






**3A MEDICAL**  
MEDICAL FACE MASK WITH EAR LOOPS  
REF 2001

CE    LOT 200803  
08.2022

 3A Medical Products Co., Ltd.  
Yu An Industrial Park 230001 Jia An  
PEOPLE'S REPUBLIC OF CHINA  
European Representative:  
Address: Effenweide 16, D-20037, Hamburg, Germany  
Luisenpark 20/21, D-20037, Hamburg, Germany  
Tel: +49-40-2011179 Fax: +49-40-2011178

**3A MEDICAL**  
MEDICAL FACE MASK WITH EAR LOOPS

Latex-free

CE     
EN14683 : 2019 TYPE IIR

REF 2001



QTY 50



## 3A Medical Face Mask

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### 1. Basic Information

**Brand:** 3A

**Model:** Latex-free Medical Face Mask

**Product Code:** 2001

**CE standard:** EN 14683:2019

**Classifications:** Type IIR (Non-Sterile)

**Material:** SPP, Melt-blown fabric

**Design:** Earloop

**Size:** 17.5cm\*9.5cm

**Expiration date:** 2 years

**Packaging:**

50pcs per box, 40 boxes per carton, 2000 pcs per carton

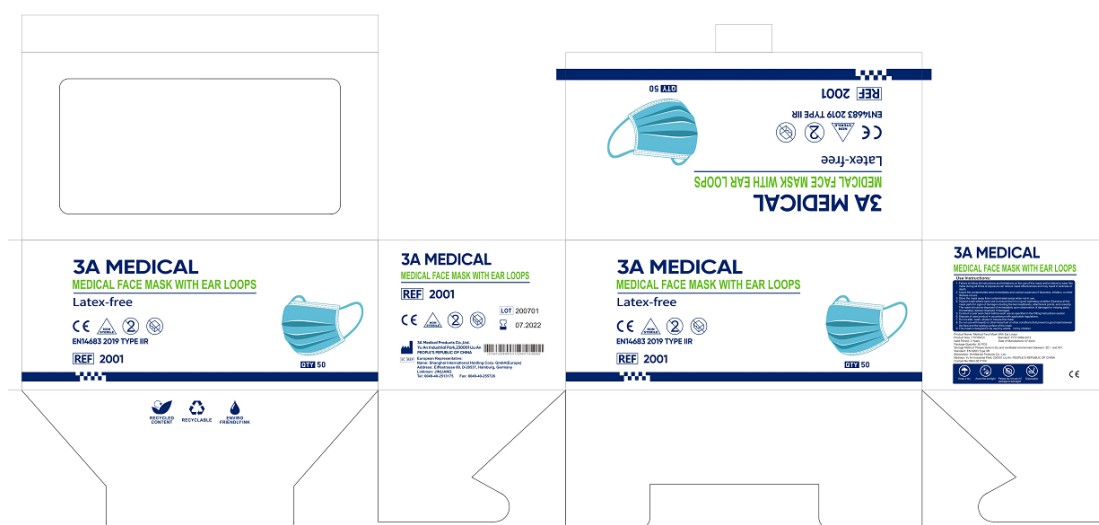
Box Size: 19\*10\*8cm

Carton Size: 52\*39.5\*34cm

Gross Weight: 9.5 kg

### 2. Pictures





## 3A MEDICAL 合格证

( CERTIFICATE OF QUALIFICATION )

产品名称：一次性使用医用口罩	产品品牌：3A Medical
Product: Medical Face Mask With Ear Loops	Brand Name: 3A Medical
产品型号：平面A型	产品代号：2001
Product Type: Plat type A	Product Code: 2001
产品规格：17.5*9.5cm	材料成份：68%SPP无纺布+32%熔喷
Product dimension: 17.5*9.5	Material Ingredient: 68%SPP +32% Melt-blow
生产批号：200701	执行标准：YY/T 0969-2013
Batch Number: 200701	Standard: YY/T 0969-2013
包装数量：50只/盒	生产日期：07. 2020
Package Quantity: 50 pieces/pack	Date of Manufacture: 07. 2020

检验日期：07. 2020  
Inspection Date: 07. 2020

有效期：储存温度-20°C-40°C，储存湿度≤80%，避光干燥的室内环境下，有效期2年。  
Expiry Date: The storage temperature is -20°C-40°C, the storage humidity is ≤80%, and it is valid for 2 years in dry indoor environment.

