Base-promoted acetal formation employing aryl salicylates

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Dedicated to Professor T. R. Govindachari on the occasion of his 85th birthday
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Abstract
New examples of base-promoted acetal formation in the reactions of aryl salicylate and salicylate-type esters 1 to 3 with aldehydes are reported.

Keywords: Oxo-acetals, aldehydes, salicylate, DABCO

Introduction
Recently we reported that aryl esters of 2-hydroxybenzoic acids (i.e. salicylates) form cyclic acetals with a variety of aldehydes (scheme 1) under base-catalyzed conditions. Key aspects of the process were that the reactions worked best under neat conditions or, where necessary, with minimal solvent to effect dissolution. This is in contrast to most methods for acetal formation which usually employ acid-catalyzed conditions.

Scheme 1

Herein we provide further examples of this process as well full experimental details for these new cases as well as the first successful reaction with an aromatic aldehyde.

Results and Discussion

Oxo-acetal formations of esters 1 to 3 were investigated. The structures of the starting materials and products are shown in Figure 1. Most reactions were carried out neat except where either, or both, the aldehyde and salicylate were solids. In those cases a very small volume of chloroform
was added to effect solution. The ratio of reactants was salicylate: aldehyde: base = 1:5:1.

![Chemical structures](image)

Figure 1

As before aliphatic aldehydes, solid paraformaldehyde and chloral all reacted well with esters 1 to 3. In our previous report we noted that the use of aromatic aldehydes did not lead to successful reaction. Interestingly we have now discovered one exception to this, namely reaction of phenyl 4-methoxysalicylate (1). Reaction with benzoaldehyde under standard conditions gave oxo-acetal 4e in 49% yield. Only in one case, that of salicylate 5 was any product formed in a reaction with an unsaturated aldehyde (cinnamaldehyde). The yield was very low (8%). Hence it can be concluded that, in general, unsaturated aldehydes (including aromatic aldehydes) are not usually good substrates for this process.

A brief investigation was carried out to examine whether any diastereoselectivity would occur in reactions with the chiral, non-racemic aldehyde, D-glyceraldehyde acetonide. Hence 2 was reacted with D-glyceraldehyde acetonide. This provided an essentially 1:1 diastereomeric mixture of acetals 7. Unfortunately this mixture proved to be inseparable.

![Chemical structures](image)

Similar reaction of 3 led to another inseparable 1:1 mixture of acetals 8.
In conclusion, this facile process represents one of the very few base-promoted methods for acetal formation. In this report, we have successfully extended the range of salicylate and salicylate-type esters which undergo this reaction.

**Experimental Section**

**General Procedures.** Melting points were determined on a Kofler hotstage and are uncorrected. Elemental microanalyses were performed by the Australian Microanalytical Service, National Analytical Laboratories, Melbourne and the University of Otago, Dunedin, New Zealand. Optical rotations were recorded on a Perkin Elmer Model 141 Polarimeter. Infrared (IR) spectra were recorded on a Perkin-Elmer 1600 Series Fourier Transform spectrophotometer (cm\(^{-1}\) scale) and refer to thin films of liquids (neat) or paraffin (Nujol) mulls of solids between NaCl plates. High resolution hydrogen-1 nuclear magnetic resonance (\(^1\)H NMR) spectra were recorded at 300 MHz on a Bruker DPX-300 spectrometer or 400 MHz on a Bruker Avance DRX 400 spectrometer. The \(^1\)H NMR spectral data refer to deuteriochloroform solutions (CDCl\(_3\)) using tetramethysilsilane (TMS) as internal reference (\(\delta\) 0.00 ppm). Carbon-13 nuclear magnetic resonance (\(^{13}\)C NMR) spectra were recorded at 75 MHz on a Bruker APX-300 spectrometer or 100 MHz on a Bruker Avance DRX 400 spectrometer. Mass spectrometry (ESI) was performed using samples in methanol on a Micromass Platform QMS Electrospray mass spectrometer. High resolution mass spectra (HRMS) for accurate mass determinations were recorded on a Bruker BioApex 47e FTMS fitted with an Analytica electrospray source using NaI for accurate mass calibration (accuracy ± 3 ppm). Low resolution mass spectra were recorded on a VG micromass 70/70F or a VG TRIO-1 mass spectrometer with an ion source temperature of 200 °C and electron impact energy of 70 eV. Esters 2 and 3 were purchased from Aldrich Chemical Company.

