PLANMECA

DECLARATION OF CONFORMITY

We

Planmeca Oy,
Asentajankatu 6,
FIN-00880 Helsinki
Finland

declare under our sole responsibility that the product

Intaoral Scanner **Planmeca Emerald**

to which this declaration relates is in conformity with following standards or other normative documents

IEC 60601-1	International standard on general safety for medical electrical equipment
IEC 60601-1-2	Medical electrical equipment. Part 1: General requirements for safety. 2. Collateral Standard: Electromagnetic compatibility - Requirements and tests.

following the provisions of **93/42/EEC Directive**.
Planmeca PlanScan is Class I device with measuring function.

The Notified Body is SGS Fimko Ltd. no 0598.

Helsinki, 2017-09-11

Olli Heikkinen
Quality Director