EC Certificate Full Quality Assurance System: Certificate US19/819943593

The management system of

Lares Research

295 Lockheed Avenue
Chico, CA, 95973, United States

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex II (excluding Section 4)

For the following products

High speed dental handpieces, low speed dental handpieces and dental handpiece couplers.

Where the above scope includes class III medical device(s), a valid EC Design Examination Certificate according to Annex II (Section 4) is a mandatory requirement for each device in addition to this certificate to place that device on the market.

This certificate is valid from 20 May 2021 until 01 April 2024
and remains valid subject to satisfactory surveillance audits.
Issue 2. Certified since 13 November 1997

Certification is based on reports numbered WW/MC 05190

Authorised by

Global Medical Devices Head of Notified Body

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