EC Certificate Full Quality Assurance System: Certificate PL19/1102977401

The management system of

Wytwórnia Zębów Sztucznych Wiedent Sp. J.

ul. Obywatelska 187/189, 94-104 Łódź, Poland

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

Wiedent ESTETIC acrylic teeth,
Estetic H heat-curing acrylic resin for crowns and bridges.

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 19 March 2021 until 11 March 2023
and remains valid subject to satisfactory surveillance audits.
Issue 3. Certified since 11 March 2003

Certification is based on reports numbered PL/WAW PL00473

Authorised by

Global Medical Devices Certification Manager

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