



HAL
open science

Tetraethyl(Pyrrolidine-2,2-diyl)Bisphosphonate

Gilles Olive, Alain Jacques

► **To cite this version:**

Gilles Olive, Alain Jacques. Tetraethyl(Pyrrolidine-2,2-diyl)Bisphosphonate. *Molecules*, 2001, 6, pp.M275. hal-00660742

HAL Id: hal-00660742

<https://hal.science/hal-00660742>

Submitted on 17 Jan 2012

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

Molecules **2001**, *6*, M275

Tetraethyl(pyrrolidine-2,2-diyl)bisphosphonate

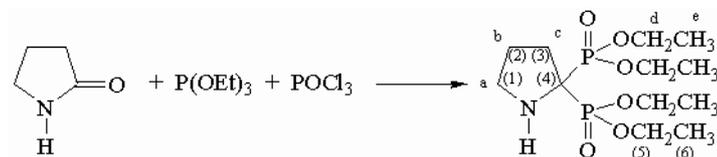
Gilles Olive ^{1*} and Alain Jacques ²

¹ Unité de physique et de chimie des hauts polymères, Université catholique de Louvain, Bâtiment Boltzmann, Place Croix du Sud, 1, B-1348 Louvain-la-Neuve, Belgium, Tel: +32 47 33 91, Fax: +32 45 15 93, e-mail: gilles.olive@excite.com

² Unité CSTR, Université catholique de Louvain, Bâtiment Lavoisier, Place Louis Pasteur, 1, B-1348 Louvain-la-Neuve, Belgium. E-mail: jacques@chim.ucl.ac.be

Received: 1 December 2001 / Accepted: 15 December 2001 / Published: 20 December 2001

Keywords: diphosphorylation, triethylphosphite, pyrrolidone, bisphosphonate.



1

The synthesis of the title compound is already described [1, 2] and we report here the fully optimized procedure. In a double walls flask, under nitrogen, phosphorus oxychloride (20 ml; 0.22 mol) is added for 1 h 00 time to a mixture at -7.5 °C of 2-pyrrolidinone (9.3 g; 0.11 mol) and triethylphosphite (35.1 g; 0.21 mol; 1.9 eq.). The reaction mixture is stirred for 1 hour at room temperature and then poured over a mixture of ice (200 g) and ammonia 30 % (400 ml). The aqueous layer is extracted with methylene chloride (3x100 ml) and then the latter is removed to obtain a yellow oil. The oil is dissolved in 100 ml of methylene chloride. An aqueous solution of hydrochloric acid (10 ml of 32 % HCl solution, 190 ml of water) is added (check that pH 1) and the aqueous layer is washed with methylene chloride (3x100 ml). A solution of sodium hydroxide (20 g of NaOH in 200 ml of water) is added up to pH 10 and the aqueous layer is extracted with methylene chloride (4x100 ml). The organic layer is dried over sodium sulfate, filtered and the removal of the solvent affords **1** (20.9 g; 58 %) as a colourless oil.

Formula: C₁₂H₂₇NO₆P₂.

Molecular weight: 343.13 g.mol⁻¹.

M.p.: -36 °C.

B.p.: 140 °C under 6.10⁻² mmHg.

R_f: 0.39 (Acetone); 0.43 (CH₂Cl₂ / EtOH 19:1).

UV (EtOH, 25 °C): λ_{max} (ε) 205 (1622), 221 (1737), 264 (212) nm (mol⁻¹.dm³.cm⁻¹).

pK_a: 3.5 - Krebs at 20 °C; 3.59 - Krebs at 37 °C; 3.47. [3]

IR (Neat): 3481 (n_{NH}); 2982 (n_{CH2} ring + n_{CH3} As.); 2932 (n_{CH2} ethoxy); 2909; 2869 (n_{CH3} Sym.); 1670 (d_{NH}); 1595; 1478; 1457 (d_{CH2} ring + ethoxy); 1392 or 1368 (d_{CHsp3} Sym.); 1324; 1293; 1243 (P=O); 1164 (P-OEt); 1098 (n_{C-N}); 1044; 968; 794; 732; 645; 580; 536 cm⁻¹.

