
mediroc

TECHNICAL DATASHEET

mediroc

KID'S FACE MASK

3-PLY EARLOOP

GENERAL INFORMATION

MANUFACTURER

Name: **STL Teknoloji Ltd. Şti.**

Add: **Catalcesme Mah. Resadiye Cad. 186 Sok. No: 1 8 Alemdag, Cekmeköy, İstanbul/TURKEY**

Authorized Representative: **Mediroc Tech LTD.**

CONFORMITY ASSESSMENT PROCEDURE

According to Regulation (EU) 2017/745 Article 52, the manufacturer follows the conformity assessment procedure relating to the EU declaration of conformity referred to in Article 19 after drawing up the technical documentation set out in Annexes II and III.

NOTIFIED BODY

No involvement of a Notified Body is needed for this Non-Sterile class I device.

PRODUCT INFORMATION

INTENDED USE

- Low-risk clinical applications that do not involve blood-borne pathogens or bodily fluids
- Enhancing infection control
- Preventing the risk of cross-contamination

DESCRIPTION

Rectangular face masks with a shapeable nose piece and two earloops present, one on each side, in order to hold mask in place.

Trade Mark: **Mediroc**

Model: **STL3PLYKIDS**

This product is ***Type II*** mask according to European Standard EU: BS EN 14683:2019

MATERIAL

Outside Layer: **Spunbond Polypropylene – SBPP (White)**
Middle Layer: **Meltblown Polypropylene – MBPP (White)**
Inner Layer: **Spunbond Polypropylene – SBPP (White)**
Nose piece: **Plastic covered iron**
Elastic Band: **Polyester**
Not formulated with Natural Rubber Latex (Latex Free)
Not formulated with DEHP
Fiberglass Free Product

MASK DIMENSIONS

Length: **175mm Width 95mm**
Length of ear loop: **170mm**
Length of nose piece: **90mm**

MANUFACTURING

This mask is made in Turkey.

REGULATION & TESTING INFORMATION

REGULATORY INFORMATION

Product CE marked as per 93/42/EEC Directive on Medical Devices.

Class 1 Medical Device - Type II - Non-Sterile

TEST METHODS

Bacterial Filtration Efficiency (BFE)

When tested in accordance with Annex B of EN 14683, the BFE of the medical face mask shall conform to the minimum value given for the relevant type in Table 1 of EN 14683.

Breathability

When tested in accordance with Annex C of EN 14683, the differential pressure of the medical face mask shall conform to the value given in relevant type in Table 1 of EN 14683.

Microbial cleanliness (Bioburden)

The bioburden of the medical face mask shall be ≤ 30 cfu/g tested. The number of masks that shall be tested is minimum 5 of the same batch/lot.

The number of masks that shall be tested is minimum 5 of the same batch/lot.

Differential Pressure

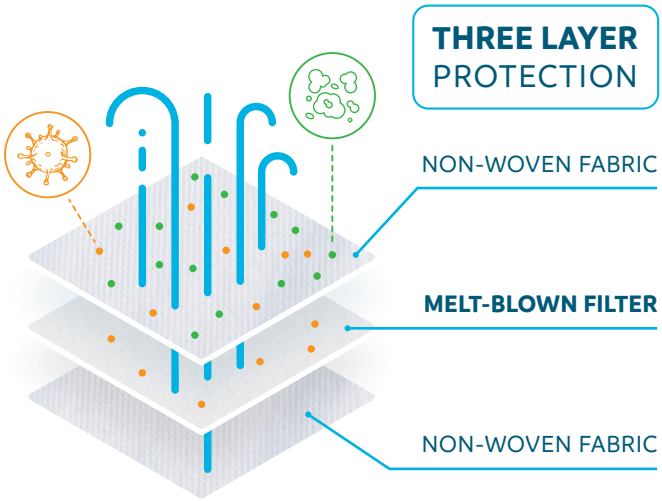
The differential pressure of the medical face mask shall be ≤ 40 cfu/g tested. The number of masks that shall be tested is minimum 5 of the same batch/lot.

TEST	TYPE I	TYPE II	TYPE IIR
Bacterial Filtiration Efficiency(BFE), (%)	≥95	≥98	**≥98**
Differantial Pressure (Pa/cm2)	<40	<40	**<60**
Splash Resistance Pressure (kPa)	Not Required	Not Required	**≥16,0**
Microbial Cleanliness (cfu/g)	≤30	≤30	**≤30**

Table – Performance Requirements for Medical Face Masks

STERILIZATION

This mask is non-sterile



STORAGE

Store in a dry and cool place, away from intense sources of heat. Keep the masks as much as practicably possible in their dispenser box. Keep dispenser boxes as much as practicably possible in their shipper box.

