medilogAR
RED Declaration of Conformity

SCHILLER
The Art of Diagnostics

Manufacturer: SCHILLER AG
Altgasse 68, 6341 Baar, Switzerland

EU Authorised Representative: SCHILLER Medizintechnik GmbH
Otto-Lilienthal-Ring 4, 85622 Feldkirchen, Germany

Radio Equipment:			
Trade Name	medilogAR		
Product Type	Digital Holter Recorder		
Conformity Assessment	Modul A (Annex II of the RED, internal production control)		
REF Number	REF #	GTIN	Description
	3.920740 (part of 1A.306000)	07613365002096	medilogAR
System numbers (System/hardware/software)	System: N/A Software: V3.0.x* Hardware: V6.2.x*		
Standards Applied	HEALTH & SAFETY (Art. 3.1(a)) EN 60601-1:2006 (IEC 60601-1: 2012) EN 62368-1:2014/AC:2015/A11:2017 EN 62479: 2010 EMC (Art. 3.1(b)) EN 60601-1-2: 2015 + A1:2021 (IEC 60601-1-2: 2020) EN 301 489-1 v2.2.3 EN 301 489-17 V3.2.2 RADIO SPECTRUM (Art. 3.2) ETSI EN 300 328 v2.2.2		

*x is a placeholder for version number for patches / bug fixes without affecting product conformity

We, hereby declare, under our sole responsibility that the radio equipment listed above to which this declaration relates, is in conformity with technical requirements of the standards listed above and the provisions of the essential requirements of the Radio Equipment Directive 2014/53/EU.

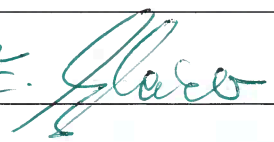
This declaration supersedes any declaration issued previously for the same product.

Signed for on behalf of: SCHILLER AG

Date of Issue: 2022-07-07
Place of Issue: Baar, Switzerland

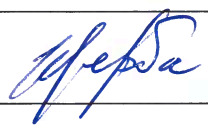
Name: ECKARD GLASER

Title / Function: HEAD OF QUALITY MANAGEMENT

Signature 

Name: VALENTINA SHCHERBA

Title / Function: HEAD OF REGULATORY AFFAIRS

Signature 

Device Dependent Declaration of Conformity Revision History

Brief Description of Change	Version	Release Date
First version	01	2022-07-07