

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.

INTERTEK ITALIA SpA
Via Miglioli, 2/A
Cernusco sul Naviglio (MI), Italy
T: +39 02 95383833
F: +39 02 95383832



PRD N° 277B

Membro degli Accordi di Mutuo Riconoscimento EA, IAF e ILAC

Signatory of EA, IAF and ILAC Mutual Recognition Agreements

Manufacturer: ACTIVE GEAR SA

Address: 1 Impasse du Nouveau Marché, 1723 Marly
, Switzerland

Authorised Representative: -

Address: -

Certificate No.: ITASLNB21014791

Category Product: II

Trade Name (Model/Product Reference):-

Article: S6507, S6508, S6509, S6510, S6511

Product type: GLOVES

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

Description:

FULL GRAIN LEATHER DRIVER GLOVE.

General Requirements: Dexterity: 5; Sizes: 7, 8, 9, 10, 11. Mechanical Risks: Abrasion resistance: 2; Blade cut resistance: 1; Tear resistance: 4; Puncture resistance: 2; Cut resistance: X.

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

Examination of the Technical File Documentation, No: FULL GRAIN LEATHER DRIVER GLOVE - S6507, S6508, S6509, S6510, S6511 - Rev.01 20/10/2021

Test Report no. See Technical File

Remark:

Note:

Issue Date 04/11/2021

Expiry Date 03/11/2026

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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Intertek Italia S.p.A. Via Miglioli, 2/A - 20063 Cernusco sul Naviglio, Milano - Italy