PACKING

Shipping case of 1800 units
10 Units are placed within 1 box and 180 boxes are placed within 1 shipping case
Box demision: **210x105x24,5mm**
Box matarial: **350gr Paper**
Shipping case demision: **640x325x500mm**
Shipping case matarial: **Carton**

SHELF LIFE

The shelf-life is 3 years after production.
The uninterrupted use duration of the device is usually less than 8 hours.

BARCODE

8 680096 103438

VISUALS

Disposable Medical

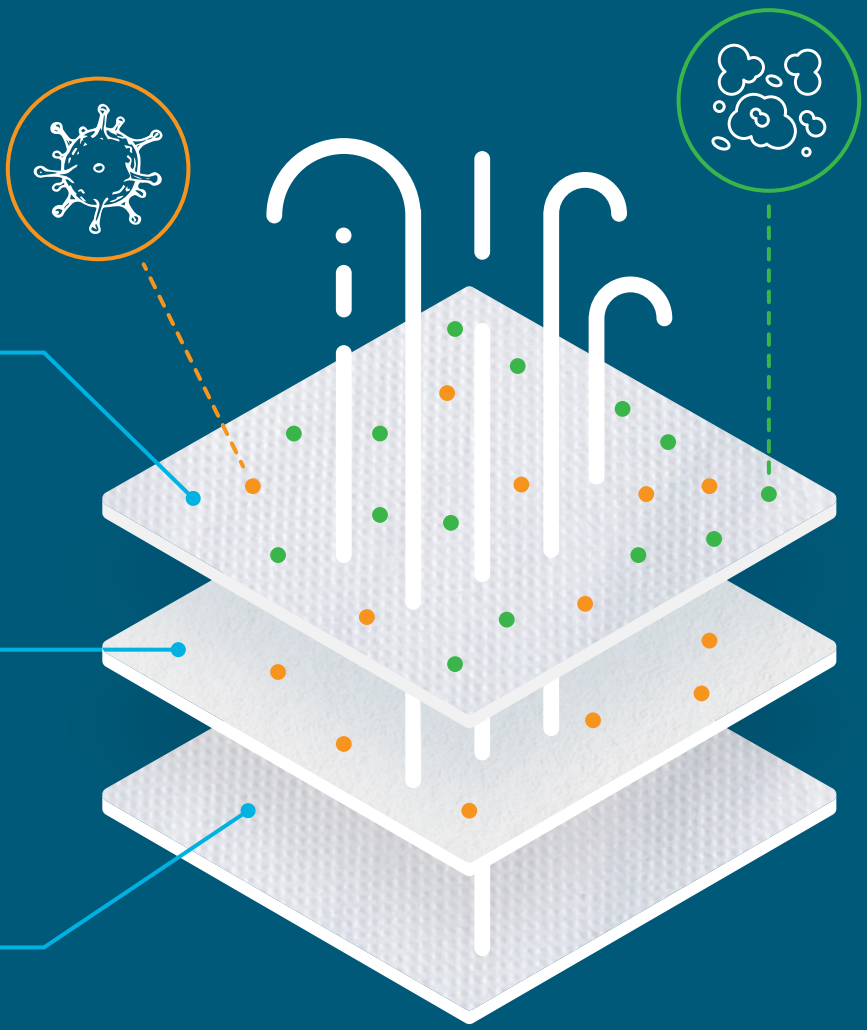
KID'S FACE MASK

DISPOSABLE MEDICAL

FACE MASK

3-PLY EARLOOP

THREE LAYER PROTECTION



TEST REPORT

Class 1 Medical Device

Standard EU: BS EN 14683:2019

ANALYSIS REPORT

Report No. : **2011194E-R1** Report Date : 02/06/2020

Applicant : STL TEKNOLOJİLTD. ŞTİ.

Address : Alemdağ, Çatalmeşe Mah. Reşadiye Cad, 186. Sk. No:18 Çekmeköy/ İstanbul/ Turkey

Sample : Dısposable Medical Face Mask

Sample Package : Carton box

Sample Amount : 100 adet

Sampling Point : -

Sampling Date : -

Sample Lot No. : -

Sample Carrying Conditions / Preservation Technique : -

Production Date : 05/2020

Packing Date : -

Expire Date : 2023

Producer Company : STL Teknoloji Ltd. Şti.

