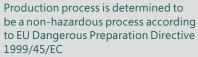


居安超高效滅殺新冠病毒細菌濾層 **Curie Ultrahigh Efficiency Viral Filter**

Efficiency 效率 VFE Virus (T4D Bacteriophage) VFE 病毒(T4D噬菌體) Killing Virus (Coronavirus COVID-19) 殺病毒率 (新冠病毒 COVID-19) BFE Bacteria (S.Aureus) BFE 細菌過濾 (金黃葡萄球菌) **BFE** Enhanced Bacteria (S.Aureus) BFE 強化濃度細菌過濾(金黃葡萄球菌) BFE Bacteria after Shelf-Life of 5 Years (S.Aureus) BFE 細菌過濾於存放5年後(金黃葡萄球菌) BFE Bacteria after 120°C for 48 hours (S.Aureus) BFE 細菌過濾於存放120度環境48小时後(金黃葡萄球菌) Killing Bacteria (S.Aureus / 60 Seconds) 殺細菌率(金黃葡萄球菌/60秒) Killing Virus (Human Orthopneumovirus H3N2) 殺病毒率 (人類呼吸道合胞病毒 H3N2) Infection Risk Complete Protection 感染風險 完全保護 intertek Nelson Labs. A Sotera Health company

Properties 特性

Test 測試	
Differential Pressure (mmH2O) 壓力差 (mmH2O)	5
Flame Spread (Second) 防火度 (秒)	5
Microbio Cleanliness (cfu) 微生物潔淨度 (cfu)	17
Shelf Life (Year) 有效保存期限 (年)	5
Formula - Food Additives A by World Health Organizat 配方成份 - 世界衛生組織核淮	ion (WHO)
Curie technology pass cher standards for baby textile p 居安技術符合以下化學安全標 兒級紡織品安全使用 EN ISO 21084:2019 EN ISO 18254:2016 EN ISO 14184:2011 JIS L 1041 DIN EN ISO 17070:2015	roducts on



64 LFBG B 82.02-08

EN ISO 14389: 2014 US CPSC-CH-C1001-09.4

根據歐盟危險制劑指令1999/45/EC規程 及其修訂版·制備被確定為無危險過程



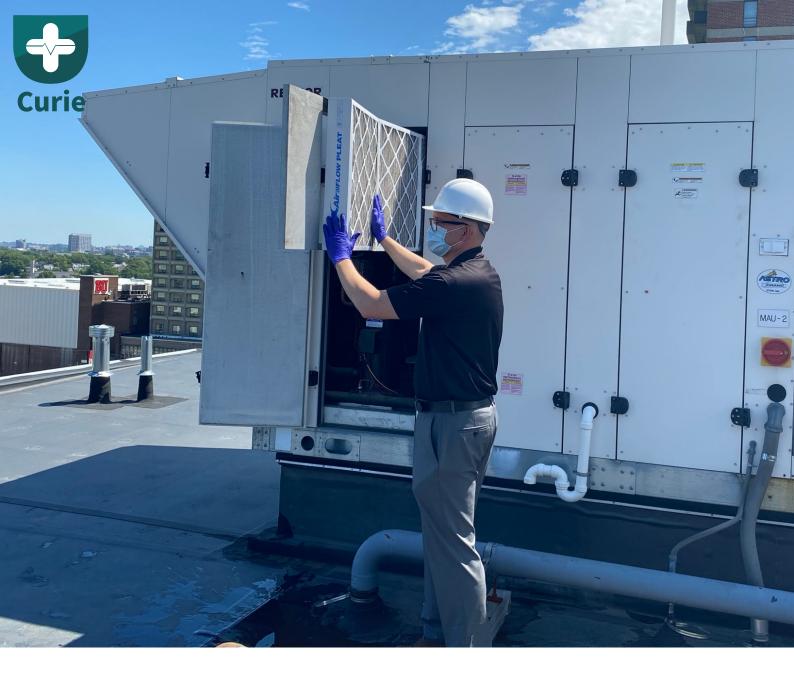












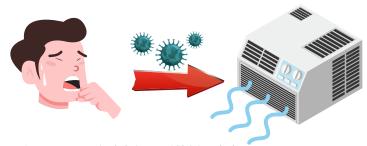
証實可以高效殺滅新冠病毒 COVID-19 (99.81%)
Proven to kill COVID-19 (99.81%)

配合超高效病毒過濾效率 VFE (>99.9a%)
Combined with Viral Filtration Efficiency VFE (>99.9a%)

比傳統靜電濾材效能高**500**倍
Over **500** times more effective than traditional electrostatic filter

不合任何毒性及金屬成份
Non toxic and non metallic composition

一次感染 Primary Infection



 呼吸
 : 釋出超過一千粒新冠病毒

 Breathing
 : > 1,000種COVID-19病毒

 咳嗽
 : 釋出超過一百萬粒新冠病毒

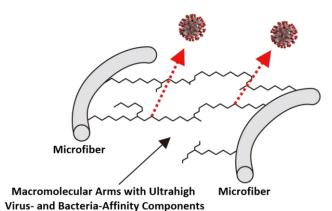
 Coughing
 : > 1,000,000種COVID-19病毒

 感染
 : 吸入超過一百粒活性新冠病毒

 Infection
 : 呼吸> 100種活躍的COVID-19病毒

濾材防護水平	穿透濾材病毒數量 No of Virus Penetrating Filter		
Type of Filter	呼吸 Breathing	咳嗽 Coughing	
VFE 95%	>50	>50,000	
VFE 99%	>10	>10,000	
Curie 居安	>1	>10	

如何攔截及殺滅新冠病毒COVID-19 How We Arrest and Kill COVID-19



居安纖維所產生的高強度梯度,與新冠狀病毒,病毒和細菌表面細菌的強韌性性的刺突蛋白正負相吸,能有效阻隔帶連續的細菌和病毒,堪稱最高等級過濾效果。

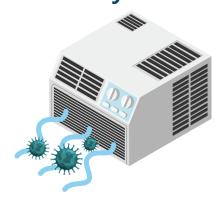
根據香港公開大學測試證明居安纖維穿透發出的極強長度,能阻隔在60秒內穿透撕裂細菌及病毒的外膜。

Spike protein over coronavirus is "strong negative charged".

The highly positive-charged fibers inside Curie's filtration material attracts and arrests "strong negative charged" protein spikes on virus and bacteria

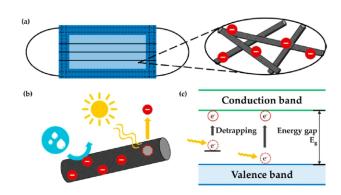
Strong attraction force can tear off protein chain over envelope of coronavirus, and achieve COVID-19 killing effect.

二次感染 Secondary Infection



- 新冠病毒能於濾材上存活7天
- Viruses can survive on the surface of traditional filter for 7 days
- 病毒和細菌數目能於使用濾材4小時後增長180倍
- Viruses can grow 180 times more after 4 hours of usage
- 瀘材上的新冠病毒進入冷氣系統造成二次感染
- COVID-19 on the filter will enter into air conditioning system
- 能高效殺滅新冠病毒的濾材,有效減少二次感染風險
- A COVID-19 killing filter can fully protect against secondary infection

傳統靜電濾材缺點 Shortcoming of Traditional Electrostatic Filter



熔噴布依靠靜電捕獲細菌和病毒,而細菌和病毒表面蛋白質帶負極,而靜電本身偏負極,兩者極性相同,捕獲效率並非為最高效。

而靜電本身帶有不穩定性,容易因炎熱天氣及潮濕環極下散失放電,在使用濾材後數小時後,過濾效率開始衰減。

Meltblown cloth relies on electrostatic to capture bacteria and viruses. Surface proteins of bacteria and viruses are negative-charged, and the electrostatic itself is negative. The polarity of the two is the same and therefore it repels – inefficient in capturing microbials.

Electrostatic itself is unstable, and it is easy to dissipate and discharge due to high temperature and humidity. Few hours after the filter used, the filtration efficiency begins to decay.

Curie Ultrahigh Efficiency Viral Filter



特強防護.殺毒殺菌.涼感透氣.耐用長效

Ultra-Protection, Anti-viral & Anti-bacterial, Light & Airy, Long-Lasting

特強防護 / Strong Protection:

防護效能達最高醫療過濾級別 (VFE / BFE > 99.9a%)·符合 **ASTM F2101 Level 3** 及 **EN14683 Type IIR** 生物危害過濾要求,用得安心。

Highest medical-grade filtration efficiency (VFE / BFE > 99.9a%), meeting biohazard filtration requirement on **ASTM F2101 Level 3** and **EN14683 Type IIR** standard, for a peace-of-mind.

殺毒殺菌 / Antiviral & Antibacterial

高效殺滅**新冠病毒 (99.81%)、病毒 (>99.9a%)** 及細菌 (>99.9a%), 殺毒殺菌時間只需 60 秒,減少二次感染風險。

Kills COVID-19 (99.81%), virus (>99.99%) and bacteria (>99.99%) in seconds, greatly reducing risk for secondary infection.

涼感透氣 / Light & Airy

超高透氣度,比同等級保護濾材透氣度高 50% 以上。

Ultra-high air permeability, 50% more air permeable vs comparable filter.

耐用長效 / Long-Lasting

攔截殺滅病毒細菌技術不依靠靜電·採用 Curie 專利強正極納米結構維持穩定效能·能於 120℃ 存放 48 小時後·維持 BFE > 99.99%·保護效能不受高溫度高濕度影響。理想儲存情況下·**有效達 5 年**之久。

Humidity and temperature proof - Curie patented strong polycationic nano structure maintain ultra filtration and disinfection efficiency, it can remain BFE > 99.99% after conditioning in 120°C for 48 hours. Effectiveness last for 5 year under optimal storage conditions.



資質 Qualification



美國專利 US Patent Number 62988900



香港專利 HK Patent Number 32020008506.8



病毒過濾效率測試 Viral Filtration Efficiency (VFE) in ASTM F2101

證實居安技術可超高效過濾病毒 (>99.9a%)

Proven that Curie technology can effectively filter virus (>99.9a%)



強化劑量細菌過濾效率測試 Bacterial Filtration Efficiency with Increased Delivery Challenge (BFE) in ASTM F2101 and EN14683

證實居安技術可超高效過濾強化劑量細菌 (99.8%)

Proven that Curie technology can effectively filter increased challenge of bacteria (99.8%)



病毒過濾效率測試 Viral Filtration Efficiency (VFE) in ASTM F2101

證實居安技術可超高效過濾病毒 (>99.9a%)

Proven that Curie technology can effectively filter virus (>99.9a%)





紡織製品抗病毒活性的測定 Determination of Antiviral Activity of Textile Products BS ISO 18184

證實居安技術可超高效殺滅新型冠狀病毒 COVID-19 (99.81%)

Proven that Curie technology can effectively kill COVID-19 Coronavirus (>99.81%)



紡織產業綜合研究所 Taiwan Textile Research Institute 空氣交換壓力差 Air Exchange Pressure in ASTM F2101 / EN14683

證實 居安技術空氣交換壓力差·符合 ASTM F2101 Level 3 及 EN14683 Type IIR 要求 Proven that air exchange pressure of Curie technology can comply standards of ASTM F2101

Level 3 and EN14683 Type IIR



細菌過濾效率測試 Bacterial Filtration Efficiency (BFE) in ASTM F2101

證實居安技術可超高效過濾細菌 (>99%)

Proven that Curie technology can effectively filter bacteria (>99%)



醫療器械無菌屏障加速老化標準 Standard Guide for Accelerated Ageing of Sterile Barrier Systems for Medical Devices in ASTM F1980-16

細菌過濾效率測試 Bacterial Filtration Efficiency (BFE) in ASTM F2101

證實居安技術在室溫 20 度存放 5 年後,可超高效過濾細菌 (>99%)

Proven that Curie technology can effectively kill bacteria (>99%)





紡織製品抗菌活性的測定 Determination of Antibacterial Activity of Textile Products BS EN ISO 20743

證實居安技術可超高效殺滅細菌 (>99%),殺菌時間少於 60 秒

Proven that Curie technology can effectively kill bacteria (>99%) · killing time is less than 60 seconds



紡織製品抗病毒活性的測定 Determination of Antiviral Activity of Textile Products BS ISO 18184

證實居安技術可超高效殺滅 H3N2 流感病毒 (>99.99%)

Proven that Curie technology can effectively kill virus (>99.99%)

紡織品烷基酚(AP)的測定 Determination of alkylphenols (AP) of Textile Products EN ISO 21084:2019

證實居安技術不含烷基酚(AP)