适用范围：用于佩戴者在不存在体液和飞溅风险的普通医疗环境下的卫生护理。  
Scope Of Application: Used for health care of the wearer in the general medical environment without the risk of body fluids and splashes.

产品注册证编号：皖械注准20202140040

Product Registration Certificate No: Anhui machinery No 20202140040

医疗器械生产许可证号：皖食药监械生产许20180032号

Medical Device production license No: Anhui Food and Drug Administration production license No. 20180032

生产厂商：安徽富美医疗科技有限公司

Manufacturer: 3A Medical Products Co., Ltd

电话：0564-3611700

Te l : 0564-3611700

生产地址：安徽省六安市裕安区城南工业园区工业路1号

Address: Yu An Industrial Park 230001 Liu AN, People' s Republic of China

### 3. Form for Medical Device Registration

Anlage 1  
(zu § 4 Abs. 1 Nr. 1 DIMDIV)  
Formularnummer 00304117

#### Allgemeine Anzeigepflicht nach §§ 25 und 30 Abs. 2 MPG General Obligation to Notify pursuant to §§ 25 and 30 (2) Medical Devices Act, MPG

#### Formblatt für Medizinprodukte, außer In-vitro-Diagnostika Form for Medical Devices except In Vitro Diagnostic Medical Devices

Zuständige Behörde / Competent authority	
Code	DE/CA05
Bezeichnung / Name	Behörde für Gesundheit und Verbraucherschutz, Referat V43
Staat / State	Land / Federal state
Deutschland	Hamburg
Ort / City	Postleitzahl / Postal code
Hamburg	20539
Straße, Haus-Nr. / Street, house no.	
Billstraße 80	
Telefon / Phone	Telefax / Fax
+49-40-428280	+49-40-427310017
E-Mail / E-mail	
medizinprodukte@bgv.hamburg.de	
Anzeige / Notification	
Registrierdatum bei der zuständigen Behörde Registration date at competent authority	Registriernummer / Registration number
Typ der Anzeige / Notification type S Erstanzeige / Initial notification E Änderungsanzeige / Notification of change E Widerrufsanzeige / Notification of withdrawal	
Frühere Registriernummer bei Änderungs- und Widerrufsanzeige Previous registration number if notification has been changed or withdrawn	
Anzeigender nach § 25 MPG / Reporter pursuant to § 25 Medical Devices Act, MPG E Hersteller / Manufacturer S Bevollmächtigter / Authorised Representative E Einführer / Importer E Verantwortlicher für das Zusammensetzen von Systemen oder Behandlungseinheiten nach § 10 Abs. 1 und 2 MPG \ Assembler of systems or procedure packs pursuant to § 10 (1) and (2) Medical Devices Act, MPG E Betrieb oder Einrichtung (aufbereiten) nach § 25 Abs. 1 MPG i. V. m. § 4 Abs. 2 MPBetriebV Institution (processing) pursuant to § 25 (1) Medical Devices Act, MPG in connection with § 4 (2) MPBetriebV E Betrieb oder Einrichtung (sterilisieren) nach § 25 Abs. 2 i. V. m. § 10 Abs. 3 MPG Institution (sterilizing) pursuant to § 25 (2) in connection with § 10 (3) Medical Devices Act, MPG	

<b>Anzeigender / Reporting organisation (person)</b>	
Code	<b>DE/0000040627</b>
Bezeichnung / Name	<b>Shanghai International Holding Corporation GmbH (Europe)</b>
Staat / State	<b>Deutschland</b>
Land / Federal state	<b>Hamburg</b>
Ort / City	<b>Hamburg</b>
Postleitzahl / Postal code	<b>20537</b>
Straße, Haus-Nr. / Street, house no. <b>Eiffestrasse 80</b>	
Telefon / Phone	<b>+49-40-2513175</b>
Telefax / Fax	<b>+49-40-255726</b>
E-Mail / E-mail <b>shholding@hotmail.com</b>	

