EC CERTIFICATION

FULL QUALITY ASSURANCE SYSTEM

Directive 93/42/EEC on Medical Devices, Annex II excluding (4)

We hereby declare that an examination of the under mentioned full quality assurance system has been carried out following the requirements of the Swedish national legislation LVFS 2003:11 to which the undersigned is subjected, transposing Annex II (with the exemption of section 4) of the Directive 93/42/EEC on medical devices. We certify that the full quality assurance system conforms with the relevant provisions of the aforementioned directive, and the result entitles the organization to use the CE 0413 marking on those products listed below.

Organization:

Mirion Technologies (Capintec), Inc

Main Site: 7 Vreeland Road, Florham Park, New Jersey 07932 United States

Product Category:

- Thyroid Uptake Systems

For further identification of the products covered, see the MDD product list/product schedule.

Certificate Number:
41313199-04

Initial Certification Date:
08 December 2000

Certificate Valid from:
9 December 2020

Certificate Expiry Date:
26 May 2024

SWEDAC ACKREDITERING
Accred. no. 1003
Certification of Management Systems
ISO/IEC 17021-1

Bob Andersson
Certification Authority MDD
Intertek Semko AB, Kista, Sweden

10 November 2020
Signed Date

Intertek Semko AB
Box 1103, SE-164 22 Kista, Sweden
Telephone +46 8 750 00 00
medtechsweden@intertek.com

The certification is subject to the organization maintaining their system in compliance with the regulations stated in this certificate, allowing regular assessments and following the contracted requirements of the Notified Body.

Intertek Semko AB is a Notified Body according to Directive 93/42/EEC on medical devices, with identification number 0413.

MDD – Product List

Products included in the Certificate No: 41313199-04
Issued to: **Mirion Technologies (Capintec), Inc**
7 Vreeland Road,
Florham Park, New Jersey 07932
United States

Product category	Type/Model designation	Class	Sterile	GMDN code (not mandatory)	Date added
Thyroid Uptake System					
	The CAPTUS® 4000e Thyroid Uptake System	IIa	No	40648	Dec 4, 2015

Signed Date: 10 November 2020
Valid Date: 9 December 2020

Intertek Semko AB
Notified Body MDD

Bob Andersson
Certification Authority MDD

This product list is only valid together with the referenced, valid EC certificate.

The GMDN codes are assigned by the manufacturer and are only provided for convenience.

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Intertek Semko AB
Torshamnsgatan 43, Box 1103, SE-164 22 Kista, Sweden
Telephone +46 8 750 00 00, Fax +46 8 750 60 30, www.intertek.se
Registered in Sweden: No SE556024059901, Registered office: As address

MDD – Decision Report

Certificate No: 41313199-04
Date: 10 November 2020
Handled by: Caroline Åman
E-mail: medtechsweden@intertek.com

Mirion Technologies (Capintec), Inc
Attn: Mary Anne Yusko
7 Vreeland Road,
Florham Park, New Jersey 07932
United States

Purpose	Assessment to issue a new certificate due to five year extension according to the national legislation for medical devices LVFS 2003:11 (Medical Device Directive 93/42/EEC), Annex II.
Activity	Certification audit was performed remote 17 August 2020 in Florham Park, New Jersey by Luis Lopes. The technical file was reviewed 18 September 2020 by Abul Kashem.at Intertek's office.
Scope of assessment	Thyroid Uptake Systems, Class IIa
Result	2 minor non conformities were noted during the audit. Presented corrective action plans have been examined and approved by us.
Certificate Valid from	9 December 2020
Conclusions/Decisions	Referring to the above a Certificate of Conformance with the national legislation for medical devices LVFS 2003:11 (Medical Device Directive 93/42/EEC), Annex II will be issued. The Certificate is valid for products specified in the "MDD – Product List".
Follow-up assessments	Follow-up assessments are going to be performed once a year.
Appeals	Any appeal against this decision will be processed by an appeals panel as Intertek. The appeal shall be submitted to Intertek Semko AB, PO-Box 1103, SE-164 22 Kista, Sweden.
Others	Any complaints, from customers and others, and corrective actions concerning your certified quality system shall be documented and retained. Upon request Intertek Semko has the right to review this documentation.

Intertek Semko AB
Notified Body MDD

Bob Andersson
Certification Authority MDD