



DATE OF MANUFACTURE

EC DECLARATION OF CONFORMITY

ACR Electronics, Inc. hereby declares under its sole responsibility that the following product is in conformity with Directive 2014/90/EU of the European Parliament and of the Council of 23 July 2016 on Marine Equipment, and has been type examined as follows and has been assessed in accordance with MED/4.18 of EU implementing regulation 2017/306. In accordance with the relevant community harmonized legislation, the product will be marked with the MED Mark of Conformity as follows:



yy = Last two digits of the year in which the mark is affixed

ACR Electronics, Inc. hereby declares under its sole responsibility that the following product is in conformity with Directive 2011/65/EU of the European Parliament and of the Council of 8 June 2011 on the restriction of the use of certain hazardous substances in electrical and electronic equipment (RoHS2). In accordance with the relevant community harmonized legislation, the product will be marked with the CE conformity marking as follows:



Product: 9 GHz Search and Rescue Transponder (SART), MED Item A.1/4.18
 Trade Names: Pathfinder PRO SART
 Model: Pathfinder PRO

Notified Bodies: TÜV SÜD BABT, Notified Body No. 0168
 Octagon House, Concorde Way, Fareham, Hampshire, PO15 5RL, UK
 EC Type Examination (Module B) Certificate No.: BABT-MED001120 Issue:01
 EC Quality System (Module D) Certificate No.: BABT-MED005722-H1 Issue 03

Regulations and Standards: IMO Resolution A.694(17) IEC 61097-1 (2007)
 IMO Resolution A.802(19) IEC 60945 Ed.4.0, 2002 incl. Corr. 1, 2008
 IMO Resolution A.530(13)
 MSC.247(83)

Manufacturer and holder of technical documents: OCEAN SIGNAL
Unit 4, Ocivan Way,
Margate
Kent, CT9 4NN

Signed for and on behalf of ACR Electronics Inc.

Signed: _____
Name: Dan Stankovic **Date:** March 16, 2017
Title: Director, Certification and Test

Document PATHFINDER PRO-002
ACR Part # Y1-15-0008C



This Declaration complies with
ISO/IEC 17050-1:2004 and
EU decision 768/2008/EC, Annex III

ACR Electronics, Inc. is registered by TUV USA to ISO 9001:2008/AS9100C



Notice to Distributors:

Please note that new MED requirements (MED MkII) call for the inclusion of the full Declaration of Conformity (see reverse) with all shipments of MED approved products. To remain in compliance, all distributors are now required to copy the Declaration of Conformity seen on the reverse of this page and include this copy with all onward shipments of applicable products. If this bulk master crate is not going to be broken down into smaller shipments, it is not necessary to make a copy of the Declaration of Conformity.