

Certificate

The LGA InterCert Zertifizierungsgesellschaft mbH, a recognized registrar for quality management systems, herewith confirms that the following company

santec *medicalprodukte gmbh*

san
tec

**Mainstraße 32
63868 Großwallstadt
Germany**

has introduced and applies a

Quality Management System

for medical devices according to

EN ISO 13485:2003 + AC:2007

Requirements for regulatory purposes

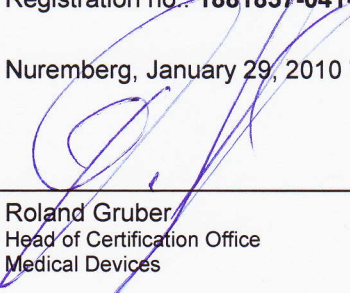
for the scopes design, development, production and distribution of sterile Tube Systems, Infusion Devices and -sets, Transfusions Systems and -sets, dynamic Bubble Traps and non absorbable surgical Suture as well as implantable Pledgets and Chordae Loops.

Proof of fulfillment of the above-mentioned standard has been provided by the certification through LGA InterCert.

This certificate is valid until August 28, 2013
Registration no.: **1881857-041-000-001**

Audit report no.: 1891693
of: September 14, 2009

Nuremberg, January 29, 2010 (*)


Roland Gruber
Head of Certification Office
Medical Devices



(*) first issue August 29, 2008, Reg. no.: 1881857-041-000 (Relocation of the company)