

天津市津津药业有限公司检验报告单

TIANJIN JINJIN PHARMACEUTICAL CO., LTD. CERTIFICATE OF ANALYSIS

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|-----------------------------|----------------|--|---------------------|-------------|
| Report No.: | 2020-004 | Analytical No.: | 2020-01-04 | Page 1 of 1 |
| Product Name: | Spironolactone | * 化验专用草 / | | |
| Batch No.: | SA1074 | Manuf. Date: | 2019.12.24 | |
| Quantity: | 30Kg | Retest Date: | 2023.12.23 | |
| Package: | drum or carton | Report Date: | 2020.1.8 | |
| Items | | Requirements | Results | |
| Appearance(Ph.Eur.) | | a white or yellowish-white powder | White powder | |
| Solubility(Ph.Eur.)(annual) | | practically insoluble in water, soluble in ethanol (96 per cent) | Complies | |
| Identification | | ommor (50 per comy | | |
| A.IR -spectrum(P | h.Eur.) | meets the requirement | Complies | |
| A.IR -spectrum(USP) | | meets the requirement | Complies | |
| B.TLC(Ph.Eur.) | | meets the requirement | Complies | |
| B. UV Absorption(USP) | | meets the requirement | Complies | |
| C. Colour Reaction(Ph.Eur.) | | meets the requirement | Complies | |
| C.Precipitate reaction(USP) | | meets the requirement | Complies | |
| Specific optical rotation | on | 2 | - p | |
| (Ph.Eur.) | | between -46° and -41° | -44.7° | |
| (USP) | ě | between -41° and -45° | -43.3° | |
| Loss on drying | | | | |
| (Ph.Eur.) | | not more than 0.5% | 0.02% | |
| (USP) | | not more than 0.5% | 0.07% | |
| Sulphated ash(Ph.Eur) | | not more than 0.1% | 0.00% | |
| Related substances(Pl | n.Eur.) | | | * |
| Impurity A | a sec | not more than 0.2% | ND | |
| Impurity F | | not more than 0.3% | ND | |
| Impurity C | | not more than 0.2% | ND | |
| Impurity D | e gi | not more than 0.15% | 0.05% | |
| Impurity E | • | not more than 0.3% | ND | |
| Impurity I | | not more than 0.5% | ND | |
| Any unspecified in | purity | not more than 0.10% | Complies | |
| Total | | not more than 0.7% | 0.11% | |
| Ordinary impurities(USP) | | not more than 2.0% | Complies | |
| Free thiol compounds() | N. C. | a blue colour develops | Complies | |
| Free mercapto compou | nds(USP) | not more than 0.10ml | Complies | |
| Assay | | | | |
| HPLC (Ph.Eur.) | | between 97.5 % and 102.0 % | 100.9% | |
| HPLC (USP) | | between 97.0 % and 103.0 % | 99.4% | |
| Residual solvent(In-hou | ise) | | | |
| Methanol | | not more than 3000ppm | бррт | |
| Tetrahydrofuran | | not more than 720ppm | Less than LOD(0.4pp | m) |
| DMF | | not more than 880ppm | Less than LOD(43ppr | |
| Particle size(In-house) | | 95% not more than 20 microns | Complies | |

CONCLUSION: THE PRODUCT CONFORMS TO Ph.Eur.9.8/USP 40

Responsible Person:

2020.1.16

Checker:

Date: 7070. 1.16