



## Application for assessment of performances of construction product

No.....  
(to be filled in by notified or authorized body)

Applicant/manufacturer (acc. to art. 1 or 2) applies for: (please tick the box in accordance with the technical specification)

- certification of constancy of performance of construction product – system 1+ (I+), 1 (I)**
- certification of conformity of factory production control – system 2+ (II+)**
- carrying out testing for assessment of performance – system 3 (III)**

pursuant to the legal act: (please tick one of the boxes)

- Regulation (EU) No. 305/2011**, of 9 March 2011 laying down harmonised conditions for the marketing of construction products and repealing Council Directive 89/106/EEC (hereinafter referred to as „CPR“)
- Act No. 133/2013 Coll.** on construction products and amendments of certain acts as amended (hereinafter referred to as „Act“)

Note: For more information, please refer to [http://www.tsus.sk/sluzby/certifikacia\\_sv\\_en.php](http://www.tsus.sk/sluzby/certifikacia_sv_en.php)

### 1 Applicant:

Trade name (acc. to the entry in the register)

Registered address:

Mailing address: (enter if different from the reg. address)

ID No.:

VAT No.:

Registration (register office and No.)

Statutory body (name, position, phone, e-mail)

Authorized employee (name, position, phone, e-mail)

Bank details:            office of the Bank:  
   IBAN:  
   BIC/SWIFT:

### 2 Manufacturer: (to be filled in when the applicant is not being the manufacturer)

Trade name (acc. to the entry in the register)

Address

ID No.:

VAT No.:

Registration (register office and No.)

Statutory body *(name, position, phone, fax, e-mail)*

Authorized employee *(name, position, phone, fax, e-mail)*

**Manufacturing plant** *(name if any, address):*

### 3 Construction product

Generic name:

Trade name:

Type(s) *(detailed identification of all its types and variants under current standards, corporate documentation, catalogue and so on., produced by the same technology):*

The product of this type is produced in \_\_\_\_\_ versions (variants).

Brands of the product covered by the tested type:

**Technical specification** *(harmonized European standard or ETA, Slovak designated standard or SK technical assessment):*

#### No. of group of products

*(acc. to Annex 1 of the Decree MDVRR SR No. 162/2013 Coll. as amended):*

**Intended use of the product in works, a way of permanent and fixed installations in works and any restriction on its use if relevant:**

#### Supplementary data *(if relevant)*

Reaction to fire class:

*(acc. to EN 13501-1:2018)*

Reaction to external fire class if relevant:

*(acc. to EN 13501-5:2016)*

Is the product made of materials for which a clearly identifiable stage in the production process results in an improvement of the reaction to fire classification (e.g. an addition of fire retardants or a limiting of organic material)?

**yes**       **no**

### 4 The product will be ready for assessment of performances starting from:

### 5 Was already the assessment of performances of the product made acc. to the CPR or the Act?

**yes** *(enter certificate No. or No. of report on testing):*

**yes**, and we apply for an amendment of the certificate *(enter certificate No. or No. of test report):*

**no**

**6 Enclosed auxiliary documents**

Information on the product (tick appropriate box(es)):

- detailed technical description and drawings, data sheets, structural analysis, calculation of thermal performance and similar if relevant;
- data on intended use;
- brief description of production process;
- data on factory production control – especially about its organization, staff and technical equipment;
- copies of reports of performed tests, certificates, audit reports and similar having some relevance to the product.

Annex No.:

A copy of the certificate of incorporation or business license current at the time of application (notarization not required) Annex No.:

When the applicant is not the manufacturer, the original or a duly certified copy of the manufacturer's authorization for the applicant indicating the extent of representation is to be enclosed. Annex No.:

**7 Declaration of the applicant/manufacturer**

- The product is developmentally completed, and data contained in this application, the documentation and other data are complete and represent the state of the product at the date of this application.
- We have not applied for the assessment of performances of the product referred to in Art. 3 to other Notified body or Authorized body, or if we did, the application has not been accepted.

**8 Additional requirements for the applicant/manufacturer**

- Submit documents needed to assess the performances of the product in Slovak language (unless otherwise agreed).
- Provide for the taking a sample, resp. provide a sample of the product for determination of product-type by testing in due time.
- Enable the performance of initial inspection of the factory and factory production control in due time.
- Provide cooperation during the assessment of performances to the extent required by the approved body.

**9 Commercial and legal relationships**

- Will be addressed in a separate contract concluded pursuant to § 591 et seq. Commercial Code, in relation to this application.

In \_\_\_\_\_ on \_\_\_\_\_

Authorized representative of the applicant

On behalf of notified or authorized body registered by:

name:

name: \_\_\_\_\_ on \_\_\_\_\_

signature: .....

signature: .....

stamp:

**Notes to fill in the application:**

*The application is filled in separately for each type of product manufactured in a Member State or abroad and for each manufacturing plant.*

*Those auxiliary documents under article 6 concerning several products applied for assessing their performance, should be attached just to only one application and in the other corresponding applications they should be just referred to.*

*The application shall be sent, together with auxiliary documents to the address indicated in the application form header. When the application is sent electronically, it must contain a scanned signature and stamp of an authorized representative of the applicant.*

**Notice:**

*Please note that the registration of the application does not confirm its integrity. The process of assessing the parameters of the product can be launched only after completion of the prescribed data and delivery of all required supporting materials acc. to article 21 para. 2 of the Act.*

**Information for the applicant/manufacture – voluntary certification, or testing**

*The assessment and verification of constancy of parameters according to the CPR or the Act covers only those (mandatory) characteristics (or at least one characteristic of them) having a relevance to the basic requirements for works and therefore referred to in the technical specifications. However, from a marketing viewpoint it is recommended to have also a voluntary certificate or at least test reports issued by an independent third party (accredited certification body or test laboratory) covering parameters of additional characteristics that might attract customers' interest in your product as opposed to your competitors having only the mandatory characteristics certified.*

*It is profitable to carry out at the same time the assessment of parameters of mandatory characteristics and of voluntary ones resulting in the issue of both certificates – mandatory and voluntary, the latter issued by the certification body for certification of products TSUS. By a less demanding procedure it is possible to obtain from the accredited TSUS test laboratory just test reports of the non-mandatory characteristics.*

*If you are interested, please indicate which non-mandatory characteristics of the product you would like to have certified or just tested.*

*The voluntary certification or testing will be carried out based on a specific contract. Detailed information are available at the address indicated in the application form header or at <http://www.tsus.sk/>.*