

Certificate of CE-Registration



This is to certify that, in accordance with either medical device Directive 93/42/EEC as amended by 2007/47/EC or Directive 98/79/EC, mdi Europa GmbH agree to perform all duties and responsibilities as the Authorized Representative for

Starkling
Gartenstraße 11
D-78573 Wurmlingen
Germany

as stipulated and demanded by the afore-mentioned Directives. The German competent authorities have allocated the medical devices of the manufacturer the following registration numbers:

| UMDNS Code | Description | Classification | Registration Number |
|------------|-------------------|----------------|---------------------|
| 10025 | Adenotomes | I | DE/CA09/0760/1601 |
| 10172 | Applicators | I | DE/CA09/0760/1598 |
| 10174 | Applicators, Clip | I | DE/CA09/0760/1592 |
| 10519 | Burs | I | DE/CA09/0760/1583 |
| 10561 | Cannulae | I | DE/CA09/0760/1581 |
| 10824 | Chisels | I | DE/CA09/0760/1632 |
| 10861 | Clamps | I | DE/CA09/0760/1629 |
| 11059 | Crushers | I | DE/CA09/0760/1626 |
| 11084 | Curets | I | DE/CA09/0760/1623 |
| 11179 | Dermatomes | I | DE/CA09/0760/1619 |
| 11254 | Dilatators | I | DE/CA09/0760/1618 |
| 11290 | Dissectors | I | DE/CA09/0760/1617 |
| 11345 | Driver/Extractors | I | DE/CA09/0760/1616 |
| 11504 | Elevators | I | DE/CA09/0760/1615 |
| 11701 | Files | I | DE/CA09/0760/1613 |
| 11774 | Forceps | I | DE/CA09/0760/1612 |

The manufacturer has provided mdi Europa with all necessary documentation, together with an appropriate Declaration of Conformity confirming that the medical devices fulfill the essential requirements of either Directive 93/42/EEC as amended by 2007/47/EC or 98/79/EC. A safety officer has been appointed for Germany and therefore is in full compliance with § 31 MPG.

Signed on 20 June 2019

Werner Sander
President

Certificate of CE-Registration



This is to certify that, in accordance with either medical device Directive 93/42/EEC as amended by 2007/47/EC or Directive 98/79/EC, mdi Europa GmbH agree to perform all duties and responsibilities as the Authorized Representative for

Starkling
Gartenstraße 11
D-78573 Wurmlingen
Germany

as stipulated and demanded by the afore-mentioned Directives. The German competent authorities have allocated the medical devices of the manufacturer the following registration numbers:

| UMDNS Code | Description | Classification | Registration Number |
|------------|---------------------|----------------|---------------------|
| 11822 | Gags | I | DE/CA09/0760/1609 |
| 11895 | Gouges | I | DE/CA09/0760/1608 |
| 11950 | Hammers, Percussion | I | DE/CA09/0760/1607 |
| 11963 | Headlights | I | DE/CA09/0760/1605 |
| 12028 | Hooks | I | DE/CA09/0760/1603 |
| 12235 | Scalpel Handles | I | DE/CA09/0760/1628 |
| 12239 | Knives | I | DE/CA09/0760/1659 |
| 12252 | Knives, Scalpel | I | DE/CA09/0760/1658 |
| 12393 | Loops | I | DE/CA09/0760/1654 |
| 12421 | Mallet | I | DE/CA09/0760/1653 |
| 12423 | Mallets, Bone | I | DE/CA09/0760/1652 |
| 12440 | Markers | I | DE/CA09/0760/1636 |
| 12549 | Mirrors Laryngeal | I | DE/CA09/0760/1638 |
| 12726 | Needle Holder | I | DE/CA09/0760/1637 |
| 12766 | Nippers, Malleus | I | DE/CA09/0760/1639 |
| 12815 | Ophthalmoskop | I | DE/CA09/0760/1640 |

The manufacturer has provided mdi Europa with all necessary documentation, together with an appropriate Declaration of Conformity confirming that the medical devices fulfill the essential requirements of either Directive 93/42/EEC as amended by 2007/47/EC or 98/79/EC. A safety officer has been appointed for Germany and therefore is in full compliance with § 31 MPG.

