EC Certificate

Full Quality Assurance System
MDD Annex II excl. 4

Registration No.:	HD 1709948-1
Manufacturer:	Nouvag AG St. Gallerstr. 23-25 9403 Goldach Switzerland
Products:	Medical devices Products included: - Suction pumps - Infiltration pumps - Motor systems for surgical and dental hand pieces - Surgical and dental hand pieces - Tube Sets, sterile - Blades, sterile - Rotating instruments - Attachments

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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TÜV Rheinland LGA Products GmbH Zertifizierstelle

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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.