**General procedure for preparation of cyclic oxo-acetals**

Unless otherwise indicated, a solution of salicylate ester, aldehyde (5 mol equivalent) and base (1.0 mol equivalent) was stirred at room temperature and allowed to react until TLC analysis indicated complete consumption of starting material. In those cases where one of the starting materials was a solid, and hence the reaction mixture was heterogeneous, a small amount of
chloroform (usually about 0.2 mL) was added to effect dissolution. The reaction mixture was then dissolved in diethyl ether (20 mL) and the ethereal solution was then washed with water (3x5 ml). The organic layer was dried (MgSO₄), filtered and concentrated under reduced pressure. Purification of the crude product was carried out either by recrystallization, flash column chromatography or preparative TLC (eluant 1:4, ethyl acetate/light petroleum). Most reactions were carried out using 500mg of the salicylate ester in each case. In the following only reaction times, chromatography details (eluant, Rf), isolated yield, physical constants and spectroscopic data are given.

**2-Hydroxy-4-methoxybenzoic acid**. To a solution of (15.4 g, 0.1 mol) of 2,4-dihydroxybenzoic acid in 20% NaOH (50 mL) was added (10.41 mL, 0.11 mol) of dimethyl sulfate. The resulting orange solution was stirred at room temperature for 20 hours before being neutralized with concentrated HCl. The mixture was extracted with ether (3x100 ml) and the aqueous layer was acidified to pH~1 with concentrated HCl, resulting in white precipitate. The precipitate was collected by filtration, washed with water and dried (under vacuum). Recrystallization from ethanol and water afforded (9.7 g, 58%) of product as cream solid. m.p. 145-146 °C (lit.⁵ 157-159 °C). IR (Nujol) 3050-2554v, 1649s, 1624s, 1576m, 1504m, 1465m, 1436s, 1348m, 1360m, 1316m, 1261m, 1250m, 1226m, 1205m, 1176m, 1150m, 1095m, 1025m, 966m, 962m cm⁻¹; ¹H NMR (200 MHz, CDCl₃) δ10.69 (s, 1H, OH), 7.82 (d, J=9.26 Hz, 1H, H5), 6.49 (d, 1H, H6), 6.47 (s, 1H, H3), 3.85 (s, CH₃); ¹³C NMR (50 MHz, CDCl₃) δ172.5 (CO), 166.9 (C-Ar), 165.3 (C-Ar), 132.6 (C-Ar), 107.9 (C5), 106.0 (C6), 101.5 (C3), 55.9 (CH₃).

**Phenyl 2-hydroxy-4-methoxybenzoate (1)**. 2-Hydroxy-4-methoxybenzoic acid (3.00 g, 17.86 mmol), phenol (1.68 g, 17.86 mmol) and POCl₃ (2.34 ml) were heated at 120-130 °C for 45 minutes. The mixture was dissolved in diethyl ether and then extracted with water. The ether layer was dried over MgSO₄ and evaporated in vacuo to gave a red oil. The crude material was purified by flash column chromatography (1:4, ethyl acetate/light petroleum). Fractions containing material of Rf=0.46 were combined and concentrated to afford the product as white solid (4.28 g, 98%). m.p. 41-42°. IR (Nujol) 2924.5s, 2854.0s, 1686.2m, 1625.0m, 1589.8m, 1459.1s, 1376.6s, 1273.9m, 1254.2m, 1194.2m, 1163.4m, 1128.2m, 1070.7m, 1030.1m, 963.2m, 856.7m, 836.4m, 773.8m, 744, 721.8m, 687.8m cm⁻¹; ¹H NMR (200 MHz, CDCl₃) δ10.73 (s, 1H, OH), 7.96 (d, J=0.84 Hz, 1H, H5), 7.23 (m, 5H, Ph), 6.55 (d, J=8.45 Hz, 1H, H6), 6.51 (s, 1H, H3), 3.87 (s, 3H, CH₃); ¹³C NMR (50 MHz, CDCl₃) δ172.5 (CO), 166.9 (C-Ar), 165.3 (C-Ar), 132.6 (C-Ar), 107.9 (C5), 106.0 (C6), 101.5 (C3), 55.9 (CH₃); MS m/z 245.14 (M+H)+.

