bsi.

By Royal Charter

EC Certificate - Production Quality Assurance

Directive 93/42/EEC on Medical Devices, Annex V

No. **CE 507707**

Issued To: **MedRx, Inc.**
1200 Starkey Road, Suite 105
Largo
Florida
33771
USA

In respect of:

The manufacture of Audiometers

Those aspects of Annex V relating to metrology in the manufacture of real ear and hearing instrument measurement devices

on the basis of our examination of the quality assurance system under the requirements of Council Directive 93/42/EEC, Annex V. The quality assurance system meets the requirements of the directive. For the placing on the market of class IIb and class III products an Annex III certificate is required.

For and on behalf of BSI, a Notified Body for the above Directive (Notified Body Number 0086):

Frank Lee, EMEA Compliance & Risk Director

First Issued: **14 September 2006** Date: **31 August 2016** Expiry Date: **13 September 2021**

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