<b>Hersteller / Manufacturer</b>	
Bezeichnung / Name	<b>3A Medical Products Co., Ltd.</b>
Staat / State	<b>CN</b>
Ort / City	<b>Liu An,</b>
Postleitzahl / Postal code	<b>230001</b>
Straße, Haus-Nr. / Street, house no. <b>Yu An Industrial Park,</b>	
Telefon / Phone	<b>+86-564-3611700</b>
Telefax / Fax	<b>+86-564-3611700</b>
E-Mail / E-mail <b>guweilin@3a-medical.cn</b>	

<b>Sicherheitsbeauftragter für Medizinprodukte nach § 30 Abs. 2 MPG 9) Safety officer for medical devices pursuant to § 30 (2) Medical Devices Act, MPG</b>	
Bezeichnung / Name	<b>Liang Jin</b>
Staat / State	<b>Deutschland</b>
Land / Federal state	<b>Hamburg</b>
Ort / City	<b>Hamburg</b>
Postleitzahl / Postal code	<b>20537</b>
Straße, Haus-Nr. / Street, house no. <b>Eiffestr.80</b>	
Telefon / Phone	<b>+49-40-2513175</b>
Telefax / Fax	<b>+49-40-255726</b>
E-Mail / E-mail <b>shholding@hotmail.com</b>	

<b>Vertreter / Deputy (optional)</b>	
	Bezeichnung / Name
	Telefon / Phone
	Telefax / Fax
	E-Mail / E-mail
	E Erstanzeige / Initial notification S Änderungsanzeige / Notification of change

Medizinprodukt (Erstmaliges Inverkehrbringen) / Medical device (First placing on the market)	
Klasse / Class	
S I	
E I - steril / sterile	
E I - mit Messfunktion / with measuring function	
E I - steril und mit Messfunktion / sterile and with measuring function	
E IIa	
E IIb	
E III	
E III - hergestellt unter Verwendung von Gewebe tierischen Ursprungs im Sinne der Verordnung (EU) Nr. 722/2012 manufactured utilising tissues of animal origin in terms of Commission Regulation (EU) No 722/2012	
E Aktives implantierbares Medizinprodukt / Active implantable medical device	
E Aktives implantierbares Medizinprodukt - hergestellt unter Verwendung von Gewebe tierischen Ursprungs im Sinne der Verordnung (EU) Nr. 722/2012 Active implantable medical device - manufactured utilising tissues of animal origin in terms of Commission Regulation (EU) No 722/2012	
App (Software auf mobilen Endgeräten)	E ja / yes    S nein / no
Nummer(n) der Bescheinigung(en) / Certificate number(s)	
Handelsname des Produktes / Trade name of the device	<b>3A</b>
Produktbezeichnung / Name of device	<b>Medical face mask</b>
Nomenklaturcode / Nomenclature code	<b>12-447</b>
Nomenklaturbezeichnung / Nomenclature term	<b>Maske</b>
Kategoriecode / Category code	<b>10</b>
Kategorie / Category	<b>Produkte zum Einmalgebrauch</b>
Kurzbeschreibung deutsch / German short description	
Kurzbeschreibung englisch / English short description	<b>Medical face mask is intend to be used for hygiene care which the wearer under general medical environment where there is no risk of bodily fluids splash.</b>

<b>Medizinprodukte (Aufbereiten) / Medical devices (Reprocessing)</b>	
	E. Semikritische Medizinprodukte / Semicritical medical devices E. Gruppe A / Group A E. Gruppe B / Group B
	E. Kritische Medizinprodukte / Critical medical devices E. Gruppe A / Group A E. Gruppe B / Group B E. Gruppe C / Group C Nummer der Bescheinigung / Certificate number
	Sterilisationsverfahren / Sterilisation procedures E. Dampfsterilisation / Steam sterilisation E. Gassterilisation / Gas sterilisation E. Strahlensterilisation / Radiation sterilisation E. andere / others Angewandtes Verfahren / Applied procedure

Ich versichere, dass die Angaben nach bestem Wissen und Gewissen gemacht wurden.  
I affirm that the information given above is correct to the best of my knowledge.

