

1. Information on ACCOUNTS	
1.1. Applicant	
Registered company name	_____
Address	_____
ZIP Code/City, Country	_____
Contact person	_____
Phone _____ e-mail _____	
1.2. Payer <input type="checkbox"/> Same as in 1.1. Applicant or enter below	
Registered company name	_____
Address	_____
ZIP Code/City, Country	_____
VAT identification number _____ VAT payer: <input type="checkbox"/> Yes <input type="checkbox"/> No	
Contact person	_____
Phone: _____ e-mail _____	
1.3. Document Holder <input type="checkbox"/> Same as in 1.1. Applicant or enter below <i>Applicant on issued documents</i>	
Registered company name	_____
Address	_____
ZIP Code/City, Country	_____
1.4. Document Recipient <input type="checkbox"/> Same as in 1.1. Applicant or enter below	
Registered company name	_____
Address	_____
ZIP Code/City, Country	_____
Contact person _____ e-mail _____	
1.5. Manufacturer <input type="checkbox"/> Same as in 1.1. Applicant or enter below	
Registered company name	_____
Address	_____
ZIP Code/City, Country	_____
2. Information on the PRODUCT	
2.1. Product	
Product name	_____
Model / type	_____
Brand name(s)	_____
Other data	_____
2.2. Places(s) of manufacture <i>Full address (Name, address, ZIP/City, Country)</i>	
1) <input type="checkbox"/> Same as in 1.5. Manufacturer	
2) _____	
3) _____	
4) _____	

3. Requested SERVICES

According to the offer _____

(offer number, date)

Fill-in the following fields if no offer was sent to you or you wish to order additional services.

3.1. Testing according to listed standards

LVD (Low Voltage Equipment)

- electric safety
- fast evaluation (a check of basic requirements)
- EMF (electromagnetic radiation)

EMC (Electromagnetic Compatibility)

- EMC
- fast evaluation (a check of basic requirements)

MD (Machinery Safety)

- machinery safety
- fast evaluation (a check of basic requirements)
- noise
- vibrations

ENV (Environmental impacts)

- vibrations
- IP
- climatic

MDD (Electrical Medical Devices)

CPR (Construction Products)

RED (Radio Equipment)

RoHS (XRF method)

According to the following standards/requirements:

3.2. Homologation

- Approval of vehicles with regard to electromagnetic compatibility according to **ECE R10**
- E-mark** (Assistance in homologation)

3.8. Assistance in acquiring foreign certificates

National approvals and other certification marks required to sell the product on individual global markets (state the country, regulator...)

- _____
- _____
- _____
- _____
- _____

3.3. Certificates

- IECEE CB** safety EMC
- IECEE CB-FCS**

Including the testing of national deviations for the following countries (mark the relevant countries for IECEE CB and IECEE CB-FCS certification mark):

- AR AT AU BE BR CA CH CN CZ DE
- DK ES FI FR GB GR HU IE IL IN
- JP KR MY MEX NL NO PL PT RS RU
- SE SG SI SK THAI TR UA US ZA

Other(s): _____

- SIQ** safety EMC

3.4. Licences

Issue of a licence for use of a mark of conformity, subject to regular inspections of a manufacturing process.

- ENEC** **CCA NTR** **CCA EMC**

Including testing of national deviations for the following countries (mark the relevant countries):

- AT BE CH DE DK ES FI FR GB GR
- HU IE CZ TR IT LU NL NO PT SE
- SI PL SK

Other(s): _____

3.5. Licences for use of SIQ marks

Issue of a licence for use of an SIQ mark of conformity, subject to regular inspections of a manufacturing process.

- SIQ**
- SIQ Bauart Geprüft / Type Approved**
- SIQ Medizin Bauart Geprüft / Medical Type Approved**

3.6. Certificates of conformity to requirements for CE marking

SIQ as an EU notified body no. 1304

- EC type examination MD (2006/42/EC)
- EC type examination NOISE (2000/14/EC)
- EC type examination RED (2014/53/EU)
- EC certificate of constancy of performance CPR (305/2011 EU Regulation)

3.7. Permission for use of SIQ marks

- SIQ Checked**

PRODUCT TESTING AND CERTIFICATION

4. Handling with SAMPLES
<p>After completion of the project and receipt of the notification, the applicant shall collect the samples delivered to SIQ. If the applicant decides to collect the samples but fails to do so within the period specified, SIQ may charge demurrage according to the valid pricelist. SIQ shall keep the samples for the period of maximum six months after completion of the project. After that period, SIQ shall destroy the samples against payment in the presence of a commission.</p> <p>The applicant agrees that after completion of the project samples shall be:</p> <p><input type="checkbox"/> destroyed in the presence of a commission</p> <p><input type="checkbox"/> stored</p> <p><input type="checkbox"/> returned to the applicant</p> <p><input type="checkbox"/> returned to the applicant via courier _____, account number _____</p>
5. Delivery of DOCUMENTS
<p>After the completion of the service and against payment, all the issued documents are sent in electronic form to the document recipient by email or at SIQ portal (https://portal.siq.si). Other means of delivery:</p> <p><input type="checkbox"/> printed documents by post (free of charge)</p> <p><input type="checkbox"/> printed documents via DHL (against payment)</p> <p><input type="checkbox"/> printed documents via courier _____, account number _____</p>
6. SUBCONTRACTING of the testing
<p>The applicant agrees that a part of testing will be carried out by one or more of the following subcontractors:</p> <p><input type="checkbox"/> SIQ Testing and Certification GmbH, Angerstraße 11, D-86807 Buchloe, Germany <i>(For the scope of the laboratory see table 23 in LP-009 published on http://www.slo-akreditacija.si/accreditation/slovenski-institut-za-kakovost-in-meroslovje-2/)</i></p> <p><input type="checkbox"/> SIQ d.o.o. Beograd, Cara Dusana 266, RS-11080 Beograd - Zemun, Serbia <i>(For the scope of the laboratory see http://www.registar.ats.rs/predmet/978/)</i></p> <p><input type="checkbox"/> The subcontractor that is stated in the offer for the provision of specific testing.</p> <p><input type="checkbox"/> _____</p>
7. Remarks
<p>Document holder, payer, document recipient, place(s) of manufacture, details of the ordered service, scope of testing, details of requested testing or certification service(s), e.g. extension of validity of documents, differences from the earlier tested/certified product, decision rule to be applied in recording test results and stating conformity with a specified requirement, any other remarks.</p> <hr/> <hr/> <hr/> <hr/> <hr/> <hr/> <hr/> <hr/> <hr/> <hr/> <hr/> <hr/> <hr/> <hr/> <hr/> <hr/>

