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Intramolecular Pauson-Khand reactions of cycloheptynedicobalt complexes

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Abstract: Cycloheptyne- $Co_2(CO)_6$ complexes bearing alkenes tethered by oxygen, sulphur, and nitrogen atoms undergo Pauson-Khand reactions to afford tricyclic compounds containing a fused 7,5- ring system.

Key words: Pauson-Khand reactions, alkynes, complexes, transition metals, Nicholas reactions

The fused 7,5- ring system an is extremely common framework within natural products, particular those of the terpene class. The guaiane, daucane, and lactarane sesquiterpenoids are only the most obvious of these systems, as many other sesquiterpenoids¹ and diterpenoids contain this structural unit. While traditional synthetic methods predominantly rely upon ring expansion protocol for access to the cycloheptane portion of these systems,² approaches based on direct cycloheptane synthesis are of increasing importance.³



Figure 1 Guaiane, daucane, and lactarane sesquiterpenoids

It has been demonstrated our group⁴ and other groups⁵ that cycloheptynedicobalt complexes are both stable and readily accessible, predominantly by Nicholas reaction chemistry. Due to the strain associated with cyclic alkynes, the $Co_2(CO)_6$ unit is central to the existence of the ring system. In addition, alkyne-Co₂(CO)₆ complexes are among the most widely employed precursors to fivemembered ring construction by Pauson-Khand reactions,^{6,7} despite recent important advances in catalytic methods and the use of other metal complexes.^{6a,d} As a result, it was our belief that the investigation of Pauson-Khand reactions on cycloheptynedicobalt complexes would be of importance for the synthesis of 7,5- ring systems. As early work in our group indicated that intermolecular Pauson-Khand reactions of cycloheptenyne-Co₂(CO)₆ complexes succeeded with a quite restricted number of alkenes,8 we concluded that intramolecular versions of the reaction were prudent. It was our intent to incorporate an alkene function with each of oxygen, sulphur, and nitrogen tethers to the alkynedicobalt unit, with particular interest in oxygen and sulphur tethers due to their capacity for reductive cleavage in the products.^{9,10} An analogous approach employing

oxygen tethered alkenes has seen success in a limited number of instances for 8,5- and 9,5- ring systems.^{11,5b, 5f} While this work was nearing completion, the preparation of 5,7,5- and 5,8,5- systems by way of intramolecular Pauson-Khand reactions of 7- and 8- membered ring cyclic ether alkyne-Co₂(CO)₆ complexes with carbon-based alkene tethers was reported by the group of Shea.¹²

We chose as our starting point homoallyl propargyl acetate **1a**, which may be obtained by ring closing metathesis of the appropriate acyclic diene, 4b,5b and the corresponding cycloheptyne complex 1b, which was prepared quantitatively by Raney[®] nickel reduction of **1a.**¹³ Complex 1a could be converted into allyl ether 2a (75% yield) and allyl thioether 2b (58% yield) by BF₃-OEt₂ (5 equiv) with allyl alcohol or allylthiol, respectively.¹⁴ In these cases, the reactions had to be performed in the presence of excess alcohol/thiol (10 equiv) in order to effectively compete with elimination of acetic acid from 1a. In the case of acetate 1b, the analogous elimination process was less of a competitive one, such that 1b could be converted into allyl alcohols 2c (84% yield) and 2d (84% yield), and allyl thioether 2e (84%) by treatment with BF₃-OEt₂ and allyl alcohol, 2-methyl-2-propen-1ol, and allylthiol, respectively. In addition, homoallyl alcohol 2f (92% yield) could be obtained by the analogous reaction with 1-buten-4-ol.

