

**To:
CERTOTTICA S.C.R.L.
Zona Industriale Villanova, 7/A
32013 Longarone BL – ITALY**

APPLICATION FOR “EU” TYPE-EXAMINATION

No. _____

for personal protective equipment (PPE)

in conformity to

**Regulation (EU) 2016/425 of the European Parliament and of
the Council of 9 March 2016**

(Module B)

(Blank spaces and boxes in this table are to be completed by CERTOTTICA staff)

Review performed by RVAL:	Date of the review: RVAL's signature and Certottica's stamp
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< Manufacturer >

Application for EU type-examination – Equipment: ... Model ...

IDENTIFICATION DATA

MANUFACTURER [1]

Company name	
Legal Headquarters	
Operative Headquarters	
Legal Representative	
Telephone	
Company Contact Person	
Role	
Telephone	
E - @ MAIL	
Manufacturing Unit	
Manufacturing Unit Headquarter	
Telephone	

[1] Any change in the above-mentioned data must be promptly passed on to the notified Body

AUTHORIZED REPRESENTATIVE OR LEGAL REPRESENTATIVE IN THE EUROPEAN UNION [1] [2]

Company Name	
Legal Headquarter	
Legal Representative	
Company Contact Person	
Role	
Telephone	

[1] Any change in the above-mentioned data must be promptly passed on to the notified Body

[2] Please enclose assignment letter signed by the manufacturer

PRODUCT – BASIC MODEL

Type of PPE	
Field of application	
Trade-mark	
Identification code /Model name	
Reference standard	
EU type-examination certificates in existence	
In the case of revision requests, include the reason for the revision	
In the case of requests for revision with renewal	<input type="checkbox"/> I wish the Certificate to be issued immediately, without waiting for the natural expiry date
	<input type="checkbox"/> I wish the Certificate to be issued on the natural expiry date
PPE category	

PRODUCT - VARIANTS

Variants of the basic model	
Identification code	

< Manufacturer >

Application for EU type-examination – Equipment: ... Model ...

CONTRACT

Between

CERTOTTICA S.C.R.L.

Loc. Villanova, 32013 Longarone BL - Italy
C.E.O. Mr. Corrado Facco

and **the Client** identified in the table “Manufacturer” and/or in the table “Authorized Representative or Legal Representative in the European Union” present in section “Identification data” (hereinafter called “the Applicant”)

INTRODUCTION

The Applicant, intends to provide the individual protection equipment described below with EU Type-Examination Certificate in accordance to the Regulation (EU) 2016/425 of the European Parliament and of the Council. CERTOTTICA is a notified body authorized to issue EU type-examination certificates for personal protection equipment according to the Regulation (EU) 2016/425 of the European Parliament and of the Council. CERTOTTICA is a body notified by the European Commission, which has assigned it identification code **2008**.

It is hereby agreed as follows:

Art. 1 - INTRODUCTION AND OBJECT OF CERTIFICATION

The preliminary statements must be considered as integral and essential part of this contract. The object of certification shall be the protection device identified in the table “Product – Basic model” and relative “Product – Variants” present in section “Identification data” hereinafter referred to as product.

Art. 2 - REFERENCE LEGISLATION

CERTOTTICA certifies the product of the Applicant, identified in the previous article, on the basis of the applicable harmonized standard, hereinafter indicated as reference law, made up by the Regulation (EU) 2016/425 of the European Parliament and of the Council for personal protection equipment.

Art. 3 - REQUIREMENTS TO OBTAIN THE EU TYPE-EXAMINATION CERTIFICATE

The Applicant has prepared the Technical File foreseen by Annex III of the reference law, in compliance with the dispositions of Art. 8 of the above-mentioned reference law and, if needed, it has supplied a sufficient number of samples as specified in the offer prepared by the relevant Sales Office as provided for by Art. 1 of this Contract. The Applicant shall consequently send to CERTOTTICA the Formal Application for “EU” type-examination, the Technical Documentation and, if applicable, the samples to test, according to point 3 of Annex V of the reference law.

Art. 4 - EXAMINATION OF THE TECHNICAL DOCUMENTATION AND TESTS ON THE SAMPLES

CERTOTTICA examines the Technical Documentation provided by the Applicant, consequently arranges and carries out the necessary tests on the product samples, if applicable, and applies all the relevant procedures foreseen by point 4 of Annex V of the reference law, with the aim to verify if the product satisfies the applicable essential health and safety requirements.

Art. 5 - ISSUING OF THE EU TYPE-EXAMINATION CERTIFICATE

In the observance of the conditions listed in the quotation, CERTOTTICA undertakes to issue and officially register the EU Type-Examination Certificate for the Applicant’s product, in compliance with point 6 of Annex V of the reference law, to the extent to which the Technical Documentation delivered upon presentation of the formal application for “EU” type-examination and the technical tests on the product prove satisfactory as regards the applicable essential health and safety requirements.

EU Type-Examination Certificate is released in double copy, one of this is forwarded to the Applicant and one is kept by CERTOTTICA.

The communication of the issuing of the EU Type-Examination Certificate shall be forwarded to the Organizations (national and/or international) to which such information is due in terms of rule of law.

Any revision of an EU Type-Examination Certificate automatically deletes and replaces the previously issued version from the date of issue (start of validity) of the revised certificate.