**7-Methoxy-4-oxo-1,3-benzodioxane (4a)**. 6 d, (Rf=0.25), 79%, colourless crystals (diethyl ether), m.p. 67-68 °C. IR (Nujol) 2922.8s, 2854.0s, 2358.4m, 1742.3m, 1624.6m, 1458.5s, 1376.8m, 1254.8m, 1180.0w, 1110.6w, 1063.6w, 1014.4w, 982.1w, 929.0w, 832.5w, 739.7w, 668.0w, 667.9w cm⁻¹; ¹H NMR (200 MHz, CDCl₃) δ7.9 (d, J=8.87 Hz, 1H, H5), 7.23 (m, 5H, Ph), 6.55 (d, J=8.45 Hz, 1H, H6), 6.51 (s, 1H, H3), 3.87 (s, 3H, CH₃); ¹³C NMR (50 MHz, CDCl₃) δ168.7 (CO), 166.2 (C-Ar), 164.4 (C-Ar), 150.1 (C-Ar), 131.7 (C5), 129.5 (2xCH-Ph), 126.2 (C6), 121.7 (2xCH-Ph), 108.0 (C3), 104.8 (C-Ph), 100.8 (CH-Ph), 55.6 (CH₃); MS m/z 245.14 (M+H)+.
2-Methyl-7-methoxy 4-oxo-1,3-benzodioxane (4b). 2 d, (Rf=0.16) (94%), white crystals (diethyl ether), m.p. 59-60 °C. IR (Nujol) 3436.0b, 2923.7s, 2853.5s, 1741.1m, 1616.4m, 1460.1s, 1376.9s, 1263.0m, 1094.0m, 1025.6m, 929.4w, 847.9w, 767.9w, 734.4w, 688.2w cm⁻¹; ¹H NMR (200 MHz, CDCl₃) δ7.89 (d, J=8.87 Hz, 1H, H5), 6.71 (dd, J=2.39, 6.44 Hz, 1H, H6), 6.48 (d, J=2.38 Hz, 1H, H8), 5.73 (q, J=5.17 Hz, 1H, H2), 3.86 (s, 3H, CH₃O), 1.73 (d, J=5.17 Hz, 3H, CH₃); ¹³C NMR (50 MHz, CDCl₃) δ166.1 (CO), 162.2 (C-Ar), 160.3 (C-Ar), 131.7 (C5), 111.2 (C6), 106.9 (C-Ar), 100.2 (C8), 98.9 (C2), 55.8 (CH₃O), 19.9 (CH₃); HRMS calc’d for C₉H₈O₄: m/z 180.0423, found 180.0421; MS m/z 180 (58), 150 (100), 122 (66).

2-Ethyl-7-methoxy 4-oxo-1,3-benzodioxane (4c). 28 h, (Rf=0.25), (86%), yellow oil. IR (neat) 3458.6w, 3079.7w, 2979.0s, 2942.1s, 2844.1m, 2364.1w, 2017.7w, 1744.1s, 1614.7s, 1584.9s, 1501.0s, 1451.3s, 1395.1s, 1332.1m, 1258.8s, 1201.6s, 1169.9s, 1133.9s, 1088.6s, 1050.7s, 1022.9s, 985.8w, 946.2w, 880.4w, 769.9s, 690.0s, 652.2s, 649.3s, 622.6s, 573.1s cm⁻¹; ¹H NMR (200 MHz, CDCl₃) δ7.9 (d, J=8.83 Hz, 1H, H5), 6.71 (dd, J=2.39, 6.33 Hz, 1H, H6), 6.50 (d, J=2.37 Hz, 1H, H8), 5.56 (m, 1H, H2), 3.87 (s, 3H, CH₃O), 2.04 (m, 2H, CH₂), 1.4 (t, J=7.46, 7.54 Hz, 3H, CH₃); ¹³C NMR (50 MHz, CDCl₃) δ165.9 (CO), 162.1 (C-Ar), 160.1 (C-Ar), 131.4 (C5), 110.9 (C6), 106.7 (C-Ar), 101.9 (C8), 100.0 (C2), 55.6 (CH₃O), 26.6 (CH₂), 6.9 (CH₃); HRMS calc’d for C₁₀H₁₀O₄: m/z 194.0579, found 194.0562; MS m/z 195.06 (M+H)+.

2-Isopropyl-7-methoxy 4-oxo-1,3-benzodioxane (4d). 4 d, (Rf=0.27), (87%), yellow oil. IR (neat) 3462.2w, 2972.3m, 2844.0m, 2362.3m, 1742.8s, 1618.3s, 1584.9s, 1501.0s, 1451.3s, 1395.1s, 1332.1m, 1258.8s, 1201.6s, 1169.9s, 1133.9s, 1088.6s, 1050.7s, 1022.9s, 985.8w, 946.2w, 880.4w, 769.9s, 690.0s, 665.2w, 649.3m, 611.6w, 573.1s cm⁻¹; ¹H NMR (200 MHz, CDCl₃) δ7.89 (d, J=8.81 Hz, 1H, H5), 6.72 (d, J=2.39 Hz, 1H, H6), 6.50 (d, J=2.37 Hz, 1H, H8), 5.36 (d, J=4.22 Hz, 1H, H2), 3.86 (s, 3H, CH₃O), 2.39 (m, 1H, CH), 1.14 (d, J=6.89 Hz, 6H, 2xCH₃); ¹³C NMR (50 MHz, CDCl₃) δ165.9 (CO), 162.3 (C-Ar), 160.4 (C-Ar), 131.4 (C5), 111.0 (C6), 106.8 (C-Ar), 104.4 (C8), 100.1 (C2), 55.6 (CH₃O), 31.6 (C-iPr), 15.9 (2xCH₃); HRMS calc. for C₁₂H₁₂O₄: m/z 222.0892, found 222.0897; MS m/z 220.09 (M+H)+.

