

DECLARATION OF CONFORMITY

MANUFACTURER

Hypro Otrokovice, s.r.o.
Přístavní 568
765 02 Otrokovice
Czech Republic, EU

MEDICAL DEVICES

Hypro-Sorb® R, O, NT, Z

bioabsorbable atelocollagen haemostatic felt

Hypro-Sorb® F

bioabsorbable barrier for guided tissue and bone regeneration

Hypro-Sorb® M

Biphasic bioabsorbable Matrix for guided tissue and bone regeneration

Hypro-Flex®

bioabsorbable atelocollagen dressing with hyaluronane

CLASSIFICATION

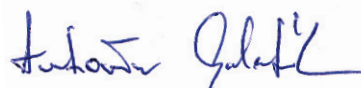
CLASS III

I, the undersigned, hereby declare that the medical devices specified above conform to the essential requirements specified in the Annex I of the Directive 93/42/EEC as amended by Directive 2007/47/EC relating on it, taking into account the product's intended use and Government order No. 336/2004 Collection of laws as amended by 245/2009 Collection of laws.

This declaration is supported by:

- EC Certificate No. 09 0627 QS/NB issued according to Annex II, Sections 3.3 and 5, of the Directive 93/42/EEC, issued in Zlin on 24th September 2009 by Institute for Testing and Certification, Notified Body No. 1023.
- EC Design-Examination Certificate No. 09 0628 CN/NB issued according to paragraph 4, Annex II of the Directive 93/42/EEC, issued in Zlin on 24th September 2009 by Institute for Testing and Certification, Notified Body No. 1023.

Otrokovice, 8th February 2011



Antonín Galatík
Company Secretary