

SQS as a conformity assessment body identification number 1250 herewith certifies the company

**Edenta Etablissement  
Industriestrasse 13  
9486 Schaanwald  
Liechtenstein**

the use of a quality assurance system in its design, development, manufacturing and distribution which fulfills the requirements set out in:

**ANNEX II**

**Directive 93/42/EEC (without section 4)**

This approval is based on the result of the report dated October 19, 2018.

The scope of validity covers the products

**rotary dental instruments, parapulpal retention pins and root posts and laboratory instruments for the dental use as well as instruments for the medical podology and footcare within the Appendix to the EC Certificate**

The following CE label can be applied to these products mentioned in the Appendix of this certificate

**CE 1250**

A condition for the validity of this certificate is a regular examination in accordance with Annex II.5 of the Directive 93/42/EEC.

Validity 01.03.2019–29.02.2024  
Issue 01.03.2019

Reg. no. 41097  
Approved Medical Responsible  
27.10.2018



F. Müller, CEO SQS



D. Taddeo, Medical Responsible



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## ANNEX II

## Directive 93/42/EEC (without section 4)

This Appendix is valid only in connection with the following certificate:

Registration Number 41097

Validity from March 1, 2019 up to and including February 29, 2024

**This approval includes the following Medical Device/s:**

Class IIa:

Product	UMDNS-Nr.
Polishers (Rubber / Arkansas stone)	16-412
Burs dental diamond (Diamond burs)	16-670
Burs dental (Tungsten carbide- and steel burs / Finishing instruments)	10-521
Burs MKG surgery (Instruments for surgery - burs)	11-341
Pin, dental retention (Parapulpal retention pins - dental)	16-700
Root canal pins (Root posts - dental)	16-202
Drilling instruments, root canal (root canal instruments - rotating)	16-411
Drilling instruments, orthopedics (Instruments podology / footcare)	17-761

Appendix issue date: March 1, 2019

