SYNTHES®

Declaration of Conformity

Synthes GmbH (Eimattstrasse 3, CH-4436 Oberdorf) hereby declares under its sole responsibility that the medical device/ device family **TRS - Trauma Recon System** (Classification IIa) meets all the provisions of the

European Council Directive 93/42/EEC, and the
Swiss Legislation on Therapeutic Products (Schweizerisches Heilmittelgesetz, HMG)
which apply.

We hereby declare the conformity of above mentioned medical device/device family according to the European Council Directive 93/42/EEC, Annex I including applicable standards as indicated in the Essential Requirements Checklist and Annex II excluding (4).

Notified Body: TÜV SÜD Product Service GmbH, Ridlerstrasse 65, 80339 Munich, Germany, identification number 0123.

The above mentioned medical device/ device family was initially CE released in the year 2008.

The list of the medical devices covered by this Declaration of Conformity is attached.

EC Certificate: G1 15 03 56032 075
Valid until: 22.12.2017

Place and Date: Oberdorf, 7 July 2015

Martin Oswald
Manager Regulatory Affairs

Martin Reinberg
WW Director of PD and Marketing Small and Large Bone Power Tools

DoC Identifier: TF10200 PWT

Attachment - Articles covered in TF10200 PWT

Article#	Article Description
05.001.201	Battery Handpiece, modular, f/TRS
05.001.241	Lid f/No. 05.001.240 (Recon Saw), f/TRS
05.001.240	Battery Handpiece, Recon Saw, w/T-Handle, f/TRS
05.001.231	Lid f/No. 05.001.201 (modular), f/TRS
05.001.227	Lid f/Battery Handpiece No. 05.001.201
05.001.202	Power Module, f/Trauma Recon System
05.001.203	Sterile Cover, f/Trauma Recon System