

EC Certificate



Full Quality Assurance System
Directive 93/42/EEC on Medical Devices, Annex II excluding (4)

Registration No.: HD 2158053-1

Manufacturer: Guilin Woodpecker Medical
Instrument Co., Ltd.
Information Industrial Park,
Guilin National High-Tech Zone
Guilin
541004 Guangxi
P.R. China

Products: Gutta Percha Obturation Device, Dental Diode Laser Device

TÜVRheinland

The Notified Body hereby declares that the requirements of Annex II, excluding section 4 of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex II, section 5 of the aforementioned directive. For placing on the market of class III devices covered by this certificate an EC design-examination certificate according to Annex II section 4 is required.

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A handwritten signature in blue ink, appearing to be 'W. Hsu'.

Dipl.-Ing. W. Hsu
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TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.