# MDD – Product List

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.) | IIa | No | - | July 1, 2009 |
| | Nomoline Adapter REF 108220 | IIa | No | - | Oct 11, 2011 |
| | Nomoline Airway Adapter Set REF 108230 | IIa | No | - | Oct 11, 2011 |
| Disposable airway adapter | IRMA Airway adapter (Adult/Pediatric) Ref 106220 | IIa | No | - | * |
| | IRMA Airway adapter Infant Ref 106260 | IIa | No | - | * |
| | EMMA Airway adapter (Adult/Pediatric) Ref 100620 | IIa | No | - | * |
| | EMMA Airway adapter Infant Ref 100660 | IIa | 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 | - | * |



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Registered in Sweden: No SE556024059901, Registered office: As address

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|---|---|---|---|---|---|
| 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.



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