

EC Certificate

mdc medical device certification GmbH

Notified Body 0483
herewith certifies that

**Ronchosan GmbH
Kirchfeldstraße 7
85653 Aying
Germany**

for the scope

fistula rasp

has introduced and applies a

Quality System

for the manufacture of the products concerned and carries out a
final inspection as specified in Annex V, Section 3.

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

**Annex V – 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 V, Section 4.

Valid from	2019-10-23
Valid until	2022-01-03
Registration no.	D1323400010
Report no.	P18-01470-132907
Stuttgart	2019-10-23



Head of Certification Body

