

EC Certificate

mdc medical device certification GmbH

Notified Body 0483
herewith certifies that

joimax[®]
helping to treat patients

joimax GmbH
Amalienbadstraße 41
RaumFabrik 61
76227 Karlsruhe
Germany

for the scope

**Instruments for the endoscopic spine surgery, TESSYS[®] Reamers and
TESSYS[®] Access Kits, Irrigation Pump with accessories, Shrill[®] System**

has introduced and applies a

Quality System

for the design, manufacture and final inspection.

The mdc audit has proven that this quality system
meets all requirements according to

Annex II – Section 3 of the Council Directive 93/42/EEC

of 14 June 1993 concerning medical devices.

The surveillance will be held as specified in Annex II, Section 5.

Valid from	2010-08-04
Valid until	2013-12-23
Registration no.	0353.15.01/0
Report no.	E 0353.15 / 2010-08-04
Stuttgart	2010-08-04



Head of Certification Body

