

報告編號(No.): HKF24300048

報告日期(DATE): 2024/03/22

頁數(PAGE): 1 of 4

### **Test Report**

台灣塑膠工業股份有限公司 (FORMOSA PLASTICS CORPORATION)

台北市內湖區南京東路六段380號A1棟10樓 ( 10F, NO. 380, SEC. 6, NANJING E. RD., NEIHU DIST., TAIPEI CITY, 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) 樣品名稱(Sample Description) 樣品型號(Style/Item No.) 樣品材質(Sample Material)	::	台灣塑膠工業股份有限公司(FORMOSA PLASTICS CORPORATION) FORMOCON(台塑鋼) FM025, FM025LCf 共聚型聚縮醛(POLYOXYMETHYLENE COPOLYMER)
	:	2024/03/13 2024/03/13 to 2024/03/22

#### 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 177.2470 (d)(2) 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 177.2470 (d)(2). 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 177.2470 (d)(1) 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 177.2470 (d)(1). Please refer to the result table(s) for the testing item(s).)

測試結果 (Test Results)

: 請見下一頁。(Please refer to next page(s).)

報告簽署(人/張伯睿 博士/部經理

Ray Chang, Ph.D. / Department Manager Signed for and on behalf of SGS Taiwan Ltd 化學實驗室-高雄/ Chemical Laboratory-Kaohsiung



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通過(PASS)

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#### 測試部位敘述(Test Part Description)

No.1 : 白色塑膠片 (WHITE PLASTIC SHEET)

#### 測試結果(Test Results)

(1)					
測試項目 (Test Item(s))	測試方法 (Method)	單位 (Unit)	RL	結果 (Result) No.1	限值 (Limit)
總萃取物 (水, 迴流, 6小時) / Total extractives (D.I. Water, reflux, 6 h)	參考美國FDA 21 CFR 177.2470. / With reference to US FDA 21 CFR 177.2470.	%(w/w)	0.02	0.0340	0.2
總萃取物 (正庚烷, 迴流, 6小時) / Net chloroform-soluble extractives (D.I. Water, 212°F, 30 min)	參考美國FDA 21 CFR 177.2470. / With reference to US FDA 21 CFR 177.2470.	%(w/w)	0.02	0.0557	0.15

(2)

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°F, 2小時) / Net chloroform-soluble extractives (D.I. Water, 150°F, 2 h)	參考美國FDA 21 CFR 177.2470 condition D. / With reference to US FDA 21 CFR 177.2470 condition D.	mg/in²	0.1	n.d.	0.5	
氯仿可萃取物 (正庚烷, 100°F, 30分鐘) / Net chloroform-soluble extractives (n-Heptane, 100°F, 30 min)	參考美國FDA 21 CFR 177.2470 condition D. / With reference to US FDA 21 CFR 177.2470 condition D.	mg/in²	0.1	n.d.	0.5	
氯仿可萃取物 (8% 乙醇, 150°F, 2小時) / Net chloroform-soluble extractives (8% Alcohol, 150°F, 2 h)	參考美國FDA 21 CFR 177.2470 condition D. / With reference to US FDA 21 CFR 177.2470 condition D.	mg/in²	0.1	n.d.	0.5	

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#### 備註(Note):

- 1. 0.1% = 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. 本實驗室之報告符合性聲明依ILAC-G8:09/2019簡單允收之二分法判定規則(w=0 · AL=TL)做為測試結果符合性聲明 判定之判定依據。(The decision rule of the statements of conformity is following the ILAC G8:09/2019 by using the simple acceptance decision rule.)

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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.)



\*\* 報告結尾 (End of Report) \*\*

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