



Sistemi s.r.l.

Manufacturer name:

AMBRA SISTEMI s.r.l.

Address:

Via Collegno 45 bis
I-10044 PIANEZZA TO

declares that following devices

Device name:

SDAL-MED

Variants:

L5, L5C, L9, V4

comply with all the requirements of the European Directive 93/42/CE, modified by the 2007/47/CE, concerning medical devices, particularly with the basic safety requirements of the Annex I, according to the EC Quality assurance System Certificate MED 28016 released by the notified body number 0476 with reference to the Annex II of the same directive.

Annexes:

⇒ EC Quality assurance System Certificate MED 28016 by the notified body number 0476

Issue date:

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Revision:

0

technical manager
Claudio Guidotti

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