Manufacturer (Name, legal form of organization, address): «Firmenname», «Strasse», «Ort»

Please list all medical devices with code number CE 0483 according to Directive 93/42/EEC on this spreadsheet.

| No. of Tech. Doc.[[1]](#footnote-2) | Product (Product Name/Product Code/Type) & intended use, manufacturing technology , sterilization method if applicable | Classification | Rule [[2]](#footnote-3) | optional  UMDNS / GMDN | In-house manu-facturing | Subcontracted manufacturing | OEM[[3]](#footnote-4) | Supplier (only for subcontracted manufacturing) |
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1. indication concerning structure of the technical documentation (per product, per product group) [↑](#footnote-ref-2)
2. 2 detailed information on applied classification rule incl. indent / paragraph [↑](#footnote-ref-3)
3. 3 Products placed on the market under your own brand, for which in the course of the conformity assessment procedure reference is made to the pre-existing certification of a device which does not or only slightly differ from the original manufacturer’s product [↑](#footnote-ref-4)