Proven that Alkylphenols (AP) is not detected from Curie technology

紡織品乙氧基化烷基酚的檢測 Detection and Determination of Alkylphenol Ethoxylates (APEO) of Textile Products EN ISO 18254:2016

證實居安技術不含乙氧基化烷基酚

Proven that Alkylphenol Ethoxylates (APEO) is not detected from Curie technology



紡織品甲醛的測定 Determination of Formaldehyde - Free and Hydrolysed Formaldehyde of Textile Products EN ISO 14184:2011 / JIS L 1041

證實居安技術不含甲醛·乎合 Type 1 - 嬰兒 < 36 個月安全標準

Proven that Formaldehyde is not detected from Curie technology, and it reach safety level for Type 1 – Baby < 36 Months

紡織品五氯苯酚含量測定 Determination of Tetrachlorophenol-, Trichlorophenol-, Dichlorophenol-, Monochlorophenol-Isomers and Pentachlorophenol Content of Textile Products DIN EN ISO 17070:2015 / 64 LFBG B 82.02-08 (Modified)

證實居安技術不含氯酚化合物及鄰苯基苯酚化合物

Proven that Chlorophenols Content and Ortho-Phenylphenol (OPP) are not detected from Curie technology

紡織品鄰苯二甲酸酯含量的測定 Determination of the Phthalate Content of Textile Products EN ISO 14389: 2014 / US CPSC-CH-C1001-09.4

證實居安技術不含鄰苯二甲酸酯

Proven that Phthalate Content are not detected from Curie technology

Curie Ultrahigh Efficiency Viral Filter



公司簡介 Company Introduction

居安為香港科技初創公司·專注創新技術開發、應用及產品化。公司以諾貝爾獎得主 Marie Curie (居里夫人)命名·希望 將居里夫人名言「我們不應害怕人生·反而要去瞭解。現在正是我們應該瞭解更多的時候·這樣我們就可以減少恐懼!」 融入公司使命·在新冠肺炎疫情肆虐下·本著這樣的精神對病原有新的認識·從而採取新方法保護民眾免於感染。

Curie Limited is a tech start-up based in Hong Kong, with a focused-on material science innovation and technology development, application and productization. Curie was named after Marie Curie, inspired by the Nobel Laureate's famous quote;

" Nothing in life is to be feared, it is only to be understood. Now is the time to understand more, so that we may fear less."

- Marie Curie

The new COVID-19 pandemic brought new meaning to these words of wisdom; Curie aimed to adapt and invent new methods of protection for the public against infection.

居安科研成果概要 Curie Filtration Technology

- 居安高效過濾技術利用獨有配方生物溶劑應用於不同材質,其產生之病毒過濾技術已獲專業認證機構證實成效卓越; Curie's Filtration Technology is applicable to different materials. This viral and bacterial filtration technology is proven by multiple lab tests by renowned certification organizations;
- 此發明擁有嶄新的病毒/細菌攔截及殺滅機制,該技術並已成功獲得美國及香港專利;
 Curie Filtration Technology kills virus and bacteria; obtained US and Hong Kong patents;
- 為香港科研、開發及生產成果;
 Research & development, design, and production in Hong Kong;
- 該技術可廣泛應用於空氣濾材及多種生活產品,透過產品為顧客提供高度保護;
 Wide-range of uses for combating COVID-19 pandemic: air filter and a other daily-use products, providing protection against viruses and bacteria;
- 公司率先將此高效病毒攔截 + 殺滅技術商業化,並以高性價比方式量產。
 Curie is the first to commercialize and mass-produce this high-efficiency, cost-effective virus killing technology
- 此高效過濾專利技術結合團隊在化學及應用材料方面的專業知識,令此科研成果成為目前世界上獨一無二的技術。
 This patented technology combines the team's expertise in chemistry and applied materials, cumulating in this "World 1st" technology.

Curie Ultrahigh Efficiency Viral Filter



居安高效過濾技術優勢 The Curie Advantage

居安高效過濾技術首階段運用在空氣濾材產品上,並經測試證實能達到高效過濾病毒細菌、特效防禦、長效保存、高透氣度等效果,充分解決傳統空氣濾材無法處理的問題。

The first application of Curie's Filtration Technology is applied to air filter products, and has been tested and proven to kill viruses and bacteria; "Ultra-Protection, Anti-viral & Anti-bacterial, Light & Airy, Long-Lasting".

	運用居安高效過濾技術的空氣濾材 Air Filter Using Curie's Ultrahigh-Efficiency Filtration Tech	傳統熔噴布空氣濾材 Traditional Air Filter Using Meltblown
過濾效果 Filtration / Killing Mechanism	高效: 根據 ASTM F2101 及 EN14683 測試結果顯示: VFE 病毒過濾效率高達 > 99.9a% BFE 細菌過濾效率高達 > 99.99% 居安纖維所產生的高強度正極,與冠狀病毒、病毒及細菌表面帶強負極性的刺突蛋白正負相吸,能有效阻隔帶負極的細菌及病毒,堪稱最高級別過濾效果。 High efficiency: According to ASTM F2101 and EN14683 test results show: VFE high as > 99.9a% BFE up to > 99.99% The highly positive-charged fibers inside Curie Filtration material attracts and arrests highly negative-charge protein spikes on virus and bacteria.	熔噴布依靠靜電捕獲細菌和病毒,而細菌和病毒表面蛋白質帶負極,而靜電本身偏負極,兩者極性相同,捕獲效率並非為最高效。而靜電本身帶有不穩定性,容易因炎熱天氣及潮濕環極下散失放電,在使用空氣濾材後數小時後,過濾效率開始衰減。 Meltblown cloth relies on electrostatic to capture bacteria and viruses. Surface proteins of bacteria and viruses are negative-charged, and the static electricity itself is negative. The polarity of the two is the same and therefore it repels – inefficient in capturing microbials. Electrostatic itself is unstable, and it is easy to dissipate and discharge due to high temperature and humidity. Few hours after the air filter used, the filtration efficiency begins to decay.
防禦原理 Principles of Protection	特效: 根據香港公開大學測試證明居安纖維底層發出的極強正極,能阻隔在 60 秒內透過撕裂細菌及病毒的外膜。 殺新冠病毒率 - 99.81% 殺流感病毒率 > 99.99%	傳統空氣濾材只能發揮不同程度阻隔功能,將細菌或病毒阻擋在空氣濾材外,但無做到殲滅的功能,細菌或病毒會開始在熔喷布纖維上數倍地增生,因此長時間使用時,容易造成二次感染。 Traditional air filter only acts as a barrier to block bacteria or viruses from the user, but they do not have the function of killing. This means bacteria or viruses will start to multiply on the meltblown fabric fibres, potentially causing secondary infection on long time usage.

Curie Ultrahigh Efficiency Viral Filter



	Killing Rate of Bacteria > 99.99%	
有效期 Effective Period	表別 長效 根據香港公開大學以符合 ASTM F1980 標準類比五年存儲條件所做的測試結果顯示,即便長期存儲,並可在存放於 120 度環境下 48 小時後,居安的過濾材料仍保持高效力,功能不減,解決了大企業、政府或公營機構需大量採購或可能面對產品失效的問題。 Long-lasting: Curie Filtration Technology material was tested against storage in an environment of 120 degrees for 48 hours, in accordance with ASTM F1980 standard simulating five-year storage conditions. After the test, Curie's filter materials still maintain high efficiency and function.,It allows long term storage that large enterprises, governments or public institutions require.	有效期短·更可能因為運送過程中的外在因素(如酷熱天氣或儲存不當)·導致熔噴布靜電大量流失·令產品出現未出售已失效的情況;亦或是現時何以民眾雖戴口罩·但疫情仍不受控的原因之一。 The effective period is short, and it is most likely due to external factors in the transportation process (such as high temperature or improper storage), electrostatic loss of the meltblown is significant over-time, causing the product to be ineffective.
透氣度 Air Permeability	高透氣度: 較傳統空氣濾材氣流阻力減少,透氣度高。 High air permeability: Compared with traditional air filter, the air resistance is reduced and the air permeability is high.	難在抗菌及透氣度之間取得平衡。 Most air filter sacrifice breathability in order to achieve the high filtration efficiency expected by users.

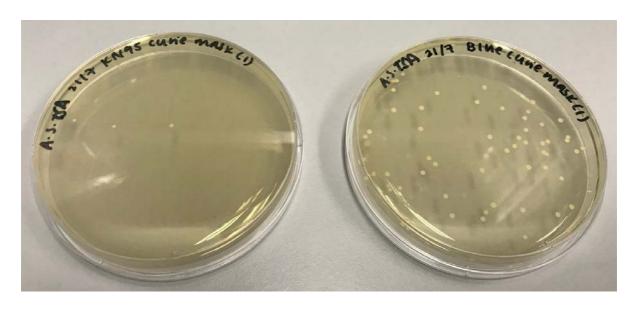
Curie Ultrahigh Efficiency Viral Filter



居安超高效滅殺新冠病毒細菌濾層與傳統熔噴布空氣濾材的比較 Comparison between Curie Air Filter and Traditional Air Filter Using Meltblown

於靜電放電後細菌過濾效率的比較

Difference on Bacterial Filtration Efficiency (BFE) after Electrostatic Discharged

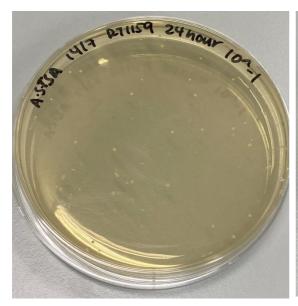


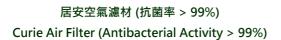
居安空氣濾材 (BFE > 99%) Curie Air Filter (BFE > 99%)

傳統熔噴布空氣濾材 (BFE < 90%) Traditional Air Filter Using Meltblown (BFE < 90%)

空氣濾材於使用 8 小時後細菌生長的比較

Difference on Bacterial Growth after 8 Hours Use of Air Filter







傳統熔噴布空氣濾材 (充滿細菌)
Traditional Air Filter Using Meltblown
(Full of Bacteria)



Sponsor: Eddie Yam Intertek Testing Services Hong Kong Ltd. 1/F, Garment Centre, 576 Castle Peak Road Kowloon, HONG KONG

Viral Filtration Efficiency (VFE) Final Report

Test Article: modified non-woven

colour: White

Style #1001

Study Number: 1280865-S01 Study Received Date: 25 Mar 2020

Testing Facility: Noison Laborate

Testing Facility: Nelson Laboratories, LLC 6280 S. Redwood Rd.

Salt Lake City, UT 84123 U.S.A.

Test Procedure(s): Standard Test Protocol (STP) Number: STP0007 Rev 16

Deviation(s): None

Summary: The VFE test is performed to determine the filtration efficiency of test articles by comparing the viral control counts upstream of the test article to the counts downstream. A suspension of bacteriophage Φ X174 was aerosolized using a nebulizer and delivered to the test article at a constant flow rate and fixed air pressure. The challenge delivery was maintained at 1.1 - 3.3 x 10³ plaque forming units (PFU) with a mean particle size (MPS) of 3.0 μ m \pm 0.3 μ m. The aerosol droplets were drawn through a six-stage, viable particle, Andersen sampler for collection. The VFE test procedure was adapted from ASTM F2101.

All test method acceptance criteria were met. Testing was performed in compliance with US FDA good manufacturing practice (GMP) regulations 21 CFR Parts 210, 211 and 820.

Test Side: Either

Test Area: ~40 cm²

VFE Flow Rate: 28.3 Liters per minute (L/min)

Conditioning Parameters: 85 ± 5% relative humidity (RH) and 21 ± 5°C for a minimum of 4 hours

Positive Control Average: 1.6 x 10³ PFU
Negative Monitor Count: <1 PFU

MPS: 2.9 µm

Study Director





James W. Luskin

Study Completion Date

1280865-501

801-290-7500 | nelsonlabs.com

sales@nelsonlabs.com

myf

FRT0007-0001 Rev 16 Page 1 of 2



Results:

Test Article Number	Percent VFE (%)	
1	>99.9ª	
2	>99.9 ^a	
3	>99.9ª	
4	>99.9 ^a	
5	>99.9ª	

^a There were no detected plaques on any of the Andersen sampler plates for this test article.

The filtration efficiency percentages were calculated using the following equation:

$$\% VFE = \frac{C - T}{C} \times 100$$

C = Positive control average

T = Plate count total recovered downstream of the test article Note: The plate count total is available upon request



myf



Sponsor: Wong Chak Ming Hong Kong Medical Supply Limited Unit A3, 2F Mai Wah Industrial Bldg. 1-7 Wah Sing Street Hong Kong, CHINA

Bacterial Filtration Efficiency (BFE) Final Report

Test Article: HKMSLMASK000 Purchase Order: HKMSLPO20200326

Study Number: 1282265-S01 Study Received Date: 28 Mar 2020

Testing Facility: Nelson Laboratories, LLC

6280 S. Redwood Rd.

Salt Lake City, UT 84123 U.S.A.

