

EC Certificate

The LGA InterCert GmbH - Notified Body 1275 according to the EC Directive concerning medical devices – herewith confirms that the following company



Amann Girrbach AG Herrschaftswiesen 1 6842 Koblach

Austria

with the additional site

Amann Girrbach GmbH Dürrenwea 40 75177 Pforzheim

Germany

applies a

Full Quality Assurance System

for the design, manufacture and final inspection of the medical devices as specified in the enclosure within the scopes: ceramic and metallic materials for production of dental prosthesis

according to

Directive 93/42/EEC, Annex II excluding (4).

In a certification audit carried out by LGA InterCert GmbH, evidence was furnished proving that the quality system of the company fulfills the requirement of the above-mentioned directive. The surveillance through the LGA InterCert GmbH is effected according to Annex II (5).

This certificate is valid until January 31, 2014 Registration no.: 1901893-006-000-002

Audit report no.: 1901893 of: December 14, 2010

Nuremberg, January 24, 2012 TE · GER,

Roland Grubes

lead of Certification Office Medical Devices

BENANNTE STELLE NOTIFIED BODY 1275

It is allowed for the company to sign the products specified in the enclosure with the adjoining sign.



ebruary 1, 2011, Reg. no.: 1901893-006-000 (Cancellation of manufacturing site)