

CARDIOVIT AT-180 RED Declaration of Conformity Rev. 01



Manufacturer:

SCHILLER AG

Altgasse 68, 6341 Baar, Switzerland

Manufacturing Site(s):

SCHILLER AG

Altgasse 68, 6341 Baar, Switzerland

EU Authorised

SCHILLER Medizintechnik GmbH

Representative:

Otto-Lilienthal-Ring 4, 85622 Feldkirchen, Germany

RED			
Trade Name	CARDIOVIT AT-180		
Product type/model	Electrocardiograph (0203)		
Conformity Assessment	Module A (Annex II of the RED, internal production control)		
REF Number	3.920570 (part of 0A.110000)		
Standards Applied	HEALTH & SAFETY (Art. 3.1(a)) EN IEC 60601-1:2020		
	EMC (Art. 3.1(b)) EN IEC 60601-1-2:2014 ETSI EN 301 489-1 V2.2.0 ETSI EN 301 489-17 V3.2.0		
	RADIO SPECTRUM (Art. 3(2)) ETSI EN 300 328 V2.2.2 ETSI EN 301 893 V2.1.1		

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: 2025-02-24

Place of Issue: Baar, Switzerland

Name: STEFAN BIGLER

Title / Function: HEAD OF REGULATORY

AFFAIRS

Signature

Name: AYNUR ASLANOVA

Title / Function: HEAD OF QUALITY

MANAGEMENT

Signature



CARDIOVIT AT-180 RED Declaration of Conformity Rev. 01



Device Dependent Declaration of Conformity Revision History

Brief Description of Change	Version	Release Date
First version	01	2025-02-24