# APPLICATION FOR PRODUCT CONFORMITY ASSESSMENT OF PRESSURE EQUIPMENT

**According to directive 97/23/EC of the european parliament and of the council as amended.**

**No.:** | | | | | | | | | |

TIC's Registry Number (do not fill in)

## APPLICANT:

| | | |
|---|---|---|
| **Company name:** | VAT No.: | |
| | Registration No.: | |
| | Registered at: | |
| Address: | Phone: | |
| | Fax: | |
| PIN Code: | e-mail: | @ |
| Banking with: | Account No.: | |
| Represented by: | | |
| Contact person authorized to conduct negotiations: | | |

## PRODUCT(s):

| |
|---|
| **Product name:**<br>Type(use page 2 of this Application form for more detailed specifications): |
| Company name and manufacturer's address (if different from that of the applicant): |

## ACCOMPANYING DOCUMENTATION:

| | |
|---|---|
| Instructions for use and assembly: | |
| Comprehensive manufacturing and design data | |
| Harmonized standards and other technical specifications used (partially or completely) in designing of the product: | |
| Technical conditions (product and manufacture characteristics, drawings, wiring diagrams, risk analysis, information for user, marking, illustrations, etc.) | |
| Test reports of accredited laboratories | |
| Other documents and information: | |

**Applicant's declaration:**

1. *We agree that product samples will not be returned after assessment.*
2. *We agree with the "Obligatory Business Conditions" of TIC and we will abide by them.*

.....
Date of Application

.....
Stamp and signature of applicant's authorized representative

3. **The Customer applies for the following activities from the Approved Body** *the activities chosen according to the pressure equipment category - Annex II of the Directive shall be marked with* ***cross****.*

| | | |
|---|---|---|
| 3.1 ☐ | **Monitoring of the final assessment** | according to Annex III of the Directive - Module A1 followed by issue of the check report ….. |
| 3.2 ☐ | **EC type-examination** | according to Annex III of the Directive - Module B Point 4 followed by issue of the EC type-examination certificate |
| 3.3 ☐ | **EC design-examination** | according to Annex III of the Directive Module B1 Point 4 followed by issue of the EC design-examination certificate |
| 3.4 ☐ | **Assessment of conformity to type** | according to Annex III of the Directive - Module C1 Point 4 (unexpected visits followed by issue of the record) |
| 3.5 ☐ | **Production quality assurance** | after EC type/design-examination according to Annex III of the Directive - Module D Points 3 and 4 (assessment of the quality system and surveillance followed by issue of the report) |
| 3.6 ☐ | **Production quality assurance** | according to Annex III of the Directive - Module D1 Points 4 and 5 (assessment of the quality system and surveillance followed by issue of the report) |
| 3.7 ☐ | **Product quality assurance** | after EC type-examination according to Annex III of the Directive -Module E Points 3 and 4 (assessment of the quality system and surveillance followed by issue of the report). |
| 3.8 ☐ | **Product quality assurance** | According to Annex III of the Directive - Module E1 Points 4 and 5 (assessment of the quality system and surveillance followed by issue of the report) |
| 3.9 ☐ | **Product verification** | after EC type/design-examination according to Annex III of the Directive -Module F Point 4 (final inspection and pressure test followed by issue of the certificate of Conformity) |
| 3.10 ☐ | **EC unit verification** | According to Annex III of the Directive - Module G Point 4 (examination of the design and testing followed by issue of the certificate of conformity) |
| 3.11 ☐ | **Full quality assurance** | according to Annex III of the Directive - Module H Points 3 and 4 (assessment of the quality system and surveillance followed by issue of the report) |
| 3.12 ☐ | **Full quality assurance with design-examination and special surveillance of the final Assessment** | according to Annex III of the Directive - Module H1 (design-examination followed by issue of the EC design-examination certificate, assessment of the quality system and surveillance followed by issue of the report) |
| 3.13 ☐ | **European approval for materials** | According to Article 11 of the Directive (inspections and tests followed by issue of the European approval for materials) |

## Detailed specification of the Pressure Equipment submitted to conformity assessment*:*

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|---|---|
| Name and description of the product: | |
| List of documentation enclosed (*figures, drawings, instruction for use & installation, technical specification etc.*) as specified in 97/23/EC directive as amended: | |
| Intended use of the product (*detailed description*): | |
| Has been this Pressure Equipment placed on the market already?<br>If yes, when has been the Pressure Equipment introduced on the market:<br>If yes, which countries have been the Pressure Equipment placed on the market in? | ☐ Yes ☐ No<br>In year |
| Is the device designed and manufactured in a controlled quality system?<br>If yes, which standards does comply the quality system with? | ☐ Yes ☐ No<br>ISO 9001:2008 |
| Is the quality system certified? | ☐ Yes ☐ No |
| Are the test reports of the accredited testing laboratories available?<br>If yes, specify the test reports in list: | ☐ Yes ☐ No |
| List of documents concerning to manufacturer's quality system (*certificates, audit reports, …*) | |
| Another documents (*information, …*): | |