Raman: 2979 (n_{CH2} ring + n_{CH3} As.); 2934 (n_{CH2} ethoxy); 2874 (n_{CH3} Sym.); 2778; 2725; 1594 (d_{NH} in plane); 1482; 1458 and 1449 (d_{CH2} ring + ethoxy); 1394 or 1369 (d_{CHsp3} Sym.); 1289; 1240 (P=O); 1190; 1162 (P-OEt); 1100 (n_{C-N}); 1027; 918; 877; 812; 757; 653; 291; 271 cm⁻¹.

¹H-NMR: δ in ppm referring to TMS (multiplicity, J in Hz); a, b, c, d and e as assigned in scheme.

Solvent	Frequency (MHz)	a	b	c	d	e
CDCl ₃	400	2.92 (t, 6.4)	1.74 (quint., 6.4)	2.16 (m)	4.07 (m)	1.20 (t, 7.0) 1.20 (t, 6.8)
C ₆ D ₆	400	2.88 (t, 6.5)	1.69 (quint., 6.5, 7.2)	2.42 (tt, 7.2 ^a , 17.7 ^b)	4.17 (m)	1.10 (t, 7.1) 1.11 (t, 7.1)
D ₂ O	500	2.98 (t, 6.4)	1.87 (quint., 6.4)	2.28 (tt, 7.2 ^a , 18.7 ^b)	4.21 (m)	1.34 (t, 7.1)
DMSO 55 °C = 328 K	500	2.92 (t, 6.9)	1.74 (quint., 7.2)	2.14 (tt, 7.2 ^a , 17.9 ^b)	4.08 (m)	1.23 (t, 7.2)
Acetone-d ₆ ^c	500	3.10 (t, 6.5)	1.91 (quint., 6.8)	2.33 (tt, 7.2 ^a , 17.7 ^b)	4.26 (m)	1.38 (t, 7.1)
CDCl ₃ ^d	500	0.87 s	0.79 s	0.58 s	1.78 s	1.85 s

^a J_{H-H}

^b J_{P-H}

^c Additional: 3.28 (ls, NH)

^d Value of T₁

¹³C-NMR: d in ppm referring to TMS (multiplicity, J_{CP} in Hz) (multiplicity, ¹J_{CH} in Hz for CDCl₃ and Acetone-d₆) ; (1), (2), (3), (4), (5) and (6) as assigned in scheme.

Solvent	Frequency (MHz)	(1)	(2)	(3)	(4)	(5)	(6)
CDCl ₃	125	47.4 (t, 4.7) (t, 139.4) ^a	26.1 (t, 3.7) (t, 133.8) ^a	30.5 (t, 3.6) (t, 135.4) ^a	61.7 (t, 152.3)	62.8 (t, 3.8) (tsext. ^c , 148.3) ^a 63.3 (t, 3.6) (tsext. ^c , 149.3) ^a	15.4 ^b (qh, 127.1) ^a 15.5 ^b (qh, 127.1) ^a
C ₆ D ₆	100	47.7 (t, 4.0)	26.5 (t, 3.1)	31.2 (t, 3.0)	62.8 (t, 151.8)	62.7 (t, 5.8) 63.4 (t, 5.8)	16.5 (t, 7.2) 16.6 (t, 5.5)
D ₂ O	125	47.0 (m)	25.4 (t, 3.1)	30.0 (m)	62.0	64.7 (q, 3.8)	15.6 (t, 2.7)
DMSO 30 °C = 303 K	125	47.0	25.6	30.5	61.4 (t, 151.3)	62.3 (t, 3.6) 62.6 ^b	16.3
DMSO 55 °C = 328 K	125	46.7 (t, 4.7)	25.3 (t, 3.7)	30.3 (t, 3.5)	61.6 (t, 151.7)	62.0 (t, 3.8) 62.3 (t, 3.4)	15.9 (t, 3.1) 16.0 (t, 2.3)
DMSO 70 °C = 343 K	125	46.9	25.6	30.6	62.1 (t, 151.5)	62.3 (t, 3.7) 62.5 (t, 3.5)	16.2
Acetone-d ₆	125	48.0 (t, 4.5) (t, 138.9) ^a	26.7 (t, 3.6) (t, 133.2) ^a	31.5 (d, 3.7) (t, 133.8) ^a	62.9 (t, 151.8)	63.2 (t, 3.7) (t, 148.04) ^a 63.6 (d, 3.4) (t, 147.9) ^a	16.8 (t, 3.8) (q, 130.5) ^a

