Vertrieb durch Micro-Medical GmbH, Am Neumarkt 2-4, 01968 Senftenberg, Tel: 03573-72031-10 oder Mail: info@micromedical.de

HYGISUN®

REF HS0501A

FFP2 Maske / FFP2 Mask

Persönliche Schutzmaske

MICRO-MEDICAL

CE 2797

EN 149: 2001+A1: 2009

CARTON DIMENSION (FFP2 - White - 20er)

HYGISUN®

1 PC/OPP	20 PCS/box	1000 PCS/carton	20000 PCS/pallet

EAN: 4260676530003

CE 2797

MICRO-MEDICAL

MICRO-MEDICAL

Bedienungsanleitung
Fitting instructions

Nehmen Sie die Maske an den Ohrschlaufen in die Hand und drücken Sie diese mit dem Bügel auf den Nasenrücken gegen Ihr Gesicht, während Sie die Ohrschlaufen hinter Ihren Ohren positionieren.
Take the mask by the ear loops in your hand and press it against your face with the strap on the bridge of your nose while you position the ear loops behind your ears.

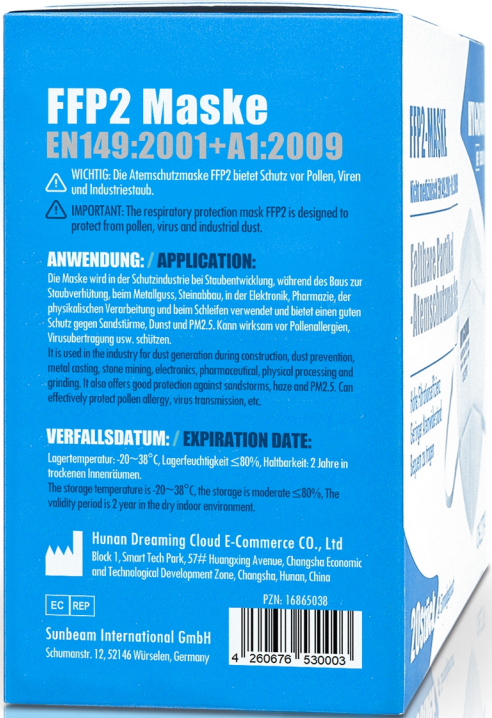
Formen Sie den Bügel mit beiden Händen in die Form Ihrer Nase.
Shape the nose clip into the shape of your nose with both hands.

Testen Sie die Passform. Nehmen Sie beide Hände über die Atemschutzmaske und atmen Sie kräftig aus. Wenn Luft um Ihre Nase strömt, ziehen Sie den Bügel fester.
Test the correct fit. Put both hands over the respirator and exhale forcefully. When air flows out around your nose, press the nose clip tighter.

HINWEIS ZUR VERWENDUNG: / NOTICE FOR USE:

- Bitte verwenden Sie dieses Produkt nicht in der Nähe einer Feuerquelle.
  Please do not use this product near fire sources.
- Da es sich bei diesem Produkt um eine Einwegmaske handelt, kann es nicht durch Waschen wiederverwendet werden.
  As this product is a disposable mask, it cannot be reused through washing.
- Von hohen Temperaturen und Luftfeuchtigkeit fernhalten und an einem sauberen Ort aufbewahren.
  Keep it away from high temperature and humidity and keep it in clean place.
- Persönliche Schutzmaske, Nicht medizinisch.
  personal protective mask, non medical.
- Verwenden Sie einzeln verpackte Produkte, sobald diese ausgepackt sind.
  Use individually packaged products as soon as they are unpacked.

MICRO-MEDICAL



MICRO-MEDICAL

MICRO-MEDICAL

MICRO-MEDICAL


Persönliche Schutzmaske
PERSONAL PROTECTIVE MASK
1Stück/ Piece
MICRO-MEDICAL

HYGISUN

QUALIFIED CERTIFICATE

产品名称	FFP2个人防护口罩(不作为医疗防护使用)
Product Name	FFP2 personal protective mask.(For personal protective only)
执行标准	EN149:2001 + A1:2009
Standard	EN149:2001 + A1:2009
产品规格	16×10.5厘米
Product Size	16×10.5CM
材质	30%熔喷布+70%无纺布
Material	30% MELT-BLOWN FABRIC+70% NON-WOVEN FABRIC
质检员	
Checker	
生产日期	
Date	QC:01
生产批号	PD:SEE ON PACKAGE
Lot code	LOT:SEE ON PACKAGE
使用周期	一次性
Usege count	Single use
有效期	2年
Period of validity	2 years
生产企业	湖南云想生活电子商务有限公司
Manufacturer	Hunan Dreaming Cloud E-Commerce CO., Ltd
生产地址	长沙经济技术开发区黄兴大道南段57号星为创芯园1栋501号
Address	Block 1, Smart Tech Park, 57# Huangxing Avenue, Changsha Economic and Technological Development Zone, Changsha, Hunan, China

QUALITY PASSED
检验 合格
HUNANDREAMINGCLOUDE-COMMERCECO.,LTD

储存条件：本品应储存温度-20~38℃，储存湿度≤80%避光干燥的室内环境下，通风良好，无腐蚀性气体的清洁环境内，如贮存不当导致发霉变质禁用。

STORAGE CONDITIONS AND METHODS: The storage temperature is -20~38℃, and the relative humidity is not more than 80%. Well ventilated and clean environment without corrosivegais, please do not use if the product gets mildewed or deteriorated due to improper storage.

本产品为个人防护用品，不作为医疗防护使用

THIS PRODUCT IS UNDER PERSONAL PROTECTIVE EQUIPMENT DIRECTIVE (PPE).FOR PERSONAL PROTECTIVE ONLY

MADE IN CHINA

MICRO-MEDICAL

HYGISUN®

ANLEITUNG

Norm:

Dieses Produkt entspricht der Norm EN149:2001 + A1:2009 für Atemschutzgeräte – Halbmaske zur Filterung zum Schutz vor Partikeln. Diese Filtermasken sind gemäß der Verordnung der Europäischen Kommission (EU) 2016/425 über PSA als Persönliche Schutzausrüstung in der Kategorie III eingestuft und entsprechend gekennzeichnet

Bestimmungsgemäße Verwendung:

Die Staubmaske ist als Kategorie FFP 2 eingestuft. Sie schützt vor Partikeln, Nebel, Rauch und Aerosolen auf Ölbasis. Die Verpackung schützt die Maske vor der Verwendung. Schützt wirksam vor Pollen. Die Maske kann nur zum persönlichen Schutz verwendet werden, nicht für medizinische Zwecke. Maske nicht bei der Brandbekämpfung und in explosionsgefährdeten Bereichen nutzen.

Dichtsitztest

1.Bedecken Sie die Maske vorsichtig mit beiden Händen ohne den Dichtsitz zu verändern.
2.stark Ausatmen;
3.Bei einer Leckage im Nasenbereich, den Nasenbügel neu anpassen. Dichtsitzprüfung wiederholen.
4.Bei einer Leckage am Maskenrand, den Sitz der Bänder überprüfen und anpassen. Dichtsitzprüfung wiederholen.

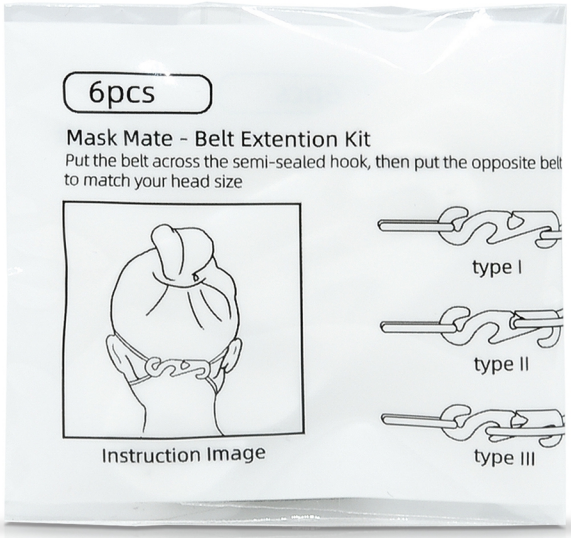
Wenn Sie KEINEN richtigen Dichtsitz erreichen können, betreten Sie NICHT den Gefahrenbereich. Informieren Sie ihren Vorgesetzten.

Warnungen und Einschränkungen:

- Vergewissern Sie sich immer, dass das Produkt:
  Geeignet ist für die Anwendung;
  Korrekt angelegt ist;
  Während des gesamten Aufenthalts im Gefahrenbereich getragen wird;
  Ersetzt wird, wenn notwendig.
- Richtige Auswahl, Schulung, Gebrauch und gegebenenfalls Reinigung sind die Voraussetzungen dafür, dass das Produkt den Anwender vor bestimmten luftgetragenen Gefahrstoffen schützt.
- Die Nichtbefolgung aller Anweisungen zur Anwendung der Maske und/oder die Fehlbenutzung während des Aufenthaltes im Gefahrenbereich kann die Gesundheit des Anwenders beeinträchtigen und zu schweren Erkrankungen oder Dauerschäden führen.
- Beachten Sie bei der Auswahl und richtigen Anwendung nationale Bestimmungen und alle mitgelieferten Informationen.
- Vor Gebrauch muss der Anwender, in Übereinstimmung mit den nationalen Regeln, in der funktionsgerechten Handhabung geschult sein.
- Dieses Produkt schützt nicht vor Gasen und Dämpfen.
- Verwenden Sie die Maske nicht in Umgebungen mit weniger als 19,5% Sauerstoff
- Verwenden Sie die Masken nicht in Umgebungen mit unbekannten Gefahrstoffen oder Konzentrationen, die die zulässigen Höchstwerte übersteigen.
- Verwenden Sie die Maske nicht, wenn Gesichtshaare im Bereich des Dichtrandes einen korrekten Dichtsitz der Maske verhindern.
- Verlassen Sie sofort den belasteten Bereich, wenn:
  a) Das Atmen schwer fällt.
  b) Schwindel oder andere Beschwerden auftreten.
  c) Die Maske beschädigt wird.
  d) Geruch oder Geschmack des Gefahrstoffs oder eine Reizung auftritt.
- Entsorgen und ersetzen Sie die Maske, wenn sie beschädigt ist, der Atemwiderstand stark erhöht ist oder am Ende einer Schicht.
- Die Maske darf niemals verändert oder repariert werden.
- Die Maske ist zum einmaligen Gebrauch vorgesehen und ist danach entsprechend der nationalen Vorgaben zu entsorgen.

