



3EC International a.s., Hraničná 18, 821 05 Bratislava, Slovakia
Notified Body No. 2265

EC CERTIFICATE

No. 2020-MDD/QS-103

issued in compliance with the Council Directive 93/42/EEC as amended,
certifies that the medical device of Class IIa,

Multifunction device for diagnostics of the vascular system
Types: VLab-4000 a VLab Basic

manufactured by company


EKOVUK MEDI s.r.o.
Na Maninách 159, 170 00 Praha 7, Czech Republic

is manufactured under conditions fulfilling the quality system requirements of Annex V, of the Directive 93/42/EEC as amended.

The Notified Body No. 2265 has performed an audit of the above device quality system. The production quality assurance has been assessed and found that it meets the requirements above. The quality system is subject to continuous surveillance according to Annex V, Sections 3.3, and 4, of the Directive 93/42/EEC as amended. The detailed description of the system, requirements and measures applied by the manufacturer are presented in the Audit Report No. 310464 and the Final protocol No. 310485/2020.

This certificate is issued under the following conditions:

It applies only to the quality system maintained in the manufacture of the above-referenced model of medical device and it does not substitute the design or type-examination procedures if requested. The certificate remains valid until the manufacturing conditions or the quality system are changed but until May 26th 2024 at the latest. The certificate validity is conditional upon positive results of regular surveillance audits and fulfilment of relevant legal and other requirements by the manufacturer.



Dr. Katarína Tomin Srdošová
Responsible to act on behalf of NB 2265

At Bratislava, on July 2nd, 2020

Certificate history:

Revision	Date of issue	Application for Conformity Assessment of MD number	Description
0	02.07.2020	310485	Initially granted certification