

Certificate of Analysis

ISO 9001

| Reference Ma | terial | \langle | |
|--|---|--|--|
| Product name | | | |
| 2-Bromo-N-[4-bromo- ylcarbonyl)phenyl]acet | | D Br | Br |
| Product code MM0064.07 | Lot number | | |
| CAS number 1694-64-0 | Appearance brown solid | | |
| Molecular weight 398.05 | Melting point 104 °C (dec.) | | |
| Molecular formula $C_{14}H_{10}Br_2N_2O_2$ | Long-term storage -18 °C, dark | | |
| / | | "as is" 6 % | |
| Date of shipment: | 06 Apr 2020 | | |
| Producer confirms that this years from the date of sh container. | is reference material (RM) meets the s ipment, provided the substance is sto | specification detailed on this Cert red under the recommended cor | ificate of Analysis for two ditions unopened in the original |
| Release by: | Date of Release: | P | Product Release |
| Dr. Sabine Schröder | Luckenwalde, 26 Mar 2020 | () Chok | |

Organisation certified to ISO 9001 | DQS 102448 and GMP (EXCiPACT $^{\mathsf{TM}})$

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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

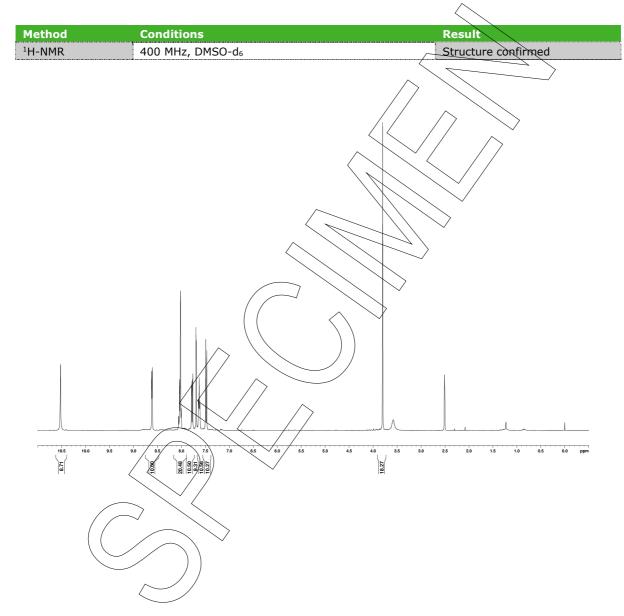
Identity Assay Final result

Revision table

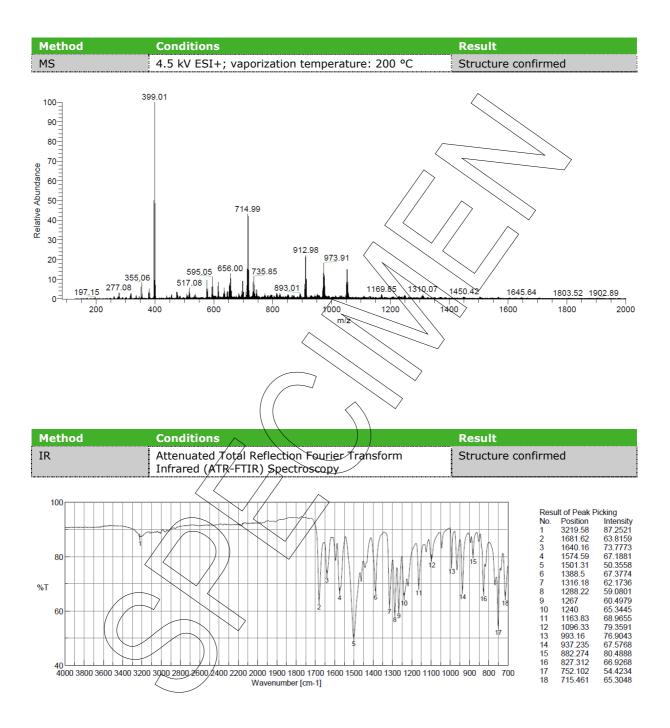


Identity

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







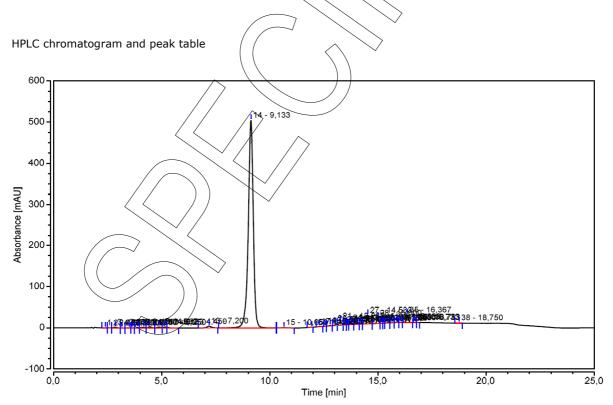


Assay

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

Purity by High Performance Liquid Chromatography (HPLC)

