



# CERTIFICATE

The Certification Body of  
**TÜV SÜD AMERICA INC.**  
Management Service Division

hereby certifies that

**Karl Storz GmbH & Co. KG**  
Mittelstraße 8  
Tuttlingen D - 78532, Germany

has implemented a Quality Management System  
in accordance with:

**ISO 13485:2003**

The scope of this Quality Management System includes:

**Design and development, manufacturing, distribution,  
installation and servicing of active medical devices, non-active  
medical devices and implants and sterile medical devices  
(products in scope: see attachment)**

This Certificate is valid until: **March 31, 2010**

Certificate Registration No: **S 951 03 2263**

Original Issue Date: **December 8, 2003**

Last Revision Date: **May 11, 2007**



  
Gary W. Minks  
VP of Regulatory Affairs



CMDCAS  
Recognized  
Registrar



# CERTIFICATE

## Products in Scope

Surgical, diagnostical and dental instruments (active and non-active)  
Rigid and flexible endoscopes for diagnostics and therapy  
Cameras, devices and auxiliary devices for imaging procedure with non ionical radiation  
Endoscopic systems  
Lithotriptors and accessories  
Motor systems and accessories  
High frequency surgery devices  
Operating room control systems and accessories  
Implantable clamps for ligation of tubings and vessels  
Bone implants (non active)  
Containers and accessories  
Laser

This Certificate is valid until: March 31, 2010

Certificate Registration No: S 951 03 2263

Original Issue Date: December 8, 2003

Last Revision Date: May 11, 2007



  
Gary V. Minks  
VP of Regulatory Affairs



CMDCAS  
Recognized  
Registrar