



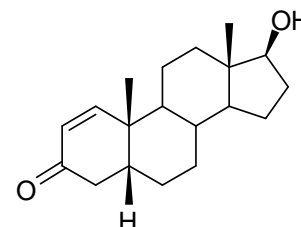
CERTIFIED REFERENCE MATERIAL CERTIFICATE OF ANALYSIS

NMIA D564: 5 β -Androst-1-en-17 β -ol-3-one

Report ID: D564.2023.01 (Ampouled 231018)

Chemical Formula: C₁₉H₂₈O₂

Molecular Weight: 288.4 g/mol



Certified value

| Batch No. | CAS No. | Mass per ampoule |
|-----------|------------|------------------|
| 99-000015 | 10529-96-1 | 994 ± 34 μg |

The uncertainty has been calculated according to ISO Guide 35 and is stated at the 95% confidence limit (k = 2).

Expiration of certification: The property values are valid till 19 October 2026, three years from the date of re-certification provided the **unopened** material is handled and stored in accordance with the recommendations below. The material as issued in the unopened container and stored as recommended below should be suitable for use beyond this date, subject to confirmation of batch stability from the issuing body. The expiry date/shelf life does not apply to sample bottles that have been opened. In such cases it is recommended that the end-user conduct their own in-house stability trials. The material will be re-tested on an annual basis to ensure that the property values are still valid. In the event a product fails the stability trial, notification will be sent to all impacted customers.

Description: The compound is supplied as a dried aliquot in a sealed ampoule under an atmosphere of argon. The CRM is intended for a single use to prepare a standard solution containing D564. The material was prepared by synthesis, and certified for identity and purity by NMIA.

Intended use: This certified reference material is suitable for use as a primary calibrator.

Instructions for use: Open the ampoule and carefully rinse the interior at least three times with a suitable organic solvent (e.g. chloroform). This will transfer 994 ± 34 μg of anhydrous 5 β -androst-1-en-17 β -ol-3-one. The mass of analyte in each ampoule is calculated from the assigned purity of the bulk and the concentration of bulk material in a stock solution used to prepare the ampoules.

Recommended storage: When not in use, this material should be stored at or below 4 °C in a closed container in a dry, dark area.

Metrological traceability: The certified purity value is traceable to the SI unit for mass (kg) through Australian national standards via balance calibration. In the mass balance approach all impurities are quantified as a mass fraction and subtracted from 100%.

Stability: This material has demonstrated stability over a minimum period of three years. The measurement uncertainty at the 95% confidence interval includes a stability component which has been estimated from annual stability trials. The long-term stability of the compound in solution has not been examined.

Homogeneity assessment: The homogeneity of the material was assessed using purity assay by GC-FID on seven randomly selected ampoules of the material. The material was judged to be sufficiently homogeneous at this level of sampling as the variation in analysis results between samples was not significantly different at a 95% confidence level from that observed on repeat analysis of the same sample.

Safety: Treat as hazardous substance. Use appropriate work practices when handling to avoid skin or eye contact, ingestion or inhalation of dust. Refer to the provided safety data sheet.

S. R. Davies

Dr Stephen R. Davies,
Team Leader,
Chemical Reference Materials, NMI.
7 November 2023

This report supersedes any issued prior to 7 November 2023.

NATA Accreditation No. 198 / Corporate Site No. 14214.

Legal notice: Terms and Conditions associated with the provision of this reference material can be found on the NMIA website.

Characterisation Report:

GC-FID: Instrument: Agilent 8890
 Column: HP-5, 30 m × 0.32 mm I.D. × 0.25 μm
 Program: 180 °C (1 min), 15 °C/min to 250 °C (5 min), 30 °C/min to 300 °C (3 min)
 Injector: 250 °C Detector Temp: 320 °C
 Carrier: Helium Split ratio: 20/1
 Relative mass fraction of the main component:
 Initial analysis: Mean = 99.5%, s = 0.009% (7 ampoules in duplicate, October 2023)

The following analytical data was obtained on the bulk material subsequently used in the preparation of the ampoules.

The identity was confirmed by a range of spectroscopic techniques, NMR, IR and MS. The certified purity value was obtained by mass balance from a combination of traditional analytical techniques, including GC-FID, thermogravimetric analysis, Karl Fischer analysis and ¹H NMR spectroscopy. The purity value is calculated as per Equation 1.

$$\text{Purity} = (100 \% - I_{\text{ORG}}) \times (100 \% - I_{\text{VOL}} - I_{\text{NVR}}) \quad \text{Equation 1}$$

I_{ORG} = Organic impurities of related structure, I_{VOL} = volatile impurities, I_{NVR} = non-volatile residue.

Supporting evidence is provided by qualitative elemental microanalysis.

