

Declaration of Conformity

We, the undersigned Manufacturer:

Company	Jotron AS
Address	Ringdalskogen 8, 3270 Larvik
Country	Norway
Telephone number	+47 33139700
Telefax number	+47 33126780
E-mail	sales@jotron.com

Certify and declare under our responsibility that the following product:

Product Description	Emergency transmitter float free
Manufacturer	Jotron AS
Brand Name	Jotron
MED - Number and item designation	MED/5.6 - 406 MHz EPIRB (COSPAS-SARSAT)
Model/Type	Tron 40S MkII, consist of; -83056 Battery Tron 40SMkII 5 years maintenance kit. -85216 FB6 Bracket w/Protective Cover -80414 HRU-kit to FB-5 and FB-6 w/H2O, HAMMAR hydrostatic release mechanism. -97821 HRU-kit to FB-4 and FBH-4 w/H2O, HAMMAR hydrostatic release mechanism. -97777 FBH-4 Bracket -101417 FB-6 Bracket Front Cover -84544 Service Kit

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	Certificate No.
IEC 61097-2 Ed.3.0	2008	MED-B-16511 Rev.02 MEDD0000296 Rev.01
IEC 60945 ed.4.0 including IEC 60945 Corrigendum 1	2002 Corr. 2008	
EN 300 066 v.1.3.1	2001-01	
IEC 63000	2018	
ISO 14001	2015	Certificate no. 901317
ISO 9001	2015	Certificate no. 900022
ISO 45001	2018	Certificate no. 907920
ISO 27001	2017	Certificate no. 904109

The product(s) also meet the requirements set out in the following SOLAS 74 Regulations and IMO Resolutions:

SOLAS 74 Reg. IV/14
SOLAS 74 Reg. X/3
SOLAS 74 Reg. IV/7
IMO Res. MSC.36(63)-(1994 HSC Code) 14
IMO Res. MSC.97(73)-(2000 HSC Code) 14
IMO Res A.662(16)
IMO Res. A.694(17)
IMO Res. A.696(17)
IMO Res. A.810(19)
IMO Res MSC/Circ 862
IMO COMSAR/Circ.32,
ITU-R M.633-4 (12/10)
ITU-R M.690-3 (03/15)

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

Directive Name	Directive number	Further identification
MED	2014/90/EU	Directive 2014/90/EU repealing Directive 96/98/EC Applies also to Implementing Regulation (EU)2022/1157
RoHS	2011/65/EU	Restriction of Hazardous Substances Applies also to Directive (EU) 2015/863

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

Notified Body number	Name and address	Directive
0470	Nemko AS, Philip Pedersens vei 11, 1366 Lysaker, Norway	MED-B
0575	DNV AS P.O.Box 300 Veritasveien 1 1322 HOVIK, Norway	MED-D

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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Address	Ringdalskogen 8, 3270 Larvik
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Telefax number	+47 33126780
Web page	jotron.com

	Drawn up in	Norway
	Date	10.11.2022
	Signature and company stamp:	Frank Løke, Certification Manager
		