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Lithium 2-(2-Lithiomethylphenyl)ethanolate from Isochroman: Easy Preparation of Substituted Benzoxepines and Functionalised Arenes

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Abstract: The reaction of isochroman (1) with an excess of lithium powder and a catalytic amount of DTBB (2.5 mol %) in THF at 20°C yields the title intermediate, which by reaction with different electrophiles (H_2O , D_2O , CO_2 and carbonyl compounds) at temperatures ranging between -78 and 20°C yields, after hydrolysis, the corresponding products 3. Diols 3h-m, derived from benzaldehyde or ketones, give upon treatment with 85% phosphoric acid the corresponding benzoxepines 5h-m. In the other cases, Friedel-Crafts products resulting from a rearrangement of the first carbenium ion formed, are obtained.

INTRODUCTION

The interest of functionalised organolithium compounds¹ in synthetic organic chemistry is focused on their capability to transfer their functionality to an electrophilic reagent, so giving directly polyfunctionalised organic structures. This type of intermediates are generally prepared by the usual ways to generate organolithium compounds², namely deprotonation, halogen-lithium exchange and mercury- or tin-lithium transmetallation, as the most important methodologies. In the case of organolithium compounds bearing an alkoxide functionality a new possibility for their preparation consists in the reductive opening of cyclic ethers³. Thus, using activated lithium (with a stoichiometric⁴ or catalytic⁵ amount of an arene) epoxides⁶, oxetanes⁷, tetrahydrofurans⁸, phthalan⁹ or dioxolanes¹⁰ (of the general type I or III) have been used as starting materials for the preparation of the corresponding oxygen-containing functionalised organolithium compounds (of the type II or IV). In this paper we describe for the first time the reductive opening of a six-membered cyclic ether, isochroman, using an arene-catalysed⁵ lithiation and the application of this process in organic synthesis.



RESULTS AND DISCUSSION

When commercially available isochroman (1) was allowed to react with an excess of lithium powder (*ca.* 1:20 molar ratio) and a catalytic amount of 4,4'-di-*tert*-butylbiphenyl (DTBB; 1:0.05 molar ratio; 2.5 mol %) in THF at ambient temperature for 45 min, a solution of the corresponding dianionic intermediate 2 was prepared, which reacted with different electrophiles (H₂O, D₂O, CO₂ and carbonyl compounds) at temperatures ranging between -78 to 20°C for 2 h to give, after hydrolysis, the corresponding products 3 (Scheme 1 and Table 1). In the absence of the catalyst lithiation times were longer (*ca.* 3 h) and yields were considerably lower.



Scheme 1. Reagents and conditions: i, Li excess, DTBB cat. (2.5 mol %), THF, 20°C, 45 min; ii, E⁺=H₂O, D₂O, CO₂, EtCHO, PrⁱCHO, BuⁱCHO, BuⁱCHO, PhCHO, Me₂CO, Et₂CO, (CH₂)₄CO, (CH₂)₅CO, PhCOMe, -78 to 20°C, 2 h; iii, H₂O, 20°C.

The hydroxyacid 3c (Table 1, entry 3) was easily cyclised to the corresponding ε -lactone 3'c (75% isolated) by means of a catalytic amount of *p*-toluenesulfonic acid at benzene reflux^{9,11} (Scheme 2).



Scheme 2. Reagents and conditions: i, TsOH cat., PhH reflux, 2 h.

We consider specially interesting the diols **3d-m** as starting materials for the corresponding oxepanes by an intramolecular dehydration. However, we could prepare these heterocycles only with the diols **3h-m** derived from benzaldehyde or ketones. Thus, by treating these precursors with 85% phosphoric acid at toluene reflux^{9,12} the corresponding benzoxepines **5h-m** were isolated (Scheme 3 and Table 2). When the more stable benzylic or tertiary carbenium ions **4h-m** are formed, they do not suffer a transposition reaction, and after nucleophilic attack of the remaining hydroxy group yield the corresponding products **5h-m**.

Entry	Electrophile E+	Producta					
		No.	E	Yield (%) ^b	Rfc		
1	H ₂ O	3a	Н	89	0.35		
2	D ₂ O	3b	D	86	0.35		
3	CO ₂	3 c	CO ₂ H	74	0.19		
4	EtCHO	3d	EtCHOH	68	0.24		
5	PriCHO	3 e	PriCHOH	80	0.28		
6	BuiCHO	3f	BuiCHOH	82	0.33		
7	ButCHO	3 g	ButCHOH	59	0.56		
8	PhCHO	3 h	PhCHOH	65	0.36		
9	Me ₂ CO	31	Me ₂ COH	44	0.23		
10	Et ₂ CO	3j	Et ₂ COH	55	0.41		
11	(CH ₂) ₄ CO	3 k	(CH ₂) ₄ COH	67	0.26		
12	(CH ₂)5CO	31	(CH ₂) ₅ COH	53	0.35		
13	PhCOMe	3m	PhC(OH)Me	44	0.38		

Table 1. Preparation of Compounds 3

^a All products **3** were >95% pure (GLC and 300 MHz ¹H NMR). ^b Isolated yield after flash chromatography (silica gel, hexane/ethyl acetate) based on the starting material **1**. ^c Silica gel, hexane/ethyl acetate: 1:1.



Scheme 3. Reagents and conditions: i, H₃PO₄, PhMe reflux, 2 h.

