



EU – DECLARATION OF CONFORMITY

1. PPE: Type **02**

2. Name and address of the manufacturer:

Louis Steitz Secura GmbH + Co. KG
Vorstadt 40
D-67292 Kirchheimbolanden

3. This declaration of conformity is issued under the sole responsibility of the manufacturer:

Louis Steitz Secura GmbH + Co. KG

4. Object of the declaration: **MED 1000 PERB**

5. The object of the declaration described in point 4 is in conformity with the relevant Union harmonisation legislation: **Regulation (EU) 2016/425**

For United Kingdom: Regulation 2016/425 on Personal Protective Equipment as brought into UK law and amended

For North Ireland: Regulation EU 2016/425 on Personal Protective Equipment

6. References to the relevant harmonised standards used, including the date of the standard, or references to the other technical specifications, including the date of the specification, in relation to which conformity is declared:

EN ISO 20345:2011

7. The notified body **PFI, Prüf- und Forschungsinstitut Pirmasens, Marie-Curie-Str. 19, 66953 Pirmasens, 0193** performed the EU type-examination (Modul B) and issued the EU type-examination certificate

2102863-02-86 Annex 01.

Included are orthopaedic inserts based on ORTHO-MED from Steitz if observed the dedicated manufacturing instruction.

Signed for and on behalf of: **Louis Steitz Secura GmbH + Co. KG**

Kirchheimbolanden, 04.07.2021

Rolf Steinacker / Testing and Certification