Sample Receiving Time : 21/05/2020 10:30:00

Analysis Beginning Time : 21/05/2020 10:45:00

Analysis Completion Time : 29/05/2020

Following analysis results were obtained from the specimen which was delivered by cargo to Çevre Laboratory;

Parameters	Unit	Finding	Tip I	Tip II	Tip IIR	LR Source	Method	Information
Differential Pressure								
DP - 1	Pa/cm²	16,73	< 40	< 40	< 60	97	EN 14683 - Annex C 122, 123, 126	
DP - 2	Pa/cm²	19,02	< 40	< 40	< 60	97	EN 14683 - Annex C 122, 123, 126	
DP - 3	Pa/cm²	16,52	< 40	< 40	< 60	97	EN 14683 - Annex C 122, 123, 126	
DP - 4	Pa/cm²	17,1	< 40	< 40	< 60	97	EN 14683 - Annex C 122, 123, 126	
DP - 5	Pa/cm²	15,92	< 40	< 40	< 60	97	EN 14683 - Annex C 122, 123, 126	
Bacterial Filtration Efficiency								
BFE - 1	%	99,6	≥95	≥98	≥98	97	EN 14683 - Annex B 122, 124, 129	
BFE - 2	%	99,6	≥95	≥98	≥98	97	EN 14683 - Annex B 122, 124, 129	
BFE - 3	%	99,6	≥95	≥98	≥98	97	EN 14683 - Annex B 122, 124, 129	
BFE - 4	%	99,9	≥95	≥98	≥98	97	EN 14683 - Annex B 122, 124, 129	

Merve BİRAH
Assistant Laboratory Responsible of Microbiology Laboratory

Approved by
02/06/2020
Ömer Yasin BALIK
Laboratory Manager

ANALYSIS REPORT

Report No. : **2011194E-R1** Report Date : 02/06/2020

Following analysis results were obtained from the specimen which was delivered by cargo to Çevre Laboratory;

Parameters	Unit	Finding	Tip I	Tip II	Tip IIR	LR Source	Method	Information
BFE - 5	%	99,6	≥95	≥98	≥98	97	EN 14683 - Annex B	122, 124, 129
Mean Positive Control Count	cfu	2395	-	-	-	-	EN 14683 - Annex B	
Negative Control Count	cfu	<1	-	-	-	-	EN 14683 - Annex B	
Mean Particle Size (MPS)	µm	2,9	-	-	-	-	EN 14683 - Annex B	
Microbial Limit - Bioburden								
Bioburden - 1	cfu/g	22	≤30	≤30	≤30	97	ISO 11737-1	120, 131
Bioburden - 2	cfu/g	16	≤30	≤30	≤30	97	ISO 11737-1	120, 131
Bioburden - 3	cfu/g	19	≤30	≤30	≤30	97	ISO 11737-1	120, 131
Bioburden - 4	cfu/g	16	≤30	≤30	≤30	97	ISO 11737-1	120, 131
Bioburden - 5	cfu/g	22	≤30	≤30	≤30	97	ISO 11737-1	120, 131

Source of Limit Ranges : 97 Medikal Yüz Maskelerinin Test Metodları ve Performans Gereksinimleri (EN 14683)

A: Acceptable NA: Not Acceptable

MU: Measurement Uncertainty

Method
EN : European Standard
ISO : International Organization for Standardization

Information
120 : Bioburden : Aerobic Bacteria and Mold-Yeast
Pozitive Controls : Bacillus atrophaeus
Extract Fluid : Peptone, Tween with Sodium Chloride
Extract Fluid Volume : 300 mL
Plating Method : Membrane Filtration
Agar Medium : Tryptic Soy Agar for Aerobic Bacteria Count and Sabouraud Dextrose Agar with Chloramphenicol for Mold and Yeast Count
Recovery Efficiency : Repetitive Rinse Method
Aerobic Bacteria : Plates are incubatede 3 days at 30-35°C, then enumerated.
Yeast - Mould : Plates are incubatede 5-7 days at 20-25°C, then enumerated.

122 : Conditioning Parameters : 85± 5 relative humidity and 21± 5 °C de minimum 4 hours

123 : Flow rate during testing : 8 L/dk

124 : Flow rate during testing : 28.3 L/dk

126 : The mask analyzed according to the results of Differential Pressure provides EN 14683 Table 1. Type I, Type II and Type IIR limits.

129 : The mask analyzed according to the results of Bacterial Filtration Efficiency (BFE) provides EN 14683 Table 1. Type I,Type II and Type IIR limits.

131 : The mask analyzed according to the results of Microbial Limit - Bioburden provides the EN 14683 Table 1. Type I, Type II and Type IIR limits.

R1 : This report supersedes 29/05/2020 date 2011194E number of report which is invalid.