Transport und Lagerung:

Die Partikelmasken haben eine Lagerdauer von 2 Jahren. Das Ende der Lagerdauer ist auf der Verpackung angegeben. Vergewissern Sie sich vor Gebrauch immer, dass das Produkt noch innerhalb der Lagerdauer liegt. Das Produkt sollte sauber, trocken und im Temperaturbereich von - 20°C bis +38°C bei einer maximalen rel. Luftfeuchtigkeit von 80% gelagert werden. Für Lagerung
und Transport die Originalverpackung verwenden. Nicht direkter Sonnenstrahlung aussetzen.



MICRO-MEDICAL

MICRO-MEDICAL

取得国外标准认证或注册的非医用口罩生产企业清单
Name List of Non-Medical Use Face Masks Companies with

序号 No.	生产企业 Company	统一社会信用代码 Uniform Social Credit Code	国外注册认证情况 Status of Certification / Authorization in Other Countires
377	[illegible] Luoyang Kaijian Technology Co.,Ltd.	[illegible]	[illegible]
378	[illegible] Xiantao Qiandeng Life Saving Equipment Co.,Ltd.	[illegible]	CE
379	湖南云想生活电子商务有限公司 Hunan Dreaming Cloud E-Commerce CO., Ltd	91430105MA4LAAUW8C	CE
380	[illegible] Linyi Meishen Medical Supplies Co., Ltd.	[illegible]	CE
381	[illegible] Nanjing Hosacy Medical Apparatus and Instruments Co.,Ltd.	[illegible]	CE
382	[illegible] Jiangsu Darying Clothing Co., Ltd	[illegible]	CE

Hunan Dreaming Cloud E-Commerce CO., Ltd
Block 1, Smart Tech Park, 57# Huangxing Avenue, Changsha Economic and Technological Development Zone,Changsha, Hunan, China

EU-KONFORMITÄTSERKLÄRUNG

Diese Konformitätserklärung wurde unter der alleinigen Verantwortung des Herstellers

Hunan Dreaming Cloud E-Commerce CO., Ltd.

Block 1, Smart Tech Park, 57 # Huangxing Avenue, Changsha Economic and Technological Development Zone, Changsha, Hunan, China

ausgestellt.

EG-Vertreter: Sunbeam International GmbH, Schumanstr.12, Würselen 52146 Deutschland

Hiermit wird erklärt, dass die folgende persönliche Schutzausrüstung (PSA)

Produktbeschreibung: HYGISUN Partikelfilter-Halbmaske

Produktmodell (e): HS0501A FFP2 NR ohne Ventil

den Bestimmungen der folgenden europäischen Verordnung entspricht:

PSA-Verordnung (Persönliche Schutzausrüstung)

Das Modell entspricht den Bestimmungen der Verordnung (EU) 2016/425, PSA zur Verwendung durch Angehörige der Gesundheitsberufe gemäß der Empfehlung der Kommission 2020/403 und der Nationalen Norm zur Umsetzung der harmonisierten europäischen Normnummer (n):

EN 149: 2001 + A1: 2009

und ist identisch mit der PSA, die Gegenstand einer EU-Typprüfung ist (Modul B der Verordnung (EU) 2016/425), auf die auf der Zertifikatsnummer verwiesen wird:

Zertifikat Nr.: CE 750475 (Ausstellungsdatum: 09/06/2021)

herausgegeben von BSI Group Niederlande BV

John M. Keynesplein 9, 1066 EP, Amsterdam, Niederlande (Notified Body No. 2797)

und entspricht den Verfahren in Modul C2 der Verordnung (EU) 2016/425 unter der Überwachung der BSI Group The Netherlands BV (Notified Body Nr. 2797), auf die auf dem vom BSI ausgestelltem Zertifikat CE 750476 (Ausstellungsdatum: 09/06/2021) verwiesen wird.

Changsha, China, 10.06.2021

OuYang Zhouya

OuYang Zhouya

(Nachname Name)

Qualitätsmanager

Hunan Dreaming Cloud E-Commerce CO., Ltd.

EU Type Examination Certificate

This is to certify that:

Sunbeam International GmbH
Schumanstr. 12
Würselen
52146
Germany

Holds Certificate Number: CE 750475

In respect of:

Respiratory protective devices - Filtering half masks to protect against particles -
To EN 149:2001+A1:2009
Model: HYGISUN HS0501A.

on the basis that BSI carried out the relevant Type Examination procedures under the requirements with the Regulation (EU) 2016/425 of the European Parliament and Council relating to Personal Protective Equipment Regulation (PPE) Annex V (Module B) and meets the relevant health and safety requirements specified in Annex II

For and on behalf of BSI, a Notified Body for the above Regulation (Notified Body Number 2797):

Drs. Dave Hagenaars, Managing Director

First Issued: 2021-06-09
Latest Issue: 2021-06-09

Effective Date: 2021-06-09
Expiry Date: 2026-06-09

PRODUCTS
RvA C 646

...making excellence a habit.™

EU Type Examination Certificate

No. CE 750475

Product Specification

Product Type:	Filtering half masks to protect against particles.
Model:	HYGISUN HS0501A.
Product description:	The particulate respirator is designed to protect against solid and non-volatile liquid particles. The masks are a single size, non-sterile, non-valved product held on the face by a pair of elasticated ear loops. The masks are intended for single shift use as denoted by the classification symbol NR.
Technical specification:	EN 149:2001+A1:2009 – Respiratory Protective Devices - Filtering half masks to protect against particles.
EN 149 classification:	FFP2 NR.

First Issued: 2021-06-09
Latest Issue: 2021-06-09

Effective Date: 2021-06-09
Expiry Date: 2026-06-09

EU Type Examination Certificate

No. CE 750475

Certificate Administration Details

Technical File reference: TCF.02.

Certificate Amendment Record:

Issue date	Comments	BSI Review Number
June 2021	First issue under PPE Regulation (EU) 2016/425. Product initially Certified as a "Covid-19" mask by BSI, Certificate CE 730303 refers.	2797:2021:3339407

Certificate validity

The Certificate holder is responsible for ensuring that the Notified Body is advised of changes to any aspect of the overall processes utilised in the manufacture of the product, failure to do so could invalidate the Certificate in respect of product manufactured following the introduction of such changes.

The validity of the Certificate for the products is also dependent on the maintenance of the EU Conformity to Type based on Internal Production Control plus supervised product checks at random intervals, Annex VII (Module C2) as referenced on BSI issued Certificate CE 750476.

First Issued: 2021-06-09
Latest Issue: 2021-06-09

Effective Date: 2021-06-09
Expiry Date: 2026-06-09

Conformity to Type based on Internal Production Control plus supervised product checks at random intervals

This is to certify that:

Sunbeam International GmbH
Schumanstr. 12
Würselen
52146
Germany

Holds Certificate Number: CE 750476

In respect of:

For the manufacture of respiratory protective devices -
Filtering half masks to protect against particles - To EN 149:2001+A1:2009.

on the basis that BSI carried out the supervised production checks at random intervals under the requirements with the Regulation (EU) 2016/425 of the European Parliament and Council relating to Personal Protective Equipment Regulation (PPE) Annex VII (Module C2)

For and on behalf of BSI, a Notified Body for the above Regulation (Notified Body Number 2797):

Drs. Dave Hagenaars, Managing Director

First Issued: 2021-06-09
Latest Issue: 2021-06-09

Effective Date: 2021-06-09
Expiry Date: 2026-06-09

PRODUCTS
RvA C 646

...making excellence a habit.™

Conformity to Type based on Internal Production Control plus supervised product checks at random intervals

No. CE 750476

Model produced by:

Hunan Dreaming Cloud E-Commerce CO., Ltd
Block 1, Smart Tech Park,
57# Huangxing Avenue,
Changsha Economic and Technological Development Zone,
Changsha,
Hunan,
China

Product details

The respiratory protective device covered by the scope of this Module C2 Certificate and the Technical Specification to which the product is manufactured are as follows:

Product type:	Respiratory protective device – Filtering half masks to protect against particles.
Model:	HYGISUN HS0501A.
Technical Specification:	EN 149:2001+A1:2009 – Respiratory Protective Devices - Filtering half masks to protect against particles.
EN 149 classifications:	FFP2 NR.

Certificate Administration Details:

Certificate Amendment Record:

Issue date	Comments	BSI Review No.
June 2021	First issue. Referenced product initially Certified as a "Covid-19" mask by BSI, with the associated BSI issued Module C2 Certificate CE 730304.	2797:21:3339408

Certificate validity

The Certificate holder is responsible for ensuring that the Notified Body is advised of changes to any aspects of the overall quality system utilized in the manufacture of the products, failure to do so could invalidate the Certificate in respect of product manufactured after the introduction of such changes.

