

测试报告

No. TAOHG1902363901

日期: 2019年06月06日 第1页,共4页

烟台白马包装有限公司 山东省烟台市福山区电信路222号

以下测试之样品是由申请者所提供及确认:一次性桌垫

SGS工作编号: QDHL1905009805CW - QD

样品接收日期: 2019年05月31日

测试周期: 2019年05月31日 - 2019年06月06日

测试要求: 根据客户要求测试

测试方法: 请参见下一页 测试结果: 请参见下一页

测试结果概要:

测试要求	结论
FDA 21 CFR 177.1520- 萃取物含量	符合
FDA 21 CFR 177.1520- 密度 (23℃)	符合
FDA 21 CFR 177.1520- 二甲苯中可溶物含量	符合

结论: 测试参数符合FDA 21 CFR 177.1520 的要求。

通标标准技术服务(青岛)有限公司 授权签名

Wang Bo, Claire 王渤

批准签署人



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测试结果:

测试样品描述:

样品编号 SGS样品ID 描述 材质

(客户提供)

SN1 TAO19-023639.001 多色印刷的透明塑料薄膜 PE

备注:

(1) mg/dm² = 毫克每平方分米

(2) mg/kg = 毫克每千克

(3) ℃= 摄氏度

(4) <= 小于

(5) MDL = 方法检测限

(6) ND = 未检出 (< MDL)

FDA 21 CFR 177.1520-萃取物含量

测试方法: 参考FDA 21 CFR 177.1520(d)(3)(ii).

常用模拟液 时间 温度 最大允许限值 样品001 结论

正已烷 2hr(s) 50℃ 5.5% (w/w) 2.1% (w/w) 符合

FDA 21 CFR 177.1520-密度 (23℃)

测试方法: 参考FDA 21 CFR 177.1520d(1).

<u>测试项目</u> <u>限值</u> <u>001</u> 密度 (23°C), g / cm³ 0.85 - 1.00 0.97 结论 符合

FDA 21 CFR 177.1520-二甲苯中可溶物含量

测试方法: 参考FDA 21 CFR 177.1520(d)(4)(ii).



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测试项目 限值 单位 MDL 001 二甲苯中可溶物含量 11.3 % (w/w) 0.5 8.4 符合 结论

检测报告仅用于客户科研、教学、内部质量控制、产品研发等目的,仅供内部参考。



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样品照片:





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