測試報告
Test Report

號碼(No.) : CY/2019/70005A　　日期(Date) : 2019/10/25　　頁數(Page) : 1 of 3

中華紙漿股份有限公司
CHUNG HWA PULP CORPORATION
台北市中正區重慶南路二段51號12樓
12F., NO. 51, SEC. 2, CHUNGCHING S. RD., ZHONGZHENG DISTRICT, TAIPEI CITY 100, 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)	:	中華紙漿股份有限公司 (CHUNG HWA PULP CORPORATION)
樣品名稱(Sample Description)	:	GREASEPROOF PAPER (防油紙)
樣品材質(Sample Material)	:	紙 (PAPER)
收件日期(Sample Receiving Date)	:	2019/07/01
測試期間(Testing Period)	:	2019/07/01 to 2019/07/10

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測試需求(Test Requested) :

客戶指定依據美國聯邦法規之藥物暨食品管理(FDA) 21 CFR 176.170(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 176.170 (Condition of use B) to conduct test. Please refer to result table for testing item(s).

測試結果(Test Results) : 請參閱下一頁 (Please refer to following pages).

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

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測試結果(Test Results)

測試部位(PART NAME)No.1　　:　　白色紙張 (WHITE PAPER)

通過(PASS)

測試項目 (Test Items)	單位 (Unit)	測試方法 (Method)	MDL	結果 (Result) No.1	限值 (Limit)
蒸發殘渣 (正庚烷, 120°F, 30 min) / Extractives residue (n-Heptane, 120°F, 30 min)	mg/in²	依據美國FDA 21 CFR 176.170 condition B (2017). / According to US FDA 21 CFR 176.170 condition B (2017).	0.2	n.d.	0.5
可溶於氯仿的萃取物 (水, 212°F, 30分鐘) / Net chloroform-soluble extractives (D.I. Water, 212°F, 30 min)	mg/in²		0.2	n.d.	0.5

備註(Note):

1. 0.1wt% = 1000ppm；mg/kg = ppm
2. MDL = Method Detection Limit (方法偵測極限值)
3. n.d. = Not Detected = below MDL (未檢出 / 低於MDL)

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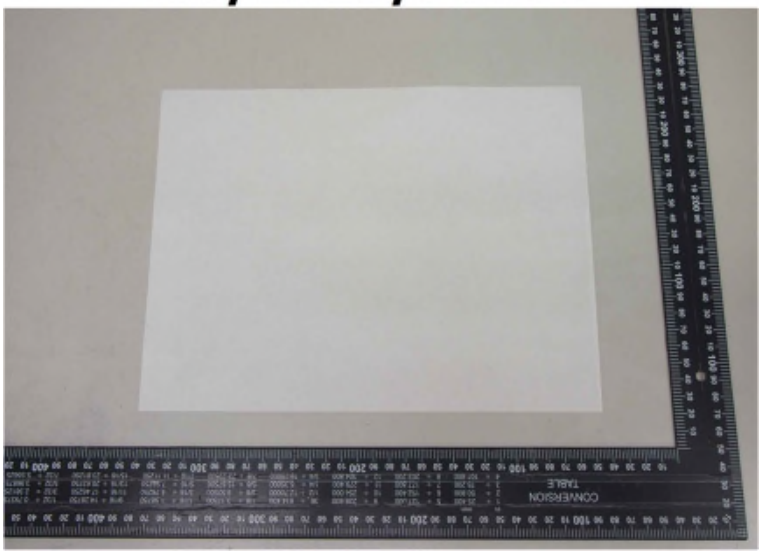
＊ 照片中如有箭頭標示，則表示為實際檢測之樣品/部位. ＊
(The tested sample / part is marked by an arrow if it's shown on the photo.)

CY/2019/70005



** 報告結尾 (End of Report) **