

EU Declaration of Conformity

We,

DEXXON ENERJİ SAN. VE TİC. A.Ş

YENİBOSNA MERKEZ MAH., 29 EKİM CAD., İSTANBUL VİZYON PARK OFİS BLOKLARI, PLAZA: 1 FLOOR : 8 BAHÇELİEVLER / İSTANBUL / TURKEY

declare under our sole responsibility that the PPE product listed below

Product Description	NON REUSABLE FFP2 NR FACE MASK
Article No	DXNMD NRFM04
Type	FFP2 NR
Manufacturer	DEXXON ENERJİ SAN. VE TİC. A.Ş
Brand Name	DEXXON

We declare that the product described above meets the requirements of the relevant provisions of the regulation as a result of the EU Type examination according to Annex 5 of the PPE Personal Protective Equipment Regulation (2016/425 / EU)

This Declaration of Conformity covers the PPE device as specified in the product list belonging to this declaration.

The product identified below complies with the general safety and performance requirements of the PPE Personal Protective Equipment Regulation (2016/425 / EU) by meeting the following standards:

Conformity Assessment Route	EN 149: 2001 + A1: 2009 Respiratory Protective Devices-Against Particles
Applicable Harmonised Standards	EN 149: 2001 + A1: 2009
Rule	Annex 5 of the PPE Personal Protective Equipment Regulation (2016/425 / EU)
Risk of the Device	The Personal Protective Equipment (PPE) has been assigned to CATIII
Classification	FFP2 NR
Base of Certificate	SZU 22 PPE 041 numbered and 08/02/2022 dated EU Type Examination Certificate ASR-PPE-C2-018/R.01 numbered and 08/02/2022 dated and inspection report

The Conformity is ensured with the following mechanism:

Complies with EU 2016/425 PPE Regulation establishing technical requirements for Category III products,

All required tests referred in above standards are conducted

Complies with other relevant harmonized legislation and community standards.

For the assessment of conformity the EU Type Examination Modul B certificate (Certificate Number : SZU 22 PPE 041) and Module C2 certificate (Certificate Number : SZU 22 PPE C 018) is issued ,after all technical evaluations for conformity to the regulation and harmonised standards conducted, by ;

SZUTEST Uygunluk Değerlendirme A.Ş. as Notified Body number 2195

The product is under surveillance of same Notified Body, NB 2195 according to the Annex III (Module C2) of the PPE Regulation (EU)2016/425,for quality assurance.

This declaration was first published on 28/01/2022 and the reference will continue to be valid unless there are significant changes in harmonized standards and factory production conditions. For detailed information about the product, see the technical file.

08.02.2022
Murat Koç
General Manager