2-Phenyl-7-methoxy 4-oxo-1,3-benzodioxane (4e). 5 d, 60 °C, (Rf=0.25), (90%), brown solid, m.p. 75-76 °C. IR (Nujol) 2920.5s, 2725.8w, 1759.8m, 1726.0m, 16180.3m, 1582.2m, 1463.0s, 1376.9s, 1315.0m, 1260.4m, 1197.7m, 1165.0m, 1074.1m, 1016.1m, 948.4m, 889.1m, 833.7m, 794.7w, 722.4m, 697.0w, 668.0w cm⁻¹; ¹H NMR (200 MHz, CDCl₃) δ7.89 (d, J=8.81 Hz, 1H, H5), 6.72 (dd, J=2.39 Hz, 1H, H6), 6.67 (d, J=2.39 Hz, 1H, H8), 5.36 (d, J=4.22 Hz, 1H, H2), 3.86 (s, 3H, CH₃O), 2.39 (m, 1H, CH), 1.14 (d, J=6.89 Hz, 6H, 2xCH₃); ¹³C NMR (50 MHz, CDCl₃) δ165.9 (CO), 162.3 (C-Ar), 160.3 (C-Ar), 131.4 (C5), 111.0 (C6), 106.8 (C-Ar), 104.4 (C8), 100.1 (C2), 55.6 (CH₃O), 31.6 (C-Pr), 15.9 (2xCH₃); HRMS calc. for C₁₅H₁₂O₄: m/z 256.0736, found 256.0739; MS m/z 256.0736, found 256.0739; MS m/z 262 (M+Na)+.
1376.9s, 1345.3m, 1277.6w, 1257.4m, 1155.9w, 1124.4w, 722.2m cm⁻¹; ¹H NMR (300 MHz, CDCl₃) δ 8.66 (s, 1H, H5), 7.95 (d, J=8.28 Hz, 1H, H6), 7.81 (d, J=8.32 Hz, 1H, H9), 7.62 (t, J=1.17, 1.27 Hz, 1H, H7), 7.48 (t, J=1.14 Hz, 1H, H8), 7.46 (s, 1H, H10), 5.74 (s, 2H, CH₂); ¹³C NMR (50 MHz, CDCl₃) δ 161.7 (CO), 153.3 (C-Ar), 137.3 (C-Ar), 133.0 (C5), 129.8 (C6), 129.6 (C9), 129.4 (C-Ar), 127.0 (C7), 125.7 (C8), 115.1 (C-Ar), 112.3 (C10), 91.4 (C2); MS m/z 200 (51), 170 (100), 142 (89), 114 (74); Anal. Calc. for C₁₂H₈O₃: C, 71.99. H, 4.03. Found: C, 71.60; H, 3.81.

2-Methyl 4-oxo-1,3-(b)-naphthodioxane (5b). 2 d, rt, K₂CO₃ in place of DABCO, (Rf=0.27), (14%), white solid, m.p. 90-91 °C. IR (Nujol) 2923.1s, 2854.1s, 1726.4m, 1636.1m, 1608.1m, 1504.4m, 1463.7s, 1270.6s, 1255.0s, 1215.3s, 1135.9m, 1084.9m, 1029.4s, 934.0s, 881.8s, 871.7s, 754.1s, 720.2m cm⁻¹; ¹H NMR (300 MHz, CDCl₃) δ 8.63 (s, 1H, H5), 7.92 (d, J=8.3 Hz, 1H, H6), 7.78 (d, J=8.3 Hz, 1H, H9), 7.60 (t, J=1.21 Hz, 1H, H8), 7.40 (s, 1H, H10), 5.8 (m, 1H, H2), 1.8 (d, J=5.2 Hz, 3H, CH₃); ¹³C NMR (50 MHz, CDCl₃) δ 162.4 (CO), 153.1 (C-Ar), 137.2 (C-Ar), 132.7 (C5), 129.6 (C6), 129.5 (C9), 129.2 (C-Ar), 127.0 (C7), 125.5 (C8), 114.5 (C-Ar), 112.0 (C10), 99.2 (C2), 20.1 (CH₃); MS m/z 214 (100), 171 (77), 143 (47), 142 (84), 115 (39), 114 (67), 113 (74), 88 (46), 87 (27), 86 (19), 76 (11), 75 (77), 74 (23), 71 (24), 63 (49), 62 (29), 57 (13).

