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# Chemical Communications

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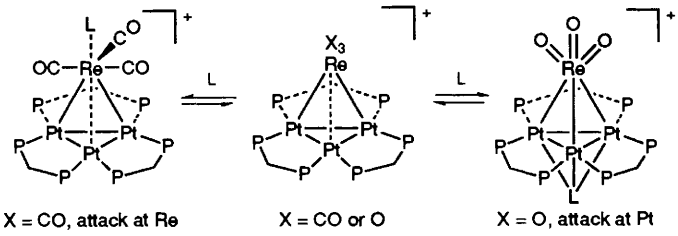
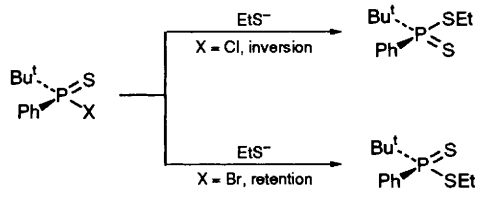
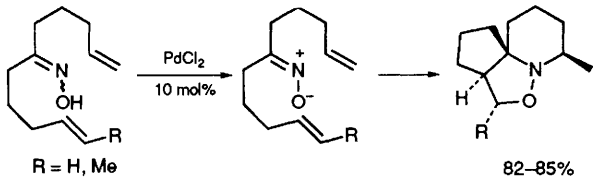


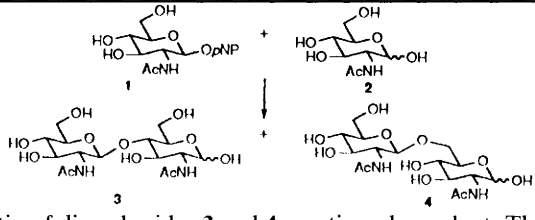




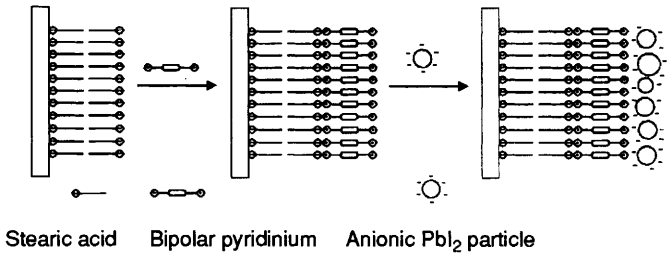





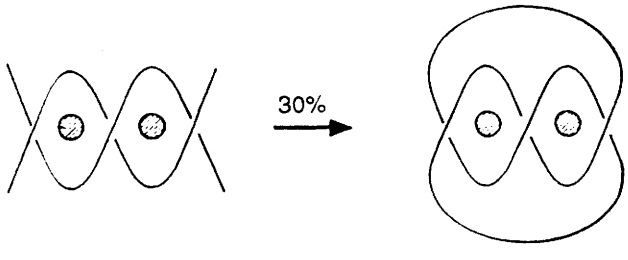



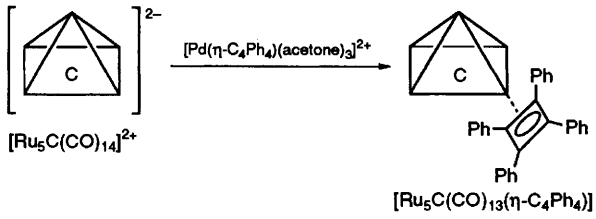
The ratio of disaccharides **3** and **4** was time-dependent. The major product in each case was isolated by selective enzymatic hydrolysis of the minor product.



