**Dicephalic surfactants**

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Two examples of dicephalic surfactants, a new class of two-headed amphiphiles, were prepared and demonstrated to exhibit unexpected aggregation behaviour.

Recently several research groups have studied the aggregation behaviour of surfactants with unusual structural features in order to further understand the relationship between the structure of the amphiphile and the type of aggregates it forms. Examples include surfactants with hyperextended1 and spiro2 hydrocarbon chains, gemini surfactants3 and super surfactants, e.g., block copolymers of polystyrene and poly(propyleneimine) dendrimers.4 Gemini surfactants, a new class of amphiphilic molecules with two hydrocarbon chains and two polar head groups connected by a linker, display aggregation behaviour that is distinctly different from that of the constituting ‘parent’ surfactants.5 The interesting properties of gemini surfactants prompted us to design and synthesise another type of amphiphilic molecule for which we propose the name dicephalic (two-headed) surfactant. Molecules of this type consist of a single hydrocarbon chain and two polar head groups (Scheme 1).‡ Both compounds can complex metal ions; the former binds transition metals to its imidazole groups, the latter calcium ions to its phosphate functions.6 Here we report on the unusual self-assembling properties of the dicephalic surfactants.

![Image](308x115 to 549x243)

The synthesis of compound 1 was initially tried by treating 1,3-dibromopropan-2-yl stearate 3 with imidazole in MeCN. Instead of the desired product 1 a mixture of compounds 4 and 6 was isolated. A mechanistic explanation is presented in Scheme 1A. Direct nucleophilic replacement of the second halogen atom in 3 by imidazole is difficult due to considerable steric hindrance as was evident from examination of CPK models.‡ During aqueous work-up a hydroxy group is probably introduced which gives 5 and after acyl migration7 the α-hydroxy stearate 6. Compound 1 could, however, be synthesised via a double nucleophilic attack of sodium imidazolide on epibromohydrin, followed by acylation of the alkoxy intermediate (Scheme 1B).

Reaction of 3 with tetraalkylammonium salts of mono- and di-benzyl phosphate, for similar steric reasons as mentioned above, did not lead to satisfactory yields of the respective desired bis-phosphates.‡ Compound 2 was prepared therefore by phosphorylation of the diiodide 7 with silver dibenzyl phosphate8 followed by catalytic hydrogenation over Pd/C and subsequent cation exchange (Scheme 1C). This indirect displacement of the halogen atoms by the dibenzyl phosphate ion is promoted by the formation of an AgI complex and the neighbouring group participation of the ester carbonyl function.

Inspection of CPK models of 1 and 2 revealed that both compounds have a very large head group section compared to their lipophilic part. According to the structure–shape concept,9 the formation of micelles may therefore be expected when these compounds are dispersed in water. Dispersion of 1 in water at pH 7.0 (0.1%, w/w) did not, however, lead to the formation of micelles, but to well defined multilayered platelets of micrometer size [Fig. 1(a)]. Powder diffraction experiments revealed a repetitive distance of 31 Å, indicating a structure in which the hydrocarbon chains are interdigitated. Electron diffraction showed several bands, indicative of the polycrystalline nature of the sample, and a strong phase transition at 58 °C (ΔH = 60 J g⁻¹) was observed by DSC. The pKₐ values of 1 (3.6 and ca. 7), make it likely that the platelets consist of a mixture of protonated and deprotonated surfactant molecules. The possibility of intermolecular hydrogen bond formation over long distances, may account for the remarkable stability of these lamellar structures.

When 0.25 equiv. of CuSO₄ were added to a dispersion of 1 in water, vesicles with diameters of 50–250 nm were formed [Fig. 1(b)]. EPR and FTIR titration experiments showed that a complex of the type Cu(Imidazole)₂⁺ had been formed,8,8 but it was not possible to differentiate between a monomeric Cu⁺₁ 1₂

![Image](Chem. Commun., 1998 743)
early by formation of a linear polymeric complex. Addition of calcium, either intramolecular by chelation or intermolecularly by formation of bundles of fibres each with a diameter of 65 Å, (aspect ratio > 2000). These fibres showed a faint phase transition at 15 °C (H(2)) in DSC. The formation of very long structures suggests that intermolecular interactions must also play an important role in stabilising these aggregates. Since at pH 7.0 the phosphate groups will be in their monoprotonated form, a network of hydrogen bonds can be formed between protonated and unprotonated P–OH groups of the stearoyloxy group or the methylene imidazole group, i.e. Cu²⁺ ions to 1 and Ca²⁺ ions to 2, gives rise to a change in molecular organisation resulting in the generation of lamellar structures in both cases. Further studies are underway and will be presented elsewhere.

Notes and References
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