

# 检验报告单

## Certificate of Analysis



### Orlistat 奥利司他

|                 |            |                         |            |
|-----------------|------------|-------------------------|------------|
| BATCH No. 批号    | ORB0141201 | MANUFACTURING DATE 生产日期 | 2015.01.04 |
| QUANTITY 数量     | 75kg       | TEST DATE 检验日期          | 2015.01.19 |
| RETEST DATE 复检期 | 2018.01.03 | REPORT DATE 报告日期        | 2015.03.24 |

| Item 项目                  | Standard 标准                                                                                                                   | Result 结果                                |
|--------------------------|-------------------------------------------------------------------------------------------------------------------------------|------------------------------------------|
| Description 性状           | White or almost white crystalline powder.<br>白色或类白色结晶性粉末                                                                      | Almost white crystalline powder 类白色结晶性粉末 |
| Solubility 溶解性           | Practically insoluble in water, freely soluble in chloroform. Very soluble in methanol and ethanol. 几乎不溶于水, 易溶于氯仿, 极易溶于甲醇和乙醇。 | Conforms 符合规定                            |
| Identification 鉴别        | IR                                                                                                                            | Conforms 符合规定                            |
| 鉴别                       | HPLC                                                                                                                          | Conforms 符合规定                            |
| Melting Point 熔点         | 42~45°C                                                                                                                       | 44.0~44.5°C                              |
| Specific Rotation 比旋度    | -31~-38°                                                                                                                      | -35°                                     |
| Water 水分                 | ≤2.0%                                                                                                                         | 0.0%                                     |
| Residual solvents 残留溶剂   | Methanol 甲醇 ≤3000ppm                                                                                                          | <59ppm                                   |
| Individual Impurity 单个杂质 | n-Heptane 正庚烷 ≤5000ppm                                                                                                        | 2956ppm                                  |
| Total Impurities 总杂质     | ≤1.0%                                                                                                                         | 0.4%                                     |
| Assay (HPLC) 含量          | ≤2.0%                                                                                                                         | 1.2%                                     |
| Bulk Density 堆密度         | ≥98.0% (on anhydrous basis)                                                                                                   | 98.7%                                    |
| Particle Size 颗粒度        | Perform and Report 检测并报告                                                                                                      | 0.31g/ml                                 |
| D10                      | Perform and Report 检测并报告                                                                                                      | 3um                                      |
| D50                      | Perform and Report 检测并报告                                                                                                      | 23um                                     |
| D90                      | Perform and Report 检测并报告                                                                                                      | 83um                                     |
| Microbial Limits 微生物限度   | Total Aerobic Microbial Count 总需氧菌 ≤1000CFU/g                                                                                 | <10CFU/g                                 |

CONCLUSION: The results conform with client standard.

结论: 本品符合客户标准。

Storage conditions: Stored in temperature between 2~8°C, away from light.

QC MANAGER 质量检验 何明霞 CHECKED BY 复核人 ANALYST 化验员 沈晓芳

FINAL BATCH DISPOSED 最终批号

Approved

Rejected

By: 李天

Version 1.1 SYB6

02/06/13

Hisun Pharmaceutical (Hangzhou) Co., Ltd.