VUTCH s.r.o.
Notified Body 3020
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A P P L I C A T I O N

☐ **For execution of EU type examination, conformity assessment of product type sample and issue of EU type examination certificate according to REGULATION OF THE EUROPEAN PARLIAMENT AND COUNCIL (EU) 2016/425 on personal protective equipment** (EU type examination - module B)

☐ **For execution of EC type examination, conformity assessment of product type sample and issue of EC type examination certificate according to DIRECTIVE OF THE EUROPEAN PARLIAMENT AND COUNCIL 2009/48/EC on the safety of toys as amended by later relevant documents** (EC type examination)

To be completed by VUTCH s.r.o., Notified Body 3020 Registered on: Registration No.:

1. Applicant

Name (company):
Home office of the company (address):

Identification number of the organization:
VAT identification number:

Bank connection - Bank name and address:

Account number (IBAN):

1.1 Represented by (statutory representative)

Name:
Position:

Phone:
E-mail:

1.2 Person authorized for negotiation

Name:
Position:

Phone:
E-mail:

2. Manufacturer

Name (company):

Address (place, country):

3. Product data

3.1 Product (name, type):

3.2 Trade name of the product (trademark):

3.3 Production classification code (KP):

3.4 Customs tariff item (informative entry):

3.5 Material composition, colour:

3.6 Standards and/or other normative documents according to which the product is to be certified:

4. Supporting documentation

4.1 Document about legal subjectivity of the applicant (excerpt from the Trade Register, trade licence) enclosure No.:

4.2 Test reports, attestations, certificates, technical reports enclosure No.:

4.3 Quality assurance system certificates enclosure No.:

4.4 Technical documentation - other documents about product safety (risk analysis, safety data sheets of used chemical substances obtained from suppliers, instructions for use, information provided by the manufacturer, etc.) enclosure No.:

5. Declaration of the manufacturer / applicant

As a type, the product is developmentally completed and the data provided in this application, the submitted technical documentation as well as other data are complete and describe the state of the product as of the date of submission of this application.
I did not ask another Notified Body to perform the EU* / EC* type test / issue the EU* / EC* type examination certificate.

5.1 The product is prepared for purchase from (date):

Notice: * delete where not applicable

6. Obligations of the manufacturer, importer:

- supply the necessary technical documentation (instructions for use, technical description of the product, drawings, diagrams, pictures, etc.) in accordance with the relevant regulation, or guidelines.
- enable the collection of samples, or take samples of the product type according to STN EN 12751, STN 01 5110, STN EN ISO 5089 and deliver the samples to Notified Body in the required time. Notified Body reserves the right to check the supplied samples of the product type and, in case of incorrect selection (collection), take them directly from the manufacturer (importer).
- to cooperate with the Notified Body on testing and certification of the product type.

7. Business and legal relations

Business and legal relations will be settled in an independent agreement, concluded according to § 591 and subsequent paragraphs of the Trade Code, linking up to this application.
The Notified Body will start to carry out the ordered works only after conclusion of the agreement and remittance of a pro-forma invoice.

8. Comments on completion of the application

A separate application shall be completed for each product type manufactured in the home country or imported on the territory of the Slovak Republic for each manufacturer and each manufacturing country.
The application and supporting documentation must be submitted in Slovak language or in English.

The delivered quantity, variety of colours and kinds of product type samples represent exhaustively the required product type.
The applicant agrees to the certification requirements and supplies any information necessary to evaluate the products to be certified.
The applicant must submit a new application in the case of significant changes after acceptance of this application by the Notified Body.

I r e q u i r e* – I d o n o t r e q u i r e* to return sample rests after testing.

In ....., on ....
Authorized Representative (name) Signature, stamp

Taken over by: Number and kind of samples:

Date:

Notice: * delete where not applicable