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Date: 30.10.20

Cust.No: 96888

Certificate of analysis

Report-No.: 113220867-99 002
Batch: 6436
Article: 82519 Dihydrotanshinone I

| Test | Unit | Limit | Testresult |
|---|------|-------------------|------------|
| Appearance, SOP 100005 | | powder | Conform |
| Color, SOP 100006 | | red | Conform |
| Solubility, SOP 105001: | | | Conform |
| Methanol | | soluble | Conform |
| Water | | sparingly soluble | Conform |
| Chloroform | | soluble | Conform |
| Identification (HPLC-HR/MS), SOP 204125 | | Conform | Conform |
| Identification (UV spectrum from HPLC-DAD analysis) according to specification, SOP 204311 | | Conform | Conform |
| Identification (IR-spectroscopy, Ph.Eur. 10.3, 2.2.24)/USP 43 NF 37 <197>), SOP 206000 | | Conform | Conform |
| Identification (1H-NMR-spectroscopy), (outsourced), SOP 206010 | | Conform | Conform |
| Identification (13C-NMR-spectroscopy), (outsourced), SOP 206020 | | Conform | Conform |
| Water content, (micro determination, coulometric titration), Ph.Eur. 10.0., 2.5.32, SOP 304291: | | | |
| Mean value | % | | < 0.2 |

Certificate of analysis

Report-No.: 113222867 - 99 002
 Batch: 6436
 Article: 82519 Dihydrotanshinone I

| Test | Unit | Limit | Testresult |
|---|------|----------|------------|
| Peakpurity, (HPLC), SOP 401367 | | Conform | Conform |
| Dihydrotanshinone I (HPLC), method 1, (% AU), SOP 440937 | % | >= 95.00 | 99.16 |
| Residual solvents, (headspace-GC), SOP 805765: Residual solvents | % | | 0.05 |
| Inorganic impurities, (ICP-MS), for reference substances, SOP 811701: | | | |
| Sodium | % | | < 0.1 |
| Potassium | % | | < 0.1 |
| Magnesium | % | | < 0.1 |
| Calcium | % | | < 0.1 |
| Aluminium | % | | < 0.1 |
| Phosphorus | % | | < 0.1 |
| Sulfur | % | | 0.2 |
| Content*, SOP 890000 | % | | 99 |

Assessment:

The above mentioned reference substance meets the specification.

*The absolute content is calculated considering the chromatographic purity, and if available, the content of water, residual solvents and inorganic impurities according to the following formula:
 Content = (100% - water content (%) - residual solvents (%) - inorganic impurities (%)) x chromatographic purity (%) / 100.

The chromatographic purity is checked regularly: the last analysis has been performed in October 2020.

The reference substance cannot be documented with an expiry date. The pack is closed and is recommended to be stored as indicated. The unopened product is guaranteed to fulfill the specifications of this analytical report for a period of 60 months. Once opened we can no longer guarantee the stability of the material.

Vestenbergsreuth, 30.10.20

Dr. Michael Schwarz
 Head of Reference Substances

This is a computer print and valid without signature. A signed certificate of analysis can be taken on request.