2-Ethyl 4-oxo-[1,3-b]-naphthodioxane (5c). 2 d, (Rf=0.35), (75%), colourless crystals (diethyl ether), m.p. 81-82 °C. IR (Nujol) 2851.1s, 2725.4w, 1762.2w, 1728.1m, 1637.4m, 1607.2w, 1460.3s, 1377.1s, 1310.2w, 1244.4m, 1207.1m, 1148.6w, 1033.0w, 947.8m, 896.4w, 873.3w, 846.7w, 817.7w, 779.3w, 722.2m cm⁻¹; ¹H NMR (300 MHz, CDCl₃) δ 8.63 (s, 1H, H5), 7.92 (d, J=8.3 Hz, 1H, H6), 7.78 (d, J=8.3 Hz, 1H, H9), 7.60 (t, J=1.22 Hz, 1H, H8), 7.40 (s, 1H, H10), 5.64 (t, J=4.92 Hz, 1H, CH), 2.12 (m, J=4.94 Hz, 2H, CH₂), 1.18 (t, J=6.92 Hz, 6H, 2xCH₃); ¹³C NMR (50 MHz, CDCl₃) δ 162.6 (CO), 153.3 (C-Ar), 137.2 (C-Ar), 132.7 (C5), 129.6 (C6), 129.5 (C9), 129.2 (C-Ar), 127.0 (C7), 114.5 (C-Ar), 112.0 (C10), 99.2 (C2), 20.1 (CH₃); MS m/z 228 (18), 170 (100), 142 (54), 114 (53), 113 (20); Anal. Calc. for C₁₄H₁₂O₃: C, 73.60. H, 5.30. Found: C, 73.13; H, 5.23.

2-Isopropyl 4-oxo-[1,3-b]-naphthodioxane (5d). 3 d, (Rf=0.42), (87%), white crystals (diethyl ether), m.p. 58-59 °C. IR (Nujol) 2923.9s, 2854.0s, 2280.6w, 1760.5m, 1639.5m, 1609.4w, 1460.3s, 1377.1s, 1344.3m, 1286.3m, 1245.5m, 1209.6m, 1135.8w, 1062.1m, 986.3m, 961.5m, 917.4m, 875.9m, 831.0w, 807.5w, 776.4m, 745.9m, 722.6m cm⁻¹; ¹H NMR (300 MHz, CDCl₃) δ 8.63 (s, 1H, H5), 7.92 (d, J=8.32 Hz, 1H, H6), 7.68 (d, J=8.39 Hz, 1H, H9), 7.59 (t, J=1.22 Hz, 1H, H7), 7.46 (t, J=1.22, 6.78 Hz, 1H, H8), 7.42 (s, 1H, H10), 5.45 (d, J=4.30 Hz, 2H, CH₂), 1.18 (t, J=6.92 Hz, 6H, 2xCH₃); ¹³C NMR (50 MHz, CDCl₃) δ 162.8 (CO), 153.4 (C-Ar), 137.4 (C-Ar), 132.7 (C5), 129.6 (C6&C9), 129.3 (C-Ar), 127.0 (C7), 125.5 (C8), 114.8 (C-Ar), 112.2 (C10), 104.9 (C2), 31.9 (CH₃), 16.1 (2xCH₃); MS m/z 242 (12), 171 (16), 170 (100), 142 (48), 114 (24); Anal. Calc. for C₁₅H₁₄O₃: C, 74.36. H, 5.82. Found: C, 74.42; H, 5.64.

2-Butyl 4-oxo-[1,3-b]-naphthodioxane (5e). 3 d, (Rf=0.39), (83%), colourless crystals (diethyl
ether), m.p. 85-86 °C. IR (Nujol) 2850.8s, 2724.7w, 1762.6m, 1637.4m, 1608.6w, 1462.8s, 1377.2s, 1270.9m, 1249.8m, 1202.0m, 1137.8w, 1087.5w, 1004.4w, 970.9m, 914.0w, 791.2w, 770.2m, 722.2s cm⁻¹; H NMR (200 MHz, CDCl₃) δ 8.63 (s, 1H, H₅), 7.93 (d, J=7.59 Hz, 1H, H₆), 7.62 (t, J=1.28, 1.30 Hz, 1H, H₇), 7.46 (d, J=5.50 Hz, 1H, H₈), 7.41 (s, 1H, H₁₀), 5.68 (t, J=5.07, 5.01 Hz, 1H, H₂), 2.09 (m, 2H, CH₂), 1.60 (m, 2H, CH₂), 1.43 (m, 2H, CH₂), 0.99 (t, J=7.08 Hz, 3H, CH₃); 13C NMR (50 MHz, CDCl₃) δ 162.5 (CO), 153.2 (C-Ar), 137.2 (C-Ar), 132.6 (C₅), 129.6 (C₆), 129.5 (C₉), 126.9 (C₇), 125.4 (C₈), 114.7 (C-Ar), 112.1 (C₁₀), 102.5 (C₂), 33.4 (CH₂), 24.9 (CH₂), 13.8 (CH₃); MS m/z 256 (14), 171 (17), 170 (100), 142 (56), 114 (50), 115 (20), 113 (15); Anal. Calc. for C₁₆H₁₆O₃: C, 74.98. H, 6.29. Found: C, 73.79; H, 6.97.

