



Notified Body No 1023  
**INSTITUTE FOR TESTING AND CERTIFICATION, Inc.**  
Zlín, Czech Republic – [www.itczlin.cz](http://www.itczlin.cz)

# EC CERTIFICATE

## No. 10 0422 QS/NB/b

Issued in compliance with the Council Directive 93/42/EEC as amended, which is implemented by the Czech Government Order No. 336/2004 (Collection of Laws) certifies that the products – medical devices of **Class III**

**Knitted vascular patch with collagen Ra ZK**  
**Knitted vascular prosthesis with collagen Ra K**  
**Knitted vascular prosthesis with collagen Ra K Incompressible**

manufactured by company

**VUP Medical, a.s.**  
**Šujanovo náměstí 3, 602 00 Brno, Czech Republic**

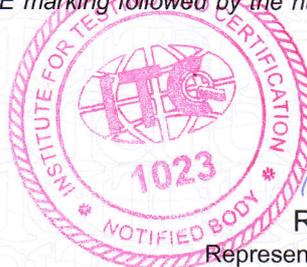
are manufactured under conditions fulfilling the quality system requirements of Annex II, Section 3.2 of the Directive 93/42/EEC and Regulation (EU) No. 722/2012, as amended.

The Notified Body No. 1023 has performed an audit of the above products quality system covering the design, manufacture and final inspection of the certified products. The quality system has been assessed, approved and is subject to continuous surveillance according to Annex II, Sections 3.3. and 5, of the Directive 93/42/EEC. The detailed description of the system parts, requirements and measures applied by the manufacturer is presented in the Final Reports No.803600811/2010, No.343602382/2014 and No.803602394/2015, which are enclosed to this Certificate.

*Condition of this Certificate use and related information:*

1. *It applies only to the quality system maintained in the manufacture of the above referenced models of the medical devices and it does not substitute the design or type-examination procedures.*
2. *The Certificate remains valid until the manufacturing conditions or the quality system are changed but until the **18<sup>th</sup> May 2017** at the latest.*
3. *The Certificate validity is conditioned by positive results of surveillance audits.*
4. *After receiving of the complementary EC Design-Examination Certificate related to the above referenced models, and fulfilling of the relevant EU legislation, the manufacturer shall affix to each medical device, of the above referenced models, the CE marking followed by the number of the Notified Body according to this example:*

**CE**  
**1023**



*Paul V*  
RNDr. Radomír Čevelík

Representative of the Notified Body No. 1023

Issued in Zlín, on 23<sup>rd</sup> March 2015

Replaces the withdrawn EC Certificate No. 10 0422 QS/NB/a issued on 4<sup>th</sup> April 2014