First Issued: 2021-06-09
Latest Issue: 2021-06-09

Effective Date: 2021-06-09
Expiry Date: 2026-06-09

Test Report 3339405.

Sunbeam International GmbH.

Introduction.

This report has been prepared by D. Key and relates to the activity detailed below:

Job/Registration Details		Client Details
Job number:	**3339405**	Sunbeam International GmbH Schumanstr. 12 Würselen 52146 Germany
Job type:	Testing samples submitted	
Start Date:	17/01/2021	
Test type:	Type	
Sample ID:	10195243	
Registration:	**CE 730303**	
Scheme:	Negative Pressure RPE	
Protocol:	PP123	
Scheme Manager:	Nathan Shipley	

The report has been approved for issue by T Wicksey – Senior Test Engineer

Approved For Issue	
	Issue Date: 22 March 2021

Objectives.

This is an independent Type Test evaluation to BS EN 149:2001+A1:2009. This report covers the gap testing from the BSI COVID-19 filtering face piece technical specification, for COVID-19 masks for use by healthcare workers.
See BSI Test Report 3220780 for the BSI COVID-19 filtering face piece technical specification test results.

Product Scope.

Respiratory protective device- Filtering half masks to protect against particles.

Report Summary.

The samples were received on 18 December 2020 and the testing was started on 17 January 2021.

The samples submitted complied with the requirements of the test work conducted.

Test Samples.

Sample ID	ER Number	Description
1 to 37	10195243	Model: HYGISUN HS0501A FFP2 NR

Description of Test Samples.

Sample Description
Model: HYGISUN HS0501A FFP2 NR. Valveless vertical fold flat particle filtering half mask with elastic earloops and removable plastic earloop clip

Test Requirements.

BS EN 149:2001 + A1:2009
Respiratory protective devices - Filtering half masks to protect against particles.

CLAUSE	REQUIREMENTS	ASSESSMENT
7	**Requirements**	-
7.1	**General**	-
7.2	**Nominal values and tolerances**	-
7.3	**Visual Inspection**	Pass (1)
7.4	**Packaging**	N/T (1)
7.5	**Material**	Pass
7.6	**Cleaning and disinfecting**	N/A (2)
7.7	**Practical performance**	N/T (3)
7.8	**Finish of parts**	Pass
7.9	**Leakage**	-
7.9.1	Total inward leakage	Pass (3)
7.9.2	Penetration of filter material	Pass (3)
7.10	**Compatibility with skin**	Pass
7.11	**Flammability**	Pass
7.12	**Carbon dioxide content of inhalation air**	N/T (3)
7.13	**Head harness**	Pass
7.14	**Field of vision**	Pass
7.15	**Exhalation valves**	N/A (4)
7.16	**Breathing resistance**	Pass (3)
7.17	**Clogging**	N/A (4)
7.18	**Demountable parts**	N/A (4)
9	**Marking**	N/T (1)
10	**Information to be supplied by the manufacturer**	N/T (1)
Appendix A - Test Panel Data		
Product Photographs		

(1) Packaging, Marking and Information not assessed as requested by BSI Product Certification
(2) Single use mask
(3) See also results from BSI COVID-19 filtering face piece technical specification testing, BSI Test Report number 3220780.
(4) Not a design feature of this product

Glossary of Terms.

Pass: Complies. Tested by BSI engineers at BSI laboratories
Pass 1: Complies. Witnessed by BSI engineers in manufacturers laboratory.
Pass 2: Complies. Tests carried out by third party lab; results accepted by BSI.
Pass*: Report resulted in uncertainty and states that Compliance is more probable than non-compliance.
Fail: Non-compliance. Product does not meet the requirements of this clause.
Fail*: Report resulted in uncertainty and states that Non-compliance is more probable than compliance.
N/T: Not Tested
N/A: Not Applicable
AR: As Received
TC: Temperature Conditioned
SW: Simulated Wear
FT: Flow Tested
MS: Mechanical strength
MMDF: Manufacturer's Minimum Design Flow

Conditions of Issue.

This Test Report is issued subject to the conditions stated in current issue of 'BSI Terms of Service'. The results contained herein apply only to the particular sample(s) tested and to the specific tests carried out, as detailed in this Test Report. The issuing of this Test Report does not indicate any measure of Approval, Certification, Supervision, Control or Surveillance by BSI of any product. No extract, abridgement or abstraction from a Test Report may be published or used to advertise a product without the written consent of BSI, who reserve the absolute right to agree or reject all or any of the details of any items or publicity for which consent may be sought.

Should you wish to speak with BSI in relation to this report, please contact Customer Services on +44 (0)8450 80 9000.

BSI
Kitemark House
Maylands Avenue
Hemel Hempstead
Hertfordshire
HP2 4SQ

UKAS TESTING
0135

Opinions and Interpretations expressed herein are outside the scope of our UKAS accreditation.
Unless otherwise stated, any results not obtained from testing in a BSI laboratory are outside the scope of our UKAS accreditation.

Test Results.

BS EN 149:2001 + A1:2009
Respiratory protective devices - Filtering half masks to protect against particles.

CLAUSE	REQUIREMENTS	ASSESSMENT
7.1	**General**	
	In all tests all samples shall meet the requirements.	-
7.2	**Nominal values and tolerances**	
	Unless otherwise specified, the values stated in this European Standard are expressed as nominal values. Except for temperature limits, values, which are not stated as maxima or minima, shall be subject to a tolerance of ±5%. Unless otherwise specified, the ambient temperature for testing shall be (16 – 32) °C, and the temperature limits shall be subject to an accuracy of ± 1°C.	-
7.3	**Visual Inspection**	
	The visual inspection shall also include the marking and the information supplied by the manufacturer.	Pass (1)
7.5	**Material**	
	Materials used shall be suitable to withstand handling and wear over the period for which the particle filtering half mask is designed to be used.	Pass
	After undergoing the conditioning described in clause 8.3.1 of the standard none of the particle filtering half masks shall have suffered mechanical failure of the facepiece or straps.	
	Three particle filtering half masks shall be tested.	Pass
	When conditioned in accordance with 8.3.1 and 8.3.2 the particle filtering half mask shall not collapse.	Pass
	Any material from the filter media released by the air flow through the filter shall not constitute a hazard or nuisance for the wearer.	Pass
	Testing shall be done in accordance with 8.2.	
7.8	**Finish of parts**	
	Parts of the device likely to come into contact with the wearer shall have no sharp edges or burrs.	Pass
	Testing shall be done in accordance with 8.2.	

(1) Marking and user information were not assessed as requested by BSI Product Certification

Test Results. (Continued)

CLAUSE	REQUIREMENTS	ASSESSMENT
7.9	**Leakage**	
7.9.1	Total inward leakage The laboratory tests shall indicate that the particle filtering half mask can be used by the wearer to protect with high probability against the potential hazard to be expected. The total inward leakage consists of three components: face seal leakage, exhalation valve leakage (if exhalation valve fitted) and filter penetration. For particle filtering half masks fitted in accordance with the manufacturer's information, at least 46 out of the 50 individual exercise results (i.e. 10 subjects x 5 exercises) for total inward leakage shall be not greater than 25% for FFP1 11% for FFP2 5% for FFP3 and, in addition, at least 8 out of the 10 individual wearer arithmetic means for the total inward leakage shall be not greater than 22% for FFP1 8% for FFP2 2% for FFP3 Testing shall be done in accordance with 8.5.	Pass (1) See Table A

Table A: Clause 7.9.1 - Total inward leakage.

Test candidate	Sample	Pre-test condition	Inward leakage (%).					
			A	B	C	D	E	Average
			Walking	Walking with head side to side	Walking with head up & down	Walking and talking	Walking	
LM2	8	TC	7.2973	3.9167	5.7087	3.2365	5.1736	5.0666
SI1	9	TC	0.2104	0.2047	0.2353	0.2276	0.1237	0.2003
KH1	10	TC	0.2112	0.2296	0.2503	0.5231	0.6973	0.3823
CB1	11	TC	3.0178	1.8510	8.0745	1.8897	5.6798	4.1025
JW1 (2)	12	TC	0.5893	0.9512	0.7366	0.4300	0.6535	0.6721

(1) Results for the remaining 'as received' samples are covered in BSI Test Report number 3220780 for the BSI COVID-19 filtering face piece technical specification testing.
(2) Earloop clip used.

Test Results. (Continued)

CLAUSE	REQUIREMENTS	ASSESSMENT
7.9.2	Penetration of filter material	
	The penetration of the filter of the particle filtering half mask shall meet the requirements of Table 1	
	A total of 9 samples of particle filtering half masks shall be tested for each aerosol. Testing in accordance with 8.11 using the Penetration test according to EN 13274-7, shall be performed on:	Pass (1) See Tables B and C
	3 samples as received,	
	3 samples after the simulated wearing treatment described in 8.3.1.	
	Testing in accordance with 8.11 using the Exposure test with a specified mass of test aerosol of 120 mg, and for particle filtering devices claimed to be re-usable additionally the Storage test, according to EN 13274-7, shall be performed:	Pass (1) See Table D and E
	for non-re-usable devices on:	
	3 samples after the test for mechanical strength in accordance with 8.3.3 followed by temperature conditioning in accordance with 8.3.2.	
	for re-usable devices on:	
	3 samples after the test for mechanical strength in accordance with 8.3.3 followed by temperature conditioning in accordance with 8.3.2 and followed by one cleaning and disinfecting cycle according to the manufacturer's instruction.	N/A (2)

Table B: Clause 8.11 - Sodium Chloride penetration test.

