

## ATTESTATION OF CONFORMITY

**Certificate No: MDD-180**

*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

**DENTA PHARMA İLAÇ MEDİKAL GIDA SANAYİ VE  
TİCARET LİMİTED ŞİRKETİ**

at the following address

Kalaba Mahallesi Ahmet Vefik Paşa Cad. No:13/B Keçiören ANKARA / TURKEY

**EN 13795-1:2019 Surgical Clothing and Drapes - Requirements and Test  
Methods - Part 1: Surgical Drapes and Gowns**

**Brand Name: DENTAPHARMA**

**Model: DGown**

(Standard Performance) are tested according to the following initial type tests by the manufacturer

For the assessment of conformity, the following documents were also reviewed:  
Laboratory test results for Microbial Penetration (wet/dry), Bioburden,  
Bursting and Tensile Strengths (wet/dry)

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 surgical gowns manufactured and designed for use to prevent the transmission of infective agents between clinical staff and patients during surgical and other invasive procedures. 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: performance level 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 09/07/2020 and valid until 08/07/2021 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that no change has been made with the product references and no change in the production process or not suspended or withdrawn for any reason.

ISTANBUL-09/07/2020



**Suat KACMAZ**  
UNIVERSAL CERTIFICATION  
Director



Verify the validity with the QR Code