| HPLC conditions: | |
|--------------------|--|
| Column | Hypersil Gold C18; 5 pm, 150 x 4.6 mm |
| Column temperature | 40 °C |
| Detector | DAD, 235/nm |
| Injector | Auto 2 μl; 0.264 mg/ml in Acetonitrile |
| Flow rate | 1.0 mk/min |
| Phase A | Water, Q.1% H ₃ PQ ₄ |
| Phase B | Acetonitrile, 0.1 % H ₃ RO ₄ |
| Gradient program | 0-8 min A/B 65/35 8-13 min A/B to 10/90 13-18 min A/B 10/90 18-20 min A/B to 65/35 20-25 min A/B 65/35 (v/v) |





Mikromol

| Area percent report - sorted by signal | | | | |
|--|----------------|---------|--------|--|
| Pk # | Retention time | Area | Area % | |
| 1 | 2.400 | 0.018 | 0.01 | |
| 2 | 2.683 | 0.017 | 0.01 | |
| 3 | 2.867 | 0.333 | 0.23 | |
| 4 | 3.167 | 0.016 | 0,01 | |
| 5 | 3.500 | 0.047 | 0.03 | |
| 6 | 3.633 | 0.017 | 0.01 | |
| 7 | 3.900 | 0.142 | 0.10 | |
| 8 | 4.167 | 0.373 | 0.25 | |
| 9 | 4.450 | 0.527 | 0.36 | |
| 10 | 4.867 | 0.469 | 0.32 | |
| 11 | 5.133 | 0.423 | 0.29 | |
| 12 | 5.250 | 0.194 | 0.13 | |
| 13 | 7.200 | 1.118 | 0.76 | |
| 14 | 9.133 | 135.077 | 91.95 | |
| 15 | 10.650 | 0.184 | 0.12 | |
| 16 | 11.933 | 0.054 | 0.04 | |
| 17 | 17.283 | 0.125 | 0.09 | |
| 18 | 12.567 | 0.026 | 0.02 | |
| 19 | 12.783 | 0.121 | 0.08 | |
| 20 | 13.050 | 0.431 | 0.29 | |
| 21 | 13.267 | 1.143 | 0.78 | |
| 22 | 13.467 | 0.158 | 0.11 | |
| 23 | 13.583 | 0.073 | 0.05 | |
| 24 | 13.750 | 0.379 | 0.26 | |
| 25 | 14.017 | 0.202 | 0.14 | |
| 26 | 14.200 | 0.046 | 0.03 | |
| 27 | 14.533 | 1.794 | 1.22 | |
| 28 | 15.000 | 1.211 | 0.82 | |



| 29 | 15.133 | 0.053 | 0.04 | |
|--------|--------|---------|--------|---|
| 30 | 15.250 | 0.011 | 0.01 | |
| 31 | 15.450 | 0.141 | 0.10 | |
| 32 | 15.650 | 0.029 | 0.02 | |
| 33 | 15.883 | 0.139 | 0.09 | |
| 34 | 16.033 | 0.048 | 0,03 | |
| 35 | 16.367 | 1.641 | 1.12 | > |
| 36 | 16.733 | 0.028 | 0.92 | |
| 37 | 16.783 | 0.019 | 0.01 | |
| 38 | 18.750 | 0.067 | 0.05 | |
| Totals | | 146.897 | 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.

Result (n = 3)

91.91 %; SD = 0.10 %

Volatile content

| Water content | | | | |
|--|---|--|--|--|
| Method / / | Karl Fischer titration | | | |
| Result (n = 3) /0.38 %; SD = 0.02 % | | | | |
| | | | | |
| Residual solvents | | | | |
| Method | ¹ H-NMR | | | |
| Result (n = 1) | No significant amounts of residual solvents were detected (< 0.05 %). | | | |
| \sim | | | | |



| Final | result |
|---------|--------|
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Assay "as is":
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91.56 %

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.

Purity (%)

100 %

The calculation of the 100% method follows the formula:

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

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 | Re | asoi | ı for revisi | on | |
|----------|-------------|-----|----------|--------------|----|-----------------------------------|
| 00 | 26 Mar 2020 | | | of the Cerl | | ate of Analysis – initial version |
| | | 7 7 | <u> </u> | <u> </u> | , | |

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