Merve BİRAH
Assistant Laboratory Responsible of Microbiology Laboratory

Approved by
02/06/2020
Ömer Yasin BALIK
Laboratory Manager

Çevre Endüstriyel Analiz Laboratuvarı Hiz. Tic. A.Ş. Merkez Mahallesi Tatlıpınar Sokak No:13 Mart Plaza Kat:2/A Kağıthane / İstanbul

ANALYSIS REPORT

Report No. **: 2011194E-R1** Report Date **:** 02/06/2020

Note

1. When request, the conformit assessment is carried out in accordance with the legal regulations and standards or the decision rules which are agreed with the customer.
2. Descriptive information about the samples / sampling in the analysis report has been declared by the customer. Our laboratory is not responsible for the legal losses.
3. Analysis report covers samples/sampling that comes to the laboratory.
4. This report and results don't not be copied and printed partially or completely without permission of Cevre Industrial Analysis Laboratory for any commercial and advertising purposes.
5. This report shall not be used official purposes related to Enviromental Regulations.
6. The test report without sign is not valid.

End of Report

Merve BİRAH
Assistant Laboratory Responsible of Microbiology Laboratory

Approved by
02/06/2020
Ömer Yasin BALIK
Laboratory Manager

Çevre Endüstriyel Analiz Laboratuvarı Hiz. Tic. A.Ş. Merkez Mahallesi Tatlıpınar Sokak No:13 Mart Plaza Kat:2/A Kağıthane / İstanbul

EU DECLARATION OF CONFORMITY & CERTIFICATION

EU DECLERATION OF CONFORMITY

Mediroc Disposable Medical Face Mask

Name and address of manufacturer	:	STL TEKNOLOJİ LTD. ŞTİ. Alemdağ, Çatalçeşme Mah. Reşadiye Cad. 186. Sok. No:18 Çekmeköy/İstanbul/Turkey
Product Name	:	Disposable Medical Face Mask
Brand Name	:	mediroc
Product Types	:	Type II, Non-Sterile
Authorized Representative	:	Mediroc Tech Ltd.

This declaration confirms that the product meets the essential requirements of following directive(s) and standart(s). The conformity was based on;

Applied Directive(s)	:	Medical Devices Directive 93/42/EEC as amended according to the Directive 2007/47/EC
Applied Standard(s)	:	EN 14683:2019 Medical Face Masks - Requiremenst and test methods
International Standards	:	ISO 13485:2016 / ISO 9001:2015(QMS)

The declaration has been carried out in accordance with individual rules and conditions. Evaluation has been carried out in accordance with:

Test Report(s) No	:	20011194E-R1
Test Conducted by	:	Cevre Industrial Analysis Laboratory
Test Lab. Adress	:	Merkez Mahallesi Tatlıponar Sokak No: 13 Mart Plaza Kat: 2/A Kağıthane/İstanbul
Issue Date	:	02/06/2020
Revision Date/No	:	-

* The undersinged herewith declarer that the above-mentioned product(s) meet the provisions of the following EC Council Directives and harmonized standards, All supporting documentations are retained under the premises of the manufacturer.

İstanbul/Turkey
05.06.2020

General Manager

STL TEKNOLOJİ LTD. ŞTİ.
Çatalmeşe Mah. Reşadiye Cad. 186 Sk. No: 18
Alemdağ - Çekmeköy - İstanbul - TÜRKİYE
Sarıgazi V.D.:773 030 1141 Ltd. Şti. No: 575043
Mersis No: 0773030116100010
http://benceiyi.com - info@benceiyi.com
Tel.: 02163145521 Fax.: 02163145523

SYNDICATE OF INTERNATIONAL SYSTEM CERTIFICATIONS

CERTIFICATE OF COMPLIANCE

This Certificate has been awarded to

STL TEKNOLOJİ LİMİTED ŞİRKETİ

ÇATALMEŞE MAH. REŞADİYE CAD. 186. SOK. NO:18
ÇEKMEKÖY / İSTANBUL / TÜRKİYE

In recognition of the organization's Managements System which complies with

EN 14683:2019+AC:2019

The scope of activities covered by this certificate is defined below

MANUFACTURE, SALES AND EXPORTS OF TEXTILE PRODUCTS, MEDICAL PROTECTIVE CLOTHING, MASK, PROTECTIVE OVERALLS, BONNET, GLOVES, OVERSHOE, APRON, SURGERY APRON FOR PATIENTS AND DOCTOR, STRETCHER COVER, DEAD BODY BAG, COLONOSCOPY SHORTS, PATIENT SHORTS, DISINFECTANT LIQUIDS, ANTIBACTERIAL SOAP AND LIQUIDS, SURFACE AND SKIN CLEANING MATERIALS, FACE PROTECTOR VISORS, SUITCASE