GC-FID: Instrument: Agilent 5890/Agilent 6890/Agilent 8890
 Column: ZB-1 or HP-5, 30 m × 0.32 mm I.D. × 0.25 μm
 Program: 180 °C (1 min), 15 °C/min to 250 °C (5 min), 30 °C/min to 300 °C (3 min)
 Injector: 250 °C Detector Temp: 320 °C
 Carrier: Helium Split ratio: 20/1
 Relative mass fraction of the main component:
 Initial analysis: Mean = 99.9% (7 sub samples, February 1999)
 Re-analysis: Mean = 99.5%, s = 0.02 (8 sub samples in duplicate, June 2006)
 Re-analysis: Mean = 99.5%, s = 0.002% (7 sub samples in duplicate, August 2014)
 Re-analysis: Mean = 99.6%, s = 0.01% (5 sub samples in duplicate, April 2015)
 Re-analysis: Mean = 99.6%, s = 0.01% (7 sub samples in duplicate, October 2023)

HPLC: Peak area percentage of total > 99.5% (3 sub samples)
 Column: Alltima C-18, 5 μm (4.6 mm × 150 mm)
 Mobile Phase: Acetonitrile/water (60:40)
 Flow Rate: 1 mL/min
 Detector: Refractive index

Karl Fischer analysis: Moisture content ≤ 0.1% mass fraction (September 2014, April 2015 & October 2023)

Thermogravimetric analysis: Volatiles content < 0.1% and non-volatile residue < 0.2% mass fraction (March 1999 & October 2005)

Spectroscopic and other characterisation data

| | | |
|---|---|--|
| GC-MS: | Parent compound: | |
| | Instrument: | HP6890/5973 |
| | Column: | HP Ultra 2, 17 m x 0.20 mm I.D. x 0.10 μ m |
| | Program: | 180 $^{\circ}$ C (1 min), 10 $^{\circ}$ C/min to 220 $^{\circ}$ C, 20 $^{\circ}$ C/min to 300 $^{\circ}$ C (3 min) |
| | Injector: | 280 $^{\circ}$ C, Split injection |
| | Transfer line temp: | 300 $^{\circ}$ C |
| | Carrier: | Helium, 1.0 mL/min |
| | Scan range: | 50-550 <i>m/z</i> |
| | <i>Bis</i> -TMS derivative: | |
| | Instrument: | HP 6890/5973 |
| Column: | HP Ultra 1, 17 m x 0.22 mm I.D. x 0.11 μ m | |
| Program: | 170 $^{\circ}$ C (0.5 min), 3 $^{\circ}$ C/min to 234 $^{\circ}$ C, 10 $^{\circ}$ C/min to 265 $^{\circ}$ C (3 min) | |
| Injector: | 280 $^{\circ}$ C, Split injection | |
| Transfer line temp: | 300 $^{\circ}$ C | |
| Carrier: | Helium | |
| Scan range: | 50-550 <i>m/z</i> | |
| The retention times of the parent compound and <i>bis</i> -TMS derivative are reported with the major peaks in the mass spectra. The latter are reported as mass/charge ratios and (in brackets) as a percentage relative to the base peak. | | |
| Parent (5.9 min): | | 288 (M^+ , 20), 270 (10), 134 (31), 122 (100), 109 (78), 78 (29) <i>m/z</i> |
| <i>Bis</i> -TMS (7.6 min): | | 432 (M^+ , 36), 417 (34), 206 (22), 194 (100), 73 (73) <i>m/z</i> |
| TLC: | Conditions: | Kieselgel 60F ₂₅₄ . Hexane/ethyl acetate (5:3) Single spot observed, R_f = 0.3 (5 sub samples) |
| IR: | Instrument: | FT-IR, Biorad WIN FTS40 |
| | Range: | 4000-400 cm^{-1} , KBr pellet |
| | Peaks: | 3426, 1673, 1450, 1273, 1056, 842, 784 cm^{-1} |
| ¹ H NMR: | Instrument: | Bruker Advance-300 |
| | Field strength: | 300 MHz |
| | Solvent: | CDCl ₃ (7.26 ppm) |
| | Key spectral data: | δ 0.77 (3H, s), 1.21 (3H, s), 3.64 (1H, t), 5.89 (1H, d), 6.83 (1H, d) ppm |
| ¹³ C NMR | Instrument: | Bruker Advance-300 |
| | Field strength: | 75 MHz |
| | Solvent: | CDCl ₃ (77.2 ppm) |
| | Spectral data: | δ 11.5, 21.2, 22.3, 23.7, 25.9, 26.7, 30.6, 35.6, 37.0, 39.0, 39.3, 41.3, 43.4, 46.8, 50.7, 82.0, 127.3, 162.0, 201.2 ppm |
| Melting point: | | 201.5 – 202.5 $^{\circ}$ C |
| Microanalysis: | Found: | C = 79.0%; H = 9.8% (February 1999) |
| | Calculated: | C = 79.1%; H = 9.8% (Calculated for C ₁₉ H ₂₈ O ₂) |