$C-5$ $C-6$ OAc $Co_2(CO)_6$	HX BF ₃ -OEt ₂ , CH ₂ Cl ₂ rt (1a), 0 °C (1b)		Co ₂ (CO) ₆	
1a, ∆°,° 1b, 5.6-H-				
10 , 0,0 11 ₂				
Acetate	Х	R	Produc	t (Yield, %)
1a	0	Н	2a	75
1a	S	Н	2b	58
1b	0	Н	2c	84
1b	0	Me	2d	84
1b	S	Н	2e	84
1b	OCH ₂	Н	2f	92

Scheme 1 Preparation of oxygen- and sulphur tethered enyne- $Co_2(CO)_6$ complexes

Our desire to include a nitrogen tethered substrate initially met with failure. Subjecting acetate **1b** to allylamine or N-allylacetamide under protic or Lewis acidic condition gave no amine incorporation with allylamine and at best trace amounts of amide incorporation with Nallylacetamide (2g). Attempts to isolate cation 3 by treatment of 1b or alcohol 1c with tetrafluoroboric acid were similarly unsuccessful. Fortunately, the use of HBF₄ with excess dimethyl sulphide rapidly gave a precipitate consistent with the formation of sulphonium salt **4**.¹⁵ Removal of solvent from **4** under vacuum and subsequent addition of allylamine and diisopropylethylamine afforded 2h, which was in turn immediately subjected to *p*-toluenesulphonic anhydride, giving **2i** in 62% yield (from 1b). While N- acetyl functions are infrequently used as tethers for intramolecular Pauson-Khand reactions,¹⁶ we desired an additional example of this Nsubstitution sequence; therefore, employing acetic anhydride in place of *p*-toluenesulphonic anhydride in reaction with unpurified 2h afforded 2g in 73% yield (from **1b**).



Scheme 2 Preparation of nitrogen tethered enyne- $Co_2(CO)_6$ complexes

With several substrates in hand, attention was turned to the Pauson-Khand reactions of **2a-f**, and **2i**. We chose **2c** for investigation of the reaction under several sets of conditions. The Sugihara amine conditions (3.5 equiv CyNH₂, ClCH₂CH₂Cl, reflux)¹⁷ indeed gave Pauson-Khand product **5c** in 65% yield, as a 5.2:1 mixture of diastereomers favouring the isomer containing methine hydrogen atoms in a *cis*- relationship (*cis*- **5c**), along with a smaller amount of Pauson-Khand/allylic reduction product^{17a} **6c** (13%). Attempted use of the Sugihara sulphide conditions (BuSMe, ClCH₂CH₂Cl, reflux)¹⁸ eliminated the byproduct **6c**, but at the expense of **5c** (51%, 2.0:1 *cis:trans*). In addition, Me₃NO (49% yield of **5c**, 1.2:1 *cis:trans*) and refluxing toluene (20% yield of **5c**, 1.1:1 *cis:trans*) were judged inferior to the Sugihara amine conditions. The Sugihara amine conditions were therefore explored with the other substrates. The corresponding allyl/homoallyl ether complex **2a** cyclized to form **5a** in excellent yield (89%) and good diastereoselectivity (*cis:trans* = 10:1). Both thioallyl complexes **2b** and **2e** cyclized successfully to give **5b** (65% yield, *cis:trans* = 3.8:1) and **5e** (66% yield, *cis:trans* = 2.0:1), respectively, although the diastereoselectivities were lower than the oxygen tethered analogues. Homoallyl ether complex **2f** similarly underwent reaction to give **5f** (73% yield, *cis:trans* = 2.0:1).¹⁹



Scheme 3 Pauson-Khand reaction of 2c under various conditions

In contrast to the above cases, both methallyl ether **2d** and N-allyl tosylate complex **2i** failed to give Pauson-Khand reaction products in synthetically useful yields under the Sugihara amine conditions, although a small amount of Pauson-Khand/allylic reduction product **6d** (14% yield) was obtained from **2d**. Therefore, several other sets of reaction protocols were examined with these substrates. Fortunately, methallyl ether complex **2d** underwent cyclization successfully under Me₃NO conditions (0 °C, 59% yield, 66% yield based on recovered **2d**) to give *cis*-**5d** exclusively. Sulphonamide tethered complex **2i** cyclized successfully under the Sugihara sulphide conditions, to give **5i** (55% yield, *cis:trans* = 1.2:1).²⁰