Sample	Pre-test condition	Continuous flow (l/min)	Penetration (%)	
			Limit	Measured
16	SW	95	6.0	0.1635
17	SW	95	6.0	0.1645
18	SW	95	6.0	0.1542

Table C: Clause 8.11 - Paraffin oil penetration test.

Sample	Pre-test condition	Continuous flow (l/min)	Penetration (%)	
			Limit	Measured
22	SW	95	6.0	1.1895
23	SW	95	6.0	1.9665
24	SW	95	6.0	1.6080

(1) Results for the remaining 'as received' samples are covered in BSI Test Report number 3220780 for the BSI COVID-19 filtering face piece technical specification testing.
(2) Not a design feature of this product.

Test Results. (Continued)

CLAUSE	REQUIREMENTS	ASSESSMENT

7.9.2 Penetration of filter material (continued)

Table D: Clause 8.11. Exposure test Sodium Chloride.

	Sample 28 MS TC	Sample 29 MS TC	Sample 30 MS TC
Flow through filter	95 l/min		
Elapsed time (minutes)	Measured penetration % (Maximum permitted penetration 6.0 %)		
5	0.239798 (1)	0.100650 (1)	0.149081 (1)
10	0.192890	0.079177	0.121274
15	0.141387	0.065270	0.096979
20	0.090230	0.052679	0.076271
25	0.056086	0.041193	0.054319
30	0.033608	0.032297	0.038793
Result	Pass	Pass	Pass

(1) The reading at which 5 subsequent sampling intervals showed a declining filter penetration. The testing was terminated without the 120mg exposure limit being reached, as permitted by BS EN 13274-7.

Test Results. (Continued)

CLAUSE	REQUIREMENTS	ASSESSMENT

7.9.2 Penetration of filter material (continued)

Table E: Clause 8.11 Paraffin oil exposure test.

	Sample 25 MS TC	Sample 26 MS TC	Sample 27 MS TC
Flow through filter	95 l/min		
Elapsed time (minutes)	Measured penetration % (Maximum permitted penetration 6.0 %)		
3	2.1110	1.5950	1.9965
5	2.2125	1.6255	2.0845
10	2.4525	1.9880	2.3595
15	2.5900	1.9280	2.4440
20	2.8495	1.9610	2.5800
25	3.0625	2.0000	2.6650
30	3.2990	2.1675	2.7670
35	3.2725	2.2115	2.9130
40	3.4220	2.2705	3.0045
45	3.6010	2.3765	3.0950
50	3.6315	2.3965	3.1885
55	3.7715	2.4610	3.2285
60	3.8520	2.5240	3.3385
(1)	4.0125	2.5060	3.3755
Result	Pass	Pass	Pass

(1) A loading of 120 mg was achieved after a period of 63 minutes, 10 seconds had elapsed.

Test Results. (Continued)

CLAUSE	REQUIREMENTS	ASSESSMENT
7.10	**Compatibility with skin**	
	Materials that may come into contact with the wearer's skin shall not be known to be likely to cause irritation or any other adverse effect to health.	Pass
	Testing shall be done in accordance with 8.4 and 8.5.	
7.11	**Flammability**	
	The material used shall not present a danger for the wearer and shall not be of a highly flammable nature.	
	When tested, the particle filtering half mask shall not burn or not continue to burn for more than 5 seconds after removal from the flame.	Pass See Table F
	The particle filtering half mask does not have to be usable after the test.	
	Testing shall be done in accordance with 8.6.	

Table F: Clause 8.6 – Flammability.

Sample	Area exposed	Comments
34 AR	Filter material, welding.	Did not ignite.
35 AR	Earloop, vertical welding.	Did not ignite.
36 TC	Filter material, welding.	Did not ignite.
37 TC	Earloop, vertical welding.	Did not ignite.

7.13	**Head harness**	
	The head harness shall be designed so that the particle filtering half mask can be donned and removed easily.	
	The head harness shall be adjustable or self-adjusting and shall be sufficiently robust to hold the particle filtering half mask firmly in position and be capable of maintaining total inward leakage requirements for the device.	Pass
	Testing shall be done in accordance with 8.4 and 8.5.	
7.14	**Field of vision**	
	The field of vision is acceptable if determined so in practical performance tests.	Pass
	Testing shall be done in accordance with 8.4.	

Test Results. (Continued)

CLAUSE	REQUIREMENTS	ASSESSMENT
7.16	**Breathing resistance**	
	The breathing resistances apply to valved and valveless particle filtering half masks and shall meet the requirements of Table 2.	
	Testing shall be done in accordance with 8.9.	
	A total of 9 valveless particle filtering half masks shall be tested: 3 as received, 3 after temperature conditioning in accordance with 8.3.2 and 3 after the test for simulated wearing in accordance with 8.3.1.	Pass* (1) (2) See Tables G, H and I
	Testing shall be done in accordance with 8.9.	
	A total of 12 valved particle filtering half masks shall be tested: 3 as received, 3 after temperature conditioning in accordance with 8.3.2, 3 after the test for simulated wearing in accordance with 8.3.1, and 3 after the flow conditioning in accordance with 8.3.4.	N/A (3)
	Testing shall be done in accordance with 8.9.	

Table G: Clause 8.9 – Breathing resistance. Inhalation resistance at a continuous flow.

Sample	Pre-test condition	Flow (l/min)	Limit (mbar)	Measured (mbar)
16	SW	30	0.7	0.54
17	SW	30	0.7	0.50
18	SW	30	0.7	0.53
31	TC	30	0.7	0.48
32	TC	30	0.7	0.47
33	TC	30	0.7	0.46

Table H: Clause 8.9 – Breathing resistance. Inhalation resistance at a continuous flow.

Sample	Pre-test condition	Flow (l/min)	Limit (mbar)	Measured (mbar)
16	SW	95	2.4	1.89
17	SW	95	2.4	1.87
18	SW	95	2.4	1.89
31	TC	95	2.4	1.79
32	TC	95	2.4	1.79
33	TC	95	2.4	1.72

(1) Results for the remaining 'as received' samples are covered in BSI Test Report number 3220780 for the BSI COVID-19 filtering face piece technical specification testing.
(2) Results for exhalation resistance are within the uncertainty of measurement, but compliance is more probable than non-compliance.
(3) Not a design feature of this product.

Test Results. (Continued)

CLAUSE	REQUIREMENTS	ASSESSMENT
7.16	**Breathing resistance (continued)**	

Table I: Clause 8.9 – Breathing resistance. Exhalation resistance at a continuous flow, measured in five orientations with the highest value recorded.

Sample	Pre-test condition	Flow (l/min)	Limit (mbar)	Measured (mbar)
16	SW	160	3.0	2.87
17	SW	160	3.0	2.96
18	SW	160	3.0	2.95
31	TC	160	3.0	2.89
32	TC	160	3.0	2.84
33	TC	160	3.0	2.79

Appendix A. – Test Panel Data

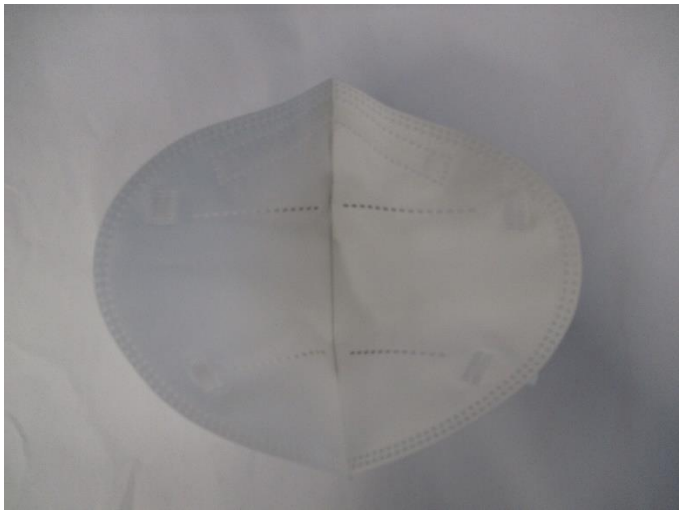
Test Candidate	Facial Dimensions (mm)					Gender
	Length of face	Width of face	Face depth	Width of mouth	Head Circumference	
JW1	116	126	122	48	570	Male
SI1	121	135	142	48	575	Male
LM2	110	148	125	47	567	Male
KH1	112	142	115	60	585	Male
CB1	117	147	130	57	566	Male

Note: All candidates were clean shaven

Product photographs.

Front view

Side view



Inside view

*** End of Report ***

Test Report No.: 244315789a 001

Client: **SUNBEAM INTERNATIONAL GMBH**

Contact Information: Schumanstr. 12, 52146 Würselen, Germany
Contact Person: Edward Zhao

Sample Description As Declared:

No. Of Sample	:	80 pcs
Product Description	:	Personal Protective Respitator Mask
Product Type	:	Single shift use only
Material	:	-
Colour	:	White
Lot No./Batch Code	:	-
Buyer Name	:	-
Trademark	:	HYGISUN
Type-identifying	:	HS0501A
Claimed Classification	:	FFP2 NR
Manufacturer	:	Hunan Dreaming Cloud E-Commerce Co., Ltd.
Country of Origin	:	-
Sales Destination (Country)	:	-
Test Type	:	Full Test
Test Specification	:	EN 149:2001 + A1:2009 Respiratory Protective Devices - Filtering Half Masks to Protect Against Particles - Requirements, Testing and Marking
Other Information	:	-

Sample Obtaining Method: Sending by customer

Delivery Condition: Apparent good, samples tested as received

Sample Receiving date: 2021-03-04 & 2021-04-21

Testing Period: 2021-03-04 to 2021-04-01 & 2021-04-21 to 2021-04-27

Place of Testing: Textiles laboratory Shanghai

For and on behalf of
TÜV Rheinland (Shanghai) Co., Ltd.