Test Procedure(s): Standard Test Protocol (STP) Number: STP0004 Rev 18

Deviation(s): None

Summary: The BFE test is performed to determine the filtration efficiency of test articles by comparing the bacterial control counts upstream of the test article to the bacterial counts downstream. A suspension of Staphylococcus aureus was aerosolized using a nebulizer and delivered to the test article at a constant flow rate and fixed air pressure. The challenge delivery was maintained at 3.5×10^3 colony forming units (CFU) with a mean particle size (MPS) of $3.0 \pm 0.3 \, \mu m$. The aerosols were drawn through a six-stage, viable particle, Andersen sampler for collection. This test method complies with ASTM F2101-19 and EN 14683:2019, Annex B; with the exception of the higher challenge level, which may represent a more severe test.

All test method acceptance criteria were met. Testing was performed in compliance with US FDA good manufacturing practice (GMP) regulations 21 CFR Parts 210, 211 and 820.

Test Side: Inside BFE Test Area: ~40 cm²

BFE Flow Rate: 28.3 Liters per minute (L/min)

Conditioning Parameters: 85 ± 5% relative humidity (RH) and 21 ± 5°C for a minimum of 4 hours

Test Article Dimensions: ~176 mm x ~160 mm

Positive Control Average: 3.5 x 10³ CFU
Negative Monitor Count: <1 CFU

MPS: 3.0 µm

The positive control average was out of specification per STP0004 Rev 18 section 6.1 which states, "The BFE positive control average shall be maintained at 1.7-3.0 x 10³ CFU." Testing with a more severe challenge to the test articles represents a worse case. The sponsor accepted the use of the higher challenge; therefore, the results are considered valid at the testing conditions that occurred.





ALL MACH OUN

James W. Luskin

20 Apr 2020 Study Completion Date



801-290-7500 | nelsonlabs.com

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brd

FRT0004-0001 Rev 22

Page 1 of 2



Results:

Test Article Number	Percent BFE (%)	
1	99.8	
2	99.8	
3	99.8	
4	99.8	
5	99.8	

The filtration efficiency percentages were calculated using the following equation:

$$\% BFE = \frac{C - T}{C} \times 100$$

C = Positive control average

T = Plate count total recovered downstream of the test article Note: The plate count total is available upon request



brd



Number: HKGT05112613-S1

Applicant: CURIE LIMITED

B3-1 G/F

SUPERLUCK INDL CTR PHASE 2

57 SHA TSUI RD TSUEN WAN NT HK

Attn: ALDRIN OR

Date: Apr 22, 2020

This is to supersede report no. HKGT05112613 dated Apr 21,

2020

Sample Description As Declared:

No. Of Sample : Several

Buyer's Name : -Agent's Name : -

Manufacturer's Name: Curie Limited

Sample Description : Curie Ultrahigh-Efficiency Viral Filter超高效病毒濾材

Colour : White Style No. : 1001 Order No. / PO No. : -Product End Uses : -

Fibre Content : Nonwoven Fabric/GMT Weight : 20g : -

Date Received/Date Test Started : Apr 15, 2020

Applicant's Provided Care Instruction/Label:

For and on behalf of

Intertek Testing Services Hong Kong Limited

Teddy Y. N. Chung

Director



Page 1 Of 4





Number: HKGT05112613-S1

Original Sample Photo:



For any queries on this report, you are welcome to contact our customer service representatives:

<u>US3</u>

Angie Yu (852) 98639123 or email to angie.yu@intertek.com

For and on behalf of

Intertek Testing Services Hong Kong Limited

Teddy Y. N. Chung

Director









Number: HKGT05112613-S1

Tests Conducted (As Requested By The Applicant)

1 Evaluation of Viral Filtration Efficiency (VFE):

Summary: The VFE test is performed to determine the filtration efficiency of test articles by comparing the viral control counts upstream of the test article to the counts downstream. A suspension of bacteriophage Φ X174 was aerosolized using a nebulizer and delivered to the test article at a constant flow rate and fixed air pressure. The challenge delivery was maintained at $1.1-3.3 \times 10^3$ plaque forming units (PFU) with a mean particle size (MPS) of $3.0 \pm 0.3~\mu$ m. The aerosols droplets were drawn through a six-stage, viable particle, Andersen sampler for collection. The VFE test procedure was adapted from ASTM F2101.

All test method acceptance criteria were met. Testing was performed in compliance with US FDA good manufacturing practice (GMP) regulations 21 CFR Parts 210, 211 and 820.

Test side: Either

Test Area: ~40 cm²

VFE Flow Rate: 28.3 Liters per minute (L/min)

Conditioning Parameters: 85 ± 5% relative humidity (RH) and 21 ± 5 °C for a minimum of 4 hours

Positive Control Average: 1.6 x 10³ PFU

Negative Monitor Count: <1 PFU

MPS: 2.9 μ m





Number: HKGT05112613-S1

Tests Conducted (As Requested By The Applicant)

Evaluation of Viral Filtration Efficiency (Cont'd)

Result:

Test Article Number	Percent VFE (%)	
1	>99.9°	
2	>99.9 ^a	
3	>99.9ª	
4	>99.9a	
5	>99.9ª	

^a There were no detected plaques on any of the Andersen sampler plates for this test article.

The filtration efficiency percentages were calculated using the following equation:

$$\%VFE = \frac{C - T}{C}x100$$

C= Positive control average

T= Plate count total recovered downstream of the test article Note: The plate count total is available upon request

Remark: The test was conducted by competent subcontractor lab.

End of Report

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This report shall not be reproduced, except in full.





CURIE LIMITED To: Attention: ALDRIN OR

Report Revision Notification Re

Report Number HKGT05112613 date APR 21, 2020

Please be informed that all the content recorded in the above captioned report will be void. This captioned report is now superseded by a revised Report, Number HKGT05112613-S1, issued on Apr 22, 2020.

Date: Apr 22, 2020

Thank you for your attention

For and on behalf of

Intertek Testing Services Hong Kong Limited

Teddy Y. N. Chung

Director





Test Report

Sample Category	Curie Ultrahigh-Efficiency Viral Filter for KV-99		
Client	Curie Limited		
Test Category	Test Entrust		
Date of Report	2020.08.18		



Report No: ATCCR20081010F

Detection Information

Client	Curie Limited		Sample Source	Inspect
Client address	Room C,23/F,Tsuen Tung Factory Building,38-40 Chai Wan KOK Street, Tsuen Wan,Hong Kong		Sample State	Normal
Date of Receives samples	2020.08.10		Date(s) of tests	2020.08.10-2020.08.18
Sample No	ATCCR20081010F-0810CP01			
Category	Test Project	Test Standard and	Method	Test Instruments
Curie Ultrahigh-Efficiency Viral Filter for KV-99	Antiviral Activity Value (COVID-19)	ISO 18184:2019 Textiles Determination of antiviral		Biosafety Cabinet

End

Remarks



Production units: Curie Limited Trademarks: Curie

Date of production: 2020.06.09

Sample model: Curie Ultrahigh-Efficiency Viral Filter for KV-99

Sample batch: 1001

Report Preparer: Authorized Signatory:

Report Reviewer: t Date of Issues Repor:

(Special Chapter for Inspection and Inspection)

Test results

Virus Types	(NO)	lg(Va _{0h}) (lgTCID ₅₀ /mL)	lg(Vb _{2h}) (lgTCID ₅₀ /mL)	$\lg(Vc_{2h})$ ($\lg TCID_{50}/mL$)
	1	6.73	6.68	3.7
COVID-19 virus MDCK cells	2	6.68	6.56	4
	3	6.7	6.57	3.9
Average Value of lgTCID ₅₀ /mL		6.70	6.61	3.88
Antiviral Activity	ntiviral Activity Value 2.72			
Antiviral Activity l	Rate (%)	99.81		



End

Beijing Shantong Medical Testing Laboratory

Declaration of Test Results

Beijing Shantong Medical Testing Lab "BSMTL" hereby declares that the test item described below has been tested by BSMTL and complies with the requirements of

ISO 18184: 2019 Textile Determination of Antiviral

The complete detail of the tests performed and the results are recorded in

Report No: ATCCR20081010F Dated: 18.08.2020

Description of item tested: Curie Ultrahigh-Efficiency Viral Filter for KV-99

Virus Tested: SARS-COV-2 / COVID-19 MDCK Cells

Summary of test results -

Antiviral Activity Value: 2.72

Antiviral Activity Rate: 99.81%

Submitted by:

is issued, namely:

Curie Limited

Room C, 23/F, Tsuen Tung Factory Building,38-40 Chai Wan Kok Street, Tsuen Wan, Hong Kong SAR

Declaration authorised by:

Title:

Date: 02/09/2020

This Declaration applied only to the particular sample tested and to the specific tests carried out as detailed in the Report referred to above.

1. This declaration does not indicate provide or imply any measure of Approval, Certification, Supervision, Control or Surveillance by BSMTL to this or any related product.

Attention is drawn to the conditions upon which this declaration

The general and specific conditions of the BSMTL Conditions of Contract for Testing, apply in all respects.

Beijing Shantong Medical Testing Laboratory Co. Ltd., Fangshan, Beijing, China



统一社会信用代码 91110111MA01A4KK4D

营业执照

(副 本) (1-1)



■ 扫描二座码登录 "国家企业信息 信息企业系统" 了解型多登记。 各家、许可、当

2

称 北京普通医学格曼实验室有限公司

*

型有限责任公司法人独资

法定代表人 杨益

经营范围

医学检验医疗服务、技术开发、技术核让、技术咨询、技术服务。(企业依法》上选择公前项目,开展经营活动。 医疗服务以及依法须经批准的项目,经相关部门批准后依 批准的内容开展经营活动;不得从事本市产业政策禁止和 限制类项目的经营活动。) 注册资本 600万元

成立日期 2018年01月24日

营业期限 2018年01月24日至 2048年01月23日

住 所 北京市房山区拱辰街道办事处学园北街11号综合 服务楼一层5106

登记机关



市场主体应当于每年1月1日至6月30日通过 国家企业信用信息公示系统报送公示年度报告。



共 人民 围

杨益 法定代表人

主要负责人

007913110111417919 登记号

30 H 2022 年 12月 02日至 该医疗机构经核准登记,准予执业



发证机关

北京市房山区卫生健康委员会

发证日期

8月 14日 2019 年

开户许可证

核准号: J1000210464502

编号: 1000-03577720

经审核, 北京善通医学检验实验室有限公司

符合开户条件,准予

开立基本存款账户。

法定代表人(单位负责人) 杨益

账号 321580100100032556



北京市卫生健康委员会

北京市卫生健康委员会关于同意 密云区医院等 11 家检测机构 开展新型冠状病毒核酸检测的通知

西城区、朝阳区、丰台区、房山区、顺义区、大兴区、密云 区卫生健康委,经济开发区,市疾控中心,市医学检验质控 中心,各相关医疗机构:

根据北京市密云区医院、北京市西城区展览路医院、北京市丰台区铁营医院、北京市朝阳区三环肿瘤医院、北京朝阳急诊抢救中心、北京朝阳中西医结合急诊抢救中心、北京市大兴区中西医结合医院、北京北亚骨科医院、北京德威铭达医学检验所、北京善通医学检验实验室中北京索真医学检验实验室等11家检测机构(以下简称11家检测机构)提交的开展新冠病毒核酸检测的申请,结合专家评估意见,经研究,现就有关事项通知如下:

- 一、同意 11 家检测机构开展新型冠状病毒核酸检测工作。
- 二、11 家检测机构要严格按照国家和本市关于开展新型 冠状病毒核酸检测、生物安全防护、生物样本资源管理的有

北京市病原微生物实验室及实验室活动备案通知书

京房山卫实验室备字[2020]第043号

北京善通医学检验实验室有限公司:

你单位于2020年06月27日提交的北京市病原微生物实验室及实验活动备案材料如下:

- 1. ☑《北京市病原微生物实验室及实验室活动备案表》;
- 2. ☑ 实验室或实验室设立单位的法人资格证明;
- 3. ☑ 实验室设立单位生物安全组织管理框架图:
- 4. ☑ 实验室布局平面图:

经本机关审查,认为申请材料齐全、符合《北京市病原微生物实验室及实验活动备案管理办法》的要求,决定予以备案。

卫生健康行政部门 (新章)

