



EC Certificate - Full Quality Assurance System

Directive 93/42/EEC on Medical Devices, Annex II excluding Section 4

No. CE 646982
Issued To: **Intra-Lock International, Inc.**
6560 West Rogers Circle
Suite #24
Boca Raton
Florida
33487
USA

In respect of:

The design and manufacture of dental implant systems, abutments, drill bits and associated accessories and systems for the preparation of autologous platelet rich fibrin (PRF) for dental, oral and maxillofacial applications

on the basis of our examination of the quality assurance system under the requirements of Council Directive 93/42/EEC, Annex II excluding section 4. The quality assurance system meets the requirements of the directive. For the placing on the market of class III products an Annex II section 4 certificate is required.

For and on behalf of BSI, a Notified Body for the above Directive (Notified Body Number 0086):

Stewart Brain, Head of Compliance & Risk -
Medical Devices

First Issued: **2016-12-21**

Date: **2017-07-12**

Expiry Date: **2022-07-17**

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Validity of this certificate is conditional on the quality system being maintained to the requirements of the Directive as demonstrated through the required surveillance activities of the Notified Body. This approval excludes all products designed and/or manufactured by a third party on behalf of the company named on this certificate, unless specifically agreed with BSI.
This certificate was issued electronically and is bound by the conditions of the contract.