citeve

Request of EC Type-Examination

Product certification – CE Marking

Manufacturer's Information:

Name .....

Address .....

E –mail .....

VAT number ..... Phone number ..... Fax number. .....

OR

Authorized Representative's Information:

Name .....

Address .....

E-mail .....

VAT number ..... Phone number ..... Fax number.....

Request of EC Type-Examination to the following Personal Protective Equipment (PPE):

PPE production plant	
Description of PPE	
Reference of model	
Certification standards	
PPE category (II or III)	
Members States of destination	

Documents and samples to be attached for the model above mentioned:

- Manufacturer's Technical File (consisting of overall and detailed plans of PPE by calculation notes and the results of prototype tests for the verification of compliance with the basic requirements and other technical specifications, if existent);

- Description of the control means and tests used in the manufacture's plant to check compliance of production PPE, including final inspection from PPE and tests, the homogeneity of production and conformity of PPE with the model described in EC Type-Examination and with the essentials exigencies from Directive 89/686/EEC.

- Information notice (this document will follow each PPE and must comply the applicable standard(s) requirements and directive 89/686/CEE).

- 1 PPE from the same reference above mentioned that the Notified Body can perform the definitive budget and decide the number of PPE samples that the manufacturer/authorized representative must sent to certification process. This PPE must be the delivered with corresponding label (marking) according certification standards.

Commitment of manufacturer/authorized representative when request this EC Type-Examination:

1. The manufacturer/authorized representative above mentioned declares that this request EC Type-Examination was only requested to Notified Body of Citeve.

2. The manufacturer/authorized representative commits that the products with EC Type-Examination must be produced according the same specifications of the samples examined by Notified Body of Citeve.

3. If the production plant is changed the manufacturer/authorized representative must inform the Notified Body.

_____, _____ of _____, _____

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Signature and Stamp