



INNGEN
Medical Supply CO.

2020

Inngen Inc.

An FDA Registered Importer



Who are we?

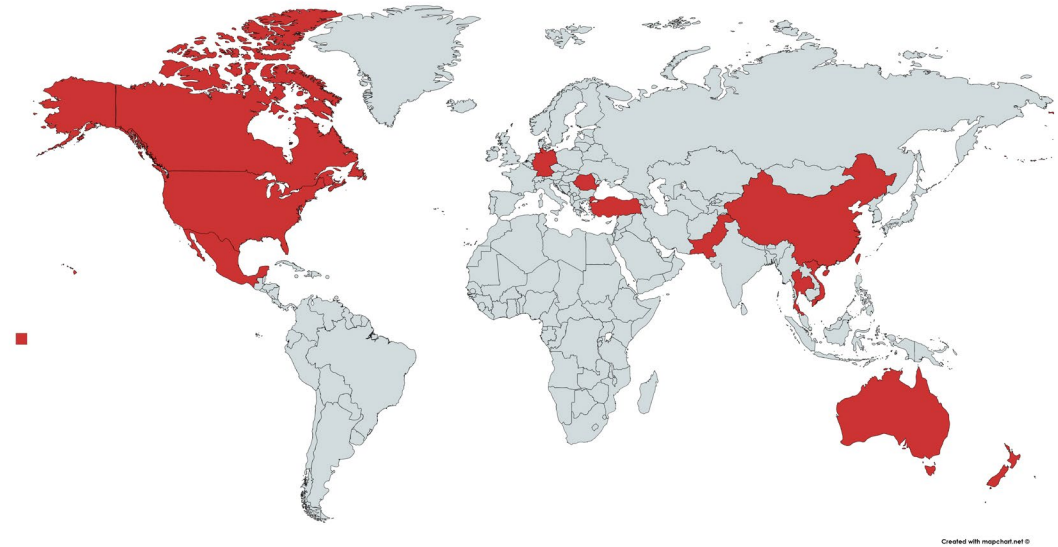
2020

A Global PPE Solution.

Inngen Inc. was founded to provide a global solution to the PPE supply problem happening as a result of COVID 19. We're an FDA registered Importer of medical supplies and have a vast network of COVID related Products.

All suppliers undergo strict onboarding due diligence that include license checks, FDA/CE Certification review, business registration review, and lab tests.

Our Supply Partners



Product Specifications

2020

Product Details

These are Latex Free. Practical barrier for a wide range of applications. Elastic cutoff available. Soft & light weight.

Output:

100,000 Available now

Payment Terms:

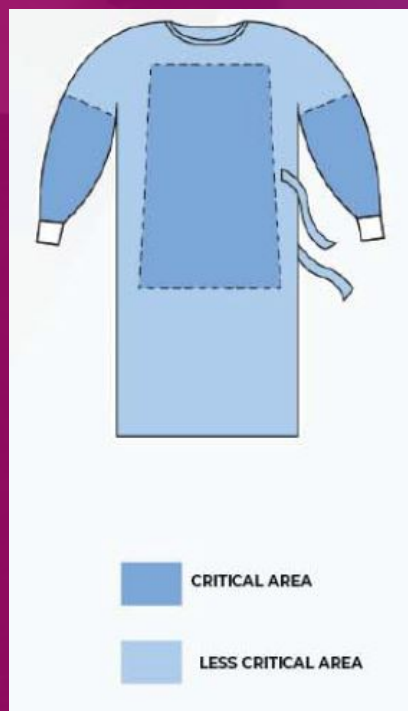
1. LOI & POF
2. Escrow/Inspect
3. Escrow is released

Surgical/Non-Surgical Isolation Gowns

Style	SMS
Weight	Level 2 - 40 GSM
Qt/Size:	100,000 Units 127*165CM (L)
Color	Blue
Packing	60PC 62X40X40CM GW:9KG
Certification:	ISO13485 CE FDA
Standard	AAMI Level 2

Isolation Gowns

Protection Zones



Pricing

Type	Price (Air Shipped)
Level 1 Isolation Gown	\$3.50
Level 2 Isolation Gown 35GSM	\$4.00
Level 3 Isolation Gown 45GSM (add .60 for sterile)	\$5.00
Level 4 Isolation Gown 45GSM + Extra protection (PE)	\$6.00
Level 4 Sterile (surgical)	\$6.60

Certifications

Establishment:
MERTOL GRUP TEKSTİL FERMUAR SANAYİ VE TİCARET A.Ş.
Gurbuz Sok. No:13/1
Güngören İstanbul, TR 34160
Registration Number: 3016945735
FEI Number*: 3016945735
Status: Active
Date Of Registration Status: 2020

Owner/Operator:
Mertol Grup Tekstil Fermuar Sanayi Ve Ticaret A.Ş.
Gurbuz Sok. No:13/1
Güngören, İstanbul TR 34160
Owner/Operator Number: 10074490

Official Correspondent:
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NVA QUALITY CERTIFICATION

NVA
NVA QUALITY CERTIFICATION

CE

EC DECLARATION OF CONFORMITY

MERTOL GRUP TEKSTİL FERMUAR SANAYİ VE ANONİM ŞİRKETİ
SANAYİ MAH. GÜRBÜZ SOK. NO:13/1 GÜNGÖREN / İSTANBUL / TÜRKİYE

In our delivered version, we declare that the product described below complies with the essential safety and health requirements of the Medical Devices Directive 93/42/EEC-General Product Safety Directive 2001/95/EEC regulations as circulating by us. This declaration will cease to be valid if the product specified below is replaced.