2021-04-30 Carmen Yan / Department Manager

Date *Name/Position*

Test Report No.: 244315789a 001

Summary of Test Results:

Clause	Item	Conclusion
7.3	Visual Inspection	P
7.4	Package	P
7.5	Material	P
7.6	Cleaning And Disinfection	N/A
7.7	Practical Performance	P
7.8	Finish Of Parts	P
7.9.1	Leakage	P
7.9.2	Penetration Of Filter Material	P
7.10	Compatibility With Skin	P
7.11	Flammability	P
7.12	Carbon Dioxide Content Of The Inhalation Air	P
7.13	Head Harness	P
7.14	Field Of Vision	P
7.15	Exhalation Valve(s)	N/A
7.16	Breathing Resistance	P
7.17	Clogging	N/A
7.18	Demountable Parts	N/A
10	Information To Be Supplied By The Manufacturer	P
9	Marking	P

Note: P = Pass F = Fail
= No Comment - = Did Not Perform
N/R = Not Request N/A = Not Applicable

Material List:

Material No.	Material	Color	Location	Remark
M001	Whole Product	White	Personal Protective Respitator Mask	Received on 2021.03.04
M001'	Whole Product	White	Personal Protective Respitator Mask	Received on 2021.04.21

Visual Inspection

Test Method: EN 149:2001+A1:2009 Clause 8.2

Clause	Item	M001
7.3	The visual inspection shall also include the marking and the information supplied by the manufacturer.	Pass
7.4	Particle filtering half masks shall be offered for sale packaged in such a way that they are protected against mechanical damage and contamination before use.	Pass
7.5	Materials used shall be suitable to withstand handling and wear over the period for which the particle filtering half mask is designed to be used.	Pass
	After undergoing the conditioning described in 8.3.1 none of the particle filtering half masks shall have suffered mechanical failure of the face piece or straps.	Pass
	When conditioned in accordance with 8.3.1 and 8.3.2 the particle filtering half mask shall not collapse.	Pass
	Any material from the filter media released by the air flow through the filter shall not constitute a hazard or nuisance for the wearer.	Pass
7.8	Parts of the device likely to come into contact with the wearer shall have no sharp edges or burrs	Pass
7.18	All demountable parts (if fitted) shall be readily connected and secured, where possible by hand.	N/A

Remark:

N/A: Due to no relevent information/material
N/R: Due to not request

Test Report No.: 244315789a 001

Practical Performance

Test Method: EN 149:2001+A1:2009 Clause 8.4 & 8.5

Clause	Item	M001
7.7	Wearing	Pass
7.7	Walking test	Pass
7.7	Work simulation test	Pass
7.10	Materials that may come into contact with the wearer's skin shall not be known to be likely to cause irritation or any other adverse effect to health	Pass
7.13	The head harness shall be designed so that the particle filtering half mask can be donned and removed easily. The head harness shall be adjustable or self-adjusting and shall be sufficiently robust to hold the particle filtering half mask firmly in position and be capable of maintaining total inward leakage requirements for the device	Pass
7.14	The field of vision is acceptable if determined so in practical performance tests	Pass

Remark:

N/A: Due to no relevent information/material
N/R: Due to not request

Test Report No.: 244315789a 001

Clause 7.9.1: Leakage

Test Method : EN 149:2001+A1:2009 Clause 8.5

Requirement : FFP2 :

At least 46 out of the 50 individual exercise results for total inward leakage ≤ 11%

At least 8 out of the 10 individual wearer arithmetic means for the total inward leakage ≤ 8%

M001								
Condition	Specimen No.	Subject	Leakage (%)					
			Walk	Head Side/Side	Head Up/Down	Talk	Walk	Mean
As received	1	BM	4.927	7.304	9.711	5.581	2.803	6.065
	2	ACH	3.824	6.874	8.145	8.664	5.217	6.545
	3	ML	4.128	6.229	8.225	7.422	3.877	5.976
	4	LLC	3.397	6.785	8.199	6.357	4.012	5.734
	5	DG	3.981	6.932	8.902	7.559	4.331	6.341
After conditioning	6	SG	4.104	5.181	10.648	7.685	3.493	6.222
	7	YL	6.247	5.487	8.375	8.247	6.027	6.877
	8	KQ	5.525	6.028	9.084	8.122	5.021	6.756
	9	KXH	6.001	6.439	9.119	8.074	5.387	7.004
	10	YY	5.743	6.009	8.911	7.936	5.111	6.742
Conclusion		Pass						

Facial Dimension Of Subject (mm)											
Subject	BM	ACH	ML	LLC	DG	SG	YL	KQ	KXH	YY	LL
Face length	135	127	120	120	130	135	115	120	130	130	121
Face width	160	159	133	140	145	155	135	135	155	165	163
Face Depth	130	122	115	115	132	132	118	115	120	143	142
Mouth Width	52	55	52	50	50	55	48	50	52	50	45

Clause 7.9.2: Penetration Of Filter Material

Test method : EN 149:2001+A1:2009 Clause 8.11

Requirement : FFP2: ≤6%

M001			
Aerosol	Condition	Specimen No.	Penetration (%)
Sodium chloride Penetration	As received	1	0.048
	As received	2	0.223
	As received	3	0.226
	Simulated wearing treatment	4	0.568
	Simulated wearing treatment	5	0.483
	Simulated wearing treatment	6	0.439
	Mechanical strength + Temperature conditioned @ Exposure test of 120mg	7	0.322
	Mechanical strength + Temperature conditioned @ Exposure test of 120mg	8	0.282
	Mechanical strength + Temperature conditioned @ Exposure test of 120mg	9	0.289
Paraffin oil Penetration	As received	10	0.566
	As received	11	0.536
	As received	12	0.521
	Simulated wearing treatment	13	0.586
	Simulated wearing treatment	14	0.623
	Simulated wearing treatment	15	0.637
	Mechanical strength + Temperature conditioned @ Exposure test of 120mg	16	0.984
	Mechanical strength + Temperature conditioned @ Exposure test of 120mg	17	2.392
	Mechanical strength + Temperature conditioned @ Exposure test of 120mg	18	1.664
Conclusion	Pass		

Clause 7.11: Flammability

Test method : EN 149:2001+A1:2009 Clause 8.6

Requirement : ≤5s

M001			
Item	Condition	Specimen No.	Test results
Afterflame time (s)	As received	1	DNI
	As received	2	DNI
	After conditioning	3	DNI
	After conditioning	4	DNI
Conclusion		Pass	

Remark:

DNI-Do not ignite

Clause 7.12: Carbon Dioxide Content Of The Inhalation Air

Test Method : EN 149:2001+A1:2009 Clause 8.7

Requirement : ≤1%

M001			
Item	Condition	Test results	
Content (%)	As received	Specimen 1	0.58
	As received	Specimen 2	0.59
	As received	Specimen 3	0.61
	As received	Mean	0.60
Conclusion		Pass	

Test Report No.: 244315789a 001

Clause 7.16: Breathing Resistance

Test Method : EN 149:2001+A1:2009 Clause 8.9

Requirement : FFP2:
Inhalation: 30l/min: ≤0.7mbar
Inhalation: 95l/min: ≤2.4mbar
Exhalation: 160l/min: ≤3.0mbar

M001'																
Flow rate (l/min)		Resistance (mbar)														
As received		Specimen 1					Specimen 2					Specimen 3				
		A	B	C	D	E	A	B	C	D	E	A	B	C	D	E
Inhalation	30	0.4	0.4	0.4	0.4	0.4	0.5	0.5	0.5	0.5	0.5	0.4	0.4	0.4	0.4	0.4
	95	1.3	1.3	1.3	1.3	1.3	1.4	1.4	1.4	1.4	1.4	1.4	1.4	1.4	1.4	1.4
Exhalation	160	2.0	2.0	2.0	2.0	2.0	2.2	2.2	2.2	2.2	2.2	2.2	2.2	2.2	2.2	2.2
Simulated wearing treatment		Specimen 4					Specimen 5					Specimen 6				
		A	B	C	D	E	A	B	C	D	E	A	B	C	D	E
Inhalation	30	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5
	95	1.5	1.5	1.5	1.5	1.5	1.4	1.4	1.4	1.4	1.4	1.5	1.5	1.5	1.5	1.5
Exhalation	160	2.4	2.4	2.4	2.4	2.4	2.3	2.3	2.3	2.3	2.3	2.3	2.3	2.3	2.3	2.3
Temperature conditioned		Specimen 7					Specimen 8					Specimen 9				
		A	B	C	D	E	A	B	C	D	E	A	B	C	D	E
Inhalation	30	0.4	0.4	0.4	0.4	0.4	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5
	95	1.4	1.4	1.4	1.4	1.4	1.5	1.5	1.5	1.5	1.5	1.5	1.5	1.5	1.5	1.5
Exhalation	160	2.3	2.3	2.3	2.3	2.3	2.4	2.4	2.4	2.4	2.4	2.2	2.2	2.2	2.2	2.2
Conclusion		Pass														

Remark : A: facing directly ahead;
B: facing vertically upwards;
C: facing vertically downwards;
D: lying on the left side;
E: lying on the right side

