

# AB Tip İnceleme Sertifikası EU Type-Examination Certificate

Belge No / Certificate No	: 89-20-02
Belgelendirme Tarihi - Bir Sonraki Belge Tarihi / Certification Date / Certificate Validity Date	: 11.12.2020-11.12.2025
Belge Geçerlilik Tarihi / Document Validity Period	: 5 yıl/ 5 years
Firma Unvanı ve Adresi / Company Name and Address	: DG Medicare GmbH Truchthari-Anger 34, 81829 München / Germany
Ürün Adı /Modeller / Product Name / Models	: DGMED2
Direktifi / Directive	: 2016/425 REGULATION
Modülü/Kategori / Module / Category	: B MODÜLÜ/ KATEGORİ III MODULE B / CATEGORY III
Test Rapor No/ları / Test Report No	: M-2020-00543
Ürün Tipi / Product Type:	

- EN 149:2001+ A1:2009 Solunumla ilgili koruyucu cihazlar - Parçacıklara karşı koruma amaçlı filtreli yarım maskeler/ Respiratory protective devices - Filtering half masks to protect against particles
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Ürünün Malzeme Bilgisi / Product Material Information: DGMED2 model ürünleri kumaş, kulak kayışı, burun klipsi ve filtre katmanı kullanılarak imal edilmiştir./ DGMED2 model products are manufactured using fabric, ear loop, nose clip, filter layer.

**Volkan AKIN**  
11.12.2020  
Karar Verici / Approver

**Okan AKEL**  
11.12.2020  
Şirket Müdürü / General manager







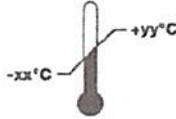

**ATTACHMENTS (89-20-02)**

To certify the PPE product at Category III level, C2 or D module is accompanied by applying one of the conformity assessment methods along with the EU Type Examination (Module B).

**Model : DGMED2**

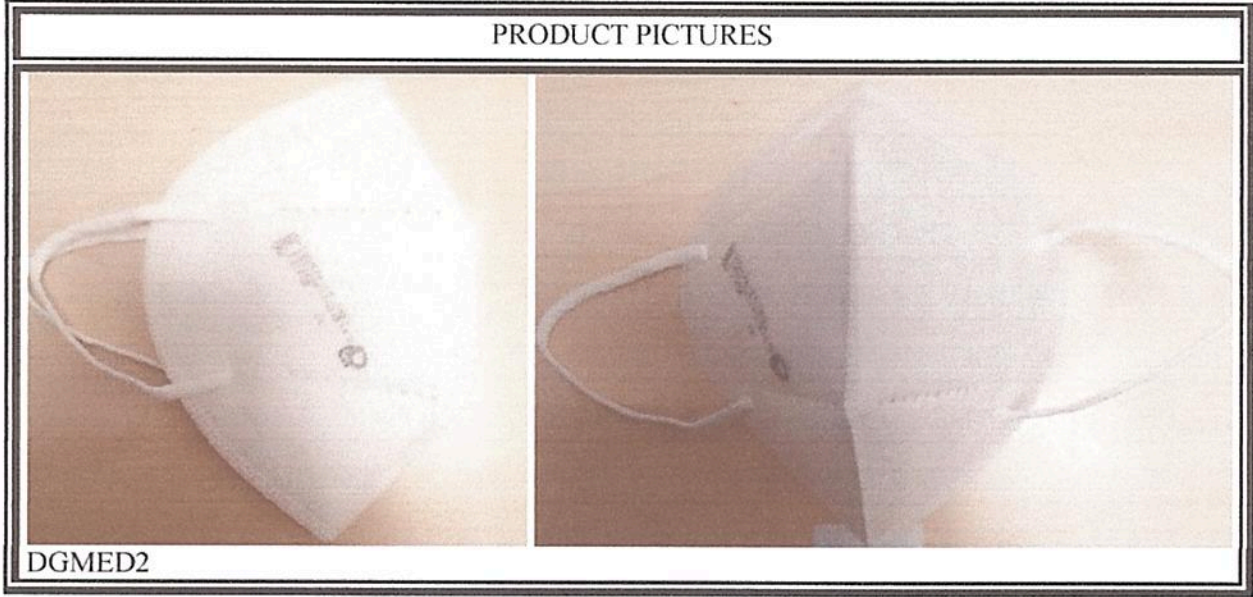
PPE SPECIFICATION	PERFORMANCE LEVELS
Classification	FFP2
Reusable / Single Shift Use	NR

PPE produced as a single unit to fit an individual user, all the necessary instructions for manufacturing such PPE on the basis of the approved basic model:

MARKING					
<b>MANUFACTURER:</b> DG Medicare GmbH					
<b>PPE TYPE :</b>					
- EN 149:2001+ A1:2009 Respiratory protective devices - Filtering half masks to protect against particles					
<b>MODEL:</b> DG MED2					
<b>PICTOGRAM AND PERFORMANCE LEVELS:</b>					
EN 149:2001+ A1:2009 FFP2 NR					
 NB 2841					
Or Condition of Storage					

MNA LABORATORIES SAN. TIC. LTD. ŞTİ declares that the above-mentioned product meets the requirements of the directive according to the EU Directive 2016/425, the safety of the product is covered by the conditions and use specified in this certificate and in the technical file.



**ATTACHMENTS (89-20-02)****DOCUMENTS IN THE TECHNICAL**

- Basic Health Safety Requirements
- Risk Assessment
- Test Reports
- Technical Report