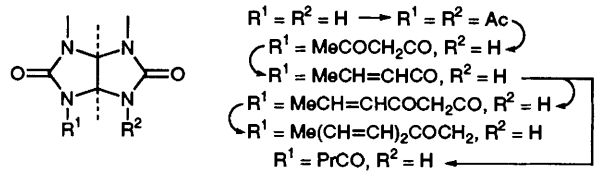




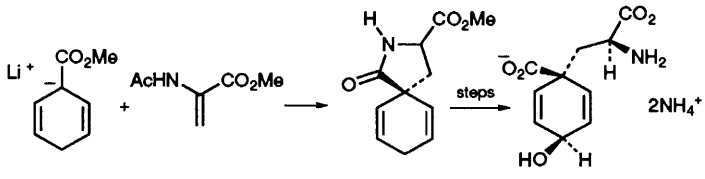





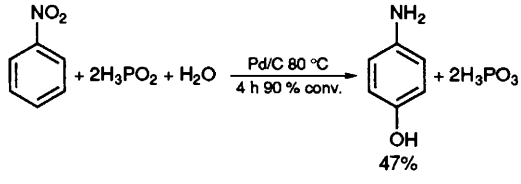


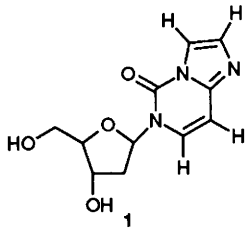




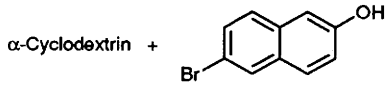





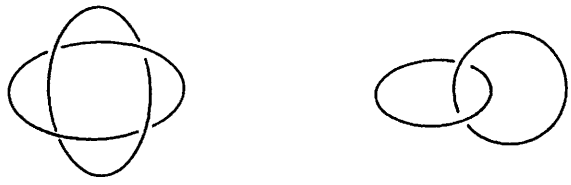




The mutagenic lesion 3,N4-etheno-2′-deoxycytidine (εdC) 1 does not form stable base pairs with A, C, G or T. A DNA duplex containing an εC : A pair adopts the B conformation with all bases *anti*.





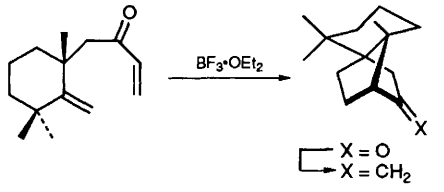

6-Bromo-2-naphthol within a 2:1 α-cyclodextrin–6-bromo-2-naphthol inclusion complex exhibits room temperature phosphorescence.



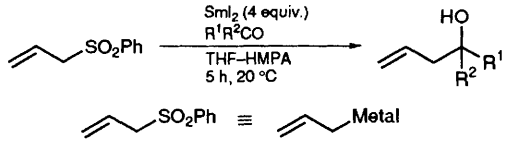
1 2

ESI-MS is able to differentiate between **1** and **2** by controlling the collision energy.

