測試報告
Test Report

報告編號(No.): HTF21C00636M01　　報告日期(DATE): 2022/01/06　　頁數(PAGE): 1 of 6

上禾伸企業有限公司 (SHANG HE SHEN ENTERPRISE CO., LTD.)
桃園市龜山區興華三街5號1樓(後段) (1F., NO. 5, XINGHUA 3RD ST., GUISHAN DIST., TAOYUAN CITY 33376, TAIWAN (R.O.C.))

以下測試樣品係由申請廠商所提供及確認 (The following sample(s) was/were submitted and identified by/on behalf of the applicant as)：

送樣廠商(Sample Submitted By)	:	上禾伸企業有限公司 (SHANG HE SHEN ENTERPRISE CO., LTD.)
樣品名稱(Sample Description)	:	THERMOPLASTIC VULCANIZATE PLASTIC (TPV塑料)
樣品型號(Style/Item No.)	:	9W560A-A7-FD
樣品材質(Sample Material)	:	TPV塑料 (THERMOPLASTIC VULCANIZATE PLASTIC)
原產國(Country Of Origin)	:	中華民國 (REPUBLIC OF CHINA)
相同材質/顏色樣品型號(Style/Item No. of Sample(s) with Same Material/Color)	:	9W550A-A7-FD，9W555A-A7-FD，9W560A-A7-FD，9W565A-A7-FD，9W570A-A7-FD，9W575A-A7-FD，9W580A-A7-FD

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收件日(Sample Receiving Date)	:	2021/12/23
測試期間(Testing Period)	:	2021/12/23 to 2022/01/04

測試需求 (Test Requested) : 請見下一頁。(Please refer to next page(s).)

測試結果 (Test Results) : 請見第三頁。(Please refer to page 3.)

Singh Hsiao / Asst. Manager
Signed for and on behalf of
SGS TAIWAN LTD.
Chemical Laboratory - Taipei

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測試需求 (Test Requested) :

(1) 客戶指定依據美國聯邦法規之藥物暨食品管理 (FDA) 21 CFR 177.1210 (使用條件A) 進行測試。測試項目請參閱測試結果表格。 (As specified by client, the sample(s) was/were tested according to U.S. Food and Drug Administration (FDA) 21 CFR 177.1210 (Condition of use A). Please refer to the result table(s) for the testing item(s).)

(2) 客戶指定依據美國聯邦法規之藥物暨食品管理 (FDA) 21 CFR 177.1210 (使用條件B) 進行測試。測試項目請參閱測試結果表格。 (As specified by client, the sample(s) was/were tested according to U.S. Food and Drug Administration (FDA) 21 CFR 177.1210 (Condition of use B). Please refer to the result table(s) for the testing item(s).)

(3) 客戶指定依據美國聯邦法規之藥物暨食品管理 (FDA) 21 CFR 177.1210 (使用條件C) 進行測試。測試項目請參閱測試結果表格。 (As specified by client, the sample(s) was/were tested according to U.S. Food and Drug Administration (FDA) 21 CFR 177.1210 (Condition of use C). Please refer to the result table(s) for the testing item(s).)

(4) 客戶指定依據美國聯邦法規之藥物暨食品管理 (FDA) 21 CFR 177.1210 (使用條件D) 進行測試。測試項目請參閱測試結果表格。 (As specified by client, the sample(s) was/were tested according to U.S. Food and Drug Administration (FDA) 21 CFR 177.1210 (Condition of use D). Please refer to the result table(s) for the testing item(s).)

(5) 客戶指定依據美國聯邦法規之藥物暨食品管理 (FDA) 21 CFR 177.1210 (使用條件F) 進行測試。測試項目請參閱測試結果表格。 (As specified by client, the sample(s) was/were tested according to U.S. Food and Drug Administration (FDA) 21 CFR 177.1210 (Condition of use F). Please refer to the result table(s) for the testing item(s).)

(6) 客戶指定依據美國聯邦法規之藥物暨食品管理 (FDA) 21 CFR 177.1210 (使用條件G) 進行測試。測試項目請參閱測試結果表格。 (As specified by client, the sample(s) was/were tested according to U.S. Food and Drug Administration (FDA) 21 CFR 177.1210 (Condition of use G). Please refer to the result table(s) for the testing item(s).)

