

# Medical Devices

## Declaration of Conformity

We, NordicNeuroLab AS declare under our sole responsibility that the products described below meet the provisions of the **Medical Device Regulations 2017/745 of 17 May 2017**, for medical device. All supporting documentation is retained under the premises of the manufacturer.

Manufacturer:	NordicNeuroLab AS
Address:	Møllendalsveien 1 N-5009 Bergen Norway
Product:	InroomViewing Device (LCD Monitor 40")
Version:	<b>3.1</b>
Single registration number (SRN):	NO-MF-000000555
Basic UDI DI:	7090042051006S
Risk Class:	<b>Class I</b>
Compliance:	Device covered by the present EU declaration is in conformity with the (EU) MDR 2017/745 and with the following standards <ul style="list-style-type: none"><li>o IEC 60601-1 (IEC 60601-1)</li><li>o IEC 60601-1-2 (IEC 60601-1-2)</li><li>o NS-EN ISO 14971:2019/A11:2021 (ISO 14971:2019)</li><li>o NS-EN ISO 13485:2016 (ISO 13485:2016)</li><li>o RoHS 2002/95/EC</li></ul>
Expiry Date:	Valid for 2 years from date of issue
 Thomas Lie Omdahl <b>Chief Executive Officer</b> NordicNeuroLab AS	 Date (Bergen, Norway)

NL-200-0553-03-EC Declaration\_MDR\_LCD 3.1- EN