Declaration of Conformity

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Date:	September - 2020

Manufacturer :	FEATHER SAFETY RAZOR CO., LTD.
	3-70, Ohyodo-Minami 3-chome, Kita-ku, Osaka, 531-0075 JAPAN
Facility :	FEATHER SAFETY RAZOR CO., LTD. MINO SITE
	600-1, Matsumori, Mino-city, Gifu, 501-3753 JAPAN
Authorized Representative :	pfm medical ag
	Wankelstr. 60, 50996 Köln, GERMANY

We, FEATHER SAFETY RAZOR CO., LTD. located at 3-70, Ohyodo-Minami 3-chome, Kita-ku, Osaka, 531-0075 JAPAN herewith declare under our sole responsibility that the product(s) stated below are in conformity with the essential requirements and other relevant requirements of EC Directive 93/42/EEC. Any alterations made without our consent shall render this declaration null and void.

Product category :	Blood Lancets
Product group :	Blood Lancets
Classification :	Class IIa *(acc. to Rule 6 Annex IX of MDD 93/42/EEC)*
Product name :	***FEATHER BLOOD LANCET***
Model No. (Ref. No.) :	02.030.00.000
Intended purpose of use :	Cutting instruments for blood collecting
Start of CE marking :	Lot No. 11013670
GMDN-Code :	Blood lancet, single-use (61579)
EC-Directive :	Medical Device Directive 93/42/EEC
Conformity Assessment Route :	Annex II excluding (4), Full quality assurance system
Standards applied :	EN ISO 13485:2016/AC:2018, EN ISO 14971:2012, EN ISO 10993-1:2009/AC2010, EN ISO 10993-5:2009, EN ISO 10993-10:2013, EN ISO 10993-18:2009, EN ISO 11137-1:2015, EN ISO 11137-2:2015, EN ISO 11737-1:2006/AC2009, EN ISO 11737-2:2009, EN ISO 11607-1:2009, EN ISO 11607-2:2006, EN ISO 15223-1:2016, EN 1041:2008, EN 62366:2008, EN ISO 14644-1:2015, JIS T 2107:2011, EN 62366:2008 AMENDMENT 1
Notified body :	TÜV SÜD Product Service GmbH ID No.0123
	Ridlerstraße 65, 80339 München, Germany
EC Certificate :	No. G1 057582 0035 Rev.01
Place, Date :	GIFU JAPAN, September 29, 2020
Signature :	Satoshi Mitsuishi
Title :	Director