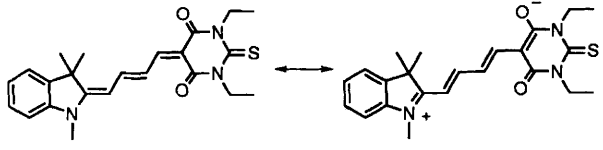






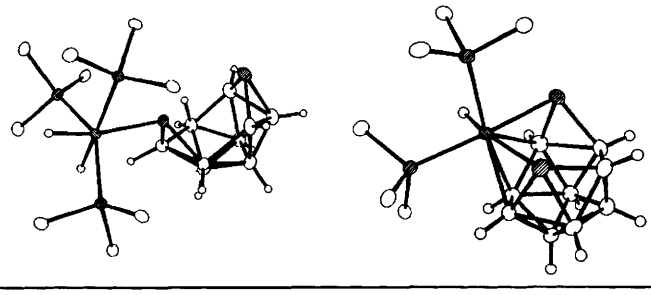





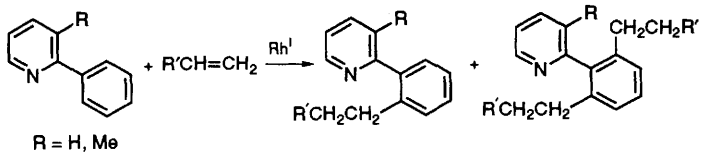



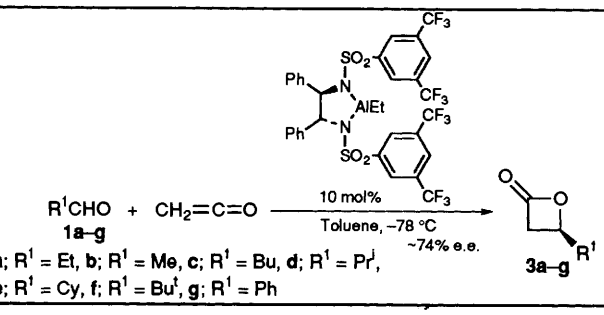
The molecular length and the endgroups determine the relative contribution of neutral and charge-separated resonance structures to the ground-state structure of merocyanines and therefore determine the magnitude and sign of their first hyperpolarisabilities.



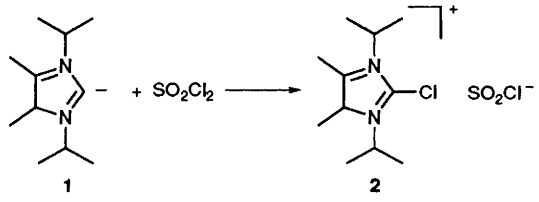




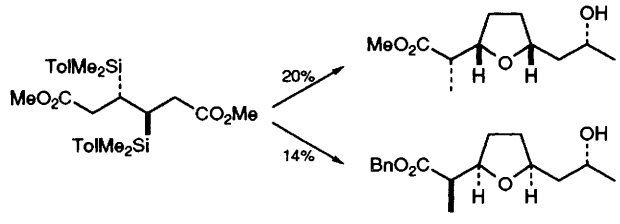




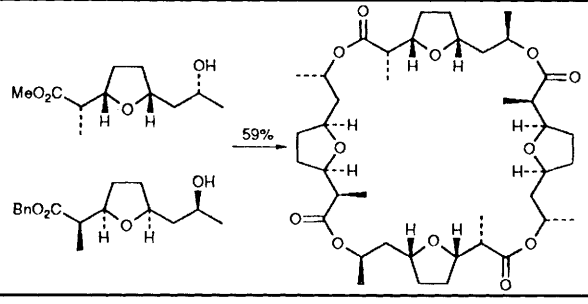


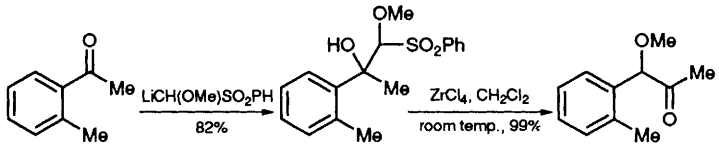




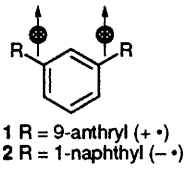



Several other examples are given

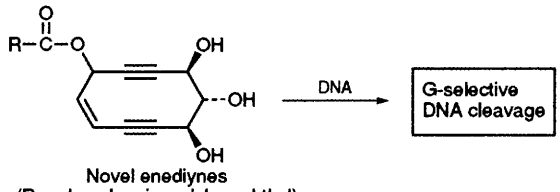


In a study of possible electrochemical control of epidermal patch medication, acetylsalicylate is now shown to be as readily released from polypyrrole as salicylate, by initially incorporating it in polymer of coarser appearance and possibly more open reticulation with hence wider exit channels.



