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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ACCREDIA
L'ENTE ITALIANO DI ACCREDITAMENTO

PRD N° 277B

Membro degli Accordi di Mutuo Riconoscimento EA, IAF e ILAC

Signatory of EA, IAF and ILAC Mutual Recognition Agreements

Manufacturer: The Jordan Al- Manara Shoes Co.Ltd
Address: Qastal Industrial, East Rail Road, Al-Jeezah, P.O.Box 340595 Amman, 11134, Jordan
Authorised Representative: -
Address: -
Certificate No.: ITASLNB21008257
Category Product: II
Trade Name (Model/Product Reference):-
Article: SU-7313, SU-5313, SU-5313 NL, SU-8313
Product type: SAFETY FOOTWEAR
Reference(s) Standard: EN ISO 20345:2011
Description:

Construction: Direct Injection; Toecap: PALADEN 604 (Metallic); Midsole: PELADEN 1180 METALIC; Last: SJ; Sole: PU/PU; Mould: ULTRA FIT; Size Range: 37-47; Category: S3 SRC.

This has been shown through satisfactory testing to: EN ISO 20345:2011

Examination of the Technical File Documentation, No: New Application the Jordan Al Manara SU-7313, SU-5313, SU-5313 NL, SU-8313 - Rev.1 05/07/2021

Test Report no. See Technical FIle

Remark:

Note:

Issue Date 27/09/2021 **Issued at:** Lastra a Signa (FI)
Expiry Date 26/09/2026 **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