2020年06月27日

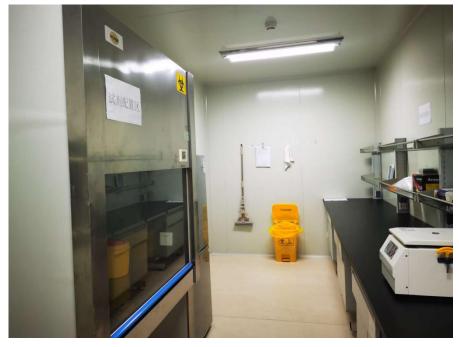
备注:此备案旨在了解你单位实验室及其实验活动基本状况,不作为审批依据。请你单位备案后,严格按照《中华人民共和国传染病法》、《病原微生物实验室生物安全管理条例》和《人间传染的高致病性病原微生物实验室和实验活动生物安全审批管理办法》等相关法律法规规定,从事相关实验活动,规范实验室管理。

北京市卫生健康委员会制定

北京善通医学检验实验室

试剂制备间





样本制备间







扩增间





样本间和灭菌间







实验室走廊



財團法人紡織產業綜合研究所 Taiwan Textile Research Institute





TEST REPORT TUCHENG

Date: Aug. 31, 2020 Date of Receipt: Aug. 24, 2020

Report No.: TFF9H598 Quantity: 1PC Page Order/Pages: P1/5 Ref. No.: NIL

Report Title: Curie Limited(U3104)

Item: Mask

Address: No. 108, Ganhe Rd., Xitun Dist., Taichung City 407, Taiwan

Test Items		Test Results	Test Methods
Sub-Micron Particulate	1	99.49	ASTM F2100-2019 9.3
Filtration Efficiency(%)	2	99.45	ASTM F2299-2017
(0.1μm PSL)	3	99.46	Flow rate:28.1
	4	99.48	(Liter/min)
	5	99.48	
	Ave.	99.47	
Air Exchange Pressure		5.3	ASTM F2100-2019 9.2
(mmH2O/cm³)	2	5.3	EN 14683:2019 Annex C
	3	4.9	
	4	5.1	
	5	5.3	
Flammability (as Received)		DNI	ASTM F2100-2019 9.5 CPSC 16 CFR 1610-2008

Note: 1mmH2O=9.8Pa.

Note: Air Exchange Pressure takes 5 pieces for testing.

Note: Flammability takes 20 samples for testing.

Note: "DNI":Did Not Ignite.

Note: Sample description is given by the client: Brandname: Curie

Model: Curie KV99 Face Mask

Hong Kong address: Unit C, 23F, Tsun Tung Factory Bldg 38-40

Chai Wan Kok St, Tsuen Wan. NT Hong Kong

Note: The test report is the test results issued by the testing institution as requested by the consignor, it shall not determine the legitimacy of the product.

Note: 1. This report is only responsible for the submitted sample(s), which will be kept for one month period.

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Authorized by president of

Taiwan Textile Research Institute

Jui - hung kao

Director,
Department of Testing and
Certification

Department of Testing and Certification, Taiwan Textile Research Institute No.6, Chengtian Rd., Tucheng Dist., New Taipei City 23674, Taiwan (R.O.C.)

Tel: +886-2-22670321 ext. 7107, 7110 Fax: +886-2-22675108, +886-2-22689839

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TEST REPORT TUCHENG

Date: Aug.07,2020	Date of Receipt:	Jul.27,2020

Report No.: TAG9G706 Quantity: 1PC Page Order/Pages: (P2/5) Ref. No.: NIL

Report Title: Curie Limited(U3104)

Mask

Address: No. 108, Ganhe Rd., Xitun Dist., Taichung City 407, Taiwan

Test Items		Test Results	Test Methods
Synthetic Blood Penetration	1	None Seen	ASTM F2100-2019 9.4
Pressure:160 mmHg	2	None Seen	ASTM F1862-2017
	3	None Seen	
	4	None Seen	
	5	None Seen	
	6	None Seen	
	7	None Seen	
	8	None Seen	
	9	None Seen	
	10	None Seen	
	11	None Seen	
	12	None Seen	5
	13	None Seen	
	14	None Seen	= °
	15	None Seen	
	16	None Seen	

Note: Sample description is given by the client: Brandname: Curie

Model: Curie KV99 Face Mask

Hong Kong address: Unit C, 23F, Tsun Tung Factory Bldg 38-40

Chai Wan Kok St, Tsuen Wan. NT Hong Kong

Note: The test report is the test results issued by the testing institution as requested by the consignor, it shall not determine the legitimacy of the product.

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TEST REPORT TUCHENG

Date: Aug. 07, 2020	Date of Receipt:	Jul.27,2020
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Report No.: TAG9G706 Quantity: 1PC Page Order/Pages: (P3/5) Ref. No.: NIL

Report Title: Curie Limited(U3104)

Mask

Address: No. 108, Ganhe Rd., Xitun Dist., Taichung City 407, Taiwan

Test Items		Test Results	Test Methods
Synthetic Blood Penetration	17	None Seen	ASTM F2100-2019 9.4
Pressure:160 mmHg	18	None Seen	ASTM F1862-2017
	19	None Seen	
	20	None Seen	
	21	None Seen	
	22	None Seen	
	23	None Seen	
	24	None Seen	
	25	None Seen	
	26	None Seen	172
	27	None Seen	100
	28	None Seen	5
	29	None Seen	
	30	None Seen	-
	31	None Seen	
	32	None Seen	

Note: Sample description is given by the client: Brandname: Curie

Model: Curie KV99 Face Mask

Hong Kong address: Unit C, 23F, Tsun Tung Factory Bldg 38-40

Chai Wan Kok St, Tsuen Wan. NT Hong Kong

Note: The test report is the test results issued by the testing institution as requested by the consignor, it shall not determine the legitimacy of the product.

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Jw - hung kao

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Tel: +886-2-22670321 ext. 7107, 7110 Fax: +886-2-22675108, +886-2-22689839





TEST REPORT TUCHENG

Date: Aug. 31, 2020 Date of Receipt: Aug. 24, 2020

Report No.: TFF9H598 Quantity: 1PC Page Order/Pages: P4/5 Ref. No.: NIL

Report Title: Curie Limited(U3104)

Item: Mask

Address: No. 108, Ganhe Rd., Xitun Dist., Taichung City 407, Taiwan

Test Items		Test Results	Test Methods
Bacterial Filtration	1	99.9	ASTM F2100-2019 9.1
Efficiency (BFE)(%)	2	99.9	ASTM F2101-2019
Staphylococcus aureus	3	> 99.9	
ATCC 6538	4	99.9	
	5	99.9	

Note: Control average: 2640 CFU.

Note: Mean particle size: $2.8 \mu m$.

Note: Testing side: outside of specimen.

Note: Testing area: 39.5 cm².

Note: Flow rate: 28.3 L/min.

Note: Sample description is given by the client: Brandname: Curie

Model: Curie KV99 Face Mask

Hong Kong address: Unit C, 23F, Tsun Tung Factory Bldg 38-40

Chai Wan Kok St, Tsuen Wan. NT Hong Kong

Note: The test report is the test results issued by the testing institution as requested by the consignor, it shall not determine the legitimacy of the product.



Note: 1. This report is only responsible for the submitted sample(s), which will be kept for one month period.

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Taiwan Textile Research Institute

Jui-hung kao

Director,
Department of Testing and
Certification

Department of Testing and Certification, Taiwan Textile Research Institute No.6, Chengtian Rd., Tucheng Dist., New Taipei City 23674, Taiwan (R.O.C.)





TEST REPORT TUCHENG

Date: Aug. 31, 2020 Date of Receipt: Aug. 24, 2020

Report No.: TFF9H598 Quantity: 1PC Page Order/Pages. P5/5) Ref. No.: NIL

Report Title: Curie Limited(U3104)

Item: Mask

Address: No. 108, Ganhe Rd., Xitun Dist., Taichung City 407, Taiwan

TFF9H598

TFF9H598

Made In Taiwan

G

Note: 1. This report is only responsible for the submitted sample(s), which will be kept for one month period.

- 2. This report cannot be reproduced in any way, except in full context, without the prior approval in writing of this Department of Testing and Certification.
- The test report should not be used for public advertisement and commercial promotion.

Authorized by president of Taiwan Textile Research Institute



Department of Testing and Certification, Taiwan Textile Research Institute No.6, Chengtian Rd., Tucheng Dist., New Taipei City 23674, Taiwan (R.O.C.)





日期 Date: 2020.08.31 收件日期 Date of Receipt:

2020.08.24

TEST REPORT

土城場區 TUCHENG

報告編號 Report No.:___

TFF9H598 W Quantity:

1件

報告頁次/頁數 (P1/5) Page Order/Pages: ____ 來文字號 空 白

Order/Pages: Ref. No.:

Kei. No.: ____ 試件類別 □罩

Item:_

報告抬頭 Report Title: 地址 40

407 台中市西屯區甘河路108號

居安有限公司(U3104)

Address:

試 驗 項 目		試驗結果	試 驗 方 法
微粒過濾效率	1	99.49	ASTM F2100-2019 9.3
(0.1 μm PSL)	2	99.45	ASTM F2299-2017
	3	99.46	流量:28.1 (Liter/min)
	4	99.48	
	5	99.48	
	平均	99,47	
空氣交換壓力差	1.4	5.3	ASTM F2100-2019 9.2
(mmH2O/cm²)	2	5.3	EN 14683:2019 Annex C
	3	4.9	3
	4	5.1	0
	5	5.3	18
現狀防火性	3 / /	DNI	ASTM F2100-2019 9.5 CPSC 16 CFR 1610-2008

註: 1mmH2O=9.8Pa。

註:空氣交換壓力差,取5個樣品測試。

註: 現狀防火性 取20個試片測試

註: "DNI"意謂不燃燒。

註:依委託者所提供來樣資料為: Brandname: Curie

Model: Curie KV99 Face Mask

香港地址: 香港新界荃灣角街 38-40 號 銓通工業大廈 23 樓C室

註: 試驗報告僅就委託者之委託事項提供試驗結果,不對產品合法性做判斷。

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Authorized by president of Taiwan Textile Research Institute

紡織產業綜合研究所檢測及驗證部 新北市土城區承天路6號 Department of Testing and Certification, Taiwan Textile Research Institute No.6, Chengtian Rd., Tucheng Dist., New Taipei City 23674, Taiwan (R.O.C.)





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日期 2020.08.07 Date:

收件日期 Date of Receipt:

2020.07.27

TEST REPORT

土城場區 TUCHENG

報告編號 Report No.:

TAG9G706 Quantity: 1件

報告頁次/頁數 (P2/5) Page Order/Pages:

來文字號 空 Ref. No .:

居安有限公司(U3104)

試件類別 □罩

Report Title:

Item:_

地址

407 台中市西屯區甘河路108號

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	To		

報告抬頭

試 驗 項 目		試驗結果	試 驗 方 法
合成血液穿透性	1	無穿透	ASTM F2100-2019 9.4
壓力:160 mmHg	2	無穿透	ASTM F1862-2017
	3	無穿透	
	4	無穿透	
	5	無穿透	
	6	無穿透	
	7	無穿透	
	8	無穿透	
	9	無穿透	3
	10	無穿透	0
	11	無穿透	18
The same of the sa	12	無穿透	
	13	無穿透	29
60	14	無穿透	
a D	15	無穿透	
	16	無穿透	B

註:依委託者所提供來樣資料為: Brandname: Curie

Model: Curie KV99 Face Mask

香港地址: 香港新界荃灣角街 38-40 號 銓通工業大廈 23 樓C室

註: 試驗報告僅就委託者之委託事項提供試驗結果,不對產品合法性做判斷

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Authorized by president of Taiwan Textile Research Institute

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日期 2020.08.07 Date:

收件日期 Date of Receipt:

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TEST REPORT

土城場區 TUCHENG

報告編號 Report No .:

TAG9G706 Quantity: 1件

報告頁次/頁數 (P3/5) Page Order/Pages:

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Ref. No .: 試件類別 □罩

Item:_

報告抬頭 Report Title: 地址

407 台中市西屯區甘河路108號

居安有限公司(U3104)

Address: -

試驗項目		試驗結果	試 驗 方 法
 合成血液穿透性	17	無穿透	ASTM F2100-2019 9.4
壓力:160 mmHg	18	無穿透	ASTM F1862-2017
	19	無穿透	
	20	無穿透	
	21	無穿透	
	22	無穿透	
	23	無穿透	
	24	無穿透	
	25	無穿透	8
	26	無穿透	00
4	27	無穿透	18
A.	28	無穿透	
	29	無穿透	29
6	30	無穿透	=10
A	31	無穿透	
	32	無穿透	

註:依委託者所提供來樣資料為: Brandname: Curie

Model: Curie KV99 Face Mask

香港地址: 香港新界荃灣角街 38-40 號 銓通工業大廈 23 樓C室

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Authorized by president of Taiwan Textile Research Institute

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日期 2020.08.31 Date:

收件日期 Date of Receipt:

2020.08.24

TEST REPORT

土城場區 TUCHENG

報告編號 TFF9H598 Report No .:

Quantity:

報告頁次/頁數 (P4/5) Page Order/Pages: 1件

來文字號 空

報告抬頭

居安有限公司(U3104)

Ref. No .: 試件類別 □罩

Item:_

Report Title: -

407 台中市西屯區甘河路108號

地址 Address: -

試驗項目		試 驗 結 果	試驗方法
細菌過濾效率(%)	1	99.9	ASTM F2100-2019 9.1
金黃色葡萄球菌	2	99.9	ASTM F2101-2019
ATCC 6538	3	> 99.9	
	4	99.9	
	5	99.9	

註: 對照組的平均菌落數: 2640 CFU。

註: 平均粒徑:2.8 μm。

註: 測試面:外側。

註: 測試面積為39.5 cm。

註: 測試流量: 28.3 L/min。

註:依委託者所提供來樣資料為: Brandname: Curie

Model: Curie KV99 Face Mask

香港地址: 香港新界荃灣角街 38-40 號 銓通工業大廈 23 樓C室

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紡織產業綜合研究所檢測及驗證部 新北市土城區承天路6號 Department of Testing and Certification, Taiwan Textile Research Institute No.6, Chengtian Rd., Tucheng Dist., New Taipei City 23674, Taiwan (R.O.C.)





