

DECLARATION OF CONFORMITY THE EC DIRECTIVE AND CE MARKING

We, Mirion Technologies (Capintec), Inc.

7 Vreeland Road, Florham Park, NJ 07932 USA

Phone (800) ASK-4CRC Fax (201) 825-1336

declare under our sole responsibility that the product CAPTUS 4000e THYROID UPTAKE SYSTEM, all models (including use with the Hewlett-Packard Officejet printer) to which this declaration relates is in conformity with the MDD 93/42/EEC as transposed into Swedish national law LVFS 2003:11. Models include 5430-30151, 5430-30152, 5430-30154, 5430-00007.

This product is in conformity with the following standards or other normative documents following:

- IEC 60601-1-2 Ed. 4.0, (2014): Medical Electrical Equipment Part 1-2: General requirements for basic safety and essential performance Collateral Standard: Electromagnetic disturbances Requirements and tests
- ANSI/AAMI ES60601-1:2005/A1:2012 Issued: 2012/08/20 Medical electrical equipment Part 1: General requirements for basic safety and essential performance
- CAN/CSA-C22.2 No. 60601-1:14: Third Edition Issued: 2014/03/01 Medical electrical equipment Part 1: General requirements for basic safety and essential performance
- IEC 60601-1-6 Issued: 2013/10/29 Ed: 3.1 Medical electrical equipment Part 1-6: General requirements for basic safety and essential performance Collateral Standard: Usability
- IEC 62366:2014, Edition 1.1 Issued 2014/01/28 Medical devices Application of usability engineering to medical devices
- IEC 62304:2006 + A1:2015 issued Medical device software Software life cycle processes

Certified to Annex II of the Directive 93/42/EEC on Medical Devices Thyroid Uptake Systems, Class IIa
Certified through Intertek Semko AB, Notified Body MDD
CE Mark as shown denotes conformance to the above statement.

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The Technical Construction File is maintained at:

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Florham Park, NJ 07932

The authorized representative located within the Community is:

Atlantico Systems Ltd.

34 Oldfield, Kingston, Galway, Republic of Ireland

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