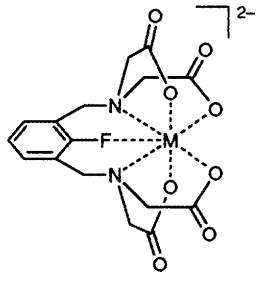

**1** R = 9-anthryl (+ •)
**2** R = 1-naphthyl (– •)






Novel enediynes
(R = phenyl, quinoxalyl, naphthyl)



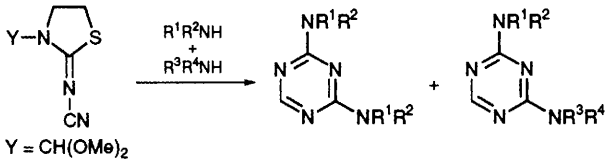





$$2[(OEP)Rh]^+ \rightarrow [(OEP)Rh]_2^{2+}$$

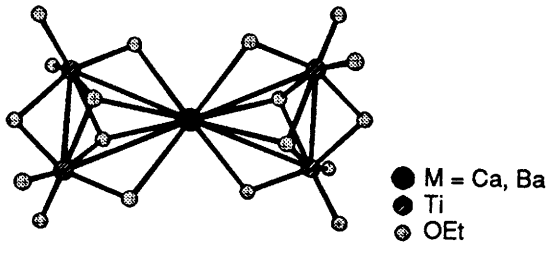
Octaethylporphyrinato rhodium cation $[(OEP)Rh]^+$ forms a tightly bound dimer, $[(OEP)Rh]_2^{2+}$ (**1**). The diamagnetism and the $^1$H NMR shifts of **1** demonstrate the loss of porphyrin ring current shifts observed for conventional aromatic porphyrin species and clearly indicate that **1** is the dimer of a porphyrin cation radical complex of rhodium(II). Relatively strong bonding between the monomer units has contributions from both a $Rh^{II}$–$Rh^{II}$ ($d_{z^2}$–$d_{z^2}$) single bond and delocalised π–π interactions between porphyrin cation radical units.

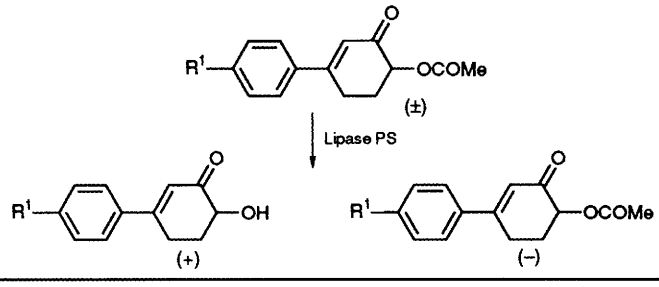




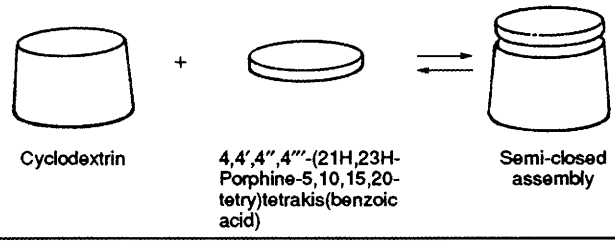

Y = $CH(OMe)_2$



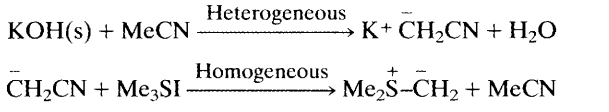








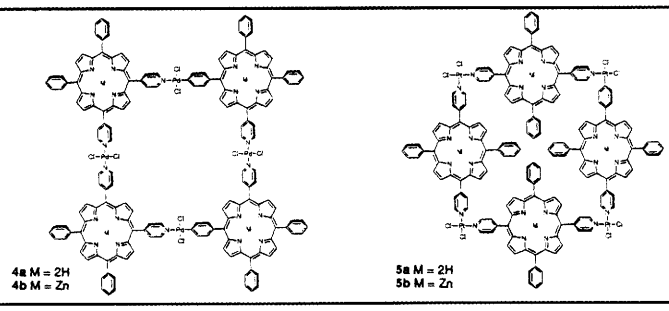




Acetonitrile acting as a solid-liquid phase-transfer catalyst



Single strands of DNA may be identified by hybridisation to a complementary probe to which is attached chelated $Eu^{3+}$. Visualisation is effected by a sensitiser localised by intercalation.





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# Journal of Materials Chemistry

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