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日期 2020.08.31 Date:

收件日期 Date of Receipt:

2020.08.24

試驗報告 TEST REPORT

土城場區 TUCHENG

報告編號 TFF9H598 Report No.:

1件

報告頁次/頁數 (P5/5) Page Order/Pages:

來文字號 空

FF90.398 Quantity: _____ 居安有限公司(U3104)

Ref. No .: 試件類別 □罩

Item:_

報告抬頭 Report Title:

407 台中市西屯區甘河路108號

地址 Address: -

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紡織產業綜合研究所檢測及驗證部 新北市土城區承天路6號 Department of Testing and Certification, Taiwan Textile Research Institute No.6, Chengtian Rd., Tucheng Dist., New Taipei City 23674, Taiwan (R.O.C.)

Tel: +886-2-22670321 ext. 7107, 7110 Fax: +886-2-22675108, +886-2-22689839

检测及检查率 古 7山 任回





TEST REPORT TUCHENG

Date: Aug. 31,2020 Date of	of Receipt:	Aug.24,2020
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Report No.: TFF9H599 Quantity: 1PC Page Order/Pages. Ref. No.: NIL

Report Title: Curie Limited(U3104)

Mask

Address: No. 108, Ganhe Rd., Xitun Dist., Taichung City 407, Taiwan

Test Items		Test Results	Test Methods
Air Exchange Pressure	1	50.7	EN 14683:2019 Annex C
(Pa/cm²)	2	49.1	
	3	49.7	
	4	51.3	
	5	48.8	

Note: Sample description is given by the client: Brandname: Curie

Model: Curie KV99 Face Mask

Hong Kong address: Unit C, 23F, Tsun Tung Factory Bldg 38–40

Chai Wan Kok St, Tsuen Wan. NT Hong Kong

Note: The test report is the test results issued by the testing institution as requested by the consignor, it shall not determine the legitimacy of the product.



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Taiwan Textile Research Institute

Jui - hung kao

Director,
Department of Testing and
Certification

Department of Testing and Certification, Taiwan Textile Research Institute No.6, Chengtian Rd., Tucheng Dist., New Taipei City 23674, Taiwan (R.O.C.)





TEST REPORT TUCHENG

Date: Aug. 31, 2020	Date of Receipt:	Aug.24,2020

Report No.: TFF9H599 Quantity: 1PC Page Order/Pages: (P2/5) Ref. No.: NIL

Report Title: Curie Limited(U3104)

Mask

Address: No. 108, Ganhe Rd., Xitun Dist., Taichung City 407, Taiwan

Test Items		Test Results	Test Methods
Synthetic Blood Penetration	1	None Seen	EN 14683:2019
Pressure:120 mmHg (16.0 kPa)	2	None Seen	ISO 22609:2004
	3	None Seen	
	4	None Seen	
	5	None Seen	
	6	None Seen	
No.	7	penetration	
	8	None Seen	
	9	None Seen	
	10	None Seen	2
	11	None Seen	
2	12	None Seen	5
63	13	None Seen	
	14	None Seen	- ·
	15	None Seen	
	16	None Seen	

Note: Sample description is given by the client: Brandname: Curie

Model: Curie KV99 Face Mask

Hong Kong address: Unit C, 23F, Tsun Tung Factory Bldg 38-40

Chai Wan Kok St, Tsuen Wan. NT Hong Kong

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Jui-hung kao

Director, Department of Testing and Certification

Department of Testing and Certification, Taiwan Textile Research Institute No.6, Chengtian Rd., Tucheng Dist., New Taipei City 23674, Taiwan (R.O.C.)





TEST REPORT TUCHENG

Date: Aug. 31, 2020 Da	e of Receipt: Aug. 24, 2020
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Report No.: TFF9H599 Quantity: 1PC Page Order/Pages: (P3/5) Ref. No.: NIL

Report Title: Curie Limited(U3104)

Item: Mask

Address: No. 108, Ganhe Rd., Xitun Dist., Taichung City 407, Taiwan

Test Items		Test Results	Test Methods
Synthetic Blood Penetration	17	None Seen	EN 14683:2019
Pressure:120 mmHg (16.0 kPa)	18	None Seen	ISO 22609:2004
	19	None Seen	
	20	None Seen	
	21	None Seen	
	22	None Seen	
No.	23	None Seen	
	24	None Seen	
	25	None Seen	
	26	None Seen	2
	27	None Seen	
2	28	None Seen	29
66	29	None Seen	
	30	None Seen	5°
	31	None Seen	
	32	None Seen	

Note: Sample description is given by the client: Brandname: Curie

Model: Curie KV99 Face Mask

Hong Kong address: Unit C, 23F, Tsun Tung Factory Bldg 38-40

Chai Wan Kok St, Tsuen Wan. NT Hong Kong

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TEST REPORT TUCHENG

Date: Aug. 31, 2020 Date of Receipt: Aug. 24, 2020

Report No.: TFF9H599 Quantity: 1PC Page Order/Pages: P4/5) Ref. No.: NIL

Report Title: Curie Limited(U3104)

Item: Mask

Address: No. 108, Ganhe Rd., Xitun Dist., Taichung City 407, Taiwan

Test Items		Test Results	Test Methods
Bacterial Filtration	1	> 99.9	EN 14683:2019 Annex B
Efficiency (BFE)(%)	2	99.9	
Staphylococcus aureus	3	99.9	
ATCC 6538	4	> 99.9	
	5	99.9	

Note: Control average: 2640 CFU.

Note: Mean particle size: $2.8~\mu\,\mathrm{m}.$

Note: Testing side: outside of specimen.

Note: Testing area: 39.5 cm².

Note: Flow rate: 28.3 L/min.

Note: Sample description is given by the client: Brandname: Curie

Model: Curie KV99 Face Mask

Hong Kong address: Unit C, 23F, Tsun Tung Factory Bldg 38-40

Chai Wan Kok St, Tsuen Wan. NT Hong Kong

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TEST REPORT TUCHENG

Date: Aug. 07, 2020	Date of Receipt:	Jul.27,2020
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Report No.: TFF9G707 Quantity: 1PC Page Order/Pages: Psf. No.: NIL

Report Title: Curie Limited(U3104)

Mask

Address: No. 108, Ganhe Rd., Xitun Dist., Taichung City 407, Taiwan

Test Items		Test Results	Test Methods
Microbial cleanliness	1	13.7	EN 14683:2019
(cfu/g)	2	10.6	EN ISO 11737-1:2018
	3	4.4	
	4	10.1	
	5	11.0	

Note: Sample description is given by the client: Brandname: Curie

Model: Curie KV99 Face Mask

Hong Kong address: Unit C, 23F, Tsun Tung Factory Bldg 38–40

Chai Wan Kok St, Tsuen Wan. NT Hong Kong

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Jui-hung Kao

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TEST REPORT TUCHENG

Date: Aug. 31, 2020 Date of Receipt: Aug. 24, 2020

Report No.: TFF9H599 Quantity: 1PC Page Order/Pages. P5/5) Ref. No.: NIL

Report Title: Curie Limited(U3104)

Item: Mask

Address: No. 108, Ganhe Rd., Xitun Dist., Taichung City 407, Taiwan

TFF9H599

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日期 2020.08.31 Date:

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試驗報告 TEST REPORT

土城場區 TUCHENG

報告頁次/頁數 (P1/5) Page Order/Pages:____ 來文字號 空 報告編號 1件 TFF9H599 Report No .: Ref. No .:

Fr96.399 Quantity:___ 居安有限公司(U3104) 試件類別 □罩 報告抬頭 Report Title: -Item:_

地址 407 台中市西屯區甘河路108號

Address: -

試驗項目		試驗結果	試驗方法
空氣交換壓力	1	50.7	EN 14683:2019 Annex C
(Pa/cm²)	2	49.1	
	3	49.7	
	4	51.3	
	5	48.8	

註:依委託者所提供來樣資料為: Brandname: Curie

Model: Curie KV99 Face Mask

香港地址: 香港新界荃灣角街 38-40 號 銓通工業大廈 23 樓C室

註: 試驗報告僅就委託者之委託事項提供試驗結果,不對產品合法性做判斷。



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日期 Date: 2020.08.31

收件日期 Date of Receipt:

2020.08.24

式 W TEST REPORT

土城場區 TUCHENG

報告編號 TFF9H599 數 Report No.: Qu

数重 _Quantity: 1件 報告

報告頁次/頁數 (P2/5) Page Order/Pages: 來文字號 空 白 Ref. No.:

試件類別 □罩 Item:_____

報告抬頭 Report Title:

407 台中市西屯區甘河路108號

居安有限公司(U3104)

地址 Address:_

試驗項目		試驗結果	試 驗 方 法
	1	無穿透	EN 14683:2019
壓力:120 mmHg (16.0 kPa)	2	無穿透	ISO 22609:2004
	3	無穿透	
	4	無穿透	
	5	無穿透	
	6	無穿透	
	7	有穿透	
N.	8	無穿透	
	9	無穿透	6
87	10	無穿透	0
	11	無穿透	38
	12	無穿透	6
हिं	13	無穿透	29
68	14	無穿透	
	15	無穿透	
	16	無穿透	

註:依委託者所提供來樣資料為: Brandname: Curie

Model: Curie KV99 Face Mask

香港地址: 香港新界荃灣角街 38-40 號 銓通工業大廈 23 樓C室

註: 試驗報告僅就委託者之委託事項提供試驗結果,不對產品合法性做判斷。

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Authorized by president of Taiwan Textile Research Institute

紡織產業綜合研究所檢測及驗證部 新北市土城區承天路6號 Department of Testing and Certification, Taiwan Textile Research Institute No.6, Chengtian Rd., Tucheng Dist., New Taipei City 23674, Taiwan (R.O.C.)





