



Product Service

EC Certificate

Production Quality Assurance

Directive 93/42/EEC on Medical Devices (MDD), Annex V

(Devices in class I in sterile conditions, sterilised systems or procedure packs)

No. G2S 10 08 57582 010

Manufacturer:**FEATHER SAFETY RAZOR CO., LTD.**

3-3-70, Ohyodo-Minami, Kita-ku,

Osaka

531-0075 JAPAN

EC-Representative:**pfm medical ag**

Wankelstr. 60

50996 Köln

GERMANY

**Product
Category(ies):**

Non-Invasive Blades, Blade Removers

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture in accordance with MDD Annex V. This quality assurance system covers those aspects of manufacture concerned with securing and maintaining sterile conditions of the respective devices / device categories and conforms to the requirements of this Directive. It is subject to periodical surveillance. See also notes overleaf.

Report No.:

ONQ73527316A

Valid until:

2015-12-25

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Hans-Heiner Junker



TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

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No. G2S 10 08 57582 010**Facility(ies):****FEATHER SAFETY RAZOR CO., LTD. HEAD OFFICE**
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