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測試部位敘述(Test Part Description)

No.1 : 白色片狀 (WHITE SHEET)

測試結果(Test Results)

(1) 通過(PASS)

測試項目 (Test Item(s))	測試方法 (Method)	單位 (Unit)	RL	結果 (Result) No.1	限值 (Limit)
氯仿可萃取物 (水, 250℉, 2小時) / Net chloroform-soluble extractives (D.I. Water, 250℉, 2 h)	參考美國FDA 21 CFR 177.1210. / With reference to US FDA 21 CFR 177.1210.	ppm	5	n.d.	50

(2) 通過(PASS)

測試項目 (Test Item(s))	測試方法 (Method)	單位 (Unit)	RL	結果 (Result) No.1	限值 (Limit)
氯仿可萃取物 (水, 212℉, 30分鐘) / Net chloroform-soluble extractives (D.I. Water, 212℉, 30 min)	參考美國FDA 21 CFR 177.1210. / With reference to US FDA 21 CFR 177.1210.	ppm	5	n.d.	50

(3) 通過(PASS)

測試項目 (Test Item(s))	測試方法 (Method)	單位 (Unit)	RL	結果 (Result) No.1	限值 (Limit)
氯仿可萃取物 (水, 沸騰, 冷卻至100℉) / Net chloroform-soluble extractives (D.I. Water, Fill boiling, cool to 100℉)	參考美國FDA 21 CFR 177.1210. / With reference to US FDA 21 CFR 177.1210.	ppm	5	n.d.	50

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(4) 通過(PASS)

測試項目 (Test Item(s))	測試方法 (Method)	單位 (Unit)	RL	結果 (Result) No.1	限值 (Limit)
氯仿可萃取物 (水, 150℉, 2小時) / Net chloroform-soluble extractives (D.I. Water, 150℉, 2 h)	參考美國FDA 21 CFR 177.1210. / With reference to US FDA 21 CFR 177.1210.	ppm	5	n.d.	50
氯仿可萃取物 (8% 乙醇, 150℉, 2小時) / Net chloroform-soluble extractives (8% Alcohol, 150℉, 2 h)	參考美國FDA 21 CFR 177.1210. / With reference to US FDA 21 CFR 177.1210.	ppm	5	n.d.	50

(5) 通過(PASS)

測試項目 (Test Item(s))	測試方法 (Method)	單位 (Unit)	RL	結果 (Result) No.1	限值 (Limit)
氯仿可萃取物 (水, 70℉, 48小時) / Net chloroform-soluble extractives (D.I. Water, 70℉, 48 h)	參考美國FDA 21 CFR 177.1210. / With reference to US FDA 21 CFR 177.1210.	ppm	5	n.d.	50

(6) 通過(PASS)

測試項目 (Test Item(s))	測試方法 (Method)	單位 (Unit)	RL	結果 (Result) No.1	限值 (Limit)
氯仿可萃取物 (水, 70℉, 24小時) / Net chloroform-soluble extractives (D.I. Water, 70℉, 24 h)	參考美國FDA 21 CFR 177.1210. / With reference to US FDA 21 CFR 177.1210.	ppm	5	n.d.	50

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備註(Note)：

1. mg/kg = ppm；0.1wt% = 1000 ppm
2. RL = Reporting Limit (報告極限值)
3. n.d. = Not Detected (未檢出) = Less than (小於) RL
4. 本報告不得分離或擷錄使用。(The report is invalid if it is partly reproduced or used.)
5. 測試結果的符合性判定不納入量測不確定度。(The statement of conformity is based on the test results, but does not include the measurement uncertainty.)
6. 本報告為 HTF21C00636 之加發報告，報告加發日期 2022年01月06日。 (This is the additional test report of HTF21C00636. The additional test report is issued on 2022/01/06.)

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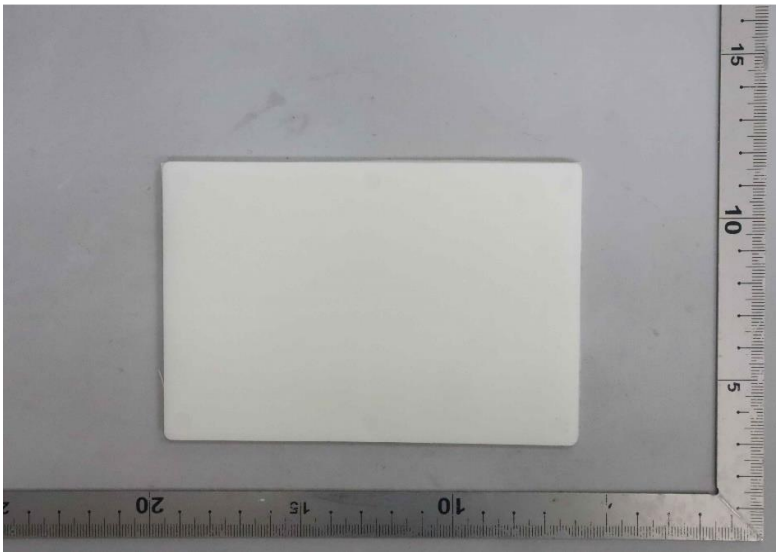
*** 照片中如有箭頭標示，則表示為實際檢測之樣品/部位。 ***
(The tested sample / part is marked by an arrow if it's shown on the photo.)

HTF21C00636



**** 報告結尾 (End of Report) ****