Art. 6 - UNICITY OF THE APPLICATION FOR “EU” TYPE-EXAMINATION

According to the dispositions provided for by point 3 of Annex V of the reference legislation, the Applicant states that for the model specified in article 1 of this Contract the same Application of “EU” type-examination is not forwarded to any other notified body, and that the model presented for examination has not been the subject of any previous EU type-examination certificate refusal decision.

Art. 7 - APPLICANT'S OBLIGATIONS

The Applicant undertakes to put all the information necessary to evaluate the product conformity to basic health and safety requirements at CERTOTTICA's disposal.

Art. 8 - CORRESPONDENCE BETWEEN THE PRODUCT UNDER EU TYPE-EXAMINATION AND SAMPLE

With the proper use of the production processes, and of examination and control methods, the Applicant undertakes to guarantee that the product launched corresponds to the sample submitted to the EU type-examination certification, of which CERTOTTICA will keep a reference sample, after the delivery of the EU Type-Examination Certificate.

Art. 9 - AMOUNTS DUE TO CERTOTTICA

For the examination of the Technical Documentation, for the execution of the Type Tests and the issuing of the EU Type-Examination Certificate, the Applicant undertakes to pay CERTOTTICA the fees stated in the offer submitted by CERTOTTICA Sales Department in accordance with the current price-list.

Art. 10 – EVALUATION REPORT

CERTOTTICA draws up an evaluation report that records the activities undertaken in accordance to the EU type-examination and their outcomes. Without prejudice to its obligations vis-à-vis the notifying authorities, CERTOTTICA shall release the content of that report, in full or in part, only with the agreement of the Applicant.

Art. 11 - EU TYPE-EXAMINATION CERTIFICATE CONDITIONS

The Applicant declares that s/he shall respect the conditions under which the EU Type-Examination Certificate is issued:

- the marking and the instructions are verified by CERTOTTICA at its choice in Italian or English. It is the Applicant's responsibility (manufacturer or its Legal Representative) to provide versions in different languages which are acceptable in the countries where the equipment is sold;
- any change to the product or to the technical file or, in the necessary cases, to the quality control manual and procedures shall timely be communicated to CERTOTTICA;
- the Applicant (manufacturer or its authorized Legal Representative) shall respect the procedures and regulations provided for by CERTOTTICA in order to issue the EU Type-Examination Certificate;
- copies of the EU Type-Examination Certificate, supplied to third parties, shall be reproduced in their entirety;
- the EU Type-Examination Certificate shall be kept by the Applicant (manufacturer or by its authorized Legal Representative) in order to show it to Control Bodies or Surveillance Authorities, on demand;
- the EU Type-Examination Certificate remains property of CERTOTTICA, which can ask for it back if some of the above-mentioned conditions are not respected;
- the validity of the EU Type-Examination Certificate is of 5 years from the issuing date.

Art. 12 - CONFIDENTIALITY

CERTOTTICA undertakes to consider all the documents and the information (know-how) it has received or that has come to its knowledge regarding the Applicant's organisation and products strictly confidential.

This clause shall remain valid and in force also after the expiry of the contract and until the confidential nature of the information and of the Know-how exchanged during its being in force persists.

All paperwork (documents, letters, notifications, etc.) and information relating to certification, since the Application for "EU" type-examination is presented, are considered confidential and access to them is governed by a specific procedure.

The only information that CERTOTTICA undertakes to communicate to all the people that require them, through specific written application to be sent by fax and/or email to the address which can be found on CERTOTTICA web site, are the information contained in the issued EU Type-Examination Certificate (without need of any authorization by the Applicant).

CERTOTTICA personnel at all levels, and external personnel involved in the supervision, tests and EU Type-Examination Certificate issuing that may gain knowledge of the information contained in these documents and other information relating to the Applicant CERTOTTICA has EU Type-Examination agreements with, are all subject to the professional secrecy act.

In the event that the law establishes that certain information should be notified to any Control Authorities which request it, CERTOTTICA will inform the Applicant regarding the information which has been disclosed.

Further to instructions from the Applicant, CERTOTTICA is irrevocably authorized to transmit the records, test reports, EU Type-Examination Certificate and any other information, to any third parties in compliance with current privacy law.

Art. 13 - ENCLOSURES

The offer, submitted by CERTOTTICA relevant Marketing Department, and approved by the Applicant, and the Regulation for the Certification of Personal Protective Equipment (PQ 16.101.004), available on CERTOTTICA site (www.certottica.it), are part of this contract.

Art. 14 - JURISDICTION

Any disputes that may arise between the parties that are directly or indirectly related to the application or interpretation of the Regulation for the Certification of Personal Protective Equipment (PQ 16.101.004) and this Application for “EU” type-examination, which cannot be amicably settled between the parties, shall be referred to the exclusive jurisdiction of the Court of Belluno.

Date: ...

THE APPLICANT
(-----)
(Legal Representative)

CERTOTTICA S.c.r.l.
Mr. Corrado Facco

The following articles are specifically approved in writing according to 1341 and following amendments:
art. 3 requirements to obtain the EU Type-Examination Certificate, art. 4 examination of the technical documentation and tests on the samples, art. 5 issuing of the EU Type-Examination Certificate, art. 8 correspondence between the product under EU Type-Examination and sample, art. 12 confidentiality, art. 14 jurisdiction.

THE APPLICANT
(-----)
(Legal Representative)

CERTOTTICA S.c.r.l.
Mr. Corrado Facco