

# Declaration of Conformity

**MANUFACTURER:** BLZ Technology (Wuhan) Co., Ltd.  
C4166, Wuhan Overseas Scholar Business Park, No.11 Dongxin Road, East  
Lake New Technology Development Zone,430074, Wuhan, China

**TRADEMARK:** N.A.

**SRN:** Not available yet

**EUROPEAN** MedPath GmbH

**REPRESENTATIVE:** Mies-van-der-Rohe-Strasse 8, 80807 Munich, Germany

**SRN:** DE-AR-000000087

**PRODUCT:** Vein Finder

**MODEL:** VS30,VS400,VS500

**BASIC UDI:** 697124654VeinFinder55

**CLASSIFICATION:** Class I (rule 13) according to Annex VIII of the Regulation(EU) 2017/745

**CONFORMITY**

**ASSESSMENT ROUTE:** Annex II + Annex III+ Annex IV of MDR

**CE CERTIFICATE NO.:** N.A.

**NAME AND ID OF THE**

**NOTIFIED BODY:** N.A.

WE HEREBY DECLARE UNDER OER SOLE RESPONSIBILITY THAT THE ABOVE MENTIONED PRODUCTS MEET THE PROVISIONS OF THE REGULATION(EU) 2017/745 ON MEDICAL DEVICES(MDR). ALL SUPPORTING DOCUMENTATTIONS ARE RETAINED UNDER THE PERMISES OF THE MANUFACTURER.

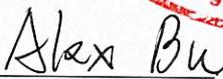
**STANDARDS** IEC 60601-1:2005+Corrigendum EN ISO 13485:2016

**APPLIED:** 1+Corrigendum 2+A1 EN ISO 14971:2012

EN 60601-1:2006+A11+A1+A12

IEC 60601-1-2:2014

EN 60601-1-2:2015

Signature:   
General Manager  
Place and Date of Issue: WUHAN, CHINA, Jan 31, 2023