The stereochemical assignments were arrived at as follows. To our surprise, the ¹H NMR spectra of majority of **5a-f** and **5i** did not give nOe enhancements between the methine hydrogen atoms for either diastereomer. Conversely, the 2D-NOESY spectrum of β -methyl substituted **5d** displayed a positive nOe interaction between the methine hydrogen and the methyl hydrogens, which was therefore indicative of a *cis*- relationship. The methine H atoms α - to the heteroatoms of **5d** and the major diastereomers of the cycloheptane products **5c**, **5f**, and **5i** also displayed very similar ¹H NMR vicinal coupling constants (dd, J = 9.9-10.9, 4.2-5.7) which were not repeated in the minor diastereomers. Finally, a small amount of the *minor* isomer of **5e** formed crystals suitable for X-ray diffraction studies, which indicated a *trans*- relationship between the two methines. The assignments for **5a-b** were based on analogy and are supported by calculations on the proposed alkene coordination intermediate, the alkene insertion transition state (MMX, PCMODEL; MM3, CAChe[®]), and on **5a-b** themselves (PM3, CAChe[®]).



Scheme 4 Intramolecular Pauson-Khand reactions of cycloheptyne Co₂(CO)₆ complexes

In summary, we have demonstrated the ability of oxygen-, sulphur-, and nitrogen tethers to enable Pauson-Khand reactions of cycloheptyne-Co₂(CO)₆ complexes, affording tricyclic systems containing a fused 7,5- structural unit. As the resultant enone C=C bond is mutual to the 7- and 5- membered rings, we view the chemistry as complementary to the work of the Moyano/Pericàs group and the group of Pérez-Castells,^{9a,21} who have employed Pauson-Khand reactions of alkynes tethered to cycloheptenes through sulphur, and oxygen and nitrogen atoms, respectively, to give 5,7,5- systems with the alkene function solely in the 5- membered ring.

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- (14)Selected compounds: (2c):IR (neat, NaCl) v_{max} 2932, 2090, 2047, 2020 cm⁻¹; ¹ H NMR δ 5.98 (m, 1H), 5.34 (dd, J = 17.1, 1.5, 1H), 5.21 (dd, J = 10.4, 1.5, 1H), 4.48 (dd, J = 10.4, 4.2, 1H), 4.35 (dd, J = 12.5, 5.1, 1H), 4.19 (dd, J = 12.5, 5.7, 1H), 3.19 (apparent dt, J = 16.7, 3.4, 1H), 2.76 (m, 1H), 2.19 (m, 1H), 2.05 (m, 1H), 1.97 (m, 1H), 1.48-1.65 (m, 2H), 1.39 (m, 1H); ¹³C NMR 203.9, 138.7, 121.0, 103.6, 102.5, 84.4, 74.2, 40.9, 39.4, 33.1, 29.7; MS m/e 380 (M-2CO⁺); HRMS m/e for $C_{16}H_{14}Co_2O_7$ calcd. (M- $2CO^+$) 379.9500, found 379.9466; (2e) IR (neat, NaCl) v_{max} 2927, 2089, 2049, 2020 cm⁻¹. ¹H NMR δ 5.88 (m, 1H), 5.15 (d, J = 17.7, 1H), 5.12 (d, J = 12.0, 1H), 3.89 (dd, J = 14.2, 4.2, 1H), 3.29 (d, J = 7.5, 2H), 3.12 (dt, J = 16.3, 3.6, 1H), 2.82 (m, 1H), 2.27 (m, 1H), 2.11 (m, 1H), 1.95 (m, 1H),1.68 (m, 1H), 1.59 (m, 1H), 1.45 (m, 1H); ¹³C NMR 200.0, 134.7, 116.9, 102.0, 99.9. 48.5, 38.0, 35.5, 35.0, 29.8, 28.8; MS m/e , 424 (M-CO⁺), 396 (M-2CO⁺); HRMS m/e for $C_{16}H_{14}Co_2O_6S$ calcd. (M-2CO⁺) 395.9271, found 395.9295. (2i) IR (neat, KBr) v_{max} 3082, 2928, 2093, 2051, 2025 cm⁻¹; ¹H NMR δ 7.72 (d, J = 8.3, 2H), 7.28 (d, J = 8.3, 2H), 5.76 (m, 1H), 5.20 (dd, J = 17.2, 1.3, 1H), 5.07 (dd, J = 10.2, 1.3, 1H), 4.96 (dd, J = 11.7, 3.2, 1H), 4.14-4.26 (m, 2H), 3.23 (br d, J = 13.5, 1H), 2.73 (m, 1H), 2.43

