

## Declaration of Conformity

We, the undersigned Manufacturer

<b>Company</b>	Jotron AS
<b>Address</b>	Ringdalskogen 8, NO-3270 Larvik
<b>Country</b>	Norway
<b>Telephone number</b>	+47 33139700
<b>Telefax number</b>	+47 33126780
<b>E-mail</b>	www.jotron.com

Certify and declare under our responsibility that the following product:

<b>Product Description</b>	Interface Control Unit; ICU
<b>Manufacturer</b>	Jotron AS
<b>Brand Name</b>	Jotron
<b>Model/Type</b>	87120 ICU

Is tested to and conforms with the essential test suites included in the following standards, which are in force within the EEA:

Standard	Issue date	Reference to report/file
EN 301 489-22 v.1.3.1	2003	Nemko Report No. E11181.00
EN 301 489-1 v.1.9.2	2008	Nemko Report No. E11181.00
IEC 60950-1, 2 <sup>nd</sup> edition+A2:2013	2013	Nemko Report No.187680, 187680 Attachment and 311173
EN 55032	2015	Based on Nemko Report No. E11181.00
EN 50581	2012	
ISO 14001	2015	Certificate number 900022/901317
ISO 9001	2015	Certificate number 900022/901317

And therefore complies with the essential requirements of the following directives:

Directive Name	Directive number	Further identification
EMC	2014/30/EU	Electromagnetic Compatibility
LVD	2014/35/EU	Low Voltage Directive
RoHS	2011/65/EU	Restriction of Hazardous Substances Directive Applies also to Directive 2015/863



The following Notified Bodies have been consulted in the Conformity Assessment procedure (whenever applicable):

Notified Body number	Name and address

The technical documentation as required by the conformity assessment procedure is kept at the following address for a period ending at least 10 years after the last product has been manufactured at the disposal of the relevant national authorities of any Member State for inspection:

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<b>Drawn up in</b>	Norway
<b>Date</b>	10.02.2020
 	
<b>Signature and Company Stamp</b>	Frank Løke Certification Manager