

EU TYPE EXAMINATION CERTIFICATE



APPROVED BODY 2575

The PPE detailed herein the criteria of an EU Type Examination in accordance with Annex V, including the applicable clauses of the Essential Health and Safety Requirements of the PPE Regulation EU 2016/425, for the category II followed by conformity to type based on internal production control (module C) set out in Annex VI.

Following an EU Declaration of Product Conformity you are hereby licensed to mark the product(s) detailed in accordance with Article 17 of the PPE Regulation EU 2016/425.

VALIDITY OF CERTIFICATE

This certificate will cease its validity at any time if needed, in particular if changes in the manufacturing process, in the raw materials or in PPE components will occur.

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PRD N° 277B

Membro degli Accordi di Mutuo Riconoscimento EA, IAF e ILAC

Signatory of EA, IAF and ILAC Mutual Recognition Agreements

Manufacturer: Vicsa Safety Comercial Limitada

Address: Panamericana Norte No.5151 Conchali Santiago - Chile

Authorised Representative: -

Address: -

Certificate No.: ITASLNB20005337

Category Product: II

Trade Name (Model/Product Reference): STEELPRO/RDL MULTIFLEX ORIGINAL IMPACT

Article: 3009710520834(M); 3009710520833(L)

Product type: GLOVES

Reference(s) Standard: EN 420:2003+A1:2009, EN 388:2016+A1:2018

Description:

General Requirements: Dexterity 5; Sizes 9, 10; Mechanical Risks: Abrasion resistance 3; Blade cut resistance 1; Tear resistance 4; Puncture resistance 1; Cut Resistance X; Impact P;

This has been shown through satisfactory testing to: EN 420:2003+A1:2009, EN 388:2016+A1:2018

Examination of the Technical File Documentation, No: MULTIFLEX ORIGINAL IMPACT - Rev 3 - 24/02/2021

Test Report no. See Technical File

Remark:

Note:

Issue Date 11/03/2021

Expiry Date 10/03/2026

Issued at: Lastra a Signa (FI)

General Manager Elena Ruffino

For and on behalf of INTERTEK ITALIA Spa

This certificate shall be issued on the following conditions:

1. This certificate refers only to the samples tested and submitted to the tests and assessments by the Body
2. The issue of the certificate does not imply that the Notified Body has carried out any surveillance or control of the manufacture.
3. The applicant shall ascertain that the manufacturing process ensures the products conformity with the approved samples as required by Regulation (EU) 2016/425



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