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Anomalous Z-selectivity in Wittig reactions of keto-stabilised ylides with ortho-heteroatom substituted benzaldehydes

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Anomalous Z-selectivity in Wittig reactions of keto-stabilised ylides with ortho-heteroatom substituted benzaldehydes

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ABSTRACT

Wittig reactions of keto-stabilised ylides with ortho-substituted benzaldehydes are found to show significantly higher Z-selectivity than analogous reactions of the same ylides with benzaldehyde itself. A cooperative effect is seen whereby the unusual Z-selectivity is further augmented if the ylide bears greater steric bulk in the α′-position. These results are consistent with our previous observations on reactions of all ylide types with aldehydes bearing a β-heteroatom. Significantly, the cooperative effect, previously limited to semi-stabilised ylides, has now been extended to stabilised ylides. Both the anomalous Z-selectivity and the cooperative effect can be rationalised within the [2+2] cycloaddition mechanism of the Wittig reaction.

1. Introduction

The Wittig reaction is perhaps the most commonly used method for the synthesis of alkenes. The many and varied applications of it and the extensive work that has been done on its mechanism can be found in various excellent reviews. A list of the very latest developments in the Wittig reaction can be found in our recent publication on the mechanism. In the latter work, we believe that we have definitively shown the unity of the mechanism of the Li-salt free Wittig reaction. By this we mean that all ylide types (unstabilised, semi-stabilised and stabilized) react by the same mechanism. Our conclusions were based on the results of Wittig reactions of β-heteroatom substituted aldehydes, which showed unexpectedly high selectivity for Z-alkene for all ylide types (see Scheme 1). Such selectivity is not seen in reactions of the same ylides with analogous aldehydes lacking the suitably placed β-heteroatom.

We showed that this change of selectivity, which was rather dramatic in many cases, could be easily accommodated within the emerging consensus on the mechanism of Li-salt free Wittig reactions: [2+2] cycloaddition of ylide and aldehyde to give oxaphosphetane (OPA) followed by cycloreversion to phosphine oxide and alkene (see Scheme 2). Therefore our results also provided strong experimental support for that mechanism. Our explanation for the heteroatom-induced increase in Z-selectivity involved an additional bonding interaction in the cis-selective cycloaddition transition state (TS), between phosphorus and the β-heteroatom, as shown in Fig. 1(a).7

Scheme 1. Highly Z-selective Wittig reactions of β-heteroatom-substituted aldehydes under conditions promoting kinetic control.

A subsidiary, but important, part of our study concerned an unexpected co-operative effect seen in the case of semi-stabilised ylides. This was also explicable on the proposed TS model, and is illustrated by the data in Scheme 3: the placement of an ortho-

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Our objectives then were to determine (i) whether there would be an increase in Z-selectivity on substitution of a heteroatom β to the aldehyde, further confirming the generality of the main effect and (ii) whether a cooperative effect (further increase in Z-selectivity) would be observed in reactions of keto-stabilised ylides with a more heavily substituted α-carbon (carbon-3 of the oxoalkylidenemethyldiphenylphosphorane). The proposed cis-selective TS for these reactions, whose existence we sought to confirm, is analogous to the one proposed previously for reactions of non-stabilised, semi-stabilised, and ester-stabilised ylides with β-heteroatom substituted aldehydes (see Fig. 1(b)). As in the previous study, we chose to focus on ylides derived from methylidiphosphinite, since kinetic control has previously been established in reactions of such ylides; but a small number of triphenylphosphine-derived cases are also included for comparison, as are some ester-stabilised ylides.

We find that there is indeed evidence for a cooperative effect with these ylides, but at a reduced level. The new enones generated by these studies proved difficult to handle and it was with great difficulty that many of the Z:E ratios were obtained; some consistently underwent decomposition and, for all the Z-isomers were extremely prone to isomerisation to the E-isomer. Therefore, despite the precautions taken to accurately measure the “kinetic Z/E ratio” of each reaction, the ratios quoted below represent only a lower bound to the true Z-selectivity of these reactions.

2. Reactions of ortho-heteroatom substituted benzaldehydes with keto-stabilised & ester-stabilized ylides

All reactions were carried out using the standard set of reaction conditions established in our previous study, with the exception that, for solubility reasons, certain reactions were carried out at higher temperatures. The ylide was generated in situ from the parent phosphonium bromide or chloride salt using non-nucleophilic NaHMDS base in THF (giving insoluble NaCl or NaBr) under an atmosphere of dry nitrogen, and then cooled to -78 °C. The aldehyde was then added dropwise to the cold ylide solution, and the reaction was stirred at low temperature for 20 minutes. Some of the reactions were quenched at low temperature by addition of acid to ensure that the reaction had occurred at that temperature. The Z/E ratios of the alkenes produced in reactions of 2-oxoalkylidenemethylidiphosphinites with selected benzaldehydes are shown in Table 1.