Ort  
City

**Hamburg**

Datum  
Date

**2020-04-30**

Name

**Liang Jin**

Unterschrift  
Signature

<b>Bearbeitungsvermerke / Processing notes</b> Nur von der zuständigen Behörde auszufüllen / To be filled in only by the competent authority			
	Bearbeiter / Person responsible		Telefon / Phone



#### 4. Declaration of Conformity



安徽富美医疗科技有限公司

3AMedical Products Co., Ltd.

地址：安徽省六安市裕安区

Add: Yu An Industrial Park, Liu An City, P.R. China

### Declaration of Conformity

Manufacturer: 3A Medical Products Co., Ltd

Address: Yu An Industrial Park, 237100, Liu An, PEOPLE'S REPUBLIC OF CHINA

European Representative:

Name: Shanghai International Holding Corp. GmbH(Hamburg)

Address: Eiffestrasse 80, 20537, Hamburg, Germany

Product name: Medical face mask

Classification: Class 1 Rule 1of Annex IX of MDD 93/42/EEC)

Type: Type IIR

Rule: According to Rule I Annex VII

We here with declare that above mentioned products meet the requirement of the (MDD 93/42/EEC) Medical device directive and the following harmonized standards:

EN14683:2019

ENISO 14971:2012

ENISO 15223-1:2016

ENISO1041:2008A1:2013

ENISO10993-1:2009/AC: 2010

ENISO10993-5:2009

ENISO10993-10:2013

The Medical face mask meets the requirement of EN14683:2019

Signature

Name: Billy Zhang

Position: Management Representative

Place: Liu An City

Date: 2020-04-17



# CERTIFICATE OF NOTIFICATION

This is to certify that, according to European Council Directive 93/42/EEC, Shanghai International Holding Corp, GmbH (Europe), performed all notification duties and responsibilities as the European authorized Representative:

**MANUFACTURER: 3A Medical Products Pty Ltd**

**Address: Yu An Industrial Park, 230001, Liu An, CN**

The manufacturer has provided Shanghai International Holding Corp, GmbH (Europe), with all the appropriate declaration according to the European Council Directive 93/42 EEC including the EC Declaration of Conformity confirming that its medical device, as stipulated here below, is fulfilling the essential requirements of the European Council Directive 93/42/EEC.

Devices: Medical Face Mask

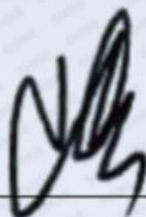
Classification: I

Model: 17.5(±5%)cm\*9.5(±5%)cm, 14.5(±5%)cm\*9.5(±5%)cm

Where the manufacturer affix the CE mark to the device listed they must ensure that all the essential requirements of European Council Directive 93/42/EEC are met.

The notification of abovementioned device has been completed by the European Authorized Representative in Germany. The Germany Competent Authority is notified of the manufacture's device and has allocated registration.

EXECUTIVE  
DIRECTOR



Shine

## 5. Full Analysis Report



**3A Medical Products Co. Ltd, Yu An  
Industrial Park, 230001 Liu An. PR  
China**

**Your notice of**  
17-03-2020

**Your reference**

**Date**  
03-04-2020

### Analysis Report 20.01605.03

Required tests :

EN 14683 (2019) + AC (2019)	EN 14683 - annex B (2019) + AC (2019)	Bacterial filtration efficiency
EN 14683 (2019) + AC (2019)	ISO 22609 (2004)	Medical face masks - Splash Test
EN 14683 (2019) + AC (2019)	EN 14683 - annex C (2019) + AC (2019)	Medical face masks - Breathability (differential pressure)
EN 14683 (2019) + AC (2019)	EN 14683 - §5.2.5 (2019) AC (2019)	Microbial cleanliness on masks

Identification number	Information given by the client	Date of receipt
T2006058	REF 2001 Lot 202001202	17-03-2020

