

**EU/RE DIRECTIVE DECLARATION OF CONFORMITY**

This is a declaration made in accordance with the requirements of Directive 2014/53/EU Of the European Parliament and of the Council of 16 April 2014 on the harmonization of the laws of the Member States relating to the making available on the market of radio equipment. The declaration of conformity is issued under the sole responsibility of the manufacturer.



**Manufacturer's Name:** NIHON KOHDEN CORPORATION  
**Business Address:** 1-31-4 Nishiochiai, Shinjuku-ku  
Tokyo 161-8560, Japan

**European Representative:** NIHON KOHDEN EUROPE GmbH  
**Address:** Raiffeisenstrasse 10, D - 61191 Rosbach, Germany

**Product Name and Model Name:** Automated External Defibrillator AED-3100

**Notified Body's Name and No.:** Intertek Semko AB Sweden No: 0413 (Module B)

**EU-Type examination Certificate No.:** SE-RED-2200432 Ed.1

**Standard Applied:** IEC 60601-1: 2005  
IEC 60601-1 Amendment 1: 2012  
IEC 60601-1-2: 2014  
IEC 60601-1-6: 2010  
IEC 60601-1-6 Amendment 1: 2013  
IEC 60601-11: 2010  
IEC 60601-12: 2014  
IEC 60601-2-4: 2010+A1:2018 (Clause 202)  
EN 301 489-1 V2.2.3  
EN 301 489-17 V3.2.4  
EN 300 328 V2.2.2  
EN 62479: 2010

**Authorized Signatory:**

Tokyo, Japan / 6 June 2022

Place and date of issue

  
Hiroko Hagiwara  
General Manager  
Clinical Development & Regulatory Affairs Division