2-Trichloromethyl 4-oxo-[1,3-b]-naphthodioxane (62f). 0.5 h, (Rf=0.30), (45%), pale yellow crystals (diethyl ether), m.p. 179-180 °C. IR (Nujol) 3433.3b, 2921.5s, 1753.6m, 1632.1m, 1609.0w, 1462.6s, 1377.1s, 1341.5m, 1270.0w, 1241.4m, 1203.9m, 1141.5m, 1099.4w, 1059.8w, 1015.3m, 910.5w, 882.2w, 864.4w, 835.4m, 765.9m, 722.2m cm⁻¹; H NMR (300 MHz, CDCl₃) δ 8.66 (s, 1H, H₅), 7.96 (d, J=8.34 Hz, 1H, H₆), 7.83 (d, J=8.25 Hz, 1H, H₉), 7.65 (t, J=6.97, 7.07 Hz, 1H, H₇), 7.58 (s, 1H, H₁₀), 7.52 (t, J=8.10, 7.02 Hz, 1H, H₈), 5.95 (s, 1H, H₂); 13C NMR (50 MHz, CDCl₃) δ 159.2 (CO), 150.9 (C-Ar), 137.5 (C-Ar), 133.1 (C₅), 130.3 (C₆), 129.7 (C₉), 129.7 (C-Ar), 127.2 (C₇), 126.3 (C₈), 113.3 (C-Ar), 112.9 (C₁₀), 100.9 (C₂), 94.4 (C₆Cl); MS m/z 316 (9), 200 (9), 199 (65), 171 (33), 170 (100), 142 (74), 115 (37), 114 (81), 113 (35), 88 (22), 86 (11), 63 (23); Anal. Calc. for C₁₃H₇O₃Cl₃: C, 49.17. H, 2.22. Found: C, 49.38; H, 2.21.

2-Cinnamal 4-oxo-[1,3-b]-naphthodioxane (5g). 3 d, (Rf=0.32), (8%), white needles (diethyl ether), m.p. 163-164 °C. IR (Nujol) 3329.6w, 2926.0s, 1729.7m, 1711.8m, 1693.9m, 1637.8m, 1536.4m, 1463.0s, 1377.1s, 1254.6m, 1207.5w, 1144.6w, 1050.0w, 1018.3w, 979.6m, 953.4m, 879.4w, 772.6w, 722.3m, 693.6w cm⁻¹; H NMR (300 MHz, CDCl₃) δ 8.67 (s, 1H, H₅), 7.94 (d, J=8.36 Hz, 1H, Ha), 7.80 (d, J=7.69 Hz, 1H, H₉), 7.60 (t, J=1.30, 7.03 Hz, 1H, H₇), 7.49 (m, 5H, Ph), 7.37 (t, J=1.64, 1.69 Hz, 1H, H₈), 7.35 (s, 1H, H₁₀), 7.06 (d, J=1.16 Hz, 1H, H₆), 6.45 (dd, J=5.24, 37.92, 5.13 Hz, 1H, Hb), 6.27 (d, J=1.16 Hz, 1H, H₂); 13C NMR (50 MHz, CDCl₃) δ 162.2 (CO), 152.9 (C₆), 159.2 (CO), 150.9 (C-Ar), 137.5 (C-Ar), 133.1 (C₅), 130.3 (C₆), 129.7 (C₉), 129.7 (C-Ar), 127.2 (C₇), 126.3 (C₈), 113.3 (C-Ar), 112.9 (C₁₀), 100.9 (C₂), 94.4 (C₆Cl); MS m/z 302 (9), 200 (9), 199 (65), 171 (33), 170 (100), 142 (74), 115 (37), 114 (81), 113 (35), 88 (22), 86 (11), 63 (23); HRMS calc. for C₂₀H₁₄O₃: m/z 302.0943, found 302.0948; MS m/z 303.1 (M+H⁺).

