Certificate of Registration

The Governing Board of
Q.A. International Certification Limited
hereby grants to:

FIZZA SURGICAL INTERNATIONAL

Registration No.: QAIC / PK / 4043 – A

(hereinafter called the Registered Company) the right to be listed in the Directory of Registered Companies in respect of the services listed below. These services shall be offered by the Registered Company at or from only the address given below in accordance with the quality management system in Compliance with the Requirements of ***ISO 13485:2016.***

Address to which this Certificate refers:

Nargate Street, Jinnah Park Colony # 2, Sialkot – Pakistan

Approved Scope to which this Certificate refers:

Manufacturers of Non-Active Reusable Surgical and Dental Instruments & Single Use Surgical Instruments.

(Please note that the above scope represents the certified activity of the named organisation and as such, the organisation may undertake additional activities that are not covered under this certification).

Signed for and on behalf of the Board

CHIEF EXECUTIVE

SCHEME MANAGER

Certificate Issue Date: *7th April 2021* - **Certificate Renewal Before:** *6th April 2022*
Date of Initial Registration: *7th April 2021* - **Re-Certification Before:** *6th April 2024*

This Certificate of Registration is granted subject to the Regulations approved by the Board.

QA INTERNATIONAL
Q.A. International Certification Ltd.
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Certificate - Zertifikat - Certificat - Certificato

CNC™
Conformity 'n Compliance
SERVICES

CERTIFICATE OF COMPLIANCE

This is to certify, that the hereunder described items of Medical Devices Directive 93/42/EEC as amended by Directive 2007/47/EC have proven their conformity to the Safety and Health requirements of the Directive.

Manufactured by: **FIZZA SURGICAL INTERNATIONAL**

Address: Nesr Gate Street, Jinnah Park, Colony no. 2,
Sialkot-51310, Pakistan

Products:

Scissors	**Extracting forceps**	**Cement spatula**
Forceps	**Curretes**	**Bone instruments**
Needle holders	**Scalars**	**Wire mesh basket and trays**
Retractors	**Root elevators**	**Orthodontic instruments**
Speculums	**Mirror and handles**	**Hollowware instruments**
Laryngoscope	**Probes**	**Crown instruments**
Dressing forceps	**Impression trays**	**Filling instruments**
Mouth gags	**Implant instruments**	**Rubber dam instruments**

Classification: Class I, re-usable, non-powered and non-measuring devices (accordingly, to the Manufacturer's declaration)

The manufacturer's technical documentation of the product(s) has been reviewed and found to comply with requirements of the above Council Directive. With drawn up an EC declaration of conformity as per Annexure VII, module A of the product, you are therefore licensed to CE mark the product(s) listed above in accordance with Article 17 of the Medical Device Directive.

This certificate shall not be reproduced except in full and remain property of CNC Services to whom must be returned on request. This certificate does not imply assessment of the series-production of the product. The holder must inform CNC Services, of any substantial changes occurred in the product or process in order to examine whether this certificate remains valid. Certificate verification available at ***https://www.cncservices.net/verification***

Conformity 'n Compliance CNC Services

CE

For CNC Services

Project Engineer

Date of Issue: 2nd April, 2021

Date of Expiry: 1st April, 2024

Certificate Number: ***EC/4970/21***

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