Signed on 20 June 2019

Werner Sander
President

Certificate of CE-Registration



This is to certify that, in accordance with either medical device Directive 93/42/EEC as amended by 2007/47/EC or Directive 98/79/EC, mdi Europa GmbH agree to perform all duties and responsibilities as the Authorized Representative for

Starkling
Gartenstraße 11
D-78573 Wurmlingen
Germany

as stipulated and demanded by the afore-mentioned Directives. The German competent authorities have allocated the medical devices of the manufacturer the following registration numbers:

| UMDNS Code | Description | Classification | Registration Number |
|------------|-----------------------|----------------|---------------------|
| 12844 | Osteotomes | I | DE/CA09/0760/1641 |
| 12849 | Otoscope | I | DE/CA09/0760/1642 |
| 13041 | Cutters, Pin and Wire | I | DE/CA09/0760/1622 |
| 13117 | Probes | I | DE/CA09/0760/1643 |
| 13126 | Proctoscope | I | DE/CA09/0760/1645 |
| 13228 | Punches | I | DE/CA09/0760/1644 |
| 13287 | Raspatories | I | DE/CA09/0760/1646 |
| 13288 | Rasps | I | DE/CA09/0760/1647 |
| 13373 | Retractor | I | DE/CA09/0760/1648 |
| 13416 | Rongeurs | I | DE/CA09/0760/1649 |
| 13448 | Saw | I | DE/CA09/0760/1650 |
| 13454 | Scalers | I | DE/CA09/0760/1627 |
| 13480 | Scissors | I | DE/CA09/0760/1630 |
| 13508 | Scoops | I | DE/CA09/0760/1633 |
| 13517 | Screwdrivers, Bone | I | DE/CA09/0760/1631 |
| 13630 | Snares | I | DE/CA09/0760/1635 |

The manufacturer has provided mdi Europa with all necessary documentation, together with an appropriate Declaration of Conformity confirming that the medical devices fulfill the essential requirements of either Directive 93/42/EEC as amended by 2007/47/EC or 98/79/EC. A safety officer has been appointed for Germany and therefore is in full compliance with § 31 MPG.

Signed on 20 June 2019

Werner Sander
President

Certificate of CE-Registration



This is to certify that, in accordance with either medical device Directive 93/42/EEC as amended by 2007/47/EC or Directive 98/79/EC, mdi Europa GmbH agree to perform all duties and responsibilities as the Authorized Representative for

Starkling
Gartenstraße 11
D-78573 Wurmlingen
Germany

as stipulated and demanded by the afore-mentioned Directives. The German competent authorities have allocated the medical devices of the manufacturer the following registration numbers:

| UMDNS Code | Description | Classification | Registration Number |
|------------|--------------------------|----------------|---------------------|
| 13645 | Spatulas | I | DE/CA09/0760/1660 |
| 13664 | Specula Nasal | I | DE/CA09/0760/1662 |
| 13665 | Specula Rectal | I | DE/CA09/0760/1663 |
| 13666 | Specula Vaginal | I | DE/CA09/0760/1664 |
| 13707 | Spreaders | I | DE/CA09/0760/1665 |
| 13730 | Sterilization Containers | I | DE/CA09/0760/1579 |
| 13823 | Strippers | I | DE/CA09/0760/1582 |
| 14066 | Tongue Depressor | I | DE/CA09/0760/1588 |
| 14154 | Trocars | I | DE/CA09/0760/1590 |
| 14255 | Tuning Forks | I | DE/CA09/0760/1595 |
| 14257 | Tweezers | I | DE/CA09/0760/1596 |
| 14291 | Urinals | I | DE/CA09/0760/1599 |
| 14427 | Kick Buckets | I | DE/CA09/0760/1604 |
| 14455 | Wire Drivers | I | DE/CA09/0760/1597 |
| 15076 | Laryngoscope Rigid | I | DE/CA09/0760/1656 |
| 15275 | Awls | I | DE/CA09/0760/1584 |

The manufacturer has provided mdi Europa with all necessary documentation, together with an appropriate Declaration of Conformity confirming that the medical devices fulfill the essential requirements of either Directive 93/42/EEC as amended by 2007/47/EC or 98/79/EC. A safety officer has been appointed for Germany and therefore is in full compliance with § 31 MPG.