2-Isopropylidene glyceral 4-oxo-[1,3-b]-naphthodioxane (7). 3 d, (Rf=0.32), (26%), colorless crystals (diethyl ether), m.p. 163-164 °C, [α]D° 14.192 (c 2.5, CHCl₃). IR (Nujol) 2926.0s, 2926.0m, 1711.8m, 1693.9m, 1668.9m, 1637.8m, 1536.4m, 1463.0s, 1377.1s, 1254.6m, 1207.5w, 1144.6w, 1050.0w, 1018.3w, 979.6m, 953.4m, 879.4w, 772.6w, 722.3m, 693.6w cm⁻¹; H NMR (300 MHz, CDCl₃) δ 8.63 (s, 1H, H₅), 7.93 (d, J=8.28 Hz, 1H, H₆), 7.79 (w, J=8.28 Hz, 1H, H₆), 7.78 (d, J=8.28 Hz, 1H, H₉), 7.61 (t, J=1.16 Hz, 1H, H₇), 7.48 (t, J=7.07 Hz, 1H, H₈), 7.44 (s, 1H, H₁₀), 5.69 (d, J=4.35 Hz, 1H, H₂), 4.52 (m, 1H, CH), 4.29 (m, 2H, CH₂), 1.50 (d, J=33.80 Hz, 6H, 2xCH₃); 13C NMR (50 MHz, CDCl₃) δ 161.3 (CO), 152.5 (C-Ar), 137.3 (C-Ar), 132.9 (C₅), 129.9 (C₆), 129.7 (C₉), 129.5 (C-
Ar), 127.1 (C7), 125.8 (C8), 114.6 (C-Ar), 112.5 (C10), 110.9 (C^2Pr), 99.8 (C2), 75.02 (CH), 64.6 (CH2), 26.3 (CH3), 25.5 (CH3); MS m/z 301.2 (M+H)+; Anal. Calc. for C_{17}H_{16}O_{5}: C, 67.99. H, 5.37. Found: C, 68.13; H, 5.30.

2-Dihydro 4-oxo-[1,3-\alpha]-naphthodioxane (6a). 1 d, (Rf=0.40), (62%), cream solid, m.p. 208-209 °C. IR (Nujol) 3329.1 m, 2855.0 s, 1753.2 m, 1729.7 m, 1712.0 m, 1694.1 m, 1661.0 m, 1631.3 m, 1578.4 m, 1536.5 m, 1513.9 m, 1463.6 s, 1377.2 s, 1352.7 m, 1282.2 m, 1254.1 w, 1220.6 m, 1151.8 m, 1102.8 m, 1048.3 m, 1029.7 m, 980.3 m, 922.1 w, 861.7 w, 820.9 m, 792.2 m, 762.2 s, 722.4 m, 698.1 w, 647.9 w cm⁻¹; \(^1\)H NMR (300 MHz, CDCl₃) δ 8.17 (d, J=7.59 Hz, 1H, H5), 7.87 (d, J=8.69 Hz, 1H, H6), 7.84 (d, J=7.95 Hz, 1H, H10), 7.66 (d, J=1.27 Hz, 1H, H7), 7.58 (m, 2H, H8&H9), 5.84 (s, 2H, CH₂); 13C NMR (50 MHz, CDCl₃) δ 161.6 (C=O), 156.9 (C-Ar), 137.2 (C-Ar), 130.1 (C5), 127.9 (C6), 126.9 (C10), 123.7 (C7), 123.0 (C8), 129.9 (C-Ar), 122.7 (C9), 109.0 (C-Ar), 91.2 (C2); HRMS calc. for C₁₂H₈O₃: m/z 200.0473, found 200.0488; MS m/z 201 (M+1)+.

2-Ethyl 4-oxo-[1,3-\alpha]-naphthodioxane (6b). 2 d, (Rf=0.42), (38%), colour less crystals (diethyl ether), m.p. 81-82 °C. IR (Nujol) 2846.7 s, 2724.6 w, 1759.2 m, 1721.2 m, 1631.2 m, 1581.3 m, 1513.9 m, 1463.9 m, 1377.0 m, 1294.0 m, 1277.0 m, 1208.9 m, 1133.8 m, 1108.3 m, 1021.8 w, 970.2 w, 823.8 m, 793.3 w, 762.5 m, 722.3 m cm⁻¹; \(^1\)H NMR (200 MHz, CDCl₃) δ 8.25 (d, J=0.62 Hz, 1H, H5), 7.87 (dd, J=8.69, 6.18 Hz, 2H, H6&H10), 7.68 (d, J=1.48 Hz, 1H, H7), 7.59 (m, 2H, H8&H9), 5.75 (t, J=4.93, 4.97 Hz, 1H, H2), 2.23 (m, 2H, CH₂), 1.25 (t, J=7.47, 7.56 Hz, 3H, CH₃); 13C NMR (50 MHz, CDCl₃) δ 162.7 (C=O), 156.8 (C-Ar), 137.3 (C-Ar), 129.9 (C5), 127.9 (C10), 126.8 (C6), 123.7 (C7), 123.1 (C9), 122.8 (C-Ar), 122.7 (C8), 108.6 (C-Ar), 102.5 (C2), 26.9 (CH₂), 7.4 (CH₃); MS m/z 228 (11), 171 (15), 170 (100), 115 (19), 114 (91), 113 (27), 88 (18), 63 (17); Anal. Calc. for C₁₄H₁₂O₃: C, 73.67. H, 5.30. Found: C, 73.87; H, 5.38.