(s, 3H), 2.11 (m, 1H), 1.92-2.04 (m, 2H), 1.73 (m, 1H), 1.35-1.60 (m, 2H); 13 C NMR 199.1, 143.1, 138.6, 135.8, 129.5, 127.3, 117.3, 102.2, 96.9, 63.5, 46.2, 37.0, 35.6, 28.9, 21.5; MS m/e , 533 (M-2CO⁺), 505 (M-3CO⁺), 421 (M-6CO⁺) ; HRMS m/e for C₂₃H₂₁Co2NO₈S calcd. (M-3CO⁺) 504.9804, found 504.9828.

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- (19) **Experimental Procedure**: A solution of **2b** (73.0 mg, 0.16 mmol) and 3.5 equivalents of cyclohexylamine (3.5 equiv, $64 \ \mu$ L) in 1,2-dichloroethane (8 mL) was heated to reflux for 20 minutes. The solution was cooled to room temperature, 20 mL diethyl ether (20 mL) was added, and the mixture was filtered through a pad of silica gel. Preparative TLC (4:1 petroleum ether:diethyl ether) of the crude material afforded **5b** (20.2 mg, 65% yield, 3.8:1 *cis:trans*).
- (5a) IR (neat, NaCl) v_{max} 3079, 2957, 1737, 1715 cm⁻¹; ¹ H (20)NMR δ 5.84 (m, 1H), 5.80 (m, 1H), 4.69 (br d, J = 10.9, 1H), 4.28 (apparent t, J = 7.1, 1H), 3.31 (m, 1H), 3.29 (d, J = 7.1, 1H), 2.95-3.08 (m, 2H), 2.63-2.71 (m, 2H), 2.23 (m, 1H), 2.16 (dd, J = 18.5, 2.7, 1H); resonances for the minor diastereomer were observed at 4.72 (apparent t, J = 6.0, 1H), 4.38 (apparent t, J = 7.4, 1H), 3.46 (dd, J = 10.9, 8.7, 1H), 2.11 (J = 18.0, 2.7, 1H); 13 C NMR 208.1 179.9, 128.7, 125.3, 72.9, 70.5, 43.2, 38.8, 33.3, 29.7, 25.1; MS m/e 176 (M⁺); HRMS m/e for C₁₁H₁₂O₂ calcd. (M⁺) 176.0837, found 176.0842. (5b) IR (neat, NaCl) v_{max} 3012, 2922, 1709, 1678 cm⁻¹; ¹ H NMR δ 5.59, (ddd, J = 12.4, 7.2, 2.1, 1H), 5.61 (m, 1H), 4.48 (br d, J = 12.0, 1H), 3.37 (m, 1H), 3.13 (dd, J = 10.9, 6.7, 1H), 3.07 (m, 1H), 2.95 (m, 1H), 2.73 (1/2 ABquartet, J = 6.3, 1H), 2.70 (1/2 ABquartet, J = 6.3, 1H), 2.57 (m, 1H), 2.43 (m, 1H), 2.22 (dd, J = 17.1, 4.1 1H); resonances for the minor diastereomer were observed at 4.20 (br d, J = 4.20, 1H), 2.66 (d, J = 6.5, 1H), 2.18 (obscured dd, 1H) ¹³C NMR 207.0, 183.4, 135.2, 126.6, 126.1, 47.9, 44.9, 42.6, 37.1, 32.3, 23.8; resonances for the minor diastereomer were observed at 129.2, 47.0, 43.2, 40.3 ; MS m/e 192 (M^+); HRMS m/e for $C_{11}H_{14}O_2$ calcd. (M⁺) 192.0603, found 192.0580. (5c) IR (neat, NaCl) v_{max} 2927, 1714, 1680 cm⁻¹; ¹ H NMR δ 4.74 (dd, J = 10.9, 5.7, 1H), 4.27 (apparent t, J = 7.7., 1H), 3.36 (dd, J = 11.3, 7.7, 1H), 3.27 (m, 1H), 2.64 (dd, J = 17.7, 6.3, 1H), 2.58 (m, 1H), 2.21 (m, 1H), 2.14 (dd, J = 17.6, 3.1, 1H), 2.09 (m, 1H), 1.99 (m, 1H), 1.93 (m, 1H), 1.56 (m, 1H), 1.30-1.49 (m, 2H); resonances for the minor diastereomer were observed at 4.35 (obscured m, 1H), 4.32 (apparent t, J =

