

A New Series of Liquid Crystalline Materials containing Boron Atoms

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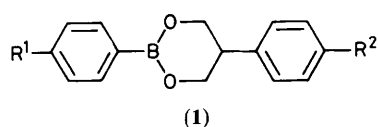
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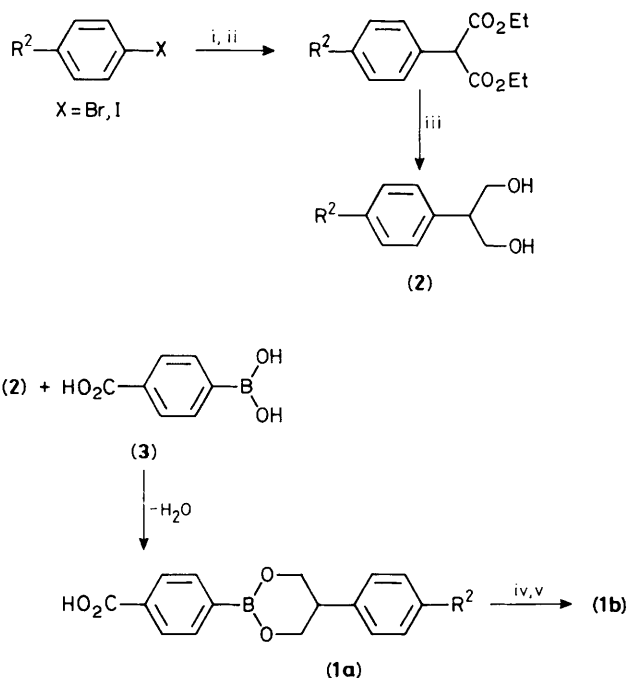
The dioxaborinane derivatives, which were synthesized by a method including a new Pd-catalysed coupling reaction, form mesomorphic phases in a wide temperature range, and provide a new series of liquid crystalline compounds containing a boron atom in the principal structure.

The majority of liquid crystalline molecules consist of carbon, oxygen, nitrogen, and hydrogen atoms. Though a few examples containing heteroatoms are known,¹ liquid crystalline materials containing metal atoms are of special interest since they are expected to have unique properties.² Previously, we have reported on lyotropic liquid-crystalline materials containing transition metals.³ Here we report the synthesis and properties of the 1,3,2-dioxaborinanes (**1**) which may be the first examples of thermotropic liquid-crystalline molecules with a boron atom in the principal structure.

The compound (**1**) was synthesized *via* the reactions shown in Scheme 1. The intermediate compound (**2**) was obtained from the coupling between the ethyl cyanoacetate anion and a 4-substituted phenyl halide, followed by esterification and reduction. The key step in the synthesis of (**2**) may be the coupling reaction which was found to proceed smoothly in the presence of a palladium catalyst in monoglyme at 70 °C.⁴ The compound (**3**) was prepared from tolylmagnesium bromide and trimethoxyborane by a two-step synthesis.⁵ Condensation



- a; R¹ = CO₂H
 b; R¹ = CN
 c; R¹ = CO₂Me
 d; R¹ = OMe



Scheme 1. Reagents: i, ⁻CH(CN)CO₂Et, Pd(PPh₃)₄; ii, EtOH-HCl-CaCl₂; iii, LiAlH₄; iv, SOCl₂, then NH₃; v, CCl₄-PPh₃/tetrahydrofuran.

Table 1. Transition temperatures^a of compounds (**1**).

(1)	R ¹	R ²	<i>T</i> (C-meso)/°C ^c	<i>T</i> (meso-I)/°C ^c
a	CO ₂ H	<i>n</i> -C ₆ H ₁₃	200(n)	260 ^b
		<i>n</i> -C ₄ H ₉ O	250 (n)	280 ^b
		<i>n</i> -C ₈ H ₁₇ O	202 (n)	273 ^b
b	CN	<i>n</i> -C ₆ H ₁₃	98 (n)	122.4
		<i>n</i> -C ₄ H ₉ O	117.0 (n)	166.0
		<i>n</i> -C ₈ H ₁₇ O	95.0 ^d	144.8
c	CO ₂ Me	<i>n</i> -C ₄ H ₉ O	156.2 (n)	158.5
		<i>n</i> -C ₈ H ₁₇ O	134.0 (s)	164.6
d	OMe	<i>n</i> -C ₄ H ₉ O	133.9 (n)	146.6
		<i>n</i> -C ₈ H ₁₇ O	110.6 ^e	132.3

^a Measured with a Mettler Thermal Microscope FP5 + FP52. ^b With decomposition. ^c C = crystal, I = isotropic, n = nematic, s = smectic.

^d *T*(C-smectic) 95.0, *T*(smectic-nematic) 113.6 °C. ^e *T*(C-smectic) 110.6, *T*(smectic-nematic) 118.7 °C.

between (**2**) and (**3**) in toluene gave (**1a**), which, on conversion of the hydroxycarbonyl group into the nitrile group yielded (**1b**).⁶ The structures of compounds (**1**) are consistent with analytical data including mass, i.r., and n.m.r. spectra. The

mesomorphic ranges of (**1**) were measured visually with an optical microscope and are summarized in Table 1.

Received, 18th September 1984; Com. 1326

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