Signed on 20 June 2019

Werner Sander
President

Certificate of CE-Registration



mdiEuropa

This is to certify that, in accordance with either medical device Directive 93/42/EEC as amended by 2007/47/EC or Directive 98/79/EC, mdi Europa GmbH agree to perform all duties and responsibilities as the Authorized Representative for

Starkling
Gartenstraße 11
D-78573 Wurmlingen
Germany

as stipulated and demanded by the afore-mentioned Directives. The German competent authorities have allocated the medical devices of the manufacturer the following registration numbers:

| UMDNS Code | Description | Classification | Registration Number |
|------------|--------------------------------------|----------------|---------------------|
| 15560 | Bougies | I | DE/CA09/0760/1587 |
| 15572 | Cup | I | DE/CA09/0760/1625 |
| 15573 | Cutters | I | DE/CA09/0760/1624 |
| 15602 | Specula | I | DE/CA09/0760/1661 |
| 15609 | Trays | I | DE/CA09/0760/1589 |
| 15710 | Forzepps, Dental | I | DE/CA09/0760/1611 |
| 15915 | Furniture Patient Room | I | DE/CA09/0760/1610 |
| 16017 | Stools | I | DE/CA09/0760/1580 |
| 16179 | Amalgam Carriers | I | DE/CA09/0760/1600 |
| 16437 | Chairs Examination/Treatment | I | DE/CA09/0760/1634 |
| 16443 | Appliers Aneurysm Clip | I | DE/CA09/0760/1594 |
| 16482 | Explorers | I | DE/CA09/0760/1614 |
| 16585 | Syringe, Cartridges | I | DE/CA09/0760/1586 |
| 16664 | Dental Hand Instruments, Orthodontic | I | DE/CA09/0760/1620 |
| 16666 | Dental Hand Instruments Surgical | I | DE/CA09/0760/1621 |
| 16779 | Tubing, Suction | I | DE/CA09/0760/1593 |

The manufacturer has provided mdi Europa with all necessary documentation, together with an appropriate Declaration of Conformity confirming that the medical devices fulfill the essential requirements of either Directive 93/42/EEC as amended by 2007/47/EC or 98/79/EC. A safety officer has been appointed for Germany and therefore is in full compliance with § 31 MPG.

Signed on 20 June 2019

Werner Sander
President

Certificate of CE-Registration



This is to certify that, in accordance with either medical device Directive 93/42/EEC as amended by 2007/47/EC or Directive 98/79/EC, mdi Europa GmbH agree to perform all duties and responsibilities as the Authorized Representative for

Starkling
Gartenstraße 11
D-78573 Wurmlingen
Germany

as stipulated and demanded by the afore-mentioned Directives. The German competent authorities have allocated the medical devices of the manufacturer the following registration numbers:

| UMDNS Code | Description | Classification | Registration Number |
|------------|------------------------|----------------|---------------------|
| 16883 | Suction Tips | I | DE/CA09/0760/1591 |
| 17673 | Approximators | I | DE/CA09/0760/1585 |
| 17858 | Saw, Bone, Rhinoplasty | I | DE/CA09/0760/1651 |
| 17954 | Hand Drills Surgical | I | DE/CA09/0760/1606 |

The manufacturer has provided mdi Europa with all necessary documentation, together with an appropriate Declaration of Conformity confirming that the medical devices fulfill the essential requirements of either Directive 93/42/EEC as amended by 2007/47/EC or 98/79/EC. A safety officer has been appointed for Germany and therefore is in full compliance with § 31 MPG.

Signed on 20 June 2019

Werner Sander
President