TEKSTİL ÜRÜNLERİ, MEDİKAL KORUYUCU KIYAFET, MASKE, KORUYUCU TULUM, BONE, ELDİVEN, GALOŞ, ÖNLÜK, HASTA VE DOKTOR İÇİN AMELİYAT ÖNLÜĞÜ, SEDYE ÖRTÜSÜ, CESET TORBASI, KOLONOSKOPİ ŞORTU, HASTA ŞORTU, DEZENKFEKTAN SIVILAR, ANTİBAKTERİYAL SABUN VE SIVILAR, YÜZEY VE CİLT TEMİZLİK MALZEMELERİ, YÜZ KORUYUCU SİPERLİK, VALİZ ÜRETİMİ, SATIŞI VE İHRACATI

Certificate Number: **SISTURAC052020104**
Date of Issue of Original Certificate: **06.05.2020**
Date of Issue of latest certificate: **27.05.2020**
Expiry Date: **05.05.2021**

Managing Director

Note: This certificate is valid only if produced with the continuation letter after the surveillance is carried out successfully.

The Organization's documentation and Implementation has been reviewed and found to comply with the relevant standard rules. This certificate of Registration is based on the evaluation of the mentioned scope given above. Organization is responsible for maintaining the responsibilities of the relevant standard rules. Any significant changes in the scope of the certification or standard referred above render this certificate invalid.

Corporate office(SIS):- Plot No. 1539, 2nd Floor, Sector-4,Gurgaon-122001, Haryana, India.
International office(SIS):- URB. Santa Ana Cal. German, Scherieber 276, San Isidro, Lima, Peru 15047.
Email us :-support@siscertifications.com, info@siscertifications.co.in. Call:- +91-9654721646
Web:- http://www.siscertifications.co.in, www.siscertifications.com
The status of this certificate can be verified on "http://www.siscertifications.co.in".

Issue No.: **02**

CERTIFICATE OF REGISTRATION

SYNDICATE OF INTERNATIONAL SYSTEM CERTIFICATIONS

This Certificate has been awarded to

STL TEKNOLOJİ LİMİTED ŞİRKETİ

ÇATALMEŞE MAH. REŞADİYE CAD. 186. SOK. NO:18
ÇEKMEKÖY / İSTANBUL / TÜRKİYE

In recognition of the organization's Management System
which complies with

ISO 9001:2015(QMS)

The scope of activities covered by this certificate is defined below

MANUFACTURE, SALES AND EXPORTS OF TEXTILE PRODUCTS, MEDICAL PROTECTIVE CLOTHING, MASK, PROTECTIVE OVERALLS, BONNET, GLOVES, OVERSHOE, APRON, SURGERY APRON FOR PATIENTS AND DOCTOR, STRETCHER COVER, DEAD BODY BAG, COLONOSCOPY SHORTS, PATIENT SHORTS, DISINFECTANT LIQUIDS, ANTIBACTERIAL SOAP AND LIQUIDS, SURFACE AND SKIN CLEANING MATERIALS, FACE PROTECTOR VISORS, SUITCASE

TEKSTİL ÜRÜNLERİ, MEDİKAL KORUYUCU KIYAFET, MASKE, KORUYUCU TULUM, BONE, ELDİVEN, GALOŞ, ÖNLÜK, HASTA VE DOKTOR İÇİN AMELİYAT ÖNLÜĞÜ, SEDYE ÖRTÜSÜ, CESET TORBASI, KOLONOSKOPİ ŞORTU, HASTA ŞORTU, DEZENKFEKTAN SIVILAR, ANTİBAKTERİYAL SABUN VE SIVILAR, YÜZEY VE CİLT TEMİZLİK MALZEMELERİ, YÜZ KORUYUCU SİPERLİK, VALİZ ÜRETİMİ, SATIŞI VE İHRACATI

Certificate Number: **SISTURQ0420202085**
Date of Issue of Original Certificate: **22.04.2020**
Date of Issue of latest certificate: **27.05.2020**
Expiry Date: **22.04.2021**

SYNDICATE OF INTERNATIONAL SYSTEM

Managing Director

Note: This certificate is valid only if produced with the continuation letter after the surveillance is carried out successfully.

The Organization's documentation and Implementation has been reviewed and found to comply with the relevant standard rules. This certificate of Registration is based on the evaluation of the mentioned scope given above. Organization is responsible for maintaining the responsibilities of the relevant standard rules. Any significant changes in the scope of the certification or standard referred above render this certificate invalid. This is an accredited certificate issued by SIS Certifications Pvt. Ltd. sanctioned for issue by International Accreditation Services , 3060 Saturn Street Suite 100 Brea, California 92821-1732, USA.