Sylvie Niessen  
Order responsible

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INRICHTING ERKEND BIJ TOEPASSING VAN DE BESLUITWET VAN 30 JANUARI 1947 / ETABLISSEMENT RECONNU PAR APPLICATION DE L'ARRÊTÉ-LOI DU 30 JANVIER 1947



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VAT BE 0459.218.289 • IBAN BE44 2100 4729 6545 • BIC GEBABEBB



Analysis Report 20.01605.03  
Date 03-04-2020  
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**Reference: T2006058 - REF 2001**  
**Lot 202001202**

**Bacterial filtration efficiency**

Date of ending the test	25-03-2020
Standard used	EN 14683 - annex B (2019) + AC (2019)
Product standard	EN 14683 (2019) + AC (2019)
Mask description	3 layers/ 18gsm SPP white/ 25gsm/meltex/ 22gsm SPP blue
Number of tested masks :	5
BFE Area tested :	± 49 cm <sup>2</sup>
Masks conditioning :	21 ± 5°C and 85 ± 5% RH
Side of the mask in contact with the bacterial challenge :	Inner side
Challenge bacterial strain used :	<i>Staphylococcus aureus</i> ATCC6538
Bacterial challenge per test :	1700 - 3000 CFU
Total test time :	1 min. delivering challenge + 1 min. without challenge (air flow continuing)
Flow rate :	28.3 l/min.
Positive control	Tests performed with no filter material in the air stream
Negative control	Test performed without challenge



### Results

B = Bacterial filtration efficiency (%)

$$B = \frac{(C - T)}{C} \times 100$$

With C = mean of the total plate counts for the positive control runs  
T = total count for the tested mask

# Mask	B (%)
1	99.9
2	99.2
3	99.8
4	99.9
5	99.8

Mean particle size of the bacterial challenge aerosol : 2.9 µm

### Controls

Mean positive controls 2636 CFU  
Negative control < 1 CFU



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Date 03-04-2020  
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**Reference: T2006058 - REF 2001**  
**Lot 202001202**

**Medical face masks - Splash Test**

Date of ending the test	26-03-2020
Standard used	ISO 22609 (2004)
Product standard	EN 14683 (2019) + AC (2019)
Mask description	3 layers/ 18gsm SPP white/ 25gsm/meltex/ 22gsm SPP blue
Number of tested masks :	32
Blood surface tension	42 ± 2 dynes/cm
Volume of the delivered blood	2 ml
Distance "canula-mask"	30 ± 1 cm
Side of the mask "impacted"	Outer side
Masks conditioning :	21 ± 5°C and 85 ± 5% RH

**Results**

**Blood pressure tested 16.0 kPa**

**Controls**

Blood visualisation on the mask	OK
Calibration procedure	OK
Control of the blood volume delivered (2 ml)	
- before the test :	OK
- after 16 masks :	OK
- after 32 masks :	OK



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Date 03-04-2020  
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INRICHTING ERKEND BIJ TOEPASSING VAN DE BESLUITWET VAN 30 JANUARI 1947 / ETABLISSEMENT RECONNU PAR APPLICATION DE L'ARRÊTÉ-LOI DU 30 JANVIER 1947

Results obtained on the set of masks

<b># Mask</b>	<b>Results : pass / fail</b>
1	Pass
2	Pass
3	Pass
4	Pass
5	Pass
6	Pass
7	Pass
8	Pass
9	Pass
10	Pass
11	Pass
12	Pass
13	Pass
14	Pass
15	Pass
16	Pass
17	Pass
18	Pass
19	Pass
20	Pass
21	Pass
22	Pass
23	Pass
24	Pass
25	Pass
26	Pass
27	Pass
28	Pass
29	Pass
30	Pass
31	Pass
32	Pass

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Performed in the microbiological lab under the responsibility of Yvette Rogister



Analysis Report 20.01605.03  
Date 03-04-2020  
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Summary P = 16.0 kPa

Number of "Pass" masks	Number of "Fail" masks
32	0

Pass = no blood detected on the observed side

Fail = blood detected on the observed side

In agreement with the customer the number of tested mask has been determined based on a single sampling plan providing an AQL of 4 % (acceptable quality limit).

