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An unusually rapid Claisen rearrangement involving ring expansion

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A Claisen rearrangement of a partially-fluorinated system involving ring expansion occurred at an unusually low temperature, 100 °C lower than a comparable system from the literature.

Paquette and co-workers showed how the Claisen rearrangement could be used to transform readily available vinylic lactones into medium-ring carbocycles during a series of landmark natural product syntheses.1 The Tebbe reaction2 fulfills a critical role in providing the vinyl ether component of the allyl vinyl rearrangement precursor; Scheme 1 summarises the sequence.

The rearrangement step requires the use of high temperatures and some special experimental precautions;3 after the Tebbe reaction, the allyl vinyl ether is sealed in base-washed tubes to minimise decomposition through enol ether protolysis and Paquette notes that following minimal enol ether purification, traces of the Tebbe reagent by-products may be present in the rearrangement medium. Given that aluminium reagents are indeed known to cause dramatic accelerations of [3,3]-Claisen rearrangements,4 some assistance to the rearrangement would not be surprising, though the nature of the active Lewis acid is not clear nor is the effect large given the reaction conditions (23 h at 175 °C in Scheme 1). Double bond migration can also compete with rearrangement forming non-productive allyl vinyl ethers.

We were interested to see if halogenation of the allylic fragment would accelerate the ring expansion and lower the reaction temperature. The literature suggests that γ,γ-diﬂuorination accelerates [3,3]-Claisen rearrangements in some cases;5 though the idea has not been tested fully and definitive theoretical work does not exist, a general consensus exists that the rehybridisation of an sp2 CF2 centre to sp3 is favoured as geminal CF2-substitution destabilises the alkene. Unambiguous accelerative effects exerted in γ,γ-diﬂuorinated systems have also been reported in other rearrangement systems including [2,3]-Wittigs,6 heteroatomatic [2,3]-rearrangements7 and oxy-Cope8 reactions.

Stork enamine chemistry allowed the syntheses of ketoesters 2a and 2b and the addition of 1-chloro-2,2-difluoroethylolithium9 generated from 1-chloro-2,2,2-trifluoroethane afforded good yields of alcohol products 3a as a mixture of diastereoisomers (Scheme 2); we assume that the major product has a trans-relationship between the sidechains.3

No lactone products were detected; what we expected was the collapse of the initial alkoxide onto the ester in at least one of the diastereoisomers. After isolating the hydroxyester, we tried to lactonise it under acidic and basic conditions but failed. Instead of hydrolysing the acid, we synthesised tert-butyl ester 3c as a mixture of diastereoisomers; a single lactone, to which we assigned structure 4a, was isolated in moderate yield after treatment of these hydroxyesters with trifluoroacetic acid in dichloromethane. Presumably, alkyl-O cleavage and carboxy protonation allows closure to the lactone; it seems less likely that the acid intercepts an allylic cation generated by protonation of, and departure of water from, the hydroxyacid. A similar successful sequence started from 1,2,2,2-tetrafluoroethane (HFC-134a) and afforded lactone 4b.10

With the lactone in hand, we attempted the Tebbe reaction† (Scheme 3) and succeeded in transforming lactone 4a completely to a new product as evinced by 19F NMR, to which we assign structure 5.

**Scheme 1** Reagents and conditions: i, vinylmagnesium bromide; ii, Cp2TiCl2(CH2)3AlMe3 (Tebbe reagent); iii, 175 °C, 23 h.

**Scheme 2** Reagents and conditions: i, ethyl acrylate or tert-butyl acrylate, dioxane, rt, 18 h; ii, 1-chloro-2,2-difluoroethenylolithium or 1,2,2-trifluoroethenylolithium, THF, −78 °C to rt; iii, CF3CO2H, THF, rt.

