



# EC Certificate

Full Quality Assurance System

Directive 98/79/EC on In Vitro Diagnostic Medical Devices (IVDD), Annex IV excluding (4, 6)  
(List A and B and devices for self-testing)

**No. V1 076216 0007 Rev. 00**

**Manufacturer:** **VPD, Bled d.o.o.**

Pot na Lisice 4  
4260 Bled  
SLOVENIA

**Facility(ies):**

VPD, Bled d.o.o.  
Pot na Lisice 4, 4260 Bled, SLOVENIA

**Product Category(ies):** **Blood glucose measuring systems for self testing  
and lipid metabolism monitoring systems for self-  
testing**

**Model(s):**

--

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device families in accordance with IVDD Annex IV. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of List A devices an additional Annex IV (4) certificate is mandatory. All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: [www.tuvsud.com/ps-cert?q=cert:V1 076216 0007 Rev. 00](http://www.tuvsud.com/ps-cert?q=cert:V1_076216_0007_Rev.00)

**Report no.:** 713206241

**Valid from:** 2021-06-09

**Valid until:** 2024-05-26

**Date,** 2021-06-08

Christoph Dicks  
Head of Certification/Notified Body