# Details of the company

|  |  |
| --- | --- |
|  | Main site (address for the certificate) |
| Company (name and legal entity) |  |
| Street |  |
| Postal code, place |  |
| Country |  |
| Homepage |  |
| E-mail address (company) |  |
| Phone (switchboard) |  |
| Fax (company) |  |
| SRN (Single Registration Number) – if known |  |
| Company VAT (TVA) ID number |  |

# Main contact person

|  |  |
| --- | --- |
| Title | Mr.  Ms.  Dr |
| First and last name |  |
| Position |  |
| Direct/mobile phone |  |
| Personal e-mail address |  |

# If the company is based in Switzerland or outside the EU (except other EFTA countries)

|  |  |  |  |
| --- | --- | --- | --- |
|  | **Address of the authorized representative based in the EU** | | |
| Company name |  | Legal Entity |  |
| Street |  | Country |  |
| Postal code |  | Place |  |
| Salutation | Mr.  Ms. | Title |  |
| First and last name |  | Position |  |
| Direct/mobile phone |  | E-mail address |  |
| The national registration number of the EU representative: | |  | |

# External consulting companies

|  |  |  |
| --- | --- | --- |
| Were you assisted in implementation or maintenance of the quality system by an external consulting company? | Yes | No |
| If yes, please provide all involved consulting companies (company name, location) | | |
|  | | |

# Short description of the company’s activities and branch:

|  |
| --- |
|  |

# Area of aimed certification

|  |  |  |
| --- | --- | --- |
|  | the whole company | the following particular parts: |

# Aimed scope of certification

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
|  | Regulation/Standard | | | Accreditation/Acceptance |
|  | Regulation (EU) 2017/745 – Medical Devices:  Annex IX, Chapter I  Annex IX, Chapter II  Annex XI, Part A  *(Please also complete the „List of Medical Devices (MDR)“ to be attached as Annex I. You will find it in the section* [*”Quotation for certification*](https://www.mdc-ce.de/downloads/mdc-documents/quotation-for-certification.html)*” of our website).* | | | ZLG[[1]](#footnote-2) |
|  | EN ISO 13485 | | | DAkkS[[2]](#footnote-3) |
| Desired scope of the certificate(s): [Activity, e.g. development, manufacturing and distribution, maintenance, etc.] + [manufactured medical devices] | |  |
|  | ISO 13485 with additional acceptance in Taiwan | | under the TCP[[3]](#footnote-4) |
|  | ISO 13485 MDSAP with additional acceptance in Australia/Brazil/Canada/Japan/USA  (*Please do also complete the questionnaire „Basic Data of MDSAP“ which you will find in the section „*[*Quotation for certification*](https://www.mdc-ce.de/downloads/mdc-documents/quotation-for-certification.html)*“ of our website.*) | | in co-operation with DQS[[4]](#footnote-5) |
|  | EN ISO 9001 | | | DAkkS2 |
| Desired scope of the certificate(s):  [Activity, e.g. development, manufacturing and distribution, etc.] + [(medical) devices] | |  |

# Information on your existing quality management system

|  |  |  |
| --- | --- | --- |
| Is your quality system certified?  *(If yes, please add copies of your certificates)* | Yes | No |
| If yes, by which Notified Body/registrar? |  | |
| If yes, when does the validity of your certificate(s) end? |  | |

# Information on the time schedule

|  |  |
| --- | --- |
| Until when can the technical documentations listed in Annex I be submitted for assessment? |  |
| When is your QM system ready for audit? *Note: before performing the audit according to Regulation (EU) 2017/745, an assessment of the Technical Documentation must be performed.* |  |
| Is there a deadline by which the audit should be completed? |  |

# Information on the language

|  |  |  |
| --- | --- | --- |
| In which language should the **audit** be conducted? | German | English |
| In which language is the **technical documentation** available? | German | English |
| In which language is the **QM documentation** available? | German | English |

