

Our Ref: CA010880

**Mr Abdul Razzaq**  
Black Smith Ltd  
13 Church Fields Avenue  
Hanworth  
Middx  
TW13 5PB  
United Kingdom

30 March 2009

Dear Mr Abdul Razzaq,

**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:- Surgiplus Instruments** located at **Manufacturers Address:- Near Civil Hospital Stadium Road Daska 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.

**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.**

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.

*Laryngoscopes/Otoscopes And Accessories*  
*Handheld Dental Mirrors And Accessories*  
*Retraction Cords/Dental Wedges/Rubber Dam/Matrix Bands*  
*Stethoscopes*  
*X-Ray Cassettes, Cassette Holders, Image Enhancers And Intensifying Screens*  
*Hospital Beds And Patient Positioning Aids*

**Custom Made Devices:**  
*None*

**Products Covered By Article 12:**  
*None*

### **Confidentiality**

Please note that in accordance with Directive 2007/47/EC as of 21st March 2010 information on the registration of persons responsible for placing devices on the market will no longer be treated as confidential and the Competent Authority will provide third parties with information on the name and address of manufacturers and authorised representatives and their devices that have been registered. However the names of individuals, their telephone numbers and email addresses will remain confidential. This will apply to all medical devices and in vitro diagnostic medical devices.

Should you have any queries regarding your registration, please do not hesitate to contact us.

Yours sincerely



Margaret Bruce-Tamakloe  
Regulatory Affairs Officer

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