

**Application no . .....**

For product assessment and for other services connected with conformity assessment of pressure equipment according to Directive 2014/68/EU of the European Parliament and of the Council ("Directive").

**1. APPLICANT**

Trade name:	
Address:	
Country:	
ID:	VAT No:
Registered in Companies Register (position, number):	
Statutory/Authorized representative: (name, position)	Contact person: (name, position)
Tel:	Tel:
E-mail:	E-mail:
Bank:	Account:

**2. PRODUCER** (do not fill in if the applicant is the producer)

Trade name:	
Address:	
Country:	
ID:	VAT No:
Statutory/Authorized representative: (name, position)	
Tel:	Contact person: (name, position)
Tel:	Tel:
E-mail:	E-mail:

**3. PRODUCT**

Product trade name:	
Type:	
Derived alternatives:	
<input type="checkbox"/> Repetitive production	<input type="checkbox"/> Custom-made in the number of _____ piece(s)

**4. THE CUSTOMER ASKS THE NOTIFIED BODY TO REALIZE FOLLOWING ACTIVITIES:**

- Internal production control plus supervised pressure equipment checks at random intervals - Annex III, point 2, (module A2).
- EU-Type examination – production type - Annex III, point 3.1, (module B).
- EU-Type examination – design type - Annex III, point 3.2, (module B).
- Conformity to type based on internal production control plus supervised pressure equipment checks at random intervals - Annex III, point 4, (module C2).
- Conformity to type based on quality assurance of the production process- Annex III, point 5, (module D).
- Quality assurance of the production process - Annex III, point 6, (module D1).
- Conformity to type based on pressure equipment quality assurance - Annex III, point 7, (module E).
- Quality assurance of final pressure equipment inspection and testing - Annex III, point 8, (module E1).
- Conformity based on unit verification - Annex III, point 10, (module G).
- Conformity based on full quality assurance - Annex III, point 11, (module H).
- Conformity based on full quality assurance plus design examination - Annex III, point 12, (module H1).

**5. THE APPLICANT ASKS TSÚ TO REALIZE FOLLOWING ACTIVITIES:**  
(outside the scope of the Notified Body activities)

- Other (specify)  
 Assistance with drawing up of the declaration of conformity.

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**Issue of test reports; language:**  
 Slovak                       English                      Bilingual:  Slovak – English

**Issue of certificates; language:**  
 Slovak                       English                      Bilingual:  Slovak – English  
 English - Russian

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**6. ACCOMPANYING DOCUMENTS**

- Technical documentation including analysis and assessment of the risk(s)- Annex no.  
*in accordance with Annex III of the Directive*
- Brief description of manufacture technology Annex no.
- Test reports, technical reports, etc. (if available) Annex no.
- Management System Certificate and last Audit report Annex no.  
*(if Management System has been certified)*
- Authorization for the Applicant containing the scope of powers given by the producer Annex no.  
*(if the application is not submitted by the producer)*

**7. DECLARATION OF APPLICANT**

The development of the product as a type is finished and all data and technical documentation presented in this conformity assessment application are complete and they represent product state on the date of the submission of this application. We hereby declare that we have not asked any other notified body for conformity assessment of the product.

**8. DUTIES OF APPLICANT**

Submit documents needed for conformity assessment in Slovak or English language as stated in point 6. Enable sampling or submit the product sample so that the conformity assessment can be completed in a given time. Provide the cooperation during the conformity assessment in the scope required by the Notified Body.

**Remarks to the filling of application:**

*The application is filled in individually for each product type; in case of various product types, these are given in annex of this application. Supporting documents in accordance with part 6, dealing with various products needed for product conformity assessment could be added to only one application and they can be referred to in other applications. Send only one exemplar of the application and supporting documents. All materials shall be sent by e-mail or by registered post at the address presented in the head of the application form. If you need more space than is given in this application, use a special annex.*

In \_\_\_\_\_ date

**Statutory/Authorized representative (name and signature):**

**Stamp**