



Notified Body No. 1023

INSTITUTE FOR TESTING AND CERTIFICATION, Inc.
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EU Type-Examination Certificate

No. 20 0312 T/NB

issued in the compliance with the Regulation (EU) 2016/425 of the European Parliament and of the Council of 9 March 2016 on personal protective equipment and repealing Council Directive 89/686/EEC, for personal protective equipment of category III:

Protective clothing and clothing components
Type: MEDIMAN 2020

Manufacturer:

LP MAN s.r.o.

K Zorníci 1654/18

957 01 Bánovce nad Bebravou

Slovenská republika

Tax Registration No: SK2023782420

This Certificate confirms that above referenced personal protective equipment (PPE) fulfils the essential health and safety requirements as they are stated in the Regulation (EU) 2016/425 of the European Parliament and of the Council, specified in detail in the harmonized technical standards:

ČSN EN ISO 13688:2014 (EN ISO 13688:2013)

ČSN EN 13034+A1:2009 (EN 13034:2005+A1:2009) [**Type 6** – coveralls with hood, **Type PB [6]** – other clothing components]

ČSN EN 14126:2004 (EN 14126:2003) [**Type B**]

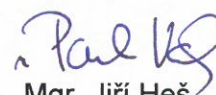
The PPE is produced in compliance with the manufacturer's technical file. The detailed product descriptions, the results of technical file examination as well as the test results including their evaluation are presented in the ITC's Evaluation Report No. 723301954/2020 that is an integral part of this Certificate.

Condition of this certificate use and related information:

1. It applies only to the type of the category III PPE submitted to test referenced above.
2. It does not imply that the Notified Body has performed any surveillance or control of PPE manufacture.
3. The manufacturer is obligated to assure that all PPEs of the respective type conform to type approved by this Certificate, which must be used only in conjunction with one of the conformity assessment procedures referred to in Article 19 (c) either of the following: conformity to type based on internal production control plus supervised product checks at random intervals (module C2) set out in Annex VII or conformity to type based on quality assurance of the production process (module D) set out in Annex VIII Regulation of the European Parliament and of the Council (EU) 2016/425.
4. The manufacturer shall inform the Notified Body of all technological changes in manufacture of the approved type and as consequence of the technical advances he shall regularly keep himself informed of any standard changes as well as modifications of testing methods conducted by Notified Body, which shall approve these changes in necessary cases by the amendment of this Certificate.
5. The manufacturer shall affix to each PPE of the approved type the conformity mark CE followed by Notified Body number, which approves the system of production control in accordance with Annex VII (module C2) or VIII (module D) the Regulation (EU) 2016/425 of the European Parliament and of the Council, according to the principles laid down in Regulation (EC) no. 765/2008.

Issued in Zlin, on 30th June 2020
Valid until: 29th June 2025




Mgr. Jiří Heš

Representative of the Notified Body No. 1023