**APPLICATION FOR CONFORMITY TO TYPE BASED ON INTERNAL PRODUCTION CONTROL PLUS SUPERVISED PRODUCT CHECKS AT RANDOM INTERVALS (Module C2)**

**ANNEX VII Clause 3. REGULATION (EU) 2016/425**

|  |  |  |  |
| --- | --- | --- | --- |
| Applicant name | Click or click here to write text | | |
| Manufacturer name | Elija un elemento. | | |
| Address | Click or click here to write text | | |
| V.A.T.: | Click or click here to write text | | |
| Telephone: | Click or click here to write text | Fax: | Click or click here to write text |
| Contact e-mail/s: | Click or click here to write text | | |
|  | | | |
| Name of authorized representative in Europe | Click or click here to write text | | |
| Address | Click or click here to write text | | |
| VAT.: | Click or click here to write text | | |
| Telephone: | Click or click here to write text | Fax: | Click or click here to write text |
| Contact e-mail/s: | Click or click here to write text | | |
|  |  | | |
| As legal representative of the company | | | |
| Position: Click or click here to write text | | | |

**I request for product checks of PPE Type according to the Regulation (EU) 2016/425, described below:**

|  |  |
| --- | --- |
| Product Name / Reference | Click or click here to write text |
| PPE certificate No | Click or click here to write text |

**Based on conformity to type based on internal production control plus supervised product checks at random intervals (Module C2)**

|  |  |
| --- | --- |
| Applicable EN standards international standards for testing and other requiremts | Click or click here to write text |

Declares that:

* The same application has not been lodged with any other notified body.
* Knows and complies with [Regulation (EU) 2016/425](https://eur-lex.europa.eu/legal-content/EN/TXT/PDF/?uri=CELEX:32016R0425&from=EN) in its entire contents, as well as Chapter II, so that, together with this application, the technical documentation described in Annex III to the Regulation(1) is provided.
* An adequate statistical sample of the manufactured PPE shall be selected by the notified body at a place agreed between the body and the manufacturer.
* The manufacturer will bear the costs of the product checks, as testing and reports according to the offer or order.
* The manufacturer will keep informed the notified body about any modification of the delivery date of the production.
* I will inform the Notified Body in possession of the technical documentation relating to the EU-type examination certificate of all modifications of the approved type and of any modifications of the technical documentation which may affect the conformity of the PPE with the essential health and safety requirements or the conditions of validity of the certificate (such modifications will require additional approval in the form of a supplement to the original EU type-examination certificate).
* The Notified Body of AITEX ensures that it has an appeal procedure, whereby the applicant, upon request, can challenge the result of a conformity assessment.

(1) Together with this application, the following documents are provided:

|  |  |  |
| --- | --- | --- |
| **List of technical documentation** | Name of the document and date of issue | Indicate with X if provided |
| Technical Documentation (2). | Click or click here to write text |  |
| User instructions | Click or click here to write text |  |
| Labelling and other marking claimed in PPE. | Click or click here to write text |  |
| Drawings and schemes for the design and manufacture of PPE and its components, sub-assemblies and circuits. | Click or click here to write text |  |
| List of materials included in the PPE | Click or click here to write text |  |
| Copy of the PPE certificate (if notified body that performed module B is not the same as the one that is going to perform module C2) | Click or click here to write text |  |
| Other documents | Click or click here to write text |  |

(2) See document on technical documentation.

And understand and have been informed that:

* (3) According to Article 12 of [Regulation (EU) 2016/425](https://eur-lex.europa.eu/legal-content/EN/TXT/PDF/?uri=CELEX:32016R0425&from=EN): An importer or distributor shall be considered a manufacturer for the purposes of this Regulation and he shall be subject to the obligations of the manufacturer set out in Article 8 where he places PPE on the market under his name or trademark or modifies PPE already placed on the market in such a way that compliance with this Regulation may be affected.
* The certificate may be withdrawn if misuse is discovered. For example, failure to pay invoices for certification support testing, and / or conformity assessment, falsification of a certificate or misleading use of the certificate in advertising will be considered as an omission. The withdrawal together with the reason, will be delivered in writing.
* The Notified Body of Aitex is designated by the number 0161. The manufacturer can only refer to that number, in documents, that exclusively fit its content as evaluated by the Notified Body of Aitex.
* Only the certificates issued by the Notified Body of AITEX, containing the original handwritten signature or the electronic signature, issued by the Technical Direction will be recognized.
* Due to the duration of the tests, the tests will start to be carried out in parallel in all the laboratories, so that in case of not complying with the parameters and not being able to certify, the customer will pay all the tests performed so far.
* Otherwise, the customer must specify the order of execution of the tests, and therefore the delivery period will be extended according to the same.
* The signature of this application form constitutes the legally enforceable Certification Agreement, whereby the applicant with sufficient charge and authority to request the evaluation of the conformity and certification of their products, undertakes to comply with all obligations defined for manufacturers and / or economic agents in the framework of the activities carried out, defined in the current legislation of application (Regulation EU No. 2016/425 regarding personal protective equipment (PPE), as well as in point 4.1.2. ISO / IEC 17065:2012. The obligations on the part of the client and the Certification Body are available at the link: [https://www.aitex.es//personal-protection-equipment-certificate/?lang=en](https://www.aitex.es/personal-protection-equipment-certificate/?lang=en)

Date: Click or click here to write text

Name, surname and title: Click or click here to write text

Authorized Signature and Company Seal: Click or click here to write text