^a Proton coupled (multiplicity, ¹J_{CH} in Hz)

^b Unresolved triplet

^c Triplet of sextuplet

Variable temperature in DMSO at 63 MHz [4]

Temp. °C	Temp. K	(1)	(2)	(3)	(4)	(5)	(6)
30	303	46.9 (t, 4.7)	25.6 (t, 3.5)	30.5 (t, 3.2)	61.4 (t, 151.7)	62.3 (t, 3.8) 62.6 (t, 3.6)	16.3 (t, 3.0)
50	323	46.9 (t, 4.8)	25.6 (t, 3.5)	30.5 (t, 3.2)	61.6 ^a	62.3 (t, 3.8) 62.5 (t, 3.6)	16.2 (t, 2.9)
60	333	46.9 (t, 4.7)	25.6 (t, 3.6)	30.5 (t, 3.2)	61.8 (t, 151.0)	62.3 (t, 3.8) 62.5 (t, 3.6)	16.2 (t, 2.7)
70	343	46.9 (t, 4.8)	25.6 (t, 3.7)	30.5 (t, 3.2)	61.9 ^a	62.3 (t, 3.8) 62.5 (t, 3.5)	16.2
90	363	46.9 (t, 4.8)	25.5 (t, 3.7)	30.5 (t, 3.2)		62.3 (t, 3.8) 62.5 (t, 3.7)	16.1 (t, 2.5)
110	383	46.9 (t, 4.8)	25.5 (t, 3.7)	30.6 (t, 3.1)		62.3 (t, 3.9) 62.5 (t, 3.7)	16.1

^a Only the central line

³¹P-NMR: d in ppm referring to external 85 % H₃PO₄

d (CDCl₃, 40 MHz): 22.5 ppm.

d (CDCl₃, 162 MHz): 24.7 ppm.

d (C₆D₆, 40 MHz): 22.8 ppm.

da (Krebs at 20 °C, 162 MHz): 16.51 ppm - db (Krebs at 20 °C, 162 MHz): 24.09 ppm. [3]

da (Krebs at 37 °C, 162 MHz): 16.53 ppm - db (Krebs at 37 °C, 162 MHz): 24.18 ppm. [3]

T₁ (CDCl₃, 202 MHz): 1.89 s.

³¹P MAS NMR: d (10 KHz spinning at 200 K): 22.39 ppm. [5]

¹⁵N-NMR: d in ppm referring to external CD₃NO₂: d (CDCl₃, 50 MHz): -341.6 (t, J_{NP} = 8.6) ppm.

d (C₆D₆, 50 MHz): -340.7 (t, J_{NP} = 8.7) ppm.

¹⁷O-NMR: d in ppm referring to external H₂O: d (DMSO, 68 MHz): 95.4 (P=O) ppm.

d (Acetone d₆, 68 MHz): 84.6 (P=O) and 53.9 (P-O) ppm.

References and Notes

1. Olive, G.; Le Moigne, F.; Mercier, A.; Rockenbauer, A.; Tordo, P. *J. Org. Chem.* **1998**, *63*, 9095-9099.
2. Olive, G.; van Genderen, M. H. P. *Magn. Reson. Chem.* **2000**, *38*(5), 379-381.
3. Pietri, S.; Miolan, M.; Martel, S.; Le Moigne, F.; Blaive, B.; Culcasi, M. *J. Biol. Chem.* **2000**, *275*(26), 19505-19512.
4. Olive, G.; Jacques, A. *To be published*.
5. Millot, Y.; Olive, G.; Magusin, P. *To be published*.

Sample Availability: Available from the author and from MDPI.

© 2001 MDPI. All rights reserved. *Molecules* website <http://www.mdpi.org/molecules/>