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CIMAC Centro Italiano Materiali di Applicazione Calzaturiera

Sede operativa
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According to the EEC instructions 89/686 dated 21st of December 1989, concerning the standardisation of legislation of all member countries with regard to individual protection systems and the relative legislative decree dated 4th of December 1992, N° 475,

A.N.C.I. servizi s.r.l. - C.I.M.A.C. section
CENTRO ITALIANO MATERIALI DI APPLICAZIONE CALZATURIERA
Authorized with the decree issued by the Ministry of Industry of the Italian Republic on the 11th of October 2000 - Community Identification N° 0465

grants:

EC TYPE-EXAMINATION CERTIFICATE
N° 0161/15826/09

for the following model of personal protective equipment:

Clog - art. " "0" GRAVITY"

Manufacturer (see notes):

MEDICAL FOOTWEAR LTD

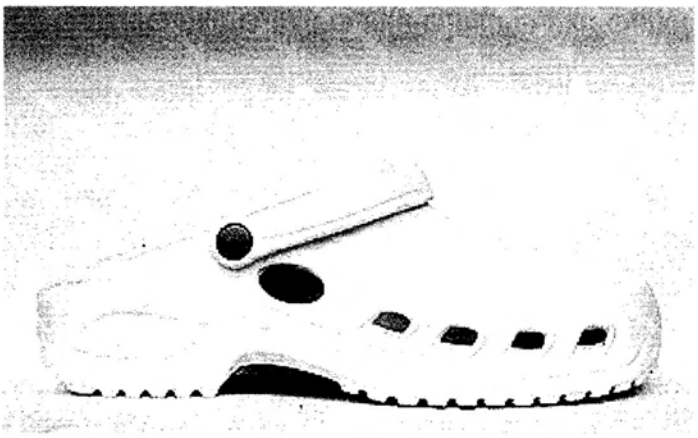
6 CORBEN MEWS
CLYSTON STREET
LONDON
SW8 4TA

Vigevano, 5th October 2009

Technical Manager of the Centre
Ing. Giuseppe Bellotti

1. Description of personal protective equipment:

PPE Category: second category
Type of PPE: occupational footwear for professional use
Design: Clog
Classification: II - All-rubber (i.e. entirely vulcanized) or all-polymeric (i.e. entirely moulded) footwear
Size range: from 35÷36 to 45÷46 (French size)
Identification of last: "0" GRAVITY
Manufacturing process: Injection
The footwear has not a removable insock



2. The tests and the examinations to verify the conformity of the article (in compliance with art. 10 of Directive 89/686/CEE) are performed applying the following harmonized standards and the Directive 91/338/CEE of 18th June 1991, 2004/21/CE of 25th February 2004 and 94/27/CE of 30th June 1994:

- EN ISO 20344:2004+A1:2007 – Personal protective equipment – Test methods for footwear.
- EN ISO 20347:2004+A1:2007 – Personal protective equipment – Occupational footwear.

3. The results of tests and examinations are contained in the following test reports:

C.I.M.A.C.	RP 20094650	dated	5th	October	2009
C.I.M.A.C.	RP 20094651	dated	5th	October	2009

4. Requirements of the personal protective equipment:

The clog art. " "0" GRAVITY" complies with:

the basic applicable requirements for occupational footwear specified in table 2 of EN ISO 20347:2004+A1:2007 standard;

the additional requirements for special applications specified in table 14 of EN ISO 20347:2004+A1:2007 standard:

- antistatic properties (A);
- energy absorption of seat region (E);
- non-cleated outsole.

The shoe complies whit what prescribed by EN ISO 20347:2004/A1:2007 standard concerning the slip-resistance of the outsole, "SRC" requirement.

5. Marking of the personal protective equipment:

The following information is provided on the footwear outsole:

- the "CE" mark
- date of manufacture (month and year)
- manufacturer's type designation: "0" GRAVITY
- manufacturer's identification mark: "MEDICAL FOOTWEAR"
- size.

6. Notes:

- The manufacturer is the one assuming the responsibility of design and fabrication of a product included in the Directive, in order to throw it in the market.
- The marking for the number and year of the reference European standard and the symbols appropriate to the protection provided should be adjacent to one another.
- This EC Type-Examination Certificate must be kept by the manufacturer, in order to produce it, on request, for the control body or safety control administration.
- The content of this EC Type-Examination Certificate is referred to the tested personal protective equipment only.
- This EC Type-Examination Certificate may be integrally duplicated; the copy must be faithful, legible (if pint size) and must contain the bold caption "TRUE COPY".
- The manufacturer should inform ANCI Servizi srl – Sezione CIMAC for any modification to the product, to the place/process of manufacturing and, if relevant (III category PPE), to the quality system if this compromises the conformity of the product to the essential requirements or other provisions of the Directive 89/686/EEC.
- The manufacturer should report all the complaints relating to the conformity of the certified product to the requirements of the reference harmonized standards and supply these reports on request of ANCI Servizi srl – Sezione CIMAC.
- The manufacturer should implement appropriate corrective actions when nonconforming product to the essential requirements of the EC certification are identified.