# Subsidiaries, branch offices, production sites and further sites which are covered by the quality system *(Please attach an annex if there are more than 5 sites)*

|  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- |
| Location | Address and legal entity | Number of Employees (no multiple assignment) | | | | | | Shift work |
| Fulltime | Part time | External sales | Marginally employed staff | Temporary workers  (permanent + non-permanent) | Trainees |
| 1 |  |  |  |  |  |  |  |  |
| 2 |  |  |  |  |  |  |  |  |
| 3 |  |  |  |  |  |  |  |  |
| 4 |  |  |  |  |  |  |  |  |
| 5 |  |  |  |  |  |  |  |  |
|  | Total |  |  |  |  |  |  |  |

# Distribution of employees in organizational areas *(Please attach an annex if there are more than 5 sites)*

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
| **Business area** *(Please add an organization chart of your company)* | Location | | | | |
| 1 | 2 | 3 | 4 | 5 |
| Design and development |  |  |  |  |  |
| Manufacturing / Production and warehouse |  |  |  |  |  |
| Repair or reprocessing of medical devices |  |  |  |  |  |
| Quality management / Quality control / Regulatory affairs |  |  |  |  |  |
| Marketing, distribution and field service |  |  |  |  |  |
| Administration and other |  |  |  |  |  |
| Total |  |  |  |  |  |

# Details on shift work *(If applicable)*

|  |  |  |
| --- | --- | --- |
| Are there any processes that cannot be audited during the regular audit period (9 am - 6 pm)? | Yes | No |
| If so, which processes/shifts are affected? | | |
|  | | |

# Details on outsourced activities

|  |  |  |
| --- | --- | --- |
| Are certain activities outsourced to external parties? *(e.g. part production)* | Yes | No |
| If yes, which activities are outsourced? | | |
|  | | |

|  |
| --- |
| ***NOTES ON DATA PROTECTION:*** *The processing of the personal data communicated in this questionnaire takes place at the request of the signatory/the submitting person for the purpose of pre-contractual measures (preparation of offers) and, if necessary, for the execution of a coming contract according to Art. 6 Para. 1 lit. b GDPR. For any further use of personal data, the consent of the data subject is regularly required. You can give such consent in the following section. Further information on data protection can be found in our data protection declaration online at* [*https://www.mdc-ce.de/privacy.html*](https://www.mdc-ce.de/privacy.html) *and in our document "Data protection information for customers and interested parties".* |
| **Consent to the use of personal data for purposes of information about services and news from the certification environment**  (If you agree with the purpose of use, please tick this accordingly. If you do not wish to give your consent, tick the field „no“. If you leave the field blank, a previously given consent remains valid).  Please inform me, independent of the certification procedure, about other aspects and news about the certification environment for medical devices, in-vitro diagnostics and health care as well as about your seminars on the subject:  Yes. gladly by post  Yes, gladly by e-mail (please include your e-mail address)  No, currently I have no interest  In accordance with data protection legislation, you have the right to information, correction, restriction and deletion of your personal data. In addition, you have the right to revoke your consent at any time and without giving reasons. If you wish to exercise any of these rights or have any further questions about data protection, please send an e-mail to [*datenschutz@mdc-ce.de*](mailto:datenschutz@mdc-ce.de) |

|  |  |  |
| --- | --- | --- |
|  |  |  |
| Place, Date |  | Signature or full name in case of electronic transmission |

**ANNEX I**

Please attach to the completed questionnaire the duly completed „List of medical devices (MDR)“ as Annex I. You will find it in the section [”Quotation for certification](https://www.mdc-ce.de/downloads/mdc-documents/quotation-for-certification.html)” on our website.

Please also enclose detailed product information such as instruction(s) for use, detailed product description(s), additional item list(s) etc. with the completed questionnaire.

If it is a recertification procedure, a "Summary of changes and scientific findings for the device" shall also be submitted in accordance with Annex VII, 4.11.

**ANNEX II**

Manufacturers of class Ir devices are also requested to attach the completed „Annex II – Questionnaire for a quotation (medical devices class Ir)”. You will find it in the section [”Quotation for certification](https://www.mdc-ce.de/downloads/mdc-documents/quotation-for-certification.html)” on our website.

1. Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten [↑](#footnote-ref-2)
2. DAkkS – Deutsche Akkreditierungsstelle GmbH [↑](#footnote-ref-3)
3. Technical Cooperation Programme on Exchange of Medical Device GMP and ISO 13485 Audit Reports [↑](#footnote-ref-4)
4. DQS Medizinprodukte GmbH [↑](#footnote-ref-5)