However, in the other cases (diols **3d-g**) we obtained always products resulting from a rearrangement of the initially formed carbocation **4**. Thus, the diol **3d** upon the same treatment as shown in Scheme 3 gave the carbenium ion **4d**, which underwent transposition to the more stable benzylic one **4'd** and final cyclisation to the cyclic ether **6d** (49% isolated; Scheme 4). Diols **3e** and **3g** behaved differently: after generating the carbocations **4e**,**g**, they rearranged to the tertiary one **4'e**,**g** (preferred than the corresponding benzylic one),

which after a Friedel-Crafts type aromatic S_E reaction gave the final isolated products 7e and 7g in 76 and 86% isolated yields, respectively (Scheme 4). Finally, the diol 3f yielded a mixture of the Friedel-Crafts product 8f (resulting from the rearranged carbenium ion 4'f) and the olefin 9f in 66% overall isolated yield (2:1 molar ratio). It is strange that neither conjugated (from 4f) or more substituted (from 4'f) olefins were obtained instead of 9f (Scheme 4).

Entry	Starting material	Producta					
		No.	R1	R ²	Yield (%) ^b	$R_{f^{c}}$	
1		5h	Ph	Н	67	0.37	
2	3i	5 i	Me	Me	72	0.28	
3	3ј	5j	Et	Et	79	0.49	
4	3k	5 k	-(CH ₂) ₄ -		69	0.40	
5	31	51	-(CH ₂) ₅ -		68	0.44	
6	3m	5m	Ph	Me	72	0.41	

 Table 2. Preparation of Benzoxepines 5

^a All products **5** were >95% pure (GLC and 300 MHz ¹H NMR). Isolated yield after flash chromatography (silica gel, hexane/ethyl acetate) based on the starting diol **3**. ^c Silica gel, hexane/ethyl acetate: 10/1.

From the results described in this paper we conclude that commercially available isochroman is an adequate precursor for functionalised arenes of the type 3 and heterocycles such as benzoxepines 5^{13} .

EXPERIMENTAL PART

General.- For general information see reference 9.

Preparation of Compounds 3. General Procedure.- To a blue suspension of lithium powder (0.125 g, 18.0 mmol) and a catalytic amount of 4,4'-di-tert-butylbiphenyl (0.047 g, 0.18 mmol) in THF (10 ml) at 20°C was added the isochroman (1) (0.250 ml, 2.0 mmol) under argon and the mixture was stirred for 45 min at the same temperature. Then, the mixture was cooled at -78°C and the corresponding electrophile (3.0 mmol; 0.5 ml in the case of water or deuterium oxide; CO_2 was bubbled for 1.5 h) was added. The mixture was stirred at the same temperature for 1 h and was hydrolysed with water. The resulting mixture was extracted with ethyl acetate. The organic layer was dried over anhydrous sodium sulfate and evaporated (15 mmHg). The residue was then purified by column chromatography (silica gel; hexane/ethyl acetate) and/or recrystallised to yield pure products **3a-m**. When the electrophile was CO_2 , after having hydrolysed the mixture with water at -78°C it was basified with 2.5 M sodium hydroxide and extracted with ethyl acetate. The aqueous layer was then acidified with 3 M hydrochloric acid and extracted with ethyl acetate. The organic layer was dried over anhydrous sodium sulfate and evaporated (15 mmHg). The resulting will acetate and evaporated (15 mmHg). The resulting mixture was then acidified with 3 M hydrochloric acid and extracted with ethyl acetate. The organic layer was dried over anhydrous sodium sulfate and evaporated (15 mmHg). The resulting residue was compound **3c** [>95% pure (GLC and 300 MHz ¹H NMR)]. Yields and R_f values are included in Table 1; other physical, analytical and spectroscopic data follow.

2-(2-Methylphenyl)ethanol (3a): v_{max} (film) 3640-3040 cm⁻¹ (OH); δ_{H} 1.95 (1H, br s, OH), 2.31 (3H, s, CH₃), 2.86 (2H, t, J =6.9, ArCH₂), 3.78 (2H, t, J =6.9, CH₂OH), 7.08-7.21 (4H, m, ArH); δ_{C} 19.3 (CH₃),



Scheme 4. Reagents and conditions: i, H₃PO₄, PhMe reflux, 2 h; ii, as in step i but 10 h.

36.3 (ArCH₂), 62.5 (CH₂OH), 125.9, 126.5, 129.5, 130.3, 136.3, 136.4 (ArC); *m/z* 136 (M+, 26%), 106 (33), 105 (100), 103 (15), 91 (23), 79 (20), 77 (25) (Found: M+, 136.0881. C₉H₁₂O requieres M, 136.0888).

2-(2-Deuteriomethylphenyl)ethanol (**3b**): ν_{max} (film) 3640-3080 cm⁻¹ (OH); δ_{H} 1.92 (1H, br s, OH), 2.29 (2H, s, CH₂D), 2.85 (2H, t, *J* =6.9, ArCH₂), 3.77 (2H, t, *J* =6.9, C *H*₂OH), 7.07-7.19 (4H, m, ArH); δ_{C} 19.1 (t, *J_{CD}* =19.3, CH₂D), 36.3 (ArCH₂), 62.5 (CH₂OH), 125.9, 126.5, 129.5, 130.3, 136.3, 136.4 (ArC); *m/z* 137 (M+, 35%), 118 (11), 107 (36), 106 (100), 105 (16), 104 (14), 92 (15), 91 (15), 80 (14), 79 (11), 78 (18), 77 (11) (Found: M+, 137.0950. C₉H₁₁DO requires M, 137.0951).