Corporate office(SIS):- Plot No. 1539, 2nd Floor, Sector-4,Gurgaon-122001, Haryana, India.
International office(SIS):- URB. Santa Ana Cal. German, Scherieber 276, San Isidro, Lima, Peru 15047.
Email us :-support@siscertifications.com, info@siscertifications.co.in. Call:- +91-9654721646
Web:- http://www.siscertifications.co.in, www.siscertifications.com
The status of this certificate can be verified on "http://www.siscertifications.co.in".

Issue No.: **02**

SIS CERT.

IAS ACCREDITED Management Systems Certification Body

IAF MEMBER OF MULTILATERAL RECOGNITION ARRANGEMENT

CERTIFICATE OF REGISTRATION

SYNDICATE OF INTERNATIONAL SYSTEM CERTIFICATIONS

CE

CE ATTESTATION OF CONFORMITY

Related Directives :
MEDICAL DEVICES 93/42/EEC---------TIBBİ CİHAZLAR DİREKTİFİ 93/42/EEC

Class Sınıf: CLASS 1 / SINIF 1, NON STERILE

Description of Product :
MEDICAL MASK
TIBBİ MASKE

Manufactured by

STL TEKNOLOJİ LİMİTED ŞİRKETİ
ÇATALMEŞE MAH. REŞADİYE CAD. 186. SOK. NO:18 ÇEKMEKÖY / İSTANBUL / TÜRKİYE

Certificate No.: SISTURCE052020705
Issue Date (Original): 06.05.2020
Issue Date(Latest): 26.05.2020
Expiry Date: 05.05.2021

CE

This Certificate is issued under the following conditions:

1.It applies only to the above referenced models of the medical devices.
2.It does not imply that the SIS has performed any surveillance or control of their manufacture.
3.The manufacture is obligated to assure conformity of all in medical devices of the respective model to type assessed by the mean of this certificate.
4.The certificate remains valid until the manufacturing condition, the quality system or relevant legislation are changed .
5.After fulfilling of the relevant EU legislation requirements, the manufacture shall affix to each medical device, of the above referenced models, the CE-marketing according to this example:

Managing Director

Note: This certificate is valid only if produced with the continuation letter after the surveillance is carried out successfully.

The Organization's documentation and Implementation has been reviewed and found to comply with the relevant standard rules. This certificate of Registration is based on the evaluation of the mentioned scope given above. Organization is responsible for maintaining the responsibilities of the relevant standard rules. Any significant changes in the scope of the certification or standard referred above render this certificate invalid.

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International office(SIS):- URB. Santa Ana Cal. German, Scherieber 276, San Isidro, Lima, Peru 15047.
Email us :-support@siscertifications.com, info@siscertifications.co.in. Call:- +91-9654721646
Web:- http://www.siscertifications.co.in, www.siscertifications.com
The status of this certificate can be verified on "http://www.siscertifications.co.in".

SIS CERT.

CERTIFICATE
of Registration

This is to Certify that the

Medical Devices – Quality Management System

of

STL TEKNOLOJİ LİMİTED ŞİRKETİ

ÇATALMEŞE MAH. REŞADİYE CAD. 186. SOK. NO:18 ÇEKMEKÖY / 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:

MANUFACTURE, SALES AND EXPORTS OF MEDICAL PROTECTIVE CLOTHING, MASK, PROTECTIVE OVERALLS, BONNET, GLOVES, OVERSHOE, APRON, SURGERY APRON FOR PATIENTS AND DOCTOR, STRETCHER COVER, DEAD BODY BAG, COLONOSCOPY SHORTS, PATIENT SHORTS, DISINFECTANT LIQUIDS, ANTIBACTERIAL SOAP AND LIQUIDS, SURFACE AND SKIN CLEANING MATERIALS, FACE SHIELDS

MEDİKAL KORUYUCU KIYAFET, MASKE, KORUYUCU TULUM, BONE, ELDİVEN, GALOŞ, ÖNLÜK, HASTA VE DOKTOR İÇİN AMELİYAT ÖNLÜĞÜ, SEDYE ÖRTÜSÜ, CESET TORBASI, KOLONOSKOPİ ŞORTU, HASTA ŞORTU, DEZENKFEKTAN SIVILAR, ANTİBAKTERİYEL SABUN VE SIVILAR, YÜZEY VE CİLT TEMİZLİK MALZEMELERİ, YÜZ KORUYUCU SİPERLİK ÜRETİMİ, SATIŞI VE İHRACATI

