



## Certificate of Analysis

### ISO Guide 34 Reference Material

#### Product Identification

Article Code: DRE-C12081000  
Article Name: 2,4'-DDT  
Formula: C<sub>14</sub>H<sub>9</sub>Cl<sub>5</sub>  
Mol. Weight: 354.49  
CAS No.: 789-02-6

Lot Number: G137925  
Expiry Date: 17.03.2023  
Storage Temperature: 20°C ± 4°C

Storage and handling: The RM should be stored in the original sealed bottle at the temperature given above. After use the bottle should be tightly closed and protected from moisture and light. The expiry date is valid for original sealed bottles under recommended storage conditions only.

Purity: 97.68% (g/g)

Expanded Uncertainty U<sub>m</sub>: 0.43% (g/g)

The uncertainty of this standard is calculated in accordance with the ISO Guide 34 and EURACHEM/CTAC Guide - Quantifying Uncertainty in Analytical Measurement, Second Edition. The expanded uncertainty is  $U(\text{exp}) = u(\text{RM}) \times k$ , where  $k$  is the coverage factor at the 95% confidence level ( $k=2$ ). Uncertainty  $u(\text{RM})$  is based on the combination of the uncertainties associated with each individual operation involved in the analysis of the product:  $u(\text{RM}) = \sqrt{u(\text{char})^2 + u(\text{bb})^2 + u(\text{sts})^2}$ ;  $u(\text{char})$  is the uncertainty of purity determination;  $u(\text{bb})$  uncertainty of homogeneity test;  $u(\text{sts})$  uncertainty of stability test long-term;  $u(\text{sts})$  uncertainty of stability test short-term.  $u(\text{ts})$  and  $u(\text{sts})$  are not included in the calculation as the stability statement is based on real evidence opposed to simulation. Minimum sample: 1 mg is recommended as the minimal sample amount. If less material is used, it is recommended to increase the certified uncertainty by a factor of two for half sample and a factor of four for a quarter of sample.

Intended use: Use this RM as calibrant for chromatography or any other analytical technique.

#### Analytical Data

Traceability of chromatography: To the International System of Units (SI).

Instrument:	GC/FID	Injector:	320°C
Detection:	FID	Initial Temp:	120°C for 4 min
Column:	Optima-5MS, 0.25 µm, 0.25 mm	End Temp:	320°C for 3 min
Inj.-Vol.:	1 µl	Gradient:	15°C/min
Flow:	1 ml/min		
Ret.Time:	14.53 min		

#### Comment

Traceability: The balances used are calibrated with weights traceable to the national standards (DKD).

Calibrated class A glassware is used for volumetric measurements.

Certificate Revision 1

Water Content: 0.01% (g/g) by Karl-Fischer-Titration ( $U(\text{exp}) = 0.03\%$  (g/g)).

Identity: EA, NMR, RT, IR

Certified on: 29.05.2017  
Certified by: M. Beck

The LGC Labor GmbH, accredited by DAkkS as indicated by the accreditation number D-RM-19883-01 & D-PL-19883-01, has shown competence based on ISO Guide 34:2009 with relevant parts of DIN EN ISO/IEC 17025:2005 for production of certified reference materials in form of organic pure substances and in form of single and multi-component solutions of organic pure substances.

<p>1</p>	<p>1</p>
<p>2</p>	<p>2</p>
<p>3</p>	<p>3</p>
<p>4</p>	<p>4</p>
<p>5</p>	<p>5</p>
<p>6</p>	<p>6</p>
<p>7</p>	<p>7</p>
<p>8</p>	<p>8</p>
<p>9</p>	<p>9</p>