Marking

Test Method: EN 149:2001+A1:2009 Clause 9

M001	
9.1 Packaging	
The following information shall be clearly and durably marked on the smallest commercially available packaging or legible through it if the packaging is transparent.	
9.1.1 The name, trademark or other means of identification of the manufacturer or supplier.	Present
9.1.2 Type-identifying marking.	Present
9.1.3 Classification The appropriate class (FFP1, FFP2 or FFP3) followed by a single space and then: "NR" if the particle filtering half mask is limited to single shift use only. Example: FFP3 NR, or "R" if the particle filtering half mask is re-usable. Example: FFP2 R D.	Present
9.1.4 The number and year of publication of this European Standard.	Present
9.1.5 At least the year of end of shelf life. The end of shelf life may be informed by a pictogram as shown in Figure 12a, where yyyy/mm indicates the year and month.	Present
9.1.6 The sentence 'see information supplied by the manufacturer', at least in the official language(s) of the country of destination, or by using the pictogram as shown in Figure 12b.	Present
9.1.7 The manufacturer's recommended conditions of storage (at least the temperature and humidity) or equivalent pictogram, as shown in Figures 12c and 12d.	Present
9.1.8 The packaging of those particle filtering half masks passing the dolomite clogging test shall be additionally marked with the letter "D". ID This letter shall follow the classification marking preceded by a single space.	N/A
9.2 Particle filtering half mask	
Particle filtering half masks complying with this European Standard shall be clearly and durably marked with the following:	
9.2.1 The name, trademark or other means of identification of the manufacturer or supplier.	Present
9.2.2 Type-identifying marking.	Present
9.2.3 The number and year of publication of this European Standard.	Present
9.2.4 Classification The appropriate class (FFP1, FFP2 or FFP3) followed by a single space and then: "NR" if the particle filtering half mask is limited to single shift use only. Example: FFP3 NR, or "R" if the particle filtering half mask is re-usable. Example: FFP2 R D.	Present
9.2.5 If appropriate the letter D (dolomite) in accordance with clogging performance. This letter shall follow the classification marking preceded by a single space.	N/A
9.2.6 Sub-assemblies and components with considerable bearing on safety shall be marked so that they can be identified.	N/A

Remark:

1. The evaluation is based on artwork.
2. N/A: Not applicable

Information To Be Supplied By The Manufacturer

Test Method: EN 149:2001+A1:2009 Clause 10

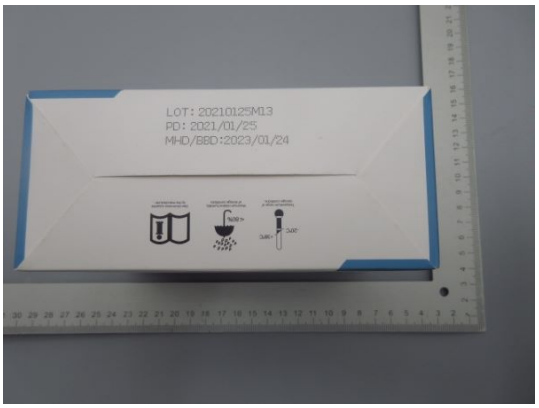
M001	
10.1 Information supplied by the manufacturer shall accompany every smallest commercial available package	Present
10.2 Information supplied by the manufacturer shall be at least in the official language(s) of the country of destination	Present
10.3 The information supplied by the manufacturer shall contain all information necessary for trained and qualified persons on	
- application/limitations	Present
- the meaning of any colour coding	N/A
- checks prior to use	Present
- donning, fitting	Present
- use	Present
- maintenance (e.g. cleaning, disinfecting), if applicable	N/A
- storage	Present
- the meaning of any symbols/pictograms used	Present
of the equipment	
10.4 The information shall be clear and comprehensible. If helpful, illustrations, part numbers, marking shall be added	Present
10.5 Warning shall be given against problems likely to be encountered, for example:	
- fit of particle filtering half mask (check prior to use)	Present
- it is unlikely that the requirements for leakage will be achieved if facial hair passes under the face seal	Present
- air quality (contaminants, oxygen deficiency)	Present
- use of equipment in explosive atmosphere	Present
10.6 The information shall provide recommendations as to when the particle filtering half mask shall be discarded	Present
10.7 For devices marked "NR", a warning shall be given that the particle filtering half mask shall not be used for more than one shift	Present

Remark:

1. The evaluation is based on artwork.
2. N/A: Not applicable

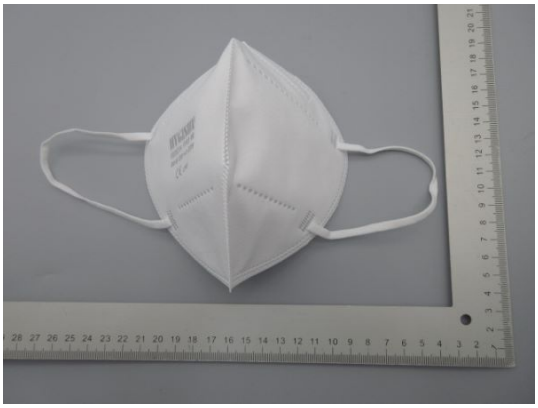
Test Report No.: 244315789a 001

Photo(s):



Photo(s):


HYGISUN
Persönliche Schutzmaske
PERSONAL PROTECTIVE MASK
15Stück/ Piece




HYGISUN
HS0501A FFP2 NR
EN149:2001+A1:2009
CE 2797

- END -

General Terms and Conditions of Business of TÜV Rheinland in Greater China

1. Scope

1.1 These General Terms and Conditions of Business of TÜV Rheinland in Greater China ("GTCB") is made between the client and one or more member entities of TÜV Rheinland in Greater China as applicable as the case may be ("TÜV Rheinland"). The Greater China hereof refers to Mainland China, Hong Kong and Taiwan.The client hereof includes :

(i) a natural person capable to form legally binding contracts under the applicable laws who concludes the contract not for the purpose of a daily use;

(ii) the incorporated or unincorporated entity duly organized, validly existing and capable to form legally binding contracts under the applicable law.

1.2 The following terms and conditions apply to agreed services including consultancy services, information, deliveries and similar services as well as ancillary services and other secondary obligations provided within the scope of contract performance.

1.3 Any standard terms and conditions of the client of any nature shall not apply and shall hereby be expressly excluded. No standard contractual terms and conditions of the client shall form part of the contract even if TÜV Rheinland does not explicitly object to them.

1.4 In the context of an ongoing business relationship with the client, this GTCB shall also apply to future contracts with the client without TÜV Rheinland having to refer to them separately in each individual case.

2. Quotations

Unless otherwise agreed, all quotations submitted by TÜV Rheinland can be changed by TÜV Rheinland without notice prior to its acceptance and confirmation by the other party.

3. Coming into effect and duration of contracts

3.1 The contract shall come into effect for the agreed terms upon the quotation letter of TÜV Rheinland or a separate contractual document being signed by both contracting parties, or upon the works requested by the client being carried out by TÜV Rheinland. If the client instructs TÜV Rheinland without receiving a quotation from TÜV Rheinland (quotation), TÜV Rheinland is, in its sole discretion, entitled to accept the order by giving written notice of such acceptance (including notice sent via electronic means) or by performing the requested services.

3.2 The contract term starts upon the coming into effect of the contract in accordance with article 3.1 and shall continue for the term agreed in the contract.

3.3 If the contract provides for an extension of the contract term, the contract term will be extended by the term provided for in the contract unless terminated in writing by either party with a six-week notice prior to the end of the contractual term.

4. Scope of services

4.1 The scope and type of the services to be provided by TÜV Rheinland shall be specified in the contractually agreed service scope of TÜV Rheinland by both parties. If no such separate service scope of TÜV Rheinland exists, then the written confirmation of order by TÜV Rheinland shall be decisive for the service to be provided.

4.2 The agreed services shall be performed in compliance with the regulations in force at the time the contract is entered into.

4.3 TÜV Rheinland is entitled to determine, in its sole discretion, the method and nature of the assessment unless otherwise agreed in writing or if mandatory provisions require a specific procedure to be followed.

4.4 On execution of the work there shall be no simultaneous assumption of any guarantee of the correctness (proper quality) and working order of either tested or examined parts nor of the installation as a whole and its upstream and/or downstream processes, organisations, use and application in accordance with regulations, nor of the systems on which the installation is based. In particular, TÜV Rheinland shall assume no responsibility for the construction, selection of materials and assembly of installations examined, nor for their use and application in accordance with regulations, unless these questions are expressly covered by the contract.

4.5 In the case of inspection work, TÜV Rheinland shall not be responsible for the accuracy or checking of the safety programmes or safety regulations on which the inspections are based, unless otherwise expressly agreed in writing.

4.6 If mandatory legal regulations and standards or official requirements for the agreed service scope change after conclusion of the contract, with a written notice to the client, TÜV Rheinland shall be entitled to additional remuneration for resulting additional expenses.

4.7The services to be provided by TÜV Rheinland under the contract are agreed exclusively with the client. A contract of third parties with the services of TÜV Rheinland, as well as making available of and justifying confidence in the work results (test reports, test results, expert reports, etc.) is not part of the agreed services. This also applies if the client passes on work results - in full or in extracts - to third parties in accordance with clause 11.4.

5. Performance periods/dates

5.1 The contractually agreed periods/dates of performance are based on estimates of the work involved which are prepared in line with the details provided by the client. They shall only be binding if being confirmed as binding by TÜV Rheinland in writing.

5.2 If binding periods of performance have been agreed, these periods shall not commence until the client has submitted all required documents to TÜV Rheinland.

5.3 Articles 5.1 and 5.2 also apply, even without express approval by the client, to all extensions of agreed periods/dates of performance not caused by TÜV Rheinland.