:: Certificate No :: TR52007H

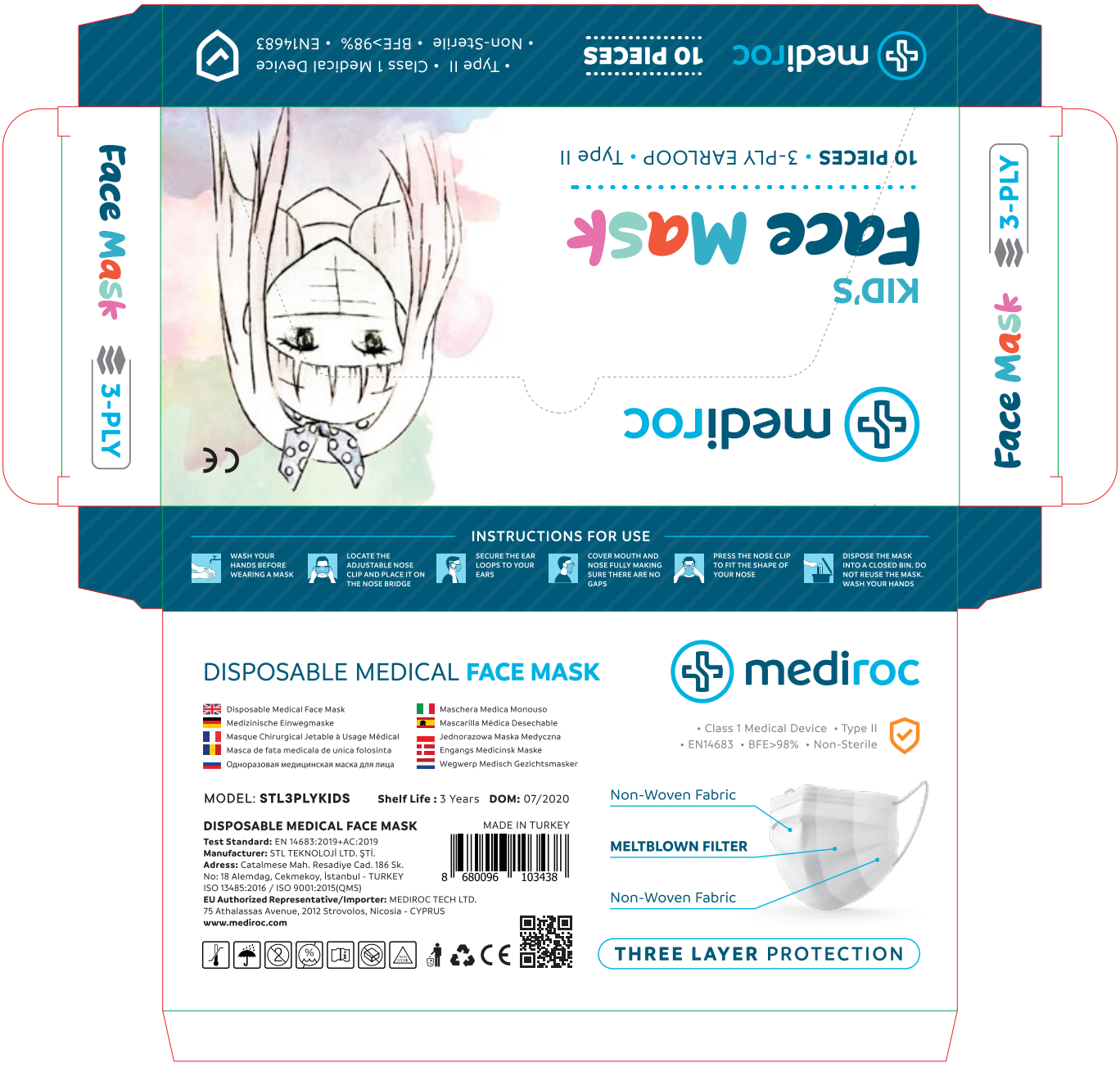
Date of initial registration	**22 April 2020**
Date of this Certificate	**22 May 2020**
Surveillance audit on or before	**21 April 2021**
Recertification Due / Certificate expiry	**21 April 2023**

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 48, 88-90 Hatton Garden, London, EC1N 8PN.

