

**ATTESTATION OF CONFORMITY****Certificate Nr: MDD - 098**

*In conformance to the European Economic Commission 93/42/EEC Medical Devices Directive on harmonisation of laws, regulations and administrative documentation of Member States on Medical Devices and European Economic Commission directive 93/68/EEC amending Medical Devices Directive dated 22 July 1993,*

the products manufactured by

**İBİŞLER TEKSTİL SANAYİ VE DIŞ TİCARET ANONİM ŞİRKETİ**

at the following address

Orhangazi Mahallesi Tunç Caddesi No:5 34358 Esenyurt / İSTANBUL

**EN 14683:2019+AC:2019 Medical Face Masks**

Brand Name : A&Z MED

Model : OLI - 2022

Type II

are tested according to the following initial type tests by the manufacturer

Technical standard EN 14683:2019+AC:2019 Medical face masks - Requirements and test methods

For the assessment of conformity, the following documents were also applied to:

Results of laboratory tests Ekoteks Testing Laboratory BFE

Results of laboratory tests Ekoteks Testing Laboratory Differential Pressure

Results of laboratory tests Ekoteks Testing Laboratory Microbial Cleanliness

UNIVERSAL CERTIFICATION has evaluated production, design, intended use, risk evaluation according to safety purpose, product itself and add-on components (if exists) and product technical drawings of the medical face masks manufactured and designed for use during the medical operations or similar medical situations with same requirements which require restriction of infectious materials to be spread to patients. With this certificate, it is approved that the product fulfils all essential requirements and the related rules of 93/42/EEC Medical Devices Directive (MDD) Class I are applied. The information on the packaging for the above listed products covers the necessary information stated in Annex I, §13, of the Medical Devices Directive (93/42/EEC) or Annex I, §23, of the Medical Device Regulation (EU) 2017/745. This information includes; reference to EN 14683 standard, type of mask (as indicated in Table 1) and other relevant information given in EN ISO 15223-1:2016 and EN 1041:2008+A1:2013. It is considered to be suitable to attach a CE mark, as seen below, on your products in accordance with the information given in this certificate with publishing an EU Declaration of Conformity.

This certificate is issued on 04/05/2020 and valid until 03/05/2021 with the conditions that no change has been made with the product references and no change in the production process or not suspended or withdrawn for any reason.

İSTANBUL -04/05/2020



Verify the validity with the QR Code

Suat KAÇMAZ  
UNIVERSAL CERTIFICATION  
Genel Müdür

# EU DECLARATION OF CONFORMITY

## PRODUCER

İBİŞLER TEKSTİL SANAYİ VE DIŞ TİCARET ANONİM ŞİRKETİ  
Orhangazi Mahallesi Tunç Caddesi No:5 34358 Esenyurt / İSTANBUL

## PRODUCT DESCRIPTION

Layered and molded medical device classified in the class I - Medical Device to be used as protection against inhalation of viruses, bacteria, other microorganisms, allergens from the environment (Type I and/or Type II)

The Producer / the Manufacturer declares on his sole responsibility that the product above is, under conditions of normal use and conditions defined by the Producer / the Manufacturer, safe and meets all the necessary legal conditions and requirements. The product, a medical device that is intended for single use and solely in accordance with the Producer's / the Manufacturer's instructions.

The Conformity is assessed especially with the following provisions:

- Government Regulation no. 93/42/EEC Medical devices establishing technical requirements for medical devices, in effective wording
- Technical standard EN 14683:2019+AC:2019 Medical face masks - Requirements and test methods
- Other relevant harmonized legislation
- Other relevant local, national and community standards
- For the assessment of conformity, the following documents were also applied to:
  - Tests for irritation and delayed-type hypersensitivity
  - Results of laboratory tests Ekoteks Testing Laboratory BFE
  - Results of laboratory tests Ekoteks Testing Laboratory Differential Pressure
  - Results of laboratory tests Ekoteks Testing Laboratory Microbial Cleanliness

## MARKING, LABELLING

Annex I, §13, of the Medical Devices Directive (93/42/EEC) or Annex I, §23, of the Medical Device Regulation (EU) 2017/745 specifies the information that should be specified on the packaging in which the medical face mask is supplied. The following information shall be supplied:

type of mask (as indicated in Table 1). EN ISO 15223-1:2016 and EN 1041:2008+A1:2013 should be considered

## MEASURES TO ENSURE CONFORMITY

The Producer / the Manufacturer declares that he has taken all necessary measures to ensure the conformity of products placed on the market with technical documentation and basic requirements for this type of product.

İlhan İBİŞ  
General Manager  
İSTANBUL 04/05/2020