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日期 2020.08.31 Date:

收件日期 Date of Receipt:

2020.08.24

TEST REPORT

土城場區 TUCHENG

報告編號 TFF9H599 Report No .:

Quantity:

居安有限公司(U3104)

1件

報告頁次/頁數 (P3/5) Page Order/Pages:

來文字號 空

Ref. No .:

試件類別 □罩

Item:_

Report Title: 地址

報告抬頭

407 台中市西屯區甘河路108號

Address: -

試驗項目		試験結果	試 驗 方 法
合成血液穿透性	17	無穿透	EN 14683:2019
壓力:120 mmHg (16.0 kPa)	18	無穿透	ISO 22609:2004
	19	無穿透	-
	20	無穿透	-
	21	無穿透	
	22	無穿透	
	23	無穿透	
N.	24	無穿透	
	25	無穿透	
8,	26	無穿透	0
	27	無穿透	38
	28	無穿透	7
	29	無穿透	29
	30	無穿透	= 6
	31	無穿透	
	32	無穿透	3

註:依委託者所提供來樣資料為: Brandname: Curie

Model: Curie KV99 Face Mask

香港地址: 香港新界荃灣角街 38-40 號 銓通工業大廈 23 樓C室

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Taiwan Textile Research Institute





日期 2020.08.31 Date:

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TEST REPORT

土城場區 TUCHENG

報告編號 TFF9H599 Report No .: Quantity:

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報告頁次/頁數 (P4/5) Page Order/Pages:

來文字號 空 Ref. No .:

試件類別 □罩 Item:_

報告抬頭 Report Title: -地址

407 台中市西屯區甘河路108號

居安有限公司(U3104)

Address: -

試驗項目		試驗結果	試 驗 方 法
細菌過濾效率(%)	1	> 99.9	EN 14683:2019 Annex B
金黃色葡萄球菌	2	99.9	
ATCC 6538	3	99.9	
	4	> 99.9	
	5	99.9	

註:對照組的平均菌落數:2640 CFU。

註: 平均粒徑: 2.8 μm。

註: 測試面:外側。

註: 測試面積:39.5 cm

註: 測試流量: 28.3 L/min。

註:依委託者所提供來樣資料為: Brandname: Curie

Model: Curie KV99 Face Mask

香港地址: 香港新界荃灣角街 38-40 號 銓通工業大廈 23 樓C室

註:試驗報告僅就委託者之委託事項提供試驗結果,不對產品合法性做判斷。



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日期 Date: 2020.08.07

收件日期 Date of Receipt:

2020.07.27

試驗報告 TEST REPORT 土城場區 TUCHENG

報告編號 TFF9G707 數量 Report No.: Quantity:__

1件 報告頁次/頁數 (P5/6) Page Order/Pages: 來文字號 空

Ref. No.: ____ 試件類別 □罩

報告抬頭 居安有限公司(U3104)

Item:_

Report Title: 40%

407 台中市西屯區甘河路108號

Address:

試驗項目		試 驗 結 果	試驗方法
微生物潔淨性	1	13.7	EN 14683:2019
(cfu/g)	2	10.6	EN ISO 11737-1:2018
	3	4.4	
	4	10.1	
	5	11.0	

註: 依委託者所提供來樣資料為: Brandname: Curie

Model: Curie KV99 Face Mask

香港地址: 香港新界荃灣角街 38-40 號 銓通工業大廈 23 樓C室

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Tel: +886-2-22670321 ext. 7107, 7110 Fax: +886-2-22675108, +886-2-22689839

謝及繼報 言瑞宏



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日期 2020.08.31 Date:

收件日期 Date of Receipt:

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試驗報告 TEST REPORT

土城場區 TUCHENG

報告編號 TFF9H599 Report No.:

1件

報告頁次/頁數 (P5/5) Page Order/Pages:

來文字號 空

Fr96.399 Quantity:___ 居安有限公司(U3104)

Ref. No .: 試件類別 □罩

報告抬頭 Report Title:

Item:_

地址

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Address: -



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Tel: +886-2-22670321 ext. 7107, 7110 Fax: +886-2-22675108, +886-2-22689839

检测及验证率 古 7山 任回





TEST REPORT TUCHENG

Date: Aug.07,2020 Date of Receipt: Jul.27,2020

Report No.: TAG9G708 Quantity: 1PC Page Order/Pages: P1/4 Ref. No.: NIL

Report Title: Curie Limited(U3104)

Item: Mask

Address: No. 108, Ganhe Rd., Xitun Dist., Taichung City 407, Taiwan

Test Items		Test Results	Test Methods
Sub-Micron Particulate	1	99.25	ASTM F2100-2019 9.3
Filtration Efficiency(%)	2	99.23	ASTM F2299-2017
$(0.1 \mu \text{m PSL})$	3	99.22	Flow rate:28.1
	4	99.16	(Liter/min)
	5	99.11	
	Ave.	99.19//	
Air Exchange Pressure		5.1	ASTM F2100-2019 9.2
(mmH2O/cm²)	2	6.1	EN 14683:2019 Annex C
0	3	5.6	
	7 4	5.1	En
	5	4.7	

Note: 1mmH2O=9.8Pa.

Note: Air Exchange Pressure takes 5 masks for testing.

Note: Sample description is given by the client: Brandname: Curie

Model: Curie KV99 3D Mask

Hong Kong address: Unit C, 23F, Tsun Tung Factory Bldg 38-40

Chai Wan Kok St, Tsuen Wan. NT Hong Kong

Note: The test report is the test results issued by the testing institution as requested by the consignor, it shall not determine the legitimacy of the product.

Note: 1. This report is only responsible for the submitted sample(s), which will be kept for one month period.

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Taiwan Textile Research Institute

Jui-hung Kao

Director,
Department of Testing and
Certification

Department of Testing and Certification, Taiwan Textile Research Institute No.6, Chengtian Rd., Tucheng Dist., New Taipei City 23674, Taiwan (R.O.C.)





TEST REPORT TUCHENG

Date: Aug. 07, 2020	Date of Receipt:	Jul.27,2020

Report No.: TAG9G708 Quantity: 1PC Page Order/Pages. Page No.: NIL

Report Title: Curie Limited(U3104)

Item: Mask

Address: No. 108, Ganhe Rd., Xitun Dist., Taichung City 407, Taiwan

Test Items		Test Results	Test Methods
Synthetic Blood Penetration	1	None Seen	ASTM F2100-2019 9.4
Pressure:160 mmHg	2	None Seen	ASTM F1862-2017
	3	None Seen	
	4	None Seen	
	5	None Seen	
	6	None Seen	
	7	None Seen	
A.	8	None Seen	
	9	None Seen	
	10	None Seen	7
	11	None Seen	
	12	None Seen	
8	13	None Seen	F
	14	None Seen	E°
	15	None Seen	
	16	None Seen	

Note: Sample description is given by the client: Brandname: Curie

Model: Curie KV99 3D Mask

Hong Kong address: Unit C, 23F, Tsun Tung Factory Bldg 38-40

Chai Wan Kok St, Tsuen Wan. NT Hong Kong

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Jui - hung kao

Director,
Department of Testing and
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Department of Testing and Certification, Taiwan Textile Research Institute No.6, Chengtian Rd., Tucheng Dist., New Taipei City 23674, Taiwan (R.O.C.)





TEST REPORT TUCHENG

Date: Aug. 07, 202	O Date of Receipt:	Jul.27,2020

Report No.: TAG9G708 Quantity: 1PC Page Order/Pages: P3/4) Ref. No.: NIL

Report Title: Curie Limited(U3104)

Mask

Address: No. 108, Ganhe Rd., Xitun Dist., Taichung City 407, Taiwan

Test Items		Test Results	Test Methods
Synthetic Blood Penetration	17	None Seen	ASTM F2100-2019 9.4
Pressure:160 mmHg	18	None Seen	ASTM F1862-2017
	19	None Seen	
	20	None Seen	
	21	None Seen	
	22	None Seen	
	23	None Seen	
R	24	None Seen	
	25	None Seen	
	26	None Seen	a n
	27	None Seen	
	28	None Seen	
8	29	None Seen	
	30	None Seen	e e
0	31	None Seen	
	32	None Seen	-

Note: Sample description is given by the client: Brandname: Curie

Model: Curie KV99 3D Mask

Hong Kong address: Unit C, 23F, Tsun Tung Factory Bldg 38-40

Chai Wan Kok St, Tsuen Wan. NT Hong Kong

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Jw - hung kao

Department of Testing and Certification

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TEST REPORT TUCHENG

Date: Aug. 07, 2020 Date of Receipt: Jul. 27, 2020

Report No.: TAG9G708 Quantity: 1PC Page Order/Pages: P4/4) Ref. No.: NIL

Report Title: Curie Limited(U3104)

Item: Mask

Address: No. 108, Ganhe Rd., Xitun Dist., Taichung City 407, Taiwan



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日期 Date: 2020.08.07 收件日期 Date of Receipt:

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式 W TEST REPORT

土城場區 TUCHENG

報告編號 Report No.:_ 報告抬頭

TAG9G708 数重 Quantity: 1件 報告頁次/頁數 (P1/4) Page Order/Pages: 來文字號 空 白 Ref. No.: _____

居安有限公司(U3104)

試件類別 口罩

Item:_

Report Title: 40°

407 台中市西屯區甘河路108號

Address:

試驗項目		試驗結果	試 驗 方 法
微粒過濾效率	1	99.25	ASTM F2100-2019 9.3
$(0.1 \mu\mathrm{m} \mathrm{PSL})$	2	99.23	ASTM F2299-2017
	3	99.22	速度:28.1 (Liter/min)
	4	99.16	
	5	99.11	
	平均	99.19	
空氣交換壓力差	7 J	5.1	ASTM F2100-2019 9.2
(mmH2O/cm²)	2	6.1	EN 14683:2019 Annex C
	3	5.6	
2 A	4	5.1	0
	5	4.7	18

註: 1mmH2O=9.8Pa。

註:空氣交換壓力差,取5個樣品測試。

註:依委託者所提供來樣資料為: Brandname: Curie

Model: Curie KV99 3D Mask

香港地址: 香港新界荃灣角街 38-40 號 銓通工業大廈 23 樓C室

註:試驗報告僅就委託者之委託事項提供試驗結果,不對產品合法性做判斷。



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Authorized by president of Taiwan Textile Research Institute

紡織產業綜合研究所檢測及驗證部 新北市土城區承天路6號 Department of Testing and Certification, Taiwan Textile Research Institute No.6, Chengtian Rd., Tucheng Dist., New Taipei City 23674, Taiwan (R.O.C.)



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日期 Date: 2020.08.07

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土城場區 TUCHENG

報告編號 Report No.: TAG9G708

数重 __Quantity: 1件 報告頁次/頁數 (P2/4) Page Order/Pages: 來文字號 空

居安有限公司(U3104)

Ref. No.: ____ 試件類別 □罩

Item:

報告抬頭 Report Title: 地址 40°

407 台中市西屯區甘河路108號

Address:

試験項目		試 驗 結 果	試 驗 方 法
合成血液穿透性	1	無穿透	ASTM F2100-2019 9.4
壓力:160 mmHg	2	無穿透	ASTM F1862-2017
	3	無穿透	
	4	無穿透	
	5	無穿透	
	6	無穿透	
	7	無穿透	
	8	無穿透	
	9	無穿透	8
	10	無穿透	00
1	11	無穿透	18
7	12	無穿透	6
	13	無穿透	29
6	14	無穿透	
	15	無穿透	
	16	無穿透	

註:依委託者所提供來樣資料為: Brandname: Curie

Model: Curie KV99 3D Mask

香港地址: 香港新界荃灣角街 38-40 號 銓通工業大廈 23 樓C室

註: 試驗報告僅就委託者之委託事項提供試驗結果,不對產品合法性做判斷。

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日期 2020.08.07 Date:

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TEST REPORT

土城場區 TUCHENG

報告編號 **TAG9G708** Report No .:

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居安有限公司(U3104)

報告頁次/頁數 (P3/4) Page Order/Pages: 1件

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Item:_

報告抬頭 Report Title: 地址

407 台中市西屯區甘河路108號

Address: -

試験項目		試驗結果	試 驗 方 法
 合成血液穿透性	17	無穿透	ASTM F2100-2019 9.4
壓力:160 mmHg	18	無穿透	ASTM F1862-2017
	19	無穿透	
	20	無穿透	
	21	無穿透	
	22	無穿透	
	23	無穿透	
	24	無穿透	
	25	無穿透	8
	26	無穿透	00
1	27	無穿透	18
The state of the s	28	無穿透	
	29	無穿透	89
6	30	無穿透	
	31	無穿透	
	32	無穿透	

註:依委託者所提供來樣資料為: Brandname: Curie

Model: Curie KV99 3D Mask

香港地址: 香港新界荃灣角街 38-40 號 銓通工業大廈 23 樓C室

註: 試驗報告僅就委託者之委託事項提供試驗結果,不對產品合法性做判斷

附記: 1.本報告僅對樣品負責,樣品保留期限為一個月。

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財團法人紡織產業綜合研究所 所長授權核發人:

Authorized by president of Taiwan Textile Research Institute

紡織產業綜合研究所檢測及驗證部 新北市土城區承天路6號 Department of Testing and Certification, Taiwan Textile Research Institute No.6, Chengtian Rd., Tucheng Dist., New Taipei City 23674, Taiwan (R.O.C.)