2-(2-Hydroxyethyl)phenylacetic Acid (3c)¹⁴: v_{max} (film) 3680-2260 (COOH), 1700 cm⁻¹ (C=O); $\delta_{\rm H}$ 2.80 (2H, t, *J*=6.9, ArCH₂), 3.62 (2H, s, CH₂COOH), 3.72 (2H, t, *J*=6.9, CH₂OH), 7.05-7.21 (4H, m, ArH), 7.37 (2H, br s, COOH, OH); $\delta_{\rm C}$ 35.4, (ArCH₂), 38.3 (CH₂COOH), 62.8 (CH₂OH), 126.7, 127.6, 130.0, 130.7, 132.5, 137.1 (ArC), 176.5 (C=O); *m/z* 162 (M⁺-H₂O, 27%), 118 (33), 117 (100), 115 (23), 105 (10), 104 (69), 103 (20), 91 (25), 78 (24), 77 (23), 65 (10), 63 (22), 52 (11), 51 (19), 50 (16).

1-[2-Hydroxyethyl)phenyl]-2-butanol (3d)¹⁴: v_{max} (film) 3680-3060 cm⁻¹ (OH); $\delta_{\rm H}$ 0.98 (3H, t, J=7.4, CH₃), 1.46-1.64 (2H, m, CH₃CH₂), 2.40-2.98 (6H, m, 2xArCH₂, 2xOH), 3.60-3.86 (3H, m, CH₂OH, CHOH), 7.14-7.25 (4H, m, ArH); $\delta_{\rm C}$ 10.1 (CH₃), 30.0 (CH₃CH₂), 35.5 (CH₂CH₂OH), 39.9 (ArCH₂), 63.3 (CH₂OH), 74.1 (CHOH), 126.5, 126.5, 129.9, 130.3, 137.4, 137.5 (ArC); *m/z* 136 (M⁺-C₃H₆O, 60%), 119 (11), 118 (54), 117 (100), 115 (29), 106 (65), 105 (97), 104 (23), 103 (26), 91 (54), 79 (13), 78 (20), 77 (32), 65 (17), 59 (77), 57 (17), 51 (13), 43 (18), 41 (35).

1-[2-(2-Hydroxyethyl)phenyl]-3-methyl-2-butanol (3e)¹⁴: v_{max} (film) 3700-3080 cm⁻¹ (OH); δ_{H} 1.02 (6H, d, $J=6.7, 2xCH_3$), 1.75-1.86 [1H, m, (CH₃)₂CH], 1.98 (2H, br s, 2xOH), 2.73 (1H, dd, J=14.0, 9.8, HCHCHOH), 2.82-2.94 (2H, m, CH₂CH₂OH), 3.00 (1H, dd, J=14.0, 6.5, HCHCHOH), 3.61 (1H, ddd, J=9.7, 5.2, 3.2, HCHOH), 3.79-3.94 (2H, m, HCHOH, CHOH), 7.16-7.23 (4H, m, ArH); δ_{C} 17.5, 18.7 (2xCH₃), 33.7 [(CH₃)₂CH], 35.5 (CH₂CH₂OH), 36.8 (CH₂CHOH), 63.4 (CH₂OH), 77.6 (CHOH), 126.5, 126.6, 130.0, 130.2, 137.6, 138.0 (ArC); *m/z* 147 (M⁺-C₃H₇-H₂O, 6%), 136 (58), 119 (24), 118 (69), 117 (98), 115 (20), 106 (100), 105 (92), 104 (26), 103 (25), 92 (11), 91 (55), 79 (17), 78 (17), 77 (31), 73 (31), 65 (12), 57 (10), 55 (35), 51 (11), 45 (13), 43 (86), 41 (69).

 $\begin{array}{l} 1\mbox{-}[2\mbox{-}(2\mbox{-}Hydroxyethyl)phenyl]\mbox{-}4\mbox{-}methyl\mbox{-}2\mbox{-}pentanol\ (3f)\mbox{1}\ v_{max}\ (film)\ 3660\mbox{-}3080\ cm\mbox{$^-1$}\ (OH);\ \delta_{H}\ 0.91\ (3H,\ d,\ J\mbox{=}6.5,\ CH_3CCH_3),\ 0.95\ (3H,\ d,\ J\mbox{=}6.5,\ CH_3CCH_3),\ 1.29\mbox{-}1.38\ (1H,\ m,\ HOCHHCHCH),\ 1.47\mbox{-}1.52\ (1H,\ m,\ HOCHHCHCH),\ 1.47\mbox{-}1.52\ (1H,\ m,\ HOCHHCHCH),\ 1.47\mbox{-}1.52\ (1H,\ m,\ HOCHHCHCH),\ 1.47\mbox{-}1.52\ (1H,\ m,\ ACH);\ \delta_{C}\ 22.0,\ 23.4,\ 24.6\ [CH(CH_3)_2],\ 35.6\ (CH_2CH_2OH),\ 40.8\ (ArCH_2),\ 46.5\ [CH_2CH(CH_3)_2],\ 63.4\ (CH_2OH),\ 70.8\ (CHOH),\ 126.6,\ 126.6,\ 130.0,\ 130.4,\ 137\mbox{-}4,\ 137\mbox{-}5\ (ArC);\ m/z\ 136\ (M^+\mbox{-}C_5H_8\mbox{-}H_2O,\ 63\%),\ 118\ (62),\ 117\ (47),\ 115\ (22),\ 106\ (71),\ 105\ (87),\ 104\ (23),\ 103\ (18),\ 91\ (38),\ 79\ (12),\ 78\ (12),\ 77\ (21),\ 69\ (24),\ 57\ (15),\ 45\ (17),\ 43\ (100),\ 42\ (16),\ 41\ (90). \end{array}$

