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The EUChemSoc Societies have taken the significant step into the future by merging their traditional journals, to form two leading chemistry journals, the *European Journal of Inorganic Chemistry* and the *European Journal of Organic Chemistry.* Three further EUChemSoc Societies (Austria, Czech Republic and Sweden) are Associates of the two journals.

COVER PICTURE

The cover picture shows the reaction between the phosphavinylidene complexes $[(\eta^5\text{-}C_5H_5)(CO)_2M{=}P{=}C(SiMe_3)_2]$ and the inversely polarized phosphaalkene $t\text{BuP}{=}C(NMe_2)_2$, which affords η^3-1,2-diphosphaallyl complexes by smooth transfer of the phosphinidene unit onto the electrophilic ligand of the precursors. The molecular structure underlines the unsymmetrical η^3-ligation of the heteroallyl ligand to the metal with the *tert*-butyl substituent *syn* to the central phosphorus atom. Treatment of the phosphavinylidene complexes with *P*-aroylphosphaalkenes leads to decomposition. In contrast, phosphavinylidene complexes and *As*-aroylarsaalkenes yield complexes featuring cyclic phosphenium ligands with an As=C bond. It is believed that the first step of this process is a formal [2+1] cycloaddition between the P=C bond and the aroylarsinidine to give a three-membered ring, which subsequently incorporates the CO unit to afford the final product. In all cases, tetrakis(dimethylamino)ethene is formed as a byproduct. Details are presented in the Microreview by L. Weber on p. 4095ff.