日期 2020.08.07 Date:

收件日期 Date of Receipt:

2020.07.27

試驗報告 TEST REPORT

土城場區 TUCHENG

報告編號 **TAG9G708** Report No.:

Quantity: 居安有限公司(U3104) 1件

報告頁次/頁數 (P4/4) Page Order/Pages: ____

來文字號 空

Ref. No .: 試件類別 □罩

Item:_

Report Title: 地址

407 台中市西屯區甘河路108號

Address: -

報告抬頭



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財團法人紡織產業綜合研究所 所長授權核發人:

Authorized by president of Taiwan Textile Research Institute

紡織產業綜合研究所檢測及驗證部 新北市土城區承天路6號 Department of Testing and Certification, Taiwan Textile Research Institute No.6, Chengtian Rd., Tucheng Dist., New Taipei City 23674, Taiwan (R.O.C.)

Tel: +886-2-22670321 ext. 7107, 7110 Fax: +886-2-22675108, +886-2-22689839

检测及验證部 古 工山 任回



政府創辦·多元創新

Government established - Diversified and innovative

TEST REPORT

Applicant: Curie Limited

Report number: IRITS202005150001

Room C, 23/F,

Date: 15 May 2020

Tsuen Tung Factory Building, 38-40 Chai Wan Kok Street,

Tsuen Wan, New Territories, Hong Kong

Attn.:

Aldrin Or

Sample Description as Declared:

No. of Sample: TWO (2) pieces of received material in zipper bag packaging

Sample Description: Curie Ultrahigh- Efficiency Viral Filter

Colour: White
Date Received: 8 May 2020

Testing Period: 9 – 14 May 2020

Tests Conducted: As requested by the Applicant, with the details as follow:

Testing Summary: The sample being tested was conditioned for a minimum of 4 hour at $21 \pm 5\,^{\circ}$ C and relative humidity of $65 \pm 5\,^{\circ}$ C. The bacterial filtration efficiency (BFE) test was performed by applying a spray of challenge bacterium *Staphylococcus aureus* in peptone water (approximately 2,200 colony forming units per spray) using a trigger sprayer. The sprayed aerosol was then drawn through the material being tested following by a tryptic soy agar plate under vacuum (flow rate: 100 Litres per minute). Number of *Staphylococcus aureus* colonies formed on the tryptic soy agar plate were counted after incubated at $37 \pm 2\,^{\circ}$ C for 48 \pm 4 hr. The BFE test procedure was modified from ASTM F2101: 2019.

For and on behalf of

Institute for Research in Innovative Technology & Sustainability

The Open University of Hong Kong

Dr. Eric Tung-po Sze

Director



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Report number: IRITS202005150001

Date: 15 May 2020

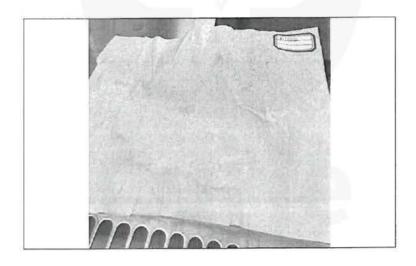
Results:

Test Sample Number

Test Sample Number	Bacterium Colonies Formed
#1	N.D.a
#2	N.D.a
Negative Control	N.D.a

^a None Detected (N.D.) – There were no detected bacterium colony of *Staphylococcus aureus* found.

Sample Photo:



<End of Test Report>



TEST REPORT

Report number: IRITS2020007030001

Date: 3 July 2020

Applicant: Curie Limited

Room C, 23/F,

Tsuen Tung Factory Building, 38-40 Chai Wan Kok Street,

Tsuen Wan, New Territories, Hong Kong

Attn.: Aldrin Or

Sample Description as Declared:

No. of Sample: TWO (2) pieces of composite material for face mask in zipper bag

packaging Curie KV99

Colour: White

Date Received: 15 June 2020 Testing Period: 16 – 24 June 2020

Tests Conducted: As requested by the Applicant, with the details as follow:

Testing Summary: The sample(s) were conditioned at an acceleration temperature of 120 $^{\circ}$ C for 48 hours, followed by pre-conditioning at a minimum of 4 hour at 21 \pm 5 $^{\circ}$ C and relative humidity of 65 \pm 5 $^{\circ}$ C. Bacterial filtration efficiency (BFE) test was then performed by spraying the samples with an aerosol of challenge bacterium *Staphylococcus aureus* in peptone water using a nebulizer. The aerosol was then drawn through the samples following by a tryptic soy agar plate under vacuum (flow rate: 100 Litres per minute). Number of *Staphylococcus aureus* colonies formed on the tryptic soy agar plate were counted after incubated at 37 \pm 2 $^{\circ}$ C for 48 \pm 4 hr. The BFE test procedure was modified from ASTM F2101: 2019.

For and on behalf of Institute for Research in Innovative Technology & Sustainability The Open University of Hong Kong

Dr. Eric Tung-po Sze

Director

科技學院 School of Science and Technology

Report number: IRITS2020007030001

Date: 3 July 2020

Results:

Test Sample Number	Bacterium Colonies Formed	Bacterial Filtration Efficiency
#1	N.D. ^a	> 99 %
#2	N.D. ^a	> 99 %
Negative Control	N.D. ^a	N/A ^b

^a None Detected (N.D.) – There were no detected bacterium colony of *Staphylococcus aureus* found

Remark: The time and temperature selected for the acceleration conditioning were based on ASTM Standard F1980-16 Appendix X1. Accelerated aging of polymers, which are equivalent to five year of room-temperature (20 $^{\circ}$ C) aging, with an aging factor Q₁₀ = 2.0.

Sample Photos:



<End of Test Report>

^b N/A – Not Applicable

TEST REPORT

Applicant: Curie Limited

Report number: IRITS2020007130001R1

Date: 23 July 2020

Room C, 23/F,

Tsuen Tung Factory Building, 38-40 Chai Wan Kok Street,

Tsuen Wan, New Territories, Hong Kong

Attn.: Aldrin Or

Sample Description as Declared:

No. of Sample: ONE (1) piece of textile material in zipper bag packaging said to be RT-

2007-T0430-DC020

Colour: White

Date Received: 21 May 2020 Testing Period: 2 – 10 July 2020

Tests Conducted: As requested by the Applicant to determine the antibacterial activity of

the sample with reference to BS EN ISO 20743: 2013 Clause 8.2 Transfer

method, with the following deviation:

 Shake-out the bacteria from specimens using peptone water instead of neutralizing solution.

For and on behalf of Institute for Research in Innovative Technology & Sustainability The Open University of Hong Kong

Dr. Eric Tung-po Sze

Director



Report number: IRITS2020007130001R1

Date: 23 July 2020

Results:

Specimen Conditions		Number of bacteria ^a (CFU per specimen)	
#1	Shake-out before incubation	0	
#2	Shake-out after incubation	0	

 $^{a}1$ millilitre of an inoculum of *Staphylococcus aureus* with concentration of 1×10^{6} CFU/ml to 3×10^{6} CFU/ml was applied onto an agar plate in the transfer method, where each specimen was set on the agar surface and weigh down with a 200 g stainless-steel cylinder for 60 s \pm 5 s to transfer the microbial content. Incubation Measurement of the number of bacteria colonies was conducted in accordance with the plate count method specified in Annex C of BS EN ISO 20743:2013.

Opinion(s) and Interpretation(s): Based on the results obtained above, the specimens demonstrated effective antibacterial property to kill bacteria during transfer phase of the experiment.

Note: This Report replaces Report number IRITS2020007130001, which has been obsoleted.

<End of Test Report>





GUANGDONG DETECTION CENTER OF MICROBIOLOGY

REPORT FOR ANALYSIS

Report №.	2020FM20686R01E
Name of Sample	Curie Ultrahigh-Efficiency Viral Filter for KV-99
Applicant	Shenzhen Qianhai e-Cycle Trading Co.,Ltd.
Test Type	Entrustment Test

Address: Building 66, No.100 Central Xian Lie Road, Guangzhou, China

Postcode: 510070

Tel: +86 20 87137666

Fax: +86 20 87137668

Website: www.gddcm.com





GUANGDONG DETECTION MICROBIOLOGY 風影器 CENTER OF

REPORT FOR ANALYSIS

Report №::2020FM20686R01E Verification Code: 03658924



Name of Sample	Curie Ultrahigh-Efficiency Viral Filter for KV-99	Test Type	Entrustment Test
Applicant	Shenzhen Qianhai e-Cycle Trading Co.,Ltd.	Address	2/F.Building B2, Yintian Industrial Area, Xixiang Street, Baoan District, Shenzhen Guangdong, China
Sample Source	Submitted for Testing by the Applicant	Sample Quantity	260cm*2m
Spec and Lot № of Sample	40g;1001	State and Characteristic	Flaky
Sample Received Date	2020-07-15	Test Completion Date	2020-07-28
Test Standard and Method	IS	O 18184: 2014 (E	Destroy Learner to Learner to Learner to Learner
the things the	Antiviral activity test		
Item Tested		Antiviral activity tes	
Item Tested Test Conclusion	The test data of the sample(s) is attac	ched to the page(s) o	Carter Ca

Editor: Chen Jingting

Verifier: Là Sujuan

Approver: Xe Xinhar





GUANGDONG DETECTION CENTER OF MICROBIOLOGY

ANALYSIS AND TEST RESULT

Report №.: 2020FM20686R01E

Virus and host cell	No.	The logarithm of infectivity titre value immediate after inoculation of the reference specimen (lgTCID ₅₀ /bottle)	The logarithm of infectivity titre value after 2h contacting with the reference specimen (lgTCID ₅₀ / bottle)	The logarithm of infectivity titre value after 2h contacting with the test specimen (lgTCID ₅₀ / bottle)
Grides Grides	erice 1	7.05	6.50	2.10
H3N2 Influenza A virus	2	6.97	6.63	2.30
Host cell: MDCK	3	7.10	6.59	2.30
lgTCID ₅₀ / bott Average	le	7.04	6.57	2.33
Logarithm of antiviral activity			4.34	
Antiviral activity rate	e (%)	Control of the Contro	Hato Telling Comment of the Comment	

(Blank below)





Report №.: 2020FM20686R01E

Notice Items

- 1. The Test report is invalid if not affixed with Authorized Stamp of Test and Paging Seal.
- 2. The Test report is invalid without signature of verifier and approver.
- 3. The Test report is invalid if being supplemented, deleted or altered.
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- 6. Any dispute of the report must be raised to the testing body within 15 days after the report is received, exceeding which the dispute will not be accepted.
- 7. For the tested sample(s) submitted by the applicant, the sample information in the test report is provided by the applicant and the laboratory is not responsible for its authenticity.



TEST REPORT

Technical Report: (5220)210-0555 August 5, 2020

Date Received: July 28, 2020 Page 1 of 8

Ken / Janus / Alice / Jacky Tsang Dakota Flat A&B 3/F, Fast Industrial Bldg. 658 Castle Peak Road, Lai Chi Kok, Kowloon, Hong Kong

Product Name: INTERLINING

Product End Use: Blouses

Color Name / Article Number:

Color Code: OFF-WHITE

H&M Order No.:

H&M Sample Type:Raw MaterialsAge Group:Child 0-3 year-oldH&M Sample Stage:D/S (Develop sample)

H&M Product Category: Knit

Date of Testing: July 28, 2020 – August 5, 2020

Number of Working Days: 7

Priority: REGULAR

Speed Order: No
No. of sample provided: /
Fiber Composition: /
Care Instruction: /

SUMMARY OF TEST RESULTS

TEST REQUESTED	CONCLUSION	REMARK
Alkylphenols (AP) Content Test	PASS	
Alkylphenol Ethoxylates (APEO) Content Test	PASS	
Formaldehyde Content Test	PASS	
Chlorophenols Content Test	PASS	
Ortho-phenylphenol (OPP) Content Test	PASS	
pH Value Test in Textile and Artificial Leather	FAIL	
Phthalates Content Test	PASS	

REMARK

If there are questions or concerns on this report, please contact:

(852) 2331 0330

hmEnquiry@hk.bureauveritas.com

BUREAU VERITAS HONG KONG LTD.

MS. YANN LO

MANAGER, ANALYTICAL

Bureau Veritas Hong Kong Ltd.
Consumer Product Services Division
Kowloon Bay Office
1/F, Pacific Trade Centre,
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Kowloon, Hong Kong
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Technical Report: **(5220)210-0555**August 5, 2020
Page 2 of 8

Photo of the Submitted Sample





August 5, 2020 Page 3 of 8

TEST RESULT

Alkylphenols (AP) Content Test

Test Method I: For Textile & Leather:

EN ISO 21084

Test Method II: For Polymers and other materials:

Organic solvent extraction, analysis with reference to EN ISO 21084.

