CERTIFICATE

for the
Quality Assurance System

DEKRA Certification

As a notified body of the European Union (Reg. No. 0124) DEKRA Certification GmbH hereby approves the Quality Assurance System applied for design, manufacture and final inspection by the company

PATH medical GmbH

Landsberger Straße 63 • 82110 Germering, Germany

Approval is based on the decision dated 18.12.2009 and the result of the report no. 51260-Z1-00 and is performed in accordance with the stipulations of

Annex II, Section 3 of the Directive 93/42/EEC

of the Council dated June 14, 1993 governing medical devices. The certification is applicable to the devices specified in the Annex. The devices in question are subjected to testing and examination in accordance with Annex II, Section 3 of the Directive 93/42/EEC. The listed devices may be affixed with the CE marking indicated below.

CE 0124

Date of the first certification:	18.12.2009	Date of the last recertification:	--.--.----
This certificate is valid until:	17.12.2014	Certificate-registration No.:	51260-16-00

English version

DEKRA Certification GmbH
Stuttgart, 18.12.2009

DEKRA Certification GmbH • D-70565 Stuttgart, Handwerkstraße 15 • DEKRA Certification

Akkreditiert durch
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
ZLG-ZQ-992.94.16