5.4TÜV Rheinland is not responsible for a delay in performance, in particular if the client has not fulfilled his duties to cooperate in accordance with clause 6.1 or has not done so in time and, in particular, has not provided TÜV Rheinland with all documents and information required for the performance of the service as specified in the contract.

5.5If the performance of TÜV Rheinland is delayed due to unforeseeable circumstances such as force majeure, strikes, business disruptions, governmental regulations, transport obstacles, etc., TÜV Rheinland is entitled to postpone performance for a reasonable period of time which corresponds at least to the duration of the hindrance plus any time period which may be required to resume performance.

6. The client's obligation to cooperate

6.1 The client shall guarantee that all cooperation required on its part, its agents or third parties will be provided in good time and at no cost to TÜV Rheinland.

6.2 Design documents, supplies, auxiliary staff, etc. necessary for performance of the services shall be made available free of charge by the client. Moreover, collaborative action of the client must be undertaken in accordance with legal provisions, standards, safety regulations and accident prevention instructions. And the client represents and warrants that:

a) it has required statutory qualifications;

b) the product, service or management system to be certified complies with applicable laws and regulations; and

c) it doesn't have any illegal and dishonest behaviours or is not included in the list of Enterprises with Serious Illegal and Dishonest Acts of People's Republic of China.

If the client breaches the aforesaid representations and warranties, TÜV Rheinland is entitled to i) immediately terminate the contract/order without prior notice; and ii) withdraw the issued testing report/certificates if any.

6.3 The client shall bear any additional cost incurred on account of work having to be redone or being delayed as a result of late, incorrect or incomplete information provided by or lack of proper cooperation from the client. Even where a fixed or maximum price is agreed, TÜV Rheinland shall be entitled to charge extra fees for such additional expense.

7. Prices

7.1 If the scope of performance is not laid down in writing when the order is placed, invoicing shall be based on costs actually incurred. If no price is agreed in writing, invoicing shall be made in accordance with the price list of TÜV Rheinland valid at the time of performance.

7.2 Unless otherwise agreed, work shall be invoiced according to the progress of the work.

7.3 If the execution of an order extends over more than one month and the value of the contract or the agreed fixed price exceeds €2,500.00 or equivalent value in local currency, TÜV Rheinland may demand payments on account or in instalments.

8. Payment terms

8.1 All invoice amounts shall be due for payment without deduction on receipt of the invoice. No discounts and rebates shall be granted.

8.2 Payments shall be made to the bank account of TÜV Rheinland as indicated on the invoice, stating the invoice and client numbers.

8.3 In cases of default of payment, TÜV Rheinland shall be entitled to claim default interest at the applicable short term loan interest rate publicly announced by a reputable commercial bank in the country where TÜV Rheinland is located. At the same time, TÜV Rheinland reserves the right to claim further damages.

8.4 Should the client default in payment of the invoice despite being granted a reasonable grace period, TÜV Rheinland shall be entitled to cancel the contract, withdraw the certificate, claim damages for non-performance and refuse to continue performance of the contract.

8.5 The provisions set forth in article 8.4 shall also apply in cases involving returned cheques, cessation of payment, commencement of insolvency proceedings against the client's assets or cases in which the commencement of insolvency proceedings has been dismissed due to lack of assets.

8.6 Objections to the invoices of TÜV Rheinland shall be submitted in writing within two weeks of receipt of the invoice.

8.7 TÜV Rheinland shall be entitled to demand appropriate advance payments.

8.8 TÜV Rheinland shall be entitled to raise its fees at the beginning of a month if overheads and/or purchase costs have increased. In this case, TÜV Rheinland shall notify the client in writing of the rise in fees. This notification shall be issued one month prior to the date on which the rise in fees shall come into effect (period of notice of changes in fees). If the rise in fees remains under 5% per contractual year, the client shall not have the right to terminate the contract. If the rise in fees exceeds 5% per contractual year, the client shall be entitled to terminate the contract by the end of the period of notice of changes in fees. If the contract is not terminated, the changed fees shall be deemed to have been agreed upon by the time of the expiry of the notice period.

8.9 Only legally established and undisputed claims may be offset against claims by TÜV Rheinland.

9. Acceptance of work

9.1 Any part of the work result ordered which is complete in itself may be presented by TÜV Rheinland for acceptance as an instalment. The client shall be obliged to accept it immediately.

9.2 If acceptance is required or contractually agreed in an individual case, this shall be deemed to have taken place two (2) weeks after completion and handover of the work, unless the client refuses acceptance within this period stating at least one fundmental breach of contract by TÜV Rheinland.

9.3 The client is not entitled to refuse acceptance due to insignificant breach of contract by TÜV Rheinland.

9.4 If acceptance is excluded according to the nature of the work performance of TÜV Rheinland, the completion of the work shall take its place.

9.5 If the client was unable to make use of the time windows provided for within the scope of a certification procedure for auditing/performance by TÜV Rheinland and the certificate is therefore to be withdrawn (e.g. performance of surveillance audits), TÜV Rheinland is entitled to immediately charge a lump-sum compensation of 10% of the order amount as compensation for expenses. The client reserves the right to prove that the TÜV Rheinland has incurred no damage whatsoever or only a considerably lower damage than the above lump sum.

9.6 Insofar as the client has undertaken in the contract to accept services, TÜV Rheinland shall also be entitled to charge lump-sum damages in the amount of 10% of the order amount as compensation for expenses if the service is not called within one year after the order has been placed. The client reserves the right to prove that the TÜV Rheinland has incurred no damage whatsoever or only a considerably lower damage than the above mentioned lump sum.

10. Confidentiality

10.1For the purpose of these terms and conditions, "confidential information" means all information, documents, images, drawings, know-how, data, samples and project documentation which one party (the "disclosing party") hands over, transfers or otherwise discloses to the other party (the "receiving party"), and the confidential information created during performance of work by TÜV Rheinland, including product testing data, defects, conformity to the technical standard and related reports. Confidential information also includes paper copies and electronic copies of such information. Confidential information is expressly not the data and know-how collected, compiled or otherwise obtained by TÜV Rheinland (non-personal) within the scope of the provision of services by TÜV Rheinland. TÜV Rheinland is entitled to store, use, further develop and pass on the data obtained in connection with the provision of services for the purposes of developing new services, improving services and analysing the provision of services.

10.2 The disclosing party shall mark all confidential information disclosed in written form as confidential before passing it onto the receiving party. The same applies to confidential information transmitted by e-mail. If confidential information is disclosed orally, the receiving party shall be appropriately informed in advance and the disclosing party shall confirm in writing the confidentiality nature of the information within five working days of oral disclosure. Where the disclosing party fails to do so within the stipulated period, the receiving party shall not take any confidentiality obligations hereunder towards such information.

10.3 All confidential information which the disclosing party transmits or otherwise discloses to the receiving party and which is created during performance of work by TÜV Rheinland:

a)may only be used by the receiving party for the purposes of performing the contract, unless expressly otherwise agreed in writing by the disclosing party;

b)may not be copied, distributed, published or otherwise disclosed by the receiving party, unless this is necessary for fulfilling the purpose of the contract or TÜV Rheinland is required to pass on confidential information, inspection reports or documentation to the government authorities, judicial court, accreditation bodies or third parties that are involved in the performance of the contract;

c)must be treated by the receiving party with the same level of confidentiality as the receiving party uses to protect its own confidential information, but never with a lesser level of confidentiality than that which is reasonably required.

10.4 The receiving party may disclose any confidential information received from the disclosing party only to those of its employees who need this information to perform the services required for the contract. The receiving party undertakes to oblige these employees to observe the same level of secrecy as set forth in this confidentiality clause.

10.5 Information for which the receiving party can furnish proof that:

a)it was generally known at the time of disclosure or has become general knowledge without violation of this confidentiality clause by the receiving party; or

b)it was disclosed to the receiving party by a third party entitled to disclose this information; or

c)the receiving party already possessed this information prior to disclosure by the disclosing party; or

d)the receiving party developed it itself, irrespective of disclosure by the disclosing party, shall not be deemed to constitute "confidential information" as defined in this confidentiality clause.

10.6 All confidential information shall remain the property of the disclosing party. The receiving party hereby agrees to immediately (i) return all confidential information, including all copies, to the disclosing party, and/or (ii) on request by the disclosing party, to destroy all confidential information, including all copies, and confirm the destruction of this confidential information to the disclosing party in writing, at any time if so requested by the disclosing party but at the latest and without special request after termination or expiry of the contract. This does not extend to include reports and certificates prepared for the client solely for the purpose of fulfilling the obligations under the contract, which shall remain with the client. However, TÜV Rheinland is entitled to make file copies of such reports, certificates and confidential information that forms the basis for preparing these reports and certificates in order to evidence the correctness of its results and for general documentation purposes required by laws, regulations and the requirements of working procedures of TÜV Rheinland.

10.7 From the start of the contract and for a period of three years after termination or expiry of the contract, the receiving party shall maintain strict secrecy of all confidential information and shall not disclose this information to any third parties or use it for itself.

11. Copyrights and rights of use, publications

11.1 TÜV Rheinland shall retain all exclusive copyrights in the reports, expert reports/opinions, test reports/results, results, calculations, presentations etc. prepared by TÜV Rheinland, unless otherwise agreed by the parties in a separate agreement. As the owner of the copyrights, TÜV Rheinland is free to grant others the right to use the work results for individual or all types of use ("right of use")

11.2 The client receives a simple, unlimited, non-transferable, non-sublicensable right of use to the contents of the work results produced within the scope of the contract, unless otherwise agreed by the parties in a separate agreement.The client may only use such reports, expert reports/opinions, test reports/results, results calculations, presentations etc. prepared within the scope of the contract for the contractually agreed purpose.