**Scheme 3** Reagents and conditions: i, Cp2TiCl2(CH2)3AlMe3, PhMe–THF, 0 to 60 °C; ii, xylene, sealed base-washed tube, 85 °C, 4 h. * Estimated yield based on clean 19F NMR and full conversion of 4a.
In contrast, the fluorinated congener 4b decomposed completely under Tebbe conditions. All attempts to purify and rigorously characterise 5 resulted in decomposition so we simply heated 5 at 85 °C. Complete consumption took place within 4 h and a new UV-active product 6 was formed. The rearrangement conditions contrast sharply with those in Scheme 1. We contend that this represents an unusually facile [3,3]-Claisen ring expansion; as the Claisen rearrangement is not usually reversible, it is hard to see how dehydrofluorination after the rearrangement could accelerate the rearrangement itself. Enol formation from the initial ketonic product followed by elimination of HF does result in the formation of a conjugated system so dienone formation is not surprising.

We are currently in pursuit of 7 in which the fluorine atoms are in the vinyl component of the Claisen precursor, so that we can compare the positional effects of fluorine atom substitution and retain both fluorine atoms in a structurally novel product.

In summary, we have shown that the rearrangement precursor can be assembled rapidly, that the fluoroallyl fragment survives Tebbe reaction conditions and that fluorine atoms at C-γ in the allyl fragment appear to lower considerably the barrier to ring expansion Claisen rearrangement.

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Notes and references

† Tebbe procedure and 19F NMR data for 5: Tebbe reagent (2.2 ml of a 0.5 M solution in toluene, 1.1 mmol) was added to a stirred solution of lactone 4a (0.236 g, 1.0 mmol) in dry THF (10 ml) at 0 °C. Fifteen min after the addition the mixture was allowed to warm to rt and stirred for a further 20 min. The mixture was then heated to 60 °C for 30 min, then cooled to 0 °C before being quenched with 15% NaOH solution (5 ml). After gas evolution had ceased, ether (10 ml) was added and the mixture was dried, filtered and evaporated. The residue was eluted through a short column of alumina with petroleum ether to give 5 (0.20 g, 89% estimated) as the only fluorine-containing product: δF (282 MHz, CDCl3) −81.8 (d, 3JF,FF 38.7, 1F); −84.9 (d, 3JF,FF 38.7, 1F). As 5 was expected to be unstable, it was taken on directly without further purification.

‡ Thermal rearrangement and selected data for 6: 5 (0.234 g, 1.0 mmol) was dissolved in xylene (5 ml) in an Ace tube that had been washed with NaOH (5 M aqueous solution). The tube was then sealed and heated to 85 °C in an oil bath. The reaction was followed by 19F NMR of aliquots until the starting material was consumed completely (4 h). Concentration and purification via silica gel column chromatography (10% ether in light petroleum) gave 6 (0.116 g, 54%) as a yellow oil; δF (300 MHz, CDCl3) 5.87 (d, 3JFF 20.2, 1H), 3.03–2.88 (m, 1H), 2.87–2.53 (m, 2H), 2.42–2.30 (m, 1H), 2.20–2.02 (m, 1H), 1.98–1.05 (m, 4H); δF (282 MHz, CDCl3) −83.9 (d, 3JFF 20.0); vmax (film/cm−1) 1662, 1627; [HRMS (Cl, M±H]+] Found: 232.0912. Calc. For C6H14F2Cl: 232.0905; m/z (Cl) 232 (M±H)+; λmax (CH3Cl) 284 nm (ε 3600 dm3 mol−1 cm−1).11

10 In this case alone, the two diastereoisomers of 3d could be separated. Only one of them lactonised (TFA in CH2Cl2) while the other decomposed completely.
11 The only similar compound we were able to find with reported UV data was 1-ethoxycarbonyl-5-oxo-1.3-cyclooctadiene ([λmax (95% EtOH) 282 nm (ε 9700 dm3 mol−1 cm−1)]; see W. C. Agosta and W. W. Lowrance, J. Org. Chem., 1970, 35, 3851.