26.5.17

Data file: 12081000-27-r001.dx

Instrument: FID 1

Sample name: 70523CY G137925

Sequence Name: 2017KW21-3b

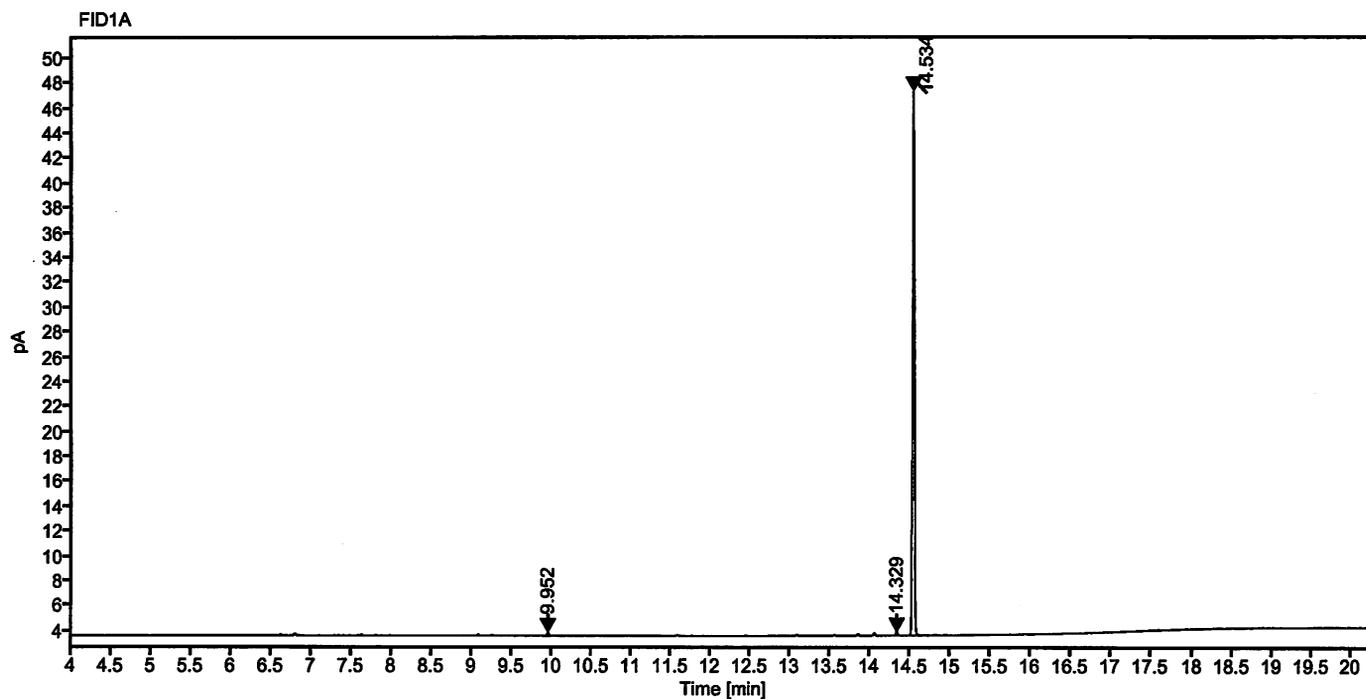
Inj. volume [µl]: 1.0

Injection date: 5/25/2017 11:55:31 AM

Acq. method: PAHK.amx

Location: 79

Sample Description 2,4'-DDT



Signal: FID1A

Nr.	RT [min]	Area [pA*s]	Height [pA]	Area%	Width [min]
1	9.952	0.39822	0.28	0.62	0.075
2	14.329	0.45337	0.30	0.70	0.098
3	14.534	63.74087	43.69	98.68	0.163
	Sum	64.59			

*P. Behr*

1. The first part of the document  
describes the general situation  
and the objectives of the study.

2. The second part of the document  
describes the methodology used  
in the study.

3. The third part of the document  
describes the results of the study  
and discusses the implications  
of the findings.

4. The fourth part of the document  
describes the conclusions of the study  
and provides recommendations  
for future research.

5. The fifth part of the document  
describes the limitations of the study  
and the strengths of the research.

6. The sixth part of the document  
describes the references used in the study  
and provides a list of the sources  
consulted.

7. The seventh part of the document  
describes the appendices of the study  
and provides additional information  
on the data and the analysis.

8. The eighth part of the document  
describes the acknowledgments of the study  
and provides a list of the people  
who assisted in the research.