11.3 The transfer of right of use of the generated work results regulated in clause 11.2. of the GTCB is subject to full payment of the remuneration agreed in favour of TÜV Rheinland.

11.4 The client may use work results only complete and unshortened. The client may only pass on the work results in full unless TÜV Rheinland has given its prior written consent to the partial passing on of work results.

11.5 Any publication or duplication of the work results for advertising purposes or any further use of the work results beyond the scope regulaed in clause 11.2 needs the prior written approval of TÜV Rheinland in each individual case.

11.6 TÜV Rheinland may revoke a once given approval according to clause 11.5 at any time without stating reasons. In this case, the client is obliged to stop the transfer of the work results immediately at his own expense and, as far as possible, to withdraw publications.

11.7 The consent of TÜV Rheinland to publication or duplication of the work results does not entitle the client to use the corporate logo, corporate design or test/certification mark of TÜV Rheinland.

12. Liability of TÜV Rheinland

12.1 Irrespective of the legal basis, to the fullest extent permitted by applicable law, in the event of a breach of contractual obligations or tort, the liability of TÜV Rheinland for all damages, losses and reimbursement of expenses caused by TÜV Rheinland, its legal representatives and/or employees shall be limited to: (i) in the case of a contract with a fixed overall fee, three times the overall fee for the entire contract; (ii) in the case of a contract for annually recurring services, the agreed annual fee; (iii) in the case of a contract expressly charged on a time and material basis, a maximum of 20,000 Euro or equivalent amount in local currency; and (iv) in the case of a framework agreement that provides for the possibility of placing individual orders, three times of the fee for the individual order under which the damages or losses have occurred. Notwithstanding the above, in the event that the total and accumulated liability calculated according to the foregoing provisions exceeds 2.5 Million Euro or equivalent amount in local currency, the total and accumulated liability of TÜV Rheinland shall be only limited to and shall not exceed the said 2.5 Million Euro or equivalent amount in local currency.

12.2 The limitation of liability according to article 12.1 above shall not apply to damages and/or losses caused by malice, intent or gross negligence on the part of TÜV Rheinland or its vicarious agents. Such limitation shall not apply to damages for a person's death, physical injury or illness.

12.3 In cases involving a fundamental breach of contract, TÜV Rheinland will be liable even where minor negligence is involved. For this purpose, a "fundamental breach" is breach of a material contractual obligation, the performance of which permits the due performance of the contract. Any claim for damages for a fundamental breach of contract shall be limited to the amount of damages reasonably foreseen as a possible consequence of such breach of contract at the time of the breach (reasonably foreseeable damages), unless any of the circumstances described in article 12.2 applies.

12.4 TÜV Rheinland shall not be liable for the acts of the personnel made available by the client to support TÜV Rheinland in the performance of its services under the contract, unless such personnel made available is regarded as vicarious agent of TÜV Rheinland. If TÜV Rheinland is not liable for the acts of the personnel made available by the client under the foregoing provision, the client shall indemnify TÜV Rheinland against any claims made by third parties arising from or in connection with such personnel's acts.

12.5 Unless otherwise contractually agreed in writing, TÜV Rheinland shall only be liable under the contract to the client.

12.6 The limitation periods for claims for damages shall be based on statutory provisions.

12.7 None of the provisions of this article 12 changes the burden of proof to the disadvantage of the client.

13. Export control

13.1When passing on the services provided by TÜV Rheinland or parts thereof to third parties in Greater China or other regions, the client must comply with the respectively applicable regulations of national and international export control law.

13.2The performance of a contract with the client is subject to the proviso that there are no obstacles to performance due to national or international foreign trade legislations or embargos and/or sanctions. In the event of a violation, TÜV Rheinland shall be entitled to terminate the contract with immediate effect and the client shall compensate for the losses incured thereof by TÜV Rheinland.

14. Data protection notice

TÜV Rheinland processes personal data of the client for the purpose of fulfilling this contract. In addition, TÜV Rheinland also processes the data for other legal purposes in accordance with the relevant legal basis. The personal data of the client will only be disclosed to other natural or legal persons if the legal requirements are met. This also applies to transfers to third countries. The personal data will be deleted immediately as soon as a corresponding reason for deletion arises. Data subjects may exercise the following rights: right of information, right of rectification, right of deletion, right of processing limitation, right of objection, right of data transferability. In addition, persons concerned by the data processing have the right to revoke their consent at any time with effect for the future, as well as the right to file a complaint with the competent data protection supervisory authority. For further details on the processing of personal data by TÜV Rheinland as the person responsible or contract processor, please refer to the respective data protection information. You can contact the Group Data Protection Officer of TÜV Rheinland by e-mail at datenschutz@de.tuv.com or by post at the following address: TÜV Rheinland AG, c/o Group Data Protection Officer, Am Grauen Stein, 51105 Cologne, Germany.

15. Test material: transport risk and storage

15.1The risk and costs for freight and transport of documents or test material to and from TÜV Rheinland as well as the costs of necessary disposal measures shall be borne by the client.

15.2Any destroyed and otherwise worthless test material will be disposed of by TÜV Rheinland for the client at the expense of the client, unless otherwise agreed.

15.3Undamaged test material shall be stored by TÜV Rheinland for four (4) weeks after completion of the test. If a longer storage period is desired, TÜV Rheinland charges an appropriate storage fee.

15.4After the expiry of the 4 weeks or any longer period agreed upon, the test material will be disposed of by TÜV Rheinland for the client for a fee in accordance with clause 15.2.

16. Termination of the contract

16.1Notwithstanding clause 3.3 of the GTCB, TÜV Rheinland and the client are entitled to terminate the contract in its entirety or, in the case of services combined in one contract, each of the combined parts of the contract individually and independently of the continuation of the remaining services with six (6) months' notice to the end of the contractually agreed term.

16.2For good causes, TÜV Rheinland may consider giving a written notice to the client to terminate the contract which includes but not limited to the following:

a) the client does not immediately notify TÜV Rheinland of changes in the conditions within the company which are relevant for certification or signs of such changes;

b) the client misuses the certificate or certification mark or uses it in violation of the contract;

c) in the event of several consecutive delays in payment (at least three times);

d) a substantial deterioration of the financial circumstances of the client occurs and as a result the payment claims of TÜV Rheinland under the contract are considerably endangered and TÜV Rheinland cannot reasonably be expected to continue the contractual relationship.

16.3In the event of termination with written notice by TÜV Rheinland for good cause, TÜVRheinland shall be entitled to a lump-sum claim for damages against the client if the conditions of a claim for damages exist. In this case, the client shall owe 15% of the remuneration to be paid until the end of the fixed contract term as lump-sum compensation. The client reserves the right to prove that there is no damage or a considerably lower damage, TÜV Rheinland reserves the right to prove a considerably higher damage in individual cases.

16.4TÜV Rheinland is also entitled to terminate the contract with written notice if the client has not been able to make use of the time windows for auditing /service provision provided by TÜV Rheinland within the scope of a certification procedure and the certificate therefore has to be withdrawn (for example during the performance of monitoring audits). Clause 16.3 applies accordingly.

17. Partial invalidity, written form, place of jurisdiction and dispute resolution

17.1 All amendments and supplements must be in writing in order to be effective. This also applies to amendments and supplements to this clause 17.1.

17.2 Should one or several of the provisions under the contract and/or these terms and conditions be or become ineffective, the contracting parties shall replace the invalid provision with a legally valid provision that comes closest to the content of the invalid provision in legal and commercial terms.

17.3 Unless otherwise stipulated in the contract, the governing law of the contract and these terms and conditions shall be chosen following the rules as below:

a)if TÜV Rheinland in question is legally registered and existing in the People's Republic of China, the contracting parties hereby agree that the contract and these terms and conditions shall be governed by the laws of the People's Republic of China.

b)if TÜV Rheinland in question is legally registered and existing in Taiwan, the contracting parties hereby agree that the contract and these terms and conditions shall be governed by the laws of Taiwan.

c)if TÜV Rheinland in question is legally registered and existing in Hong Kong, the contracting parties hereby agree that the contract and these terms and conditions shall be governed by the laws of Hong Kong.

17.4 Any dispute in connection with the contract and these terms and conditions or the execution thereof shall be settled friendly through negotiations.

Unless otherwise stipulated in the contract, if no settlement or no agreement in respect of the extension of the negotiation period can be reached within two months of the arising of the dispute, the dispute shall be submitted:

a)in the case of TÜV Rheinland in question being legally registered and existing in the People's Republic of China, to China International Economic and Trade Arbitration Commission (CIETAC) to be settled by arbitration under the Arbitration Rules of CIETAC in force when the arbitration is submitted. The arbitration shall take place in Beijing, Shanghai, Shenzhen or Chongqing as appropriately chosen by the claiming party.

b)in the case of TÜV Rheinland in question being legally registered and existing in Taiwan, to Chinese Arbitration Association Taipei Branch to be arbitrated in accordance with its then current Rules of Arbitration. The arbitration shall take place in Taipei.

c)in the case of TÜV Rheinland being legally registered and existing in Hong Kong, to Hong Kong International Arbitration Centre (HKIAC) to be settled by arbitration under the HKIAC Administered Arbitration Rules in force when the Notice of Arbitration is submitted in accordance with these rules. The arbitration shall take place in Hong Kong.

The decision of the relevant arbitration tribunal shall be final and binding on both parties. The arbitration fee shall be borne by the losing party.