Occupational Safety Research Institute, v.v.i. Jeruzalémská 1283/9, 110 00 Praha 1, Czech Republic Notified Body 1024



EU TYPE-EXAMINATION CERTIFICATE No. 1024/E-011/2023

This EU type-examination certificate is issued to:

Manufacturer: Western Ukrainian Center Medservis, LLC

79029, Ukraine, Lviv, Antonovich str., 128

Identification number: 804

PPE product: Personal protective equipment to protect the body against X rays

OBERIH, model Protective x-ray headgear

It is certified that the manufacturer's technical file and above-mentioned PPE product have been assessed and found to be in accordance with the essential health and safety requirements of Regulation (EU) 2016/425 of the European parliament and of the Council on personal protective equipment and repealing Council Directive 89/686/EEC, as recorded in Report No.1024/ZZ-002/2023, which is an integral part of this Certificate.

When examined the model was found to meet all of the relevant requirements of the appropriate harmonized standard(s):

ČSN EN 61331-3 ed.2:2015 Protective devices against diagnostic medical X-radiation – Part 3:

Protective clothing, eyewear (idt. IEC 61331-3:2014)

ČSN EN ISO 13688:2014 Protective clothing – General requirements (idt EN ISO 13688:2013)

The certification was performed according to the certification scheme of Regulation (EU) 2016/425 Module B. Marking and instructions have been assessed in the English language only.

Certificate is valid until 2028, February 14

For and on behalf of Occupational Safety Research Institute

Notified Body No. 1024:

Date of Issue: 2023-02-14

Ustav bezarati i subjekt 1030 e Conosti pracije pracij

Ing. Jiří Tilhon, Ph.D., LL.M.

This certificate was issued in Czech and English versions. Both versions have the same validity.

Description and picture of the PPE product:



Personal protective equipment to protect the body against X rays OBERIH, model Protective x-ray headgear for protection against X-Ray ionization in professional application according to instruction for using.

The product is classified in category III and shall only be used in conjunction with conformity assessment procedure Module C2 within the meaning of Regulation (EU) 2016/425.



Graphical appearance of CE mark. In the event of the involvement of a notified body in the stage of manufacturing check (category III, Module C2 or D) its distinguishing number shall be added to CE mark.