 2017-01-30
Valid until: 2022-01-29
First Issued: 2017-01-30
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RNDr. Radomír Čevelík
Representative of the Notified Body No. 1023