|  |  |
| --- | --- |
| Applicant Body Name |  |
| Applicant Address |  |
| Phone |  | Tax Office |  |
| e-mail |  | Tax No |  |
| Web Address |  | Name and Position of the contact person |  |

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| --- |
| The preffered conformity to type process for products under Category II:[ ]  PPE Regulation Annex VII, (Module C2), conformity to type based on internal production control plus supervised product checks at random intervals[ ]  PPE Regulation Annex VIII, (Module D), conformity to type based on quality assurance of the production process |

 Information on the EU Type Examination Certificates of the products in this application:

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| Product Type and Model | Issued Notified Body | Certificate No | Issue Date | End Date |
|  |  |  |  |  |

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| Declarations (Do not forget to check the boxes) |
| [ ]  I declare that, I did not apply for Module C2 or Module D service to another Notified Body for the Products listed above [ ]  I declare that, I will follow the requirements of “Universal Certification” certification process based on EU 2016/425 reguation and follow the requirements of the EU 2016/425 regulation[ ]  I declare that, I will cooperate and accept any visit or audit requests from Universal Certification and accompany to sampling of adequate variety and quantity of products for examination and show necessary ease for the products within this application.[ ]  I declare that i will not start mass production of the Products listed above and will not refer or use CE mark with Universal Certification Notified Body number until successful completion of conformity to type examination or audit processes.[ ]  I declare that I will annex any result of conformity to type gathered from another Notified Body for the products within this application (Expired conformity to type certificate and reports, any negative result for conformity to type examination result etc)  |

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| --- | --- | --- |
| Place  | Date | Seal / Authorised Name and Signature |

* The following annexes shall be provided with the application
1. EU Type Examination Certificate, Annexes and Test Reports
2. Technical documentation provided for EU Type Examination Certificate (as defined in 2016/425/EU Annex 3)
3. Official entitiy registration documents
4. Authorised signature samples
5. List of inspection and test equipments
6. Any certificate for quality management system or similar
7. List of inspection and test equipments, Number of Manufacturing Lines, Manufacturing Capacity (i.e piece per month)
8. Sub contractor agreement (if applicable)
9. Authorisation letter (if applicable)
10. Trademark registration (if applicable)
11. Yearly production plans