2-Isopropyl 4-oxo-[1,3-\alpha]-naphthodioxane (6c). 2 d, (Rf=0.40), (32%), colourless crystals (diethyl ether), m.p. 69-70 °C. IR (Nujol) 2924.4 s, 1757.9 m, 1720.6 m, 1633.1 m, 1513.7 m, 1462.8 s, 1377.0 s, 1294.0 m, 1277.0 m, 1248.5 m, 1217.1 m, 1133.8 m, 1111.0 m, 1051.7 m, 1020.1 m, 984.2 m, 957.7 m, 867.0 w, 820.4 m, 790.1 m, 724.3 m, 649.3 w cm⁻¹; \(^1\)H NMR (200 MHz, CDCl₃) δ 8.23 (d, J=0.85 Hz, 1H, H5), 7.88 (dd, J=8.66, 7.67 Hz, 2H, H6&H10), 7.68 (d, J=1.48 Hz, 1H, H7), 7.59 (m, 2H, H8&H9), 5.75 (t, J=4.93, 4.97 Hz, 1H, H2), 2.41 (m, 1H, CH), 1.26 (d, J=5.27 Hz, 6H, 2xCH₃); \(^1\)C NMR (50 MHz, CDCl₃) δ 162.8 (C=O), 156.9 (C-Ar), 137.3 (C-Ar), 129.9 (C5), 127.9 (C10), 126.8 (C6), 123.7 (C7), 123.1 (C9), 122.8 (C-Ar), 122.7 (C8), 108.6 (C-Ar), 104.9 (C2), 31.9 (CH-^3Pr), 16.3 (CH₃), 16.2 (CH₃); MS m/z 242 (9), 171 (15), 170 (100), 115 (19), 114 (91), 113 (27), 88 (18), 63 (17); Anal. Calc. for C₁₅H₁₄O₃: C, 74.36. H, 5.82. Found: C, 74.14; H, 5.71.

2-Butyl 4-oxo-[1,3-\alpha]-naphthodioxane (6d). 2 d, (Rf=0.35), (38%), colourless crystals (diethyl ether), m.p. 60-62 °C. IR (Nujol) 2851.0 s, 1757.7 s, 1633.1 m, 1582.1 m, 1514.0 m, 1463.2 s, 1376.9 s, 1278.6 m, 1208.9 m, 1138.6 w, 1108.3 w, 1021.8 w, 970.2 w, 823.8 m, 793.3 w, 762.5 m, 722.3 m cm⁻¹; \(^1\)H NMR (200 MHz, CDCl₃) δ 8.24 (d, J=0.66 Hz, 1H, H5), 7.87 (dd, J=8.67, 5.94 Hz, 2H, H6&H10), 7.84 (d, J=1.51 Hz, 1H, H7), 7.63 (m, 2H, H8&H9), 5.78 (t, J=5.1, 5.14 Hz, 1H, H2), 2.20 (m, 2H, CH₂), 1.66 (m, 2H, CH₂), 1.48 (m, 2H, CH₂), 1.00 (t, J=7.17, 7.21 Hz, 3H,
2-Isopropylidene glyceral 4-oxo-[1,3-α]-naphthodioxane (8). 1 d, (Rf = 0.22), (65%), colourless crystals (diethyl ether), m.p. 105-106 °C, [α]D 0 4.9 (c 3, CHCl₃). IR (Nujol) 3329.6 m, 2922.4 s, 2853.8 s, 1729.4 m, 1711.7 m, 1694.1 m, 1680.9 m, 1536.4 m, 1462.7 s, 1377.0 s, 1291.0 w, 1254.0 m, 1164.6 w, 1054.3 w, 979.6 m, 763.7 w, 722.2 m, 702.0 w cm⁻¹; ¹H NMR (200 MHz, CDCl₃) δ 8.22 (d, J=0.90 Hz, 1H, H5), 7.89 (m, 2H, H6 & H10), 7.72 (d, J=5.43 Hz, 1H, H7), 7.63 (m, 2H, H8 & H9), 5.83 (d, J=4.09 Hz, 1H, H2), 4.62 (m, 1H, CH), 4.38 (m, 2H, CH₂), 1.55 (d, J=28.04 Hz, 6H, 2xCH₃); ¹³C NMR (50 MHz, CDCl₃) δ 161.4 (C=O), 137.4 (C-Ar), 130.2 (C5), 128.1 (C10), 127.1 (C6), 123.7 (C7), 123.3 (C9), 123.1 (C-Ar), 122.7 (C8), 111.0 (C-Ar), 108.8 (C-Ar), 99.8 (C2), 95.4 (C-²Pr), 74.9 (CH), 64.6 (CH₂), 26.5 (CH₃), 25.2 (CH₃); HRMS calc. for C₁₇H₁₆O₅: m/z 300.0998, found 300.1008; MS m/z 301.1 (M+H)⁺.

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References