Tested Item(s): I001 White fabric with transparent adhesive

Maximum Limit: 100 mg/kg (sum)

Tooted Homes	Test	Result	Ä.,.		Canalasian
Tested Item(s)	Method	Detected Analyte(s)	Conc.	Unit	Conclusion
I001	I	ND	ND	mg/kg	PASS

Note:

ND = Not detected ">" = More than Conc. = Concentration

ppm = part(s) per million = mg/kg mg/kg = milligram(s) per kilogram

Reporting Limit (mg/kg): Sum (OP & NP): 10

Remark:

- The list of alkylphenols is summarized in table of Appendix.

Alkylphenol Ethoxylates (APEO) Content Test

Test Method I : For Textile and Other materials:

With reference to EN ISO 18254-1, analysis by Liquid Chromatograph Mass

Spectrometer (LC-MS)

Test Method II : For Leather:

With reference to EN ISO 18218-1 and EN ISO 18254-1

Tested Item(s): I001 White fabric with transparent adhesive

Maximum Limit: Others: 100 mg/kg (Sum)
Recycled materials: 1000 mg/kg (Sum)

Tostad Itam(s)	Test	Result			Canalusian
Tested Item(s)	Method	Detected Analyte(s)	Conc.	Unit	Conclusion
I001	I	ND	ND	mg/kg	PASS

Note:

ND = Not detected ">" = More than Conc. = Concentration

ppm = part(s) per million = mg/kg mg/kg = milligram(s) per kilogram

Reporting Limit (mg/kg): Sum (OPEOs & NPEOs): 20

Remark:

- The list of alkylphenol ethoxylates is summarized in table of Appendix.



August 5, 2020 Page 4 of 8

TEST RESULT

Formaldehyde Content Test

Test Method I : Textiles & Other Materials: EN ISO 14184-1

Test Method II : Leather: ISO 17226:2 and/or ISO 17226-1

Test Method III : Glue & 0-3 years old products: JIS L 1041: 2011 Method A

Test Method IV : Carpets and mats: GB 18587 Grade B.

Tested Item(s): 1001 White fabric with transparent adhesive

Maximum Limit:	Type I	For baby < 36 months: 16 mg/kg						
	Type II	Others: 75 mg/kg	Others: 75 mg/kg					
	Type III	Carpets and mats : < 0.050 mg/m ² /h						
Tested Item(s)	Туре	Test Method	Result	Unit	Conclusion			
1001	I	III	ND	mg/kg	PASS			

Note:

ND = Not detected ">" = More than



August 5, 2020 Page 5 of 8

TEST RESULT

Chlorophenols Content Test

Test Method: With reference to ISO 17070:2015 or 64 LFBG B 82.02-08 (Modified). Potassium

hydroxide extraction, derivatisation and analysis by Gas Chromatograph Mass

Spectrometer (GC-MS).

Tested Item(s) : I001 White fabric with transparent adhesive

Maximum Limit:	0.5 mg/kg (Each)				
Tested Item(s)	Resu	C			
	Detected Analyte(s)	Conc.	Unit	Conclusion	
I001	ND	ND	mg/kg	PASS	

Note:

ND = Not detected ">" = More than

ppm = part(s) per million = mg/kg mg/kg = milligram(s) per kilogram

Reporting Limit (mg/kg): Each: 0.5

Remark:

The list of chlorophenols is summarized in table of Appendix.

Ortho-phenylphenol (OPP) Test

Test Method: 1 M KOH extraction, 16 hours at 90 degrees C, derivatization and analysis

§ 64 LFGB B 82.02-08 or DIN EN ISO 17070:2015.

Tested Item(s) : I001 White fabric with transparent adhesive

Maximum Limit:	1000 mg/kg		
Tested Item(s)	Result	Unit	Conclusion
I001	ND	mg/kg	PASS

Note:

ND = Not detected ">" = More than

ppm = part(s) per million = mg/kg mg/kg = milligram(s) per kilogram

Reporting Limit (mg/kg): 100



August 5, 2020 Page 6 of 8

TEST RESULT

pH Value Test in Textile and Artificial Leather

Test Method: ISO 3071, extraction with potassium chloride solution.

Tested Item(s): 1001 White fabric with transparent adhesive

Maximum Allowable Limit		pH 4.0- pH 7.5				
Parameter	Unit	Result				
Test Item(s)		I001		-		
pH (Trial 1)	9 89	8.3		-		
pH (Trial 2)		8.3) = 0	-		
Mean of Trial 1 & 2	(4):	8.3	1=1	_		
Conclusion		FAIL	· · · · · · · · · · · · · · · · · · ·	2		

Phthalates Content Test

Test Method I : Textile: CPSC-CH-C1001-09.4 and EN ISO 14389: 2014.

Test Method II : Others: CPSC-CH-C1001-09.4, analysis by Gas Chromatograph Mass Spectrometer

(GC-MS).

Tested Item(s): I001 White fabric with transparent adhesive

Maximum Limit: 500 mg/kg (Each) 1000 mg/kg (Sum)

Tested Item(s)	Test	Resi	ult		Conclusion
resteu rtem(s)	method	Detected Analyte(s)	Conc.	Unit	Conclusion
1001	I	ND	ND	mg/kg	PASS

Note:

ND = Not detected ">" = More than Conc. = Concentration

ppm = part(s) per million = mg/kg mg/kg = milligram(s) per kilogram

Reporting Limit (mg/kg): Each: 50

Remark:

- The list of phthalates is summarized in table of Appendix.

END



Technical Report: **(5220)210-0555**August 5, 2020
Page 7 of 8

APPENDIX

No.	Name of Analytes	CAS-No.	No.	Name of Analytes	CAS-No.
î	Octylphenol (OP)	Various (140-66-9, 27193-28-8, 1806-26-4)	2	Nonylphenol (NP)	Various (25154-52-3 104-40-5, 84852-15-3, 11066-49-2'

No.	Name of Analytes	CAS-No.	No.	Name of Analytes	CAS-No.
1	Octylphenolethoxylates (OPEOs)	Various (9002-93-1, 9036-19-5, 68987-90-6)	2	Nonylphenolethoxylates (NPEOs)	Various (9016-45-9, 26027-38-3, 127087-87-0 37205-87-1, 68412-54-4)

No.	Name of Analytes	CAS-No.	No.	Name of Analytes	CAS-No.
1	Pentachlorophenol (PCP) and its salts and esters	87-86-5	3	Trichlorophenol (TriCP) and its salts and esters: 2,3,5- Trichlorophenol; 2,3,6- Trichlorophenol & 2,4,5- Trichlorophenol; 2,4,6- Trichlorophenol; 3,4,5- Trichlorophenol & 2,3,4- Trichlorophenol	933-78-8; 933-75-5 & 95-95-4; 88-06-2; 609-19-8 & 15950-66-0
2	Tetrachlorophenol (TeCP) and its salts and esters: 2,3,5,6- Tetrachlorophenol 2,3,4,6- Tetrachlorophenol 2,3,4,5- Tetrachlorophenol	935-95-5 58-90-2 4901-51-3	-	-	



Technical Report: **(5220)210-0555**August 5, 2020
Page 8 of 8

APPENDIX

No.	Name of Analytes	CAS-No.	No.	Name of Analytes	CAS-No.
1	Butyl benzyl phthalate (BBP)	85-68-7	12	Di-iso-pentyl phthalate (DIPP)	605-50-5
2	Dibutyl phthalate (DBP)	84-74-2	13	n-Pentyl-iso-pentyl phthalate (NPIPP)	776297-69-9
3	Di-2-ethylhexyl phthalate (DEHP)	117-81-7	14	Di-cyclohexyl phthalate (DCHP)	84-61-7
4	Di-iso-butyl phthalate (DIBP)	84-69-5	15	Di-iso-octyl phthalate (DIOP)	27554-26-3
5	Di-n-octyl phthalate (DNOP)	117-84-0	16	Di-n-pentyl phthalate (DPP)	131-18-0
6	Di-iso-nonyl phthalate (DINP)	28553-12-0	17	Dihexyl phthalate, branched and linear	68515-50-4
7	Di-iso-decyl phthalate (DIDP)	26761-40-0	18	Dimethyl phthalate (DMP)	131-11-3
8	Bis(2-methoxyethyl) phthalate (DMEP)	117-82-8	19	1,2-benzenedicarboxylic acid, dipentylester, branched and linear	84777-06-0
9	Di-n-hexyl phthalate (DHP)	84-75-3	20	1,2-benzenedicarboxylic acid, di- C7-11-branched and linear, alkyl esters (DHUNP)	68515-42-4
10	Diethyl phthalate (DEP)	84-66-2	21	1,2-benzenedicarboxylic acid, di- C6-8-branched alkyl esters, C7- rich (DIHP)	71888-89-6
11	Di-n-propyl phthalate (DPRP)	131-16-8	22	1,2-Benzenedicarboxylic acid, di- C6-10-alkyl esters or mixed decyl and hexyl and octyl diesters with ≥ 0.3% of dihexyl phthalate; 1,2- Benzenedicarboxylic acid, mixed decyl and hexyl and octyl diesters; 1,2-Benzenedicarboxylic acid, di-C6-10-alkyl esters	68515-51-5 68648-93-1



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62/988,900

166693

03/12/2020

WIPC2002

CONFIRMATION NO. 9401 UPDATED FILING RECEIPT

Date Mailed: 07/09/2020

Law Offices of Sergei Orel, LLC 2125 Center Avenue, Suite 616 Fort Lee, NJ 07024

Receipt is acknowledged of this provisional patent application. It will not be examined for patentability and will become abandoned not later than twelve months after its filing date. Any correspondence concerning the application must include the following identification information: the U.S. APPLICATION NUMBER, FILING DATE, NAME OF FIRST INVENTOR, and TITLE OF INVENTION. Fees transmitted by check or draft are subject to collection.

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Inventor(s)

Jianliang GONG, Yau Ma Tei, HONG KONG; Chun Yin OR, Cheung Sha Wan, HONG KONG;

Applicant(s)

Jianliang GONG, Yau Ma Tei, HONG KONG; Chun Yin OR, Cheung Sha Wan, HONG KONG;

Power of Attorney:

Michael Yablonsky--40407

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Permission to Access Search Results: Yes

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The country code and number of your priority application, to be used for filing abroad under the Paris Convention,

is US 62/988,900

Projected Publication Date: None, application is not eligible for pre-grant publication

Non-Publication Request: No Early Publication Request: No

page 1 of 3

** MICRO ENTITY **
Title

Air Filtration System and Manufacturing Method Therefor

Statement under 37 CFR 1.55 or 1.78 for AIA (First Inventor to File) Transition Applications: No

PROTECTING YOUR INVENTION OUTSIDE THE UNITED STATES

Since the rights granted by a U.S. patent extend only throughout the territory of the United States and have no effect in a foreign country, an inventor who wishes patent protection in another country must apply for a patent in a specific country or in regional patent offices. Applicants may wish to consider the filing of an international application under the Patent Cooperation Treaty (PCT). An international (PCT) application generally has the same effect as a regular national patent application in each PCT-member country. The PCT process **simplifies** the filing of patent applications on the same invention in member countries, but **does not result** in a grant of "an international patent" and does not eliminate the need of applicants to file additional documents and fees in countries where patent protection is desired.

Almost every country has its own patent law, and a person desiring a patent in a particular country must make an application for patent in that country in accordance with its particular laws. Since the laws of many countries differ in various respects from the patent law of the United States, applicants are advised to seek guidance from specific foreign countries to ensure that patent rights are not lost prematurely.

Applicants also are advised that in the case of inventions made in the United States, the Director of the USPTO must issue a license before applicants can apply for a patent in a foreign country. The filing of a U.S. patent application serves as a request for a foreign filing license. The application's filing receipt contains further information and guidance as to the status of applicant's license for foreign filing.

Applicants may wish to consult the USPTO booklet, "General Information Concerning Patents" (specifically, the section entitled "Treaties and Foreign Patents") for more information on timeframes and deadlines for filing foreign patent applications. The guide is available either by contacting the USPTO Contact Center at 800-786-9199, or it can be viewed on the USPTO website at http://www.uspto.gov/web/offices/pac/doc/general/index.html.

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Title 35, United States Code, Section 184

Title 37, Code of Federal Regulations, 5.11 & 5.15

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08 June, 2020

WORLD IP CONSULTANCY Unit B, 3/F, Cheong Yu Bldg No. 143-151 Castle Peak Rd, Yuen Long HONG KONG

Application for Grant of a Short-term Patent **Under Application No. 32020008506.8**

We refer to your application for a short-term patent lodged on 02 June, 2020.

The above application is found to have satisfied the minimum requirements as laid down in section 114(2) of the Patents Ordinance. The accorded date of filing is 02 June, 2020.

In general, you will receive our further letter for the application at least five months after the date of this letter. If you do not receive any letter from us after this period, please contact us at 2961 6901.

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