7.8, 1H), 3.35 (dd, J = 11.4, 8.3, 1H), 2.26 (dd, J = 16.6, 4.7, 1H), 2.55 (obscured m, 1H); ¹³C NMR 208.4, 186.0, 137.7, 79.6, 71.9, 44.8, 41.0, 34.0, 27.6, 26.5, 23.5; resonances for the minor diastereomer were observed at 206.8, 182.2, 135.4, 77.1, 71.4, 43.6, 39.0, 29.5, 27.5, 24.6, 23.4 MS m/e 178 (M⁺), 150 (M-CO⁺); HRMS m/e for $C_{11}H_{14}O_2$ calcd. (M-CO⁺) 150.1045, found 150.1038. (5d) IR (neat, NaCl) v_{max} 2928, 1713, 1677 cm⁻¹; ¹ H NMR δ 4.79 (ddd, J = 10.9, 5.7, 1.6, 1H), 3.93 (d, J = 8.0, 1H), 3.54 (d, J = 8.0, 1H), 2.55 (apparent dt, J = 16.1, 4.2, 1H), 2.39 (1/2 ABquartet, J = 17.1, 1H), 2.35 (1/2 ABquartet, J = 17.1, 1H), 2.21 (m, 1H), 2.05 (m, 1H), 1.98 (m, 1H), 1.90 (m, 1H), 1.55 (m, 1H), 1.40 (m, 1H), 1.33 (m, 1H), 1.31 (s, 3H); ¹³C NMR 208.1, 185.7, 136.2, 76.1, 48.6, 47.4, 33.8, 27.5, 24.9, 24.5, 23.4 ; MS m/e 192 (M⁺); HRMS m/e for C₁₂H₁₆O₂ calcd. (M⁺) 192.1150, found 192.1143. (5e) IR (neat, NaCl) ν_{max} 2928, 1703, 1665 cm $^{-1};$ 1 H NMR δ 4.14 (br d, J = 11.6, 1H), 3.31 (m, 1H), 3.04 (dd, J = 10.6, 6.7, 1H), 2.69 (dd, J = 17.0, 6.0, 1H), 2.65 (m, 1H), 2.60 (apparent t, J = 11.3, 1H), 2.18 (dd, J = 16.8, 4.3, 1H), 2.04 (m, 1H), 1.85-1.97 (m, 2H), 1.53 (m, 2H), 1.47 (m, 1H), 1.14 (m, 1H); resonances for the minor diastereomer were observed at 4.19 (br d, J = 11.3, 1H), 2.65 (J = 18.5, 6.2, 1H), 2.58 (m, 1H), 2.23 (m, 1H), 1.41 (m, 1H); ¹³C NMR 207.4, 184.9, 139.0, 48.5, 42.6, 36.1, 31.2, 30.9, 27.0, 23.0; resonances for the minor diastereomer were observed at 208.1, 181.0, 139.5, 48.2, 46.8, 41.2, 37.4, 35.3, 28.5, 23.5 MS m/e 194 (M⁺); HRMS (ESI) m/e for $C_{11}H_{14}OS$ calcd. (M+Na⁺) 217.0658, found 217.0650. (5f) IR (neat, NaCl) v_{max} 