1-[2-(2-Hydroxyethyl)phenyl]-3,3-dimethyl-2-butanol (**3g**): mp 90-91°C (pentane/dichloromethane); v_{max} (KBr) 3660-3040 cm⁻¹ (OH); $\delta_{\rm H}$ 1.00 [9H, s, (CH₃)₃C], 2.25 (2H, br s, 2xOH), 2.65 (1H, dd, *J*=13.9, 10.6, ArCHH), 2.79-2.88 (2H, m, CH₂CH₂OH), 2.96 (1H, dd, *J*= 13.9, 2.1, ArCHH), 3.44 (1H, dd, *J*=10.6, 2.1, CHOH), 3.74-3.90 (2H, m, CH₂OH), 7.15-7.25 (4H, m, ArH); $\delta_{\rm C}$ 25.7 [(CH₃)₃C], 34.2 (CH₂CH₂OH), 35.1 [(CH₃)₃C], 35.4 (ArCH₂), 63.5 (CH₂OH), 80.7 (CHOH), 126.5, 126.6, 130.0, 130.2, 137.7, 138.5 (ArC); *m/z* 204 (M⁺-H₂O, 1%), 136 (100), 129 (16), 119 (28), 118 (96), 117 (73), 115 (23), 106 (88), 105 (91), 104 (19), 103 (16), 92(10), 91 (51), 87 (23), 79 (18), 78 (15), 77 (30), 69 (27), 65 (14), 57 (43), 45 (21), 43 (29), 41 (67). Anal. Calcd. for C₁₄H₂₂O₂: C, 75.63; H, 9.97. Found : C, 75.41; H, 10.05.

2-[2-(2-Hydroxyethyl)phenyl]-1-phenylethanol (**3h**)¹⁴: v_{max} (film) 3640-3080 cm⁻¹ (OH); $\delta_{\rm H}$ 2.57 (2H, br s, 2xOH), 2.74-2.90 (2H, m, CH₂CH₂OH), 2.94 (1H, dd, J=14.0, 4.6, ArHCH), 3.06 (1H, dd, J=14.0, 8.7, ArHCH), 3.66-3.79 (2H, m, CH₂OH), 4.83 (1H, dd, J=8.7, 4.6, CHOHPh), 7.11-7.34 (9H, m, ArH); $\delta_{\rm C}$ 35.3 (CH₂CH₂OH), 42.3 (ArCH₂), 63.3 (CH₂OH), 75.2 (CHOH), 125.7, 126.4, 126.7, 127.5, 128.3, 129.9, 130.4, 136.8, 137.5, 144.1 (ArC); *m/z* 165 (M⁺-C₆H₅, 3%), 136 (67), 118 (38), 117 (32), 115 (21), 107 (77), 106 (39), 105 (57), 104 (27), 103 (22), 91 (31), 79 (100), 78 (28), 77 (98), 51 (15).

 $\begin{array}{l} 1\-\[2-(2-Hydroxyethyl)phenyl]\-\[2-methyl-2-propanol\ (3i)\]^{14}\!: v_{max}\ (film)\ 3680\-\[3000 cm\]^1\ (OH);\ \delta_H\ 1.23\ (6H,\ s,\ 2xCH_3),\ 2.55\ (2H,\ br\ s,\ 2xOH),\ 2.85\ (2H,\ s,\ CH_2COH),\ 2.97\ (2H,\ t,\ J=6.7,\ CH_2CH_2OH),\ 3.80\ (2H,\ t,\ J=6.7,\ CH_2OH),\ 7.12\-\7.26\ (4H,\ m,\ ArH);\ \delta_C\ 29.6\ (2xCH_3),\ 35.5\ (ArCH_2),\ 45.2\ (CH_2COH),\ 63.2\ (CH_2OH),\ 71.4\ (COH),\ 125.9,\ 126.7,\ 129.7,\ 131.9,\ 136.4,\ 138.0\ (ArC);\ m/z\ 161\ (M^+\-CH_3\-H_2O,\ 5\%),\ (CH_2OH),\ 5\%\),\ 126.7\ (CH_2OH),\ 125.9\ (CH_$

143 (12), 136 (53), 118 (37), 117 (58), 115 (18), 106 (46), 105 (66), 104 (19), 103 (13), 91 (32), 79 (10), 78 (16), 77 (21), 65 (10), 59 (100), 43 (54), 41 (18).

