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CERTIFICATE OF COMPLIANCE

REGISTRATION NUMBER - QMS/1397

THIS IS TO CERTIFY THAT

ASSURE SURGICAL INTERNATIONAL

MOHALLAH MASHRIQI, MURAD PUR, SIALKOT 51310 PAKISTAN

HAS BEEN ASSESSD AGAINST THE APPLICABLE REQUIREMENTS OF DIRECTIVE 93/42/EEC AS UPDATED DIRECTIVE 2007/47/EC FOR CLASS I MEDICAL DEVICES, AND FOUND TO MEET THE REQUIREMENTS OF DIRECTIVES SATISFACTORILY

PRODUCTS

MANUFACTURE OF SURGICAL INSTRUMENTS

AND SKILL WAS EXERCISE IN CARRYING OUT THIS DUE CARE WHILE ALL ASSESSMENT, HOWEVER REGULAR CONFORMANCE ALL THE REGULATORY TO IS THE SOLE RESPONSIBILITY OF THE REQUIREMENTS MANUFACTURER. DOCUMENT AND LEGAL CANNOT USED THIS NOT BE



SCHEME MANAGER

ISSUE DATE: EXPIRY DATE: 08-02-2016 07-02-2017

NUMBER OF SITES: 01 SITES

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CERTIFICATE

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MOHALLAH MASHRIQI, MURAD PUR, SIALKOT 51310 PAKISTAN

THE MANUFACTURING SCOPE OF REGISTRATION IS AS FOLLOW

MANUFACTURE OF SURGICAL INSTRUMENTS

HAS BEEN ASSESSD AND CERTIFIED AS MEETING THE REQUIREMENTS OF

ISO 13485:2003

WHILE ALL DUE CARE AND SKILL WAS EXERCISE IN CARRYING OUT REGULAR CONFORMANCE TO ALL THE REGULATORY ASSESSMENT, HOWEVER SOLE RESPONSIBILITY REQUIREMENTS THE OF THE MANUFACTURER. DOCUMENT LEGAL CANNOT AND THIS NOT BE USED

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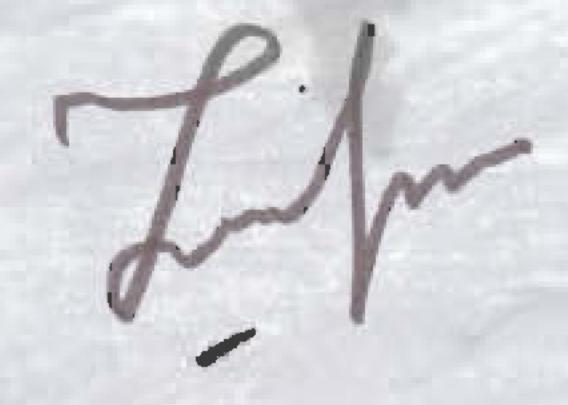
THE MANUFACTURING SCOPE OF REGISTRATION IS AS FOLLOW

MANUFACTURE OF SURGICAL INSTRUMENTS

HAS BEEN ASSESSD AND CERTIFIED AS MEETING THE REQUIREMENTS OF

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DUE CARE AND SKILL WAS EXERCISE IN CARRYING ASSESSMENT, HOWEVER REGULAR CONFORMANCE TO ALL THE REGULATORY RESPONSIBILITY SOLE THE REQUIREMENTS MANUFACTURER. THE LEGAL NOT DOCUMENT THIS AND CANNOT BE AS SUCH USED



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