Teslim edilen versiyonumuzda aşağıda açıklanan ürünün, temel güvenlik ve sağlığını koruma amaçlarıyla Tıbbi Cihazlar Direktifi 93/42/EEC Genel Ürün Güvenliği Direktifi 2001/95/EEC yönetmeliklerinin temel güvenlik ve sağlık gerekliliklerine uygun olduğunu beyan ederiz. Aşağıda bilgileri belirtilen ürünün değiştirilmesi halinde bu beyan geçerliliğini yitirecektir.

Description Of The Product / Ürünün Tanımı : OVERALLS (DISPOSABLE)
TULUM (TEK KULLANIMLIK)

Product Type / Ürün Tipi : NON STERILE

Product Commercial Brand / Marka : MERTOL

Applicable EC Directives / Geçerli AT Direktifleri : GENERAL PRODUCT SAFETY DIRECTIVE 2001/95/EEC
MEDICAL DEVICES DIRECTIVE 93/42/EEC
GENEL ÜRÜN GÜVENLİĞİ DİREKTİFİ 2001/95/EEC
TIBBİ CİHAZLAR DİREKTİFİ 93/42/EEC

Applicable Harmonised Standards / Geçerli Uyumlaştırılmış Standartlar : TS EN 14325, TS EN 464, EN ISO 13485

Applicable National Technical Standards and Specifications / Uygulanabilir Ulusal Teknik Standartlar ve Özellikler : CLASS-I / NONSTERILE
Sınıflandırma : 20060210
Certificate Number / Sertifika Numarası : MERTOL GRUP
Certificate Code / Sertifika Kodu : 01.06.2020
Certificate Issue Date / Sertifika Yayın Tarihi : 01.06.2021
Certificate Validity Date / Sertifikanın Geçerlilik Tarihi : 01.06.2021
Company Official / Firma Yetkilisi : (Authorized Signature and Title) / (Yetkili İmza ve Unvan)

MERTOL GRUP TEKSTİL FERMUAR SANAYİ VE ANONİM ŞİRKETİ declares that the Medical Devices Directive 93/42/EEC-General Product Safety Directive 2001/95/EEC has fulfilled the applicable requirements and responsibility has been taken for the above-described product group. The product groups described above have been controlled depending on internal production controls.

MERTOL GRUP TEKSTİL FERMUAR SANAYİ VE ANONİM ŞİRKETİ olarak yukarıda tanımlanmış olan ürün grupları için Tıbbi Cihazlar Direktifi 93/42/EEC Genel Ürün Güvenliği Direktifi 2001/95/EEC' nin uygulanabilir gerekliliklerine yerine getirilip ve sorumluluğun altına alındığını beyan ederiz. Yukarıda tanımlanmış ürün grupları, iç üretim kontrolüne bağlı olarak kontrol edilmektedir.

www.mertolgrup.com.tr info@nvaquality.com

CERTIFICATE of Registration



This is to Certify that the
Medical Devices – Quality Management System

of
**MERTOL GRUP TEKSTİL FERMUAR SANAYİ
VE ANONİM ŞİRKETİ**
SANAYİ MAH. GÜRBÜZ SOK. NO:13/1 GÜNGÖREN /
İSTANBUL / TÜRKİYE

has been independently assessed and is compliant
with the requirements of

ISO 13485:2016

This Certificate is applicable to the following product or service ranges:

PRODUCTION, SALES AND EXPORT OF SINGLE USE FACE MASK,
SURGICAL AND MEDICAL SINGLE USE GOWNS, SINGLE
USE OVERALLS, SINGLE USE GLOVES

TEK KULLANIMLIK YÜZ MASKESİ, CERRAHİ VE MEDİKAL TEK
KULLANIMLIK ÖNLÜK, TEK KULLANIMLIK TULUM, TEK
KULLANIMLIK ELDİVEN ÜRETİMİ, SATIŞI VE İHRACATI

:: Certificate No :: TR52578H

Date of initial registration 08 June 2020

Date of this Certificate 08 June 2020

Surveillance audit on or before 07 June 2021

Recertification Due / Certificate expiry 07 June 2023

This Certificate is property of Staunchly Management & System Services Ltd. and remains valid
subject to satisfactory surveillance audits.



Director
STAUNCHLY MANAGEMENT & SYSTEM SERVICES LTD.
Suite 4E, B9-90 Hatton Garden, London, EC1N 8PN,
Phone: +44 345 680 0190

Email: info@staunchlyservices.com Web: www.staunchlyservices.com

For precise and updated information concerning the present certificate mail to: info@staunchlyservices.com

This Certificate is the property of Staunchly Management & System Services Private Limited and shall be returned immediately when demanded

2020



THANK YOU!

Email: Orders@inngeninc.com