2-Ethyl-1-[2-(2-hydroxyethyl)phenyl]-2-butanol (3j)¹⁴: v_{max} (film) 3640-3080 cm⁻¹ (OH); $\delta_{\rm H}$ 0.92 (6H, t, J=7.5, 2xCH₃), 1.46 (2H, q, J=7.5, 1xCH₃CH₂), 1.55 (2H, q, J=7.5, 1xCH₃CH₂), 1.90 (2H, br s, 2xOH), 2.83 (2H, s, ArCH₂), 3.00 (2H, t, J=6.5, CH₂CH₂OH), 3.85 (2H, t, J=6.5, CH₂OH), 7.15-7.26 (4H, m, ArH); $\delta_{\rm C}$ 8.0 (2xCH₃), 30.7 (2xCH₃CH₂), 35.6 (CH₂CH₂OH), 40.8 (ArCH₂), 63.4 (CH₂OH), 75.2 (COH), 125.9, 126.7, 129.7, 132.0, 136.1, 138.6 (ArC); *m/z* 193 (M⁺-C₂H₅, 3%), 175 (13), 136 (50), 129 (13), 118 (41), 117 (43), 115 (18), 106 (47), 105 (18), 104 (12), 91 (30), 87 (100), 79 (11), 78 (11), 77 (18), 69 (18), 57 (62), 45 (80), 43 (24), 41 (28).

1-{[2-(2-Hydroxyethyl)phenyl]methyl}cyclohexanol (31): mp 104-105°C (pentane/dichloromethane); v_{max} (KBr) 3600-3040 cm⁻¹ (OH); $\delta_{\rm H}$ 1.19-1.59 (10H, m, 5xring CH₂), 2.12 (2H, br s, 2xOH), 2.83 (2H, s, ArCH₂COH), 3.00 (2H, t, *J*=6.6, CH₂CH₂OH), 3.83 (2H, t, *J*=6.6, CH₂OH), 7.14-7.22 (4H, m, ArH); $\delta_{\rm C}$ 21.9, 25.7, 35.7, 37.6 (5xring CH₂, CH₂CH₂OH), 44.9 (CH₂COH), 63.3 (CH₂OH), 71.7 (COH), 125.8, 126.7, 129.7, 132.1, 135.8, 138.3 (ArC); *m/z* 136 (M⁺-C₆H₁₀O, 36%), 129 (15), 118 (35), 117 (42), 115 (20), 106 (55), 105 (35), 104 (30), 103 (17), 99 (100), 91 (35), 81 (84), 79 (21), 78 (18), 77 (26), 65 (12), 57 (13), 55 (47), 53 (14), 43 (34), 42 (16), 41 (57). Anal. Calcd. for C₁₅H₂₂O₂: C, 76.88; H, 9.46. Found : C, 76.11; H, 9.83.

 $\begin{array}{l} 1\-\[2-(2-Hydroxyethyl)phenyl]\-\2-phenyl\-\2-propanol\ (3m)\^{14}:\ v_{max}\ (film)\ 3660\-\3080\ cm\^{-1}\ (OH);\ \delta_{H}\ 1.59\ (3H, s, CH_3),\ 2.40\ (2H, br\ s,\ 2xOH),\ 2.84\ (2H,\ t,\ J=6.5,\ CH_2CH_2OH),\ 3.14\ (2H,\ s,\ CH_2COH),\ 3.78\ (2H,\ t,\ J=6.5,\ CH_2OH),\ 6.88\-\7.40\ (9H,\ m,\ ArH);\ \delta_{C}\ 29.6\ (CH_3),\ 35.3\ (CH_2CH_2OH),\ 46.2\ (CH_2COH),\ 63.4\ (CH_2OH),\ 75.0\ (COH),\ 124.9,\ 125.8,\ 126.6,\ 126.9,\ 128.0,\ 129.6,\ 131.9,\ 135.5,\ 138.3,\ 147.9\ (ArC);\ m/z\ 178\ (M^+\-C_6H_6,\ 1\%),\ 136\ (14),\ 121\ (58),\ 106\ (10),\ 105\ (13),\ 77\ (15),\ 43\ (100).\end{array}$

Preparation of Compound 3'c. - A solution of the hydroxyacid 3c (1mmol) in benzene (50 ml) in the presence of a catalytic amount of *p*-toluenesulfonic acid (0.001 g) was heated in a 120°C oil bath temperature at a Dean-Stark apparatus for 15 h, then the benzene was removed by distillation and the resulting residue was hydrolysed with a saturated solution of sodium carbonate and extracted with ethyl acetate. The organic layer was dried over anhydrous sodium sulfate and evaporated (15 mmHg). The residue was then purified by column chromatography (silica gel; hexane ethyl acetate) and recrystallised to yield the tittle compound 3'c (75% yield). 4,5-Dihydro-3-benzoxepin-2(1H)-one (3'c)¹⁵: mp 83-84°C (hexane/dichloromethane)[lit. mp 92°C]; v_{max} (KBr) 1720 cm⁻¹ (C=O); $\delta_{\rm H}$ 3.29 (2H, t, J=5.5, ArCH₂CH₂), 4.03 (2H, s, ArCH₂C=O), 4.60 (2H, t, J=5.5, CH₂O), 7.07-7.22 (4H, m, ArH); $\delta_{\rm C}$ 33.8 (ArCH₂CH₂), 40.3 (CH₂C=O), 65.7 (CH₂O), 126.9, 127.6, 128.5, 130.3, 130.8, 134.7 (ArC), 172.3 (C=O); *m/z* 162 (M⁺, 56%), 118 (42), 117 (100), 115 (22), 105 (13), 104 (53), 103 (23), 91 (17), 78 (18), 77 (15), 63 (11), 51 (14).