2928, 1702, 1648 cm⁻¹; ¹ H NMR δ 4.23 (dd, J = 9.9, 4.2, 1H), 4.06 (m, 1H), 3.66 (apparent dt, J = 2.1, 11.9, 1H), 2.77 (m, 1H), 2.62 (dd, J = 18.7, 6.3, 1H), 2.26-2.28 (2H), 1.97-2.15 (m, 3H), 1.81-1.91 (m, 2H), 1.28-1.75 (m, 4H); resonances for the minor diastereomer were observed at 4.26 (br d, obscured, 1H), 4.09 (m, 1H), 3.74 (ddd, J = 11.4, 9.3, 4.6, 1H), 3.01 (m, 1H), 2.70 (m, 1H), 2.67 (dd, J = 18.2, 6.3, 1H), 2.70 (m, 1H), 1.41 (m, 1H), 1.14 (m, 1H); ¹³C NMR 207.5, 170.4, 139.2, 78.6, 66.8, 41.0, 37.8, 34.0, 32.2, 25.9, 22.3, 21.3; resonances for the minor diastereomer were observed at 206.5, 179.3, 137.5, 63.6, 41.1, 32.3, 31.9, 28.2, 27.5, 26.6; MS m/e 192 (M⁺); HRMS (ESI) m/e for C₁₁H₁₄O₂ calcd. (M+Na⁺) 215.1048, found 215.1050. (5i) IR (neat, NaCl) v_{max} 2923, 1714, 1680 cm⁻¹; ¹ H NMR δ 7.78 (d, J = 8.2, 2H) and 7.72 (d, J = 8.2, 2H), 7.34 (d, J = 8.2, 2H) and 7.38 (d, J = 8.2, 2H), 4.46 (ddd, J = 10.9, 5.4, 1.5 1H), 4.00 (dd, J = 12.0, 7.8, 1H), 3.96 (apparent t, J = 7.8, 1H), 3.57 (br d, J = 9.8, 1H), 3.12 (m, 1H), 2.94 (apparent t, J = 11.5), 2.73 (m, 1H), 2.62 (dd, J = 16.9, 6.0, 1H), 2.50-2.54 (m, 3H), 2.43 (s, 3H) and 2.45 (s, 3H), 2.39-2.50 (m, 3H), 2.18 (m, 1H), 1.96-2.10 (m, 4H), 1.79-1.93 (m, 2H), 1.43-1.64 (m, 4H), 1.24-1.39 (m, 2H), 1.13 (m, 1H); ¹³C NMR 207.2, 205.4, 180.9, 178.5, 144.3, 143.8, 139.2, 137.7, 136.0, 131.7, 130.0, 129.9, 128.0, 127.1, 62.6, 61.4, 54.2, 52.7, 41.6, 40.9, 40.4, 39.3, 35.7, 30.9, 30.3, 28.1, 27.1, 27.0, 25.9, 23.5, 22.8, 21.6; MS m/e 194 (M⁺); HRMS m/e for $C_{12}H_{18}O_2$ calcd. (M⁺) 194.1301, found 194.1295.

(21) Rosillo, M.; Casarrubios, L.; Domínguez, G; Pérez-Castells, J. Org. Biomol. Chem. 2003, 1, 1450. _____