PACKAGING



INSTRUCTIONS FOR USE 1) Wash your hands before wearing a mask. 2) Locate the adjustable nose clip and place it on the nose bridge. 3) Secure the ear loops to your ears. 4) Cover mouth and nose fully making sure there are no gaps. 5) Press the nose clip to fit the shape of your nose. 6) Dispose the mask into a closed bin. Do not reuse the mask. Wash your hands. **ИНСТРУКЦИИ ПО ИСПОЛЬЗОВАНИЮ** 1) Вымойте руки перед тем, как надеть маску. 2) Поместите регулируемый зажим для носа на переносицу. 3) Закрепите ушные петли в удобном положении на ушах. 4) Убедитесь, что маска полностью покрывает область рта и носа. 5) Нажмите на зажим для носа так, чтобы он плотно прилегал. 6) После использования - утилизируйте маску и вымойте руки. Не используйте маску повторно. **MODE D'EMPLOI** 1) Lavez-vous les mains avant de porter un masque. 2) Localisez le clip de nez réglable et placez-le sur le haut du nez. 3) Fixez les élastiques à vos oreilles. 4) Couvrez complètement la bouche et le nez en vous assurant qu'il n'y a pas d oubli. 5) Appuyez sur le pince nez pour l'adapter a la forme de votre nez. 6) Jeter le masque dans une poubelle fermée. Ne pas réutiliser le masque. Lavez-vous les mains après. **INSTRUCCIONES DE USO** 1) Lavese las manos antes de usar una máscara. 2) Ubique la pinza nasal y coloquela en el puente de la nariz. 3) Asegure el arnes a ambos lados de tus oidos. 4) Cubrase la boca y la nariz, asegurando de que no hay huecos. 5) Presione el clip de la pinza nasal para adaptarlo a la forma de su nariz. 6) Deseche la máscara en un cesto cerrado. No reutilice la máscara. Lavese las manos. **ISTRUZIONI PER L'USO** 1) Lavarsi le mani prima di indossare una maschera. 2) Individuare il nasello regolabile e posizionarlo sul ponte del naso. 3) Fissare i passanti per le orecchie. 4) Coprire completamente la bocca e il naso assicurandosi che non ci siano spazi vuoti. 5) Premere il ferma nasale per adattarlo alla forma del naso. 6) Disporre la maschera in un contenitore chiuso. Non riutilizzare la maschera. Lavarsi le mani. **INSTRUKCJA UŻYCIA** 1) umyj ręce przed założeniem maski. 2) umieść zacisk maski na nosie. 3) umocuj maskę na uszach. 4) usta i nos powinny być całkowicie zakryte. 5) użyj zacisku na nosie, aby dopasować maskę do kształtu twojego nosa. 6) użytą maskę wyrzuć do kosza. Nie używaj maski ponownie. Umyj ręce. **GEBRAUCHSANWEISUNG** 1) Waschen Sie Ihre Hände bevor Sie eine Maske anziehen. 2) Suchen Sie den verstellbaren Nasenclip und platzieren Sie ihn auf der Nasenbrücke. 3) Befestigen Sie die Maske mit den Gummibändern an Ihren Ohren. 4) Decken Sie Mund und Nase vollständig ab und stellen Sie sicher, dass kein Luftspalt vorhanden ist. 5) Passen Sie den Nasenclip der Form Ihrer Nase an. 6) Entsorgen Sie die Maske in einem geschlossenen Behälter. Verwenden Sie die Maske nicht mehrmals. Waschen Sie die Hände. **GEBRUIKSAANWIJZING** 1) Was je handen voordat je een masker draagt. 2) Lokaliseer de neusklem en plaats deze streep op de neusbrug. 3) Pak masker bij de elastieken en plaats deze om de oren. 4) De onderkant van je masker spreid je nu open tot onder je kin zodat neus, mond en kin volledig zijn afgedekt en het masker goed aansluit. 5) Druk de neusklem aan zodat het vorm van je neus aanneemt. 6) Gooi het masker in de afvalbak en raak deze niet onnodig aan. Was opnieuw je handen. **BRUGSANVISNING** 1) Vask hænder eller brug håndsprit før du tager ansigtsmaske på. 2) Find den justerbare næseclip og sæt den på næsen. 3) Fastgør ørestroppene til ørerne. 4) Sørg for, at munden og næsen er fuldstændig dækket af masken. 5) Tilpas næseclippen til næsen. 6) Efter brug smides masken væk – helst i en lukket affaldsbøtte. Masken må ikke genbruges. Vask hænder efter brugen af masken. **INSTRUCTIUNI DE FOLOSIRE** 1) Spalati-va mâinile înainte de a folosi masca. 2) Localizați clema de nas reglabilă și atașați o pe podul nasului. 3) Fixati bucle de elastic după urechе. 4) Acoperiți gura și nasul complet. Asigurați-vă ca nu exista spațiu liber. 5) Apăsați clema pentru a se potrivi cu forma nasului. 6) Aruncați masca într-un Coș și închideți coșul.Nu refolosiți. Spalativa pe mâini.

THANK YOU!