Treatment of Diols 3d-m with Phosphoric Acid. Isolation of Compounds 5-9. General Procedure.-To a solution of the corresponding diol 3d-m (1 mmol) in toluene (5 ml) was added 85% phosphoric acid (0.4 ml). The reaction mixture was heated at 110°C for 10 h, then the toluene was removed by distillation and the resulting residue was hydrolysed with water and extracted with ethyl acetate. The organic layer was dried over anhydrous sodium sulfate and evaporated (15 mmHg). The resulting residue was purified by column chromatography (silica gel; hexane ethyl acetate) to yield pure products 5-9. Yields and physical data of compounds 5h-m (R_f) are included in Table 2; analytical and spectroscopic data follow.

1,2,4,5-Tetrahydro-2-phenyl-3-benzoxepine (**5**h): v_{max} (film) 3040, 3000, 1600, 750, 690 cm⁻¹ (ArH); $\delta_{\rm H}$ 2.76 (1H, dd, J = 15.2, 4.6, HCHCH₂O), 2.90 (1H, d, J = 15.2, HCHCH₂O), 3.41 (1H, d, J = 15.2, ArHCH), 3.48 (1H, dd, J = 15.2, 9.8, ArHCH), 3.60-3.68 (1H, m, HCHO), 4.26-4.32 (1H, m, HCHO), 4.45 (1H, d, J = 9.8, CHOPh), 7.10-7.43 (9H, m, ArH); $\delta_{\rm C}$ 39.5 (CH₂CH₂O), 47.2 (CH₂CHOPh), 70.0 (CH₂O), 82.0 (CHO), 125.8, 126.4, 126.5, 127.4, 128.4, 129.3, 129.7, 140.1, 141.5, 143.7 (ArC); *m/z* 224 (M⁺, 9%), 118 (73), 117 (100), 115 (23), 105 (11), 91 (21), 78 (11), 77 (20), 51 (13) (Found: M⁺, 224.1196. C₁₆H₁₆O requires M, 224.1201).

1,2,4,5-Tetrahydro-2,2-dimethyl-3-benzoxepine (5i): v_{max} (film) 3040, 3000, 750 cm⁻¹ (ArH); $\delta_{\rm H}$ 1.16 (6H, s, 2xCH₃), 2.94 (2H, t, J=5.0, ArCH₂CH₂), 2.95 [2H, s, CH₂CO(CH₃)₂], 3.80 (2H, t, J=5.0, CH₂O), 6.98-7.15 (4H, m, ArH); $\delta_{\rm C}$ 26.8 (2xCH₃), 38.9 (ArCH₂CH₂), 49.3 (CH₂CO), 62.6 (CH₂O), 73.0 (CO), 126.2, 126.4, 128.7, 130.0, 138.8, 140.9 (ArC); m/z 176 (M⁺, 9%), 118 (34), 117 (100), 115 (23), 91 (22), 77 (12), 51 (12), 43 (34), 41 (11) (Found: M⁺, 176.1207. C₁₂H₁₆O requires M, 176.1201).

2,2-Diethyl-1,2,4,5-tetrahydro-3-benzoxepine (5j): v_{max} (film) 3020, 3000, 750 cm⁻¹ (ArH); $\delta_{\rm H}$ 0.84 (6H, t, J=7.5, 2xCH₃), 1.33-1.50 (4H, m, 2xCH₃CH₂), 2.92 (2H, s, ArCH₂), 2.93 (2H, t, J=4.9, CH₂CH₂O), 3.77 (2H, t, J=4.9, CH₂O), 7.01-7.24 (4H, m, ArH); $\delta_{\rm C}$ 7.5 (2xCH₃), 27.7 (2xCH₃CH₂), 38.9 (CH₂CH₂O), 45.6 (ArCH₂), 61.9 (CH₂O), 76.9 (CO), 126.1, 126.3, 128.7, 130.1, 138.5, 141.0 (ArC); *m*/z 204 (M⁺, 6%), 119 (16), 118 (75), 117 (100), 115 (13), 91 (15), 57 (12) (Found: M⁺, 204.1504. C₁₄H₂₀O requires M, 204.1514).

Spirocyclopentane-2-[1,2,4,5-tetrahydro-3-benzoxepine] (5k): v_{max} (film) 3040, 3000, 750 cm⁻¹ (ArH); $\delta_{\rm H}$ 1.43-1.74 (8H, m, 4xring CH₂), 2.95 (2H, t, *J*=4.8, CH₂CH₂O), 3.01 (2H, s, ArCH₂), 3.75 (2H, t, *J*=4.8, CH₂O), 7.00-7.23 (4H, m, ArH); $\delta_{\rm C}$ 23.3, 36.6, 38.9, 47.3 (4xring CH₂, 2xArCH₂), 63.5 (CH₂O), 84.9 (CO), 126.1, 126.3, 128.7, 129.9, 139.2, 141.1 (ArC); *m/z* 202 (M⁺, 5%), 119 (14), 118 (55), 117 (100), 115 (41), 103 (11), 91 (29), 78 (11), 67 (10), 55 (15), 41 (13) (Found: M⁺, 202.1351. C₁₄H₁₈O requires M, 202.1358).

