

測試報告 號碼(No.): CY/2020/50387

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, 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) :

日期(Date): 2020/06/09

EASY SEAL BOARD (益利咖) 樣品名稱(Sample Description) • 樣品材質(Sample Material) 紙 (PAPER) • 收件日期(Sample Receiving Date) 2020/05/29 • 測試期間(Testing Period) 2020/05/29 to 2020/06/09 :

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). :





頁數(Page): 1 of 3

PIN CODE: 4893019B

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測試結果(Test Results)

測試部位(PART NAME)No.1 : 白色紙(與食物接觸面)(WHITE PAPERBOARD (FOOD CONTACT SIDE))

通過(PASS)

	上述 2世 (1 AGS)				
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 (2020). / According to US FDA 21 CFR 176.170 condition B (2020).	0.2	n. d.	0.5
蒸發殘渣(水, 212°F, 30分鐘)/ Extractives residue (D.I. Water, 212 °F, 30 min)	mg/in ²	依據美國FDA 21 CFR 176.170 condition B (2020). / According to US FDA 21 CFR 176.170 condition B (2020).	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)
- 4. 符合性结果之判定係以測試結果與限值做比較。/ The statement of compliance conformity is based on comparison of testing results and limits.

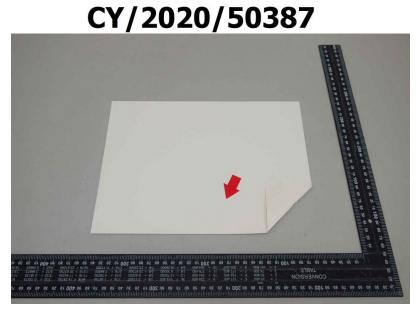


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* 照片中如有箭頭標示,則表示為實際檢測之樣品/部位. * (The tested sample / part is marked by an arrow if it's shown on the photo.)



** 報告結尾 (End of Report) **

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