

Certificate of Analysis 9001 **Reference Material Product name** 2-Amino-5-nitrothiazole **Product code** Lot number MM3279.01 1009621 **CAS number** Appearance H_2 121-66-4 brown solid Molecular weight **Melting point** 145.14 202 °C (dec) **Molecular formula** Long-term storage $C_3H_3N_3O_2S$ 2 to 8 °C, dark Assay "as is" **99.2 %** Date of shipment: 02 Sep 2019 Producer confirms that this reference material (RM) meets the specification detailed on this Certificate of Analysis for one year from the date of shipment, provided the substance is stored under the recommended conditions unopened in the original container. **Release by:** Date of Release: Product Release Dia Dr. Sabine Schröder Luckenwalde, 16 Jul 2019 Page 1/7

Organisation certified to ISO 9001 | DQS 102448 and GMP (EXCIPACT)

Producer: LGC GmbH Louis-Pasteur-Str. 30 D-14943 Luckenwalde Germany www.lgcstandards.com



Product information

For laboratory use only. Not suitable for human or animal consumption.

Before usage of the RM, it should be allowed to warm to room temperature. No drying required, as the certified value is already corrected for the content of water and other volatile materials.

The product quality is controlled by regularly performed quality control tests (retests).

Further content

Identity Assay Final result

Revision table



Identity

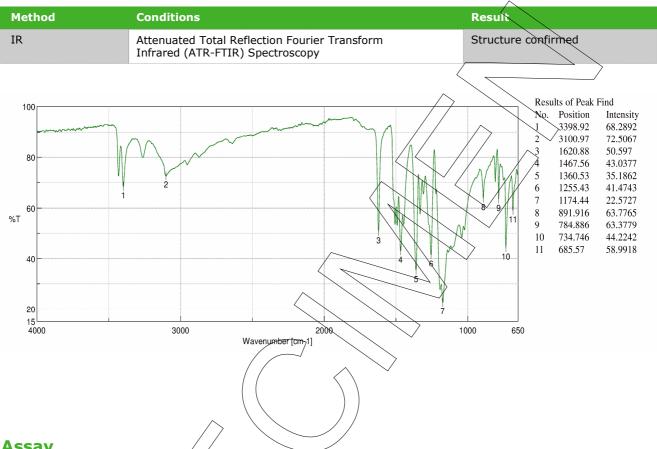
The identity of the reference material was established by following analyses.

| Method | Conditions | Resuit |
|--------------------|------------------------------|---------------------|
| ¹ H-NMR | 400 MHz, DMSO-d ₆ | Structure confirmed |
| | | |
| | | |
| 9.5 9.0 | | |
| | | |



| Method | Conditions | Result |
|---|--|--------------------------------------|
| MS | 3.5 kV ESI+; capillary temperature: 269 °C Theoretical value: 146.00187 | Structure confirmed |
| | | |
| Apex 7587 Scan: #14 100 90 80 70 70 80 70 70 80 70 70 80 70 70 70 70 70 70 70 70 70 7 | 9 RT: 1.58 min NL: 2.43E+007 | FTMS + p ESI Full ms [80.00-1000.00] |
| لىليات قارلقان قار سات بەر 1. قار ل o | 9517 9.98879 261,13064 391,28406 | m/z |
| 79 100 | 200 300 400 600 600 600 600 | 700 800 900 1.010 |
| | | |
| 4 | | |
| | | |





Assay

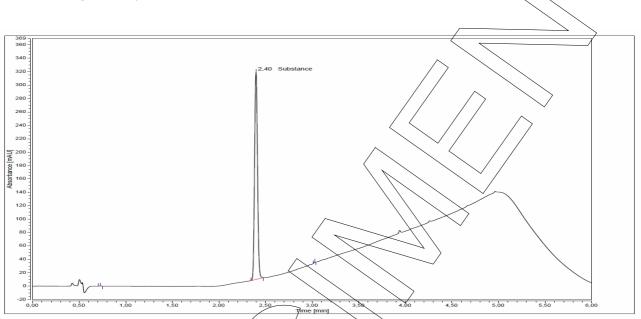
The assay of the reference material was assessed by following analyses.

Purity by High Performance Liquid Chromatography (HPLC)

| HPLC Conditions: | |
|--------------------|--|
| Column | Kinetex Phenyl-Hexyl; 1.7 µm, 100 x 2.1 mm |
| Column temperature | 40 °C |
| Detector | DAD, 240 nm |
| Injector | Auto 2 µl; 0.080 mg/ml in Acetonitrile/Water 50/50 (v/v) |
| Flow rate | 0.5 ml/min |
| Phase A | Water, 0.1 % HCOOH |
| Phase B | Acetonitrile, 0.1 % HCOOH |
| Gradient program | 0-1 min A/B 98/2 |
| | 1-4 min A/B to 2/98 |
| | 4-5 min A/B to 98/2 |
| | 5-6 min A/B 98/2 (v/v) |

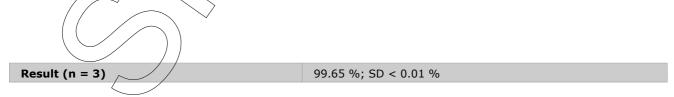


HPLC chromatogram and peak table



| Area percent report - sorted by signal | | | | |
|--|----------------|----------|--------|--|
| Pk # | Retention time | Area | Area % | |
| 1 | 0.728 | 0.0120 | 0.11 | |
| 2 | 2.398 | ~11.3243 | 99.65 | |
| 3 | 3.027 | 0.0282 | 0.25 | |
| Totals | | 11.3645 | 100.00 | |

The content of the analyte was determined as ratio of the peak area of the analyte and the cumulative areas of the purities, added up to 100 %. System peaks were ignored in calculation.





| Volatile content | | |
|-----------------------|------------------------|--|
| Water content | | |
| Method | Karl Fischer titration | |
| Result (n = 3) | 0.43 %; SD = 0.03 % | |
| | | |

| Residual solvents | | | | | | \sim | |
|-------------------|--|-------------------------|-------------------|------|--------|-----------|----|
| Method | ¹ H-NMR | $\langle \cdot \rangle$ | $\langle \rangle$ | / / | | / 7 | |
| Result (n = 1) | No significant amounts of residual sol | vent | s were | dete | cted (| < ⁄0.05 % |). |

Final result

Assay "as is": 99.22 %

The assay "as is" is assessed by 100% method (mass balance) and is equivalent to the assay based on the not anhydrous and not dried substance respectively.

The calculation of the 100% method follows the formula:

Assay (%) = (100 % - volatile contents(%))

Purity (%)

Volatile contents are considered as absolute contributions and purity is considered as relative contribution. Inorganic residues are excluded by additional tests.

| Revision table | | | |
|----------------|--|---------------------|--|
| Revision | Date | Reason for revision | |
| 00 | 16 Jul 2019 Release of the Certificate of Analysis - initial version | | |

Product warranties for the RM are set out in the terms and conditions of purchase.