



Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
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ZLG-BS-244.10.08



Product Service

EC Certificate

Production Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex V
(Devices in Class IIa, IIb or III)

No. G2 063838 0015 Rev. 02

Manufacturer:

Medax S.r.l. Unipersonale

Via R. Piva, 1/A
46025 Poggio Rusco (MN)
ITALY

Facility(ies):

Medax S.r.l. Unipersonale
Via Sandro Pertini, 4, 41039 S. Possidonio (MO), ITALY

**Product
Category(ies):**

Reusable biopsy guns. Veress needles, thoracentesis and
paracentesis kits. Vertebroplasty needles. Bone marrow
aspiration and biopsy needles. Soft tissue biopsy needles.
Punch needles for skin biopsy. Intraosseous infusion
needles. Local anesthesia needles.
Device for pre-operative localisation of non-palpable lesions
of the breast.

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex V. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class IIb and III devices an additional Annex III certificate is mandatory. See also notes overleaf.

Report No.:

ITA1321318C

Valid from:

2020-02-05

Valid until:

2024-05-26

Date,

2020-01-29

Christoph Dicks
Head of Certification/Notified Body