Notified Body 2369

Application for EU Type Examination and Conformity assessment of Personal Protective Equipment according to Regulation (EU) 2016/425

Fill out CB	Review date:	Registration date:	Registration number:

1. Applicant

Name of company: .....

Address: .....

VAT No: ..... IBAN: ..... SWIFT: .....

Banking connection

Name and address of the bank: ..... account number: .....

Authorized person	Contact person
Name:	Name: .....
Position:	Position: .....
Telephone:	Telephone: .....
E – mail:	E – mail: .....
Website:	

2. Product

Product name: .....

Type of product (Model): .....

Type series: .....

.....

Manufacturer (if different from applicant): .....

.....

(Name, Address, Country)

Place of manufacturing (if different from applicant):

.....

.....

(Address)

3. The applicant asks the notified body 2369 to realize following activities:

☐ **EU-Type examination** - Annex V Regulation (Module B) *

☐ **Conformity to type based on internal production control plus supervised product checks at random intervals**- Annex VII (Module C2)

4. Technical requirements

Technical requirements (technical regulation or standard) under which conformity of product type samples is to be assessed:

.....

Note: * For category III PPE is required to choose module C2 together with the module B. In a separate module B is impossible to affix the CE marking with Notified Body Number for the product/s.

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5. Accompanying documentation for the application

Specification	Document ID and Date
Product specifications and product variants with photo	
Material sheets of materials used on the product with manufacturer's indication	
Test results – test reports and final reports	
Technical documentation as required by Annex III Regulation	

6. Declaration of the 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 and have been the Testing and Certification Regulation recognized and acknowledged.

After obtaining the certificate, the applicant undertakes:

- always to fulfil the certification requirements, including implementing appropriate changes when they are communicated by the certification body,
- if the certification applies to ongoing production, the certified product continues to fulfil the product requirements,
- to make all necessary arrangements for:
  1) the conduct of the evaluation and surveillance (if required), including provision for examining documentation and records, and access to relevant equipment, location(s), area (s), personnel, and applicant's subcontractors,
  2) investigate of complaints,
  3) the participation of observers, if applicable,
- to make claims regarding certification consistent with the scope of certification,
- do not to use its product certification in such a manner as to bring the certification body into disrepute and do not to make any statement regarding its product certification that the certification body may consider misleading or unauthorized,
- upon suspension, witdrawal, or termination of certification do not to use of all advertising matter that contains any reference thereto and to take action as required by the certification scheme (e.g. the return of certification documents) and to take any other required measure,
- to provide copies of the certification documents to others, the documents shall be reproduced in their entirety or specified in the certification scheme,
- in making reference to its product certification in communications media such as documents, brochures or advertising, to comply with the requirements of the certification body or as specified by the certification scheme,
- to comply with any requirements that may be prescribed in thecertification scheme relating to the use of marks of conformity, and on information related to the product,
- to keep records of all complaints made known to it relating to the compliance with the certification requirements and to make these records avaible to the certification body when requested, and to take appropriate action with respect to such complaints and any deficiencies found in products that affect compliance with requirements for certification and to document the actions taken,
- to inform the certification body, without delay, of changes that may affect its ability to conform with the certification requirements.

7. Duties of applicant

Submit documents needed for conformity assessment in Slovak or English language as stated in point 5. 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.

8. Applicant's consent *

The applicant has been notified and agrees to ensure that the tests are carried out in a subcontracting manner:

yes **no**

9. Trade-legal relations

An application initialed by both parties for certification activities is considered legally enforceable and takes into account the responsibility of the certifikation body and its client.

……..……

Place, date

…… ……

Name authorized person **Signature and Stamp**

Fill out CB

Registration number and date of receipt of the sample:	**Amount of sample:**

Note**: *** choose one of the options

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