



Medicines and Healthcare products  
Regulatory Agency

Hannibal House  
Elephant and Castle, London SE1 6TQ

No. 191/2552  
Royal Thai Embassy  
Islamabad, Pakistan  
0-1 SEP 2009

General enquiries  
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Certified genuine signature of  
Raja M. Sarwar Khan  
*(Signature)*  
**(Mr. Pramote Pramotsab)**  
Second Secretary/Consul

Direct line  
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Mr D Iqbal  
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40 Thorn Cliff Road  
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ATTESTED

*(Signature)*

This Ministry is not responsible  
for the content of this document

Our ref: CA 007203

30 September 2003

Dear Mr D Iqbal,

**Raja M. Sarwar Khan**  
Asstt. Director  
M/o Foreign Affairs  
Islamabad

31 AUG 2009

**MEDICAL DEVICES REGULATIONS 2002: REGULATION 19**  
**Registration of Persons Placing General Medical Devices on the Market**

Thank you for informing the Competent Authority of the details of **Manufacturers Name:- Gobble Surgical**) located at **Manufacturers Address:- P. O. Gohad Pur Sialkot Pakistan** for whom you are acting as the authorised representative and for supplying the medical device information.

Your registration has been recorded based on your declaration that you have determined that the device(s) fall within the definition of "medical device", and that you have classified it/them as falling within Regulation 19 taking into account the intended purpose(s) and mode(s) of action. In accepting your registration, I should make clear that the Competent Authority does not examine each individual notification and therefore cannot and does not necessarily endorse these determinations. Neither does this letter represent any form of accreditation or approval by the UK Competent Authority.

Your registration is based upon your declaration on the RG2 form and means that:

**For Manufacturers of Class I medical devices, Assemblers, and Sterilisers**

You should now be operating under the Medical Devices Directive and the above Regulations for the products you asked us to register, by fully complying with the essential requirements, CE marking those products or labelling them as such.

**For Manufacturers of Custom-made devices**

You should be ready to claim compliance with the Directive and Regulations and should be manufacturing custom-made devices in accordance with their requirements.



Head Office  
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An Executive Agency of the Department of Health

ชำระค่าธรรมเนียมแล้ว  
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No. 192/2552 01 SEP 2009

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Islamabad, Pakistan

If you stop placing devices on the market or if you are not complying with the Regulations you should inform us so that we can amend our records. You should be aware that it is an offence to place on the market CE marked devices that do not comply with the regulations.

genuine signature of  
RATA M SARWAR KHAN

Pranote Pramonsab  
Second Secretary/Consul

The information you provided has been recorded against the reference number shown at the top of this letter, which we ask you to quote in all future correspondence and communications.

Please inform us of any changes to:

- the company information
- additional generic groups of devices (not individual products within an existing generic group)
- discontinuation of a generic group of devices.

Please use RG2, the Registration form, to tell us about any of these changes.

Thank you for registering the following generic groups of devices:

**Class I Devices:**

*Surgical Instruments (Re-Usable And Non-Powered)*

*Dental Instruments (Re-Usable & Non-Powered)*

**Custom Made Devices:**

None

**Products Covered By Article 12:**

None

ATTESTED

Raja M. Sarwar Khan

Asstt Director

M/o Foreign Affairs

Islamabad

Should you have any queries regarding your registration please contact us on the telephone number given at the top of this letter.

ATTESTED

Yours sincerely

Jasu Patel

Jasu Patel

Raja M. Sarwar Khan

Asstt Director  
M/o Foreign Affairs  
Islamabad

16 JUL 2007

UMAIR BUTT  
COUNTERSIGNED

The State of Commerce & Industry  
Public Relations Officer

DISTRICT OFFICER HEALTH  
SIALKOT