 $\begin{array}{l} \textit{Spirocyclohexane-2-[1,2,4,5-tetrahydro-3-benzoxepine] (51): v_{max} (film) 3040, 3000, 750 cm^{-1}$ (ArH); δ_{H} 1.23-1.62 (10H, m, 5xring CH_2), 2.91 (2H, s, ArCH_2), 2.93-2.96 (2H, m, CH_2CH_2O), 3.76-3.79 (2H, m, CH_2O), 6.99-7.13 (4H, m, ArH); δ_{C} 21.7, 25.9, 34.8, 39.0 (5xring CH_2, CH_2CH_2O), 48.2 (ArCH_2CO), 61.4 (CH_2O), 73.3 (CO), 126.1, 126.3, 128.7, 130.0, 138.3, 140.9 (ArC); m/z 216 (M⁺, 19%), 119 (27), 118 (100), 117 (96), 115 (24), 91 (18), 41 (12) (Found: M⁺, 216.1510. C_{15}H_{20}O requires M, 216.1514). \end{array}$

 $\begin{array}{l} 1,2,4,5\text{-}Tetrahydro-2\text{-}methyl-2\text{-}phenyl-3\text{-}benzoxepine} \ (\textbf{5m}): \nu_{max} \ (film) \ 3020, \ 3000, \ 1590, \ 750, \ 700 \ cm^{-1} \\ (ArH); \ \delta_{H} \ 1.40 \ (3H, \ s, \ CH_{3}), \ 2.99\text{-}3.03 \ (2H, \ m, \ CH_{2}CH_{2}O), \ 3.22 \ (1H, \ d, \ \textit{J=14.8}, \ \textit{HCHCO}), \ 3.50 \ (1H, \ d, \ \textit{J=14.8}, \ \textit{HCHCO}), \ 3.79\text{-}3.97 \ (2H, \ m, \ CH_{2}O), \ 7.01\text{-}7.51 \ (9H, \ m, \ ArH); \ \delta_{C} \ 29.1 \ (CH_{3}), \ 38.1 \\ (CH_{2}CH_{2}O), \ 47.9 \ (CH_{2}CO), \ 62.7 \ (CH_{2}O), \ 77.4 \ (CO), \ 125.8, \ 126.1, \ 126.5, \ 128.1, \ 128.9, \ 130.8, \ 137.7, \ 140.6, \ 146.4 \ (ArC); \ m/z \ 238 \ (M^{+}, \ 5\%), \ 118 \ (49), \ 117 \ (100), \ 115 \ (19), \ 115 \ (19), \ 91 \ (19), \ 78 \ (12), \ 77 \ (17), \ 51 \ (12), \ 43 \ (17) \ (Found: \ M^{+}, \ 238.1364. \ C_{17}H_{18}O \ requires \ M, \ 238.1358). \end{array}$

3,4-Dihydro-4-propyl-1H-2-benzopyran (6d): (49% yield) v_{max} (film) 3020, 3000, 740 cm⁻¹ (ArH); $\delta_{\rm H}$ 0.96 (3H, t, J=7.3, CH₃), 1.49 (2H, sextet, J=7.3, CH₃CH₂), 1.73-1.90 (2H, m, ArCHCH₂), 2.68 (1H, dt, J=16.2, 3.8, HCHCH₂O), 2.97 (1H, ddd, J=16.2, 9.5, 5.4, HCHCH₂O), 3.76 (1H, ddd, J=11.2, 9.5, 3.8, CH₂HCHO), 4.13 (1H, ddd, J=11.2, 5.4, 3.8, CH₂HCHO), 4.75 (1H, dd, J=7.9, 3.5, CHO), 7.05-7.19 (4H, m, ArH); $\delta_{\rm C}$ 14.1 (CH₃), 18.5 (CH₃CH₂), 29.2 (CHCH₂), 38.1 (ArCH₂), 63.1 (CH₂O), 75.6

(CHO), 124.8, 126.0, 126.0, 128.8, 133.9, 138.6 (ArC); m/z 176 (M⁺, 2%), 133 (100), 115 (12), 105 (15), 103 (10), 79 (10), 77 (17), 43 (13), 41 (22) (Found: M⁺, 175.8951. C₁₂H₁₆O requires M, 176.1201).

2,3-Dihydro-4-(2-hydroxyethyl)-1,1-dimethyl-1H-indene (7e): (76% yield) v_{max} (film) 3680-3080 cm⁻¹ (OH); $\delta_{\rm H}$ 1.25 (6H, s, 2xCH₃), 1.74 (1H, br s, OH), 1.92 [2H, d, J=7.2, C(CH₃)₂CH₂], 2.82-2.88 (4H, m, 2x ArCH₂), 3.82 (2H, t, J=7.2, CH₂OH), 6.98-7.14 (3H, m, ArH); $\delta_{\rm C}$ 28.4 [C(CH₃)₂CH₂], 28.7 (2xCH₃), 36.6, 40.9 (2xArCH₂), 44.0 [C(CH₃)₂], 62.5 (CH₂OH), 120.2, 126.8, 126.9, 134.0, 141.7, 152.9 (ArC); *m*/z 190 (M⁺, 42%), 176 (21), 175 (100), 159 (14), 157 (54), 143 (12), 142 (24), 141 (17), 130 (11), 129 (61), 128 (51), 127 (16), 117 (10), 115 (16), 91 (20), 77 (17), 65 (11), 63 (13), 51 (11), 41 (17) (Found: M⁺, 190.1356. C₁₃H₁₈O requires M, 190.1358).

