

測試報告

頁數(Page): 1 of 3 號碼(No.): CY/2018/80049 日期(Date): 2018/08/10

Test Report

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, CHONGCING S. RD., JHONGZHENG 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):

中華紙漿股份有限公司 (CHUNG HWA PULP CORPORATION) 送樣廠商(Sample Submitted By)

樣品名稱(Sample Description) CHP PAPER STRAW (紙吸管)

樣品型號(Style/Item No.) **PEDRO**

樣品批號(Lot No.)

樣品材質(Sample Material) 紙 (PAPER)

製造日期(Manufacturing Date)

原產國(Country of Origin) 台灣 (TAIWAN) 收件日期(Sample Receiving Date) 2018/08/01

2018/08/01 TO 2018/08/10 測試期間(Testing Period)

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E)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 E) 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).



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測試結果(Test Results)

測試部位(PART NAME)No.1 : 白色吸管 (WHITE STRAW)

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,70°F, 30分鐘)/ Extractives residue(n-Heptane, 70°F, 30 min)	mg/in²	依據美國FDA 21 CFR 176.170 condition E (2017). / According to US FDA 21 CFR 176.170 condition E (2017).	0. 2	n. d.	0.5
蒸發殘渣(水, 120°F, 24小時)/ Extractives residue (D.I. Water, 120 °F, 24 hr)	mg/in²		0.2	0. 323	0.5
可溶於氯仿的萃取物(50% 乙醇,120°F, 24小時)/ Net chloroform-soluble extractives(50% Alcohol,120°F,24 hr)	mg/in²		0. 2	n. d.	0.5
可溶於氯仿的萃取物 (8% 乙醇, 120°F, 24 小時) / Net chloroform-soluble extractives (8% Alcohol, 120°F, 24 hr)			0. 2	n. d.	0.5

備註(Note):

- 1. 0.1 wt% = 1000 ppm; 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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號碼(No.): CY/2018/80049 日期(Date): 2018/08/10 頁數(Page): 3 of 3

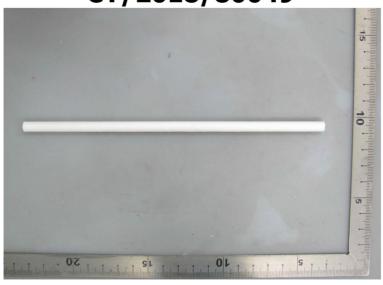
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* 照片中如有箭頭標示,則表示為實際檢測之樣品/部位. * (The tested sample / part is marked by an arrow if it's shown on the photo.)

CY/2018/80049



** 報告結尾 (End of Report) **

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