

Products included in the certificate no: 41314263
 Issued to: **Masimo Sweden AB**
 Svärdvägen 15
 SE-182 33 Danderyd
 Sweden

Product category	Type/Model designation	Class	Sterile	GMDN code (not mandatory)	Date added
Mainstream multi-gas sensor	IRMA CO2 Ref 200101	Iib	No	-	*
	IRMA CO2 Bluepoint Ref 200104	Iib	No	-	July 1, 2009
	IRMA CO2 Breas Ref 200105	Iib	No	-	Mar 28, 2011
	IRMA AX+ Ref 200601	Iib	No	-	*
	IRMA AX+ Max Ref 200603	Iib	No	-	*
	IRMA AX+ Bluepoint Ref 200604	Iib	No	-	Sept 8, 2010
Sidestream multi-gas sensor	ISA OR+ Ref 800401	Iib	No	-	July 1, 2009
	ISA AX+ Ref 800601	Iib	No	-	July 1, 2009
Sampling line	Nomoline REF 108210 (Connected to ISA side-stream multi-gas sensor.)	Ila	No	-	July 1, 2009
	Nomoline Adapter REF 108220	Ila	No	-	Oct 11, 2011
	Nomoline Airway Adapter Set REF 108230	Ila	No	-	Oct 11, 2011
Disposable airway adapter	IRMA Airway adapter (Adult/Pediatric) Ref 106220	Ila	No	-	*
	IRMA Airway adapter Infant Ref 106260	Ila	No	-	*
	EMMA Airway adapter (Adult/Pediatric) Ref 100620	Ila	No	-	*
	EMMA Airway adapter Infant Ref 100660	Ila	No	-	*
Mainstream multi-gas monitor for pocket PC	VEO Multigas Monitor PC, AX+ Ref 300602	Iib	No	-	*
Mainstream multi-gas monitor	Max Multigas Analyzer Ref 500100	Iib	No	-	*

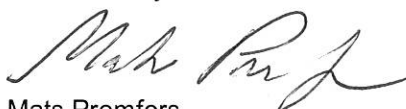
Product list for certificate no: 41314263
 Date: April 28, 2014
 Page 1 of 2

Product category	Type/Model designation	Class	Sterile	GMDN code (not mandatory)	Date added
Mainstream emergency Capnometer	EMMA (kPa) Ref 605100	IIb	No	-	*
	EMMA (mmHg) Ref 605102	IIb	No	-	*
	EMMA Capnometer (kPa) Ref 606100	IIb	No	36552	Oct 18, 2013
	EMMA Capnometer (mmHg) Ref 606102	IIb	No	36552	Oct 18, 2013
Sidestream Capnometer	ISA CO2 Ref 800101	IIb	No	-	July 1, 2009
	ISA CO2 ROOT Ref 800106	IIb	No	-	April 19, 2013

* Product added before July 1, 2009

Date of Issue: April 28, 2014

Intertek Semko AB
Notified Body MDD



Mats Premfors
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.

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

Product list for certificate no: 41314263
Date: April 28, 2014
Page 2 of 2