2,3-Dihydro-4-(2-hydroxyethyl)-1,1,2-trimethyl-1H-indene (7g): (86% yield) v_{max} (film) 3600-3080 cm⁻¹ (OH); $\delta_{\rm H}$ 0.95 (3H, s, CH₃CCH₃), 1.06 (3H, d, J=6.9, CH₃CH), 1.26 (3H, s, CH₃CHCH₃), 2.03-2.15 (1H, m, CH₃CH), 2.45 (1H, dd, J=15.5, 10.1, ArHCHCHCH₃), 2.83 (2H, t, J=6.8, CH₂CH₂OH), 2.96 (1H, dd, J= 15.5, 7.5, ArHCHCHCHC₃), 2.97 (1H, br s, OH), 3.81 (2H, t, J=6.8, CH₂OH), 6.97-7.13 (3H, m, ArH); $\delta_{\rm C}$ 13.9 (CH₃CH), 23.1, 26.6 [(CH₃)₂C], 36.6 (CH₂CH₂OH), 36.8 (ArCH₂), 45.0 (CH₃CH), 45.4 [(CH₃)₂C], 62.6 (CH₂OH), 120.5, 126.7, 126.8, 133.7, 140.9, 153.7 (ArC); *m/z* 204 (M⁺, 33%), 189 (66), 172 (19), 171 (82), 157 (10), 156 (27), 143 (67), 142 (16), 141 (42), 130 (11), 129 (44), 128 (87), 127 (19), 115 (67), 105 (22), 91 (36), 79 (14), 78 (11), 77 (44), 65 (22), 63 (23), 55 (23), 53 (21), 52 (13), 51 (37), 45 (11), 43 (64), 41 (100) (Found: M⁺, 204.1510. C₁₄H₂₀O requires M, 204.1514).

1,2,3,4-Tetrahydro-5-(2-hydroxyethyl)-1,1-dimethylnaphthalene (**8f**): (44% yield) v_{max} (film) 3640-3080 cm⁻¹ (OH); $\delta_{\rm H}$ 1.28 (6H, s, 2xCH₃), 1.63 [2H, t, J=5.8, C(CH₃)₂CH₂], 1.70 (1H, br s, OH), 1.73-1.86 (2H, m, CH₂CH₂CH₂Ar), 2.71 (2H, t, J=6.4, ArCH₂), 2.85 (2H, t, J=6.9, CH₂CH₂OH), 3.81 (2H, t, J=6.9, CH₂OH), 6.97 (1H, dd, J= 7.4, 1.2, 1xArH), 7.10 (1H, t, J=7.4, 1xArH), 7.25 (1H, dd, J= 7.4, 1.2, 1xArH); $\delta_{\rm C}$ 19.5 (CH₂CH₂CH₂CH₂Ar), 27.3 [C(CH₃)₂CH₂], 32.0 (2xCH₃), 34.1 [C(CH₃)₂], 36.3, 38.7 (2xArCH₂), 62.4 (CH₂OH), 125.3, 125.5, 126.7, 134.6, 135.9, 146.3 (ArC); *m*/z 204 (M⁺, 41%), 190 (10), 189 (75), 173 (13), 172 (13), 171 (100), 159 (10), 156 (10), 148 (10), 143 (29), 142 (17), 141 (26), 131 (26), 130 (13), 129 (54), 128 (63), 127 (12), 119 (14), 117 (17), 115 (42), 105 (16) (Found: M⁺, 204.1507. C₁₄H₂₀O requires M, 204.1514).

(*E*)-2-[2-(4-Methyl-2-pentenyl)phenyl]ethanol (**9f**): (22% yield) v_{max} (film) 3620-3080 cm⁻¹ (OH); $\delta_{\rm H}$ 0.97 (6H, d, *J*=6.7, 2xCH₃), 1.47 (1H, br s, OH), 2.13-2.35 [1H, m, (CH₃)₂CH], 2.91 (2H, t, *J*=6.9, CH₂CH₂OH), 3.36 (2H, d, *J*=5.6, CH₂CH=CH), 3.83 (2H, t, *J*=6.9, CH₂OH), 5.40 (1H, dd, *J*=15.5, 6.0, CH₂CH=CH), 5.50 (1H, dt, *J*=15.5, 5.6, CH₂CH=CH), 7.14-7.23 (4H, m, ArH); $\delta_{\rm C}$ 22.5 (2xCH₃), 31.0 [(CH₃)₂CH], 35.8, 35.9 (2xArCH₂), 63.2 (CH₂OH), 125.5, 126.2, 126.7, 129.8, 129.9, 136.3, 139.1, 139.3 (ArC, CH=CH); *m*/z 204 (M⁺, 17%), 171 (13), 144 (13), 143 (100), 141 (10), 135 (15), 133 (12), 130 (21), 129 (37), 128 (51), 127 (14), 119 (17), 117 (63), 116 (13), 115 (59), 105 (23), 104 (17), 103 (12), 91 (43), 78 (15), 77 (19), 69 (17), 65 (17), 63 (11), 55 (15), 53 (11), 51 (15), 43 (20), 41 (56) (Found: M⁺, 204.1516. C₁₄H₂₀O requires M, 204.1514).

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