Certificate Number: PPE19161376

EU TYPE-EXAMINATION CERTIFICATE

This is to certify that INSPEC International B.V., Notified Body 2849, has evaluated the Personal Protective Equipment type(s) in respect of the product detailed on this certificate and deemed it(them) to be in compliance with Annex V (Module B) of the Personal Protective Equipment Regulation (EU) 2016/425 and the applicable Essential Health & Safety Requirements.

Manufacturer:

Majestic Safety Products & Services B.V.
Jan Campertlaan 6
3201 AX Spijkenisse
P.O. Box 80
3200 AB Spijkenisse
The Netherlands

Compliance with the applicable Essential Health & Safety Requirements has been demonstrated as above, including examination in accordance with the harmonised standard below:

EN 352-2:2002

Product description: Hearing protection – Earplugs;

Models: OXXA Sana; 8011-I, 8011-R, 8021

Date of initial certification: 15 May 2019
Date of current issue: 13 September 2023
Period of validity: 17 July 2023 - 17 July 2028

13.09.23
Certificate Signatory

Product details

Model identification: OXXA Sana: 8011-I, 8011-R and 8021 (Aural type, Disposable)

Variants:

Model	Description
OXXA Sana 8011-I	Earplugs without interconnecting cord
OXXA Sana 8011-R	Earplugs without interconnecting cord
OXXA Sana 8021	Earplugs with interconnecting cord

Technical file reference: TF18161019 & TF19161376

Category: III (three)

Performance levels:

Frequency	125	250	500	1000	2000	4000	8000
Mean Attenuation (dB)	33.0	33.3	36.1	37.1	36.1	41.8	38.5
Standard Deviation (dB)	6.7	7.6	7.4	4.7	4.8	3.6	3.9
Assumed Protection (dB)	26.3	25.7	28.7	32.4	31.3	38.2	34.6

H= 33dB M=31dB L=28dB SNR = 34 dB

Options: None

Sizes: Nominal diameter, 6-12mm

Accessories: None

Certificate amendment record

Date	Description
15/05/2019	Initial issue
13/09/2023	Renewed following Simplified Review and Amendment to model names

Conditions attached to the issue of this certificate:

1. This certificate alone, if Category II (two) PPE, forms INSPECs permission to the manufacturer to use the 'CE' conformity mark for compliant products to be placed on the European Union internal market. In this case, the manufacturer may affix the conformity mark to each PPE or on a document supplied with the PPE, and draw up a written EU declaration of conformity for each PPE model referencing this certificate as per Article 17.

2. The manufacturer / authorised representative shall undertake to fulfil the obligations arising out of the Personal Protective Equipment Regulation (EU) 2016/425, and with INSPEC's Regulations governing this Module.

3. The manufacturer / authorised representative shall inform INSPEC without delay of any planned changes to the product, technical file or manufacturer information which may affect the validity of this certificate, before any such change is made.

4. Marking and instructions have been assessed in the English language only. It is the manufacturer's / authorised representative's responsibility to obtain and supply language versions acceptable to the country where the product is to be sold.

5. For category III product, the manufacturer must obtain and maintain an approval decision to Module C2 or Module D prior to placing product on the European Union internal market.

6. This certificate remains the property of INSPEC and may be withdrawn if any of the conditions attached to its issue are not complied with.

7. This certificate may be copied or reproduced by the certificate holder, complete and without omissions or additions, and in accordance with INSPEC's terms of business.

8. The manufacturer shall not use its product certification in any manner as to bring INSPEC into disrepute, nor make any statements regarding the product certification that INSPEC considers misleading or unauthorised.

9. Upon suspension, withdrawal or termination of this certificate, the manufacturer must discontinue all advertising matter that references the product certification and take action to cease production of products.