If 29 masks or more over 32 obtain a "Pass" result the 4% AQL is reached.





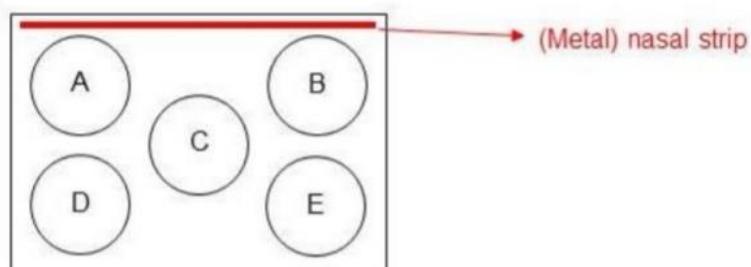
Analysis Report 20.01605.03  
Date 03-04-2020  
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**Reference:** T2006058 - REF 2001  
Lot 202001202

**Medical face masks - Breathability (differential pressure)**

Date of ending the test	24-03-2020
Standard used	EN 14683 - annex C (2019) + AC (2019)
Product standard	EN 14683 (2019) + AC (2019)
Mask description	3 layers/ 18gsm SPP white/ 25gsm/meltex/ 22gsm SPP blue
Number of tested masks :	5
Number of areas per mask	5 (see figure)
Dimension of the areas :	Disc whose diameter is 2.5 cm
Surface areas :	4.9 cm <sup>2</sup>
Flow rate :	8 l/min.
Direction of the air flow :	From the inside of the mask to the outside
Masks conditioning :	21 ± 5°C and 85 ± 5% RH

Figure : Distribution of the areas in the mask





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INRICHTING ERKEND BIJ TOEPASSING VAN DE BESLUITWET VAN 30 JANUARI 1947 / ETABLISSEMENT RECONNU PAR APPLICATION DE L'ARRÊTÉ-LOI DU 30 JANVIER 1947

### Results      $\Delta P$

	Mask 1	Mask 2	Mask 3	Mask 4	Mask 5
Area A	52.6	23.0	52.6	35.2	40.3
Area B	51.1	30.6	59.1	39.7	39.7
Area C	53.0	34.8	45.4	43.4	38.3
Area D	46.7	35.7	48.1	44.2	41.4
Area E	44.2	29.7	39.5	39.7	41.4
<b>Average <math>\Delta P</math> (Pa/cm<sup>2</sup>)</b>	<b>49.5</b>	<b>30.8</b>	<b>48.9</b>	<b>40.4</b>	<b>40.2</b>

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Performed in the microbiological lab under the responsibility of Yvette Rogister



**Reference: T2006058 - REF 2001**  
**Lot 202001202**

**Microbial cleanliness on masks**

Date of ending the test 30-03-2020  
 Standard used EN 14683 - §5.2.5 (2019) AC (2019)  
 Product standard EN 14683 (2019) + AC (2019)

Number of tested masks 5  
 Extraction liquid Peptone 1g/l, NaCl 5g/l & Tween 20 2g/l  
 Extraction volume 300 ml  
 Extraction time 5 min.  
 Counting technique Membrane filtration  
 Filtration volume 100 ml  
 Culture media TSA (Tryptic Soy Agar)  
 SDA (Sabouraud Dextrose Agar with chloramphenicol)

Incubation conditions 3 days at 30°C (TSA)  
 7 days at 20-25°C (SDA)

**Results**

# Mask	Mask weight (g)	CFU*/mask		Microbial cleanliness	
		<i>Aerobic microbial count (bacteria)</i>	<i>Fungi count (SDA)</i>	$\Sigma$ CFU/mask	$\Sigma$ CFU/g
1	4.25	33	<3	< 36	< 9
2	4.22	3	<3	< 6	< 2
3	4.33	3	<3	< 6	< 2
4	4.35	36	<3	< 39	< 9
5	4.37	3	<3	< 6	< 2