Acetonyldimenyldiphosphinite (6, R = CH, in the diagram accompanying Table 1) was found to be insoluble in THF below -50 °C and attempted Wittig reactions with it at temperature below that (with low temperature quenching) yielded no alkene product. Therefore its reactions were carried out at -45 °C or 20 °C. The reactions of this ylide with benzaldehyde (2a) showed high E-selectivity (Table 1 entries 1 & 2), while its reactions with ortho-heteroatoom substituted benzaldehydes (2c–d) showed a moderate increase in Z-selectivity (up to 30 or 40% Z-isomer produced, see Table 1 entries 3 & 4). Thus the aldehyde
ortho-heteroatom-induced increase in Z-selectivity previously observed in reactions of non-stabilised, semi-stabilised and ester-stabilised ylides is also observed in reactions of keto-stabilised ylides. The magnitude of the shift in the energy of the cis-selective TS is not as great as was generally observed in our previously disclosed results.\(^7\)

Table 1. Z/E ratio\(^a\) for reactions of selected 2-oxoalkylidemethyldiphenylphosphoranes 6-8 (generated in situ from the corresponding phosphonium salts 3-5)\(^b\) with selected benzaldehydes 2a-d:

<table>
<thead>
<tr>
<th>Entry</th>
<th>Ylide R</th>
<th>Aldehyde Y</th>
<th>Temp °C</th>
<th>Z/E ratio</th>
</tr>
</thead>
<tbody>
<tr>
<td>1(^i)</td>
<td>CH₃</td>
<td>H</td>
<td>-45</td>
<td>19:81</td>
</tr>
<tr>
<td>2(^i)</td>
<td>CH₃</td>
<td>H</td>
<td>20</td>
<td>20:80</td>
</tr>
<tr>
<td>3(^i)</td>
<td>CH₃</td>
<td>Cl</td>
<td>-45</td>
<td>33:67</td>
</tr>
<tr>
<td>4(^i)</td>
<td>CH₃</td>
<td>Br</td>
<td>20</td>
<td>40:60</td>
</tr>
<tr>
<td>5(^i)</td>
<td>CH₂Cl</td>
<td>H</td>
<td>-78</td>
<td>12:88</td>
</tr>
<tr>
<td>6(^i)</td>
<td>t-Bu</td>
<td>H</td>
<td>-78</td>
<td>17:83</td>
</tr>
<tr>
<td>7(^i)</td>
<td>CH₂Cl</td>
<td>Cl</td>
<td>-78</td>
<td>50:50</td>
</tr>
<tr>
<td>8(^i)</td>
<td>CH₂Cl</td>
<td>Br</td>
<td>-78</td>
<td>50:50</td>
</tr>
<tr>
<td>9(^i)</td>
<td>t-Bu</td>
<td>Br</td>
<td>-78</td>
<td>48:52</td>
</tr>
<tr>
<td>10(^i)</td>
<td>t-Bu</td>
<td>I</td>
<td>-78</td>
<td>50:50</td>
</tr>
</tbody>
</table>

\(^a\) Z/E ratio determined by \(^1\)H NMR analysis of the crude product obtained after aqueous work-up of the reaction mixture. See ESI for full details.

\(^b\) Phosphonium salt counter-ion Z = Cl except where otherwise noted.

\(^c\) Ylides (6-8) were generated at 20°C from the parent phosphonium salts (3-5) in dry THF by treatment with NaHMDS. The ylide solution was then brought to the reaction temperature indicated, and the appropriately substituted benzaldehyde was added dropwise. After stirring for 20 minutes, the reaction was quenched as indicated in footnotes d-f below.

\(^d\) Quenched with 5% aqueous HCl after 20 minutes at the reaction temperature indicated.

\(^e\) Reactions at 20 °C stirred for 4 hours, then quenched with 5% aqueous HCl.

\(^f\) Removed from cooling bath after 20 minutes, and stirred for 12 hours at room temperature before being quenched with 5% aqueous HCl.

\(^g\) Counter-ion Z = Br.

Ylides with one or more substituents on carbon-3 of the 2-oxoalkylidene group (7 and 8) gave rise to increased E-selectivity in their reactions with benzaldehyde (Table 1 entries 5 & 6) but with ortho-heteroatom-stabilised benzaldehydes 2bd-d (Table 1 entries 7-10) they gave considerably increased Z-selectivity by comparison, indicating the operation of the same cooperative effect previously observed in reactions of semi-stabilised ylides with benzaldehydes.\(^7\) The unsubstituted acetonyle (6) reacts with benzaldehyde (2a) with reasonably high E-selectivity, and reacts with ortho-heteroatom substituted benzaldehydes (2c-d) with somewhat increased Z-selectivity. By comparison, the shift in Z-selectivity from the highly E-selective reactions of 7 and 8 (which have greater steric bulk at carbon-3 of the 2-oxoalkylidene moiety) with benzaldehyde to the reactions of the same ylides with ortho-heteroatoms substituted benzaldehydes 2b-d is much greater.

The Z/E ratios for alkenes produced in reactions of keto-stabilised ylides derived from triphenylphosphine are shown in Table 2. These were carried out using pre-formed ylide (i.e. not generated in situ) at -78 °C but not quenched until they had been allowed to warm to room temperature. Thus the actual reaction temperature is not certain for these reactions.

Table 2. Z/E ratio\(^a\) for alkenes produced in the reactions of selected acetyldienetriphenylphosphoranes 16 and 17\(^c\) and (tert-butoxycarbonyl)ethylidene triphenylphosphorane 18\(^d\) with selected benzaldehydes 2a and 2d-f:

<table>
<thead>
<tr>
<th>Entry</th>
<th>Ylide R</th>
<th>Aldehyde Y</th>
<th>Enone Z/E ratio</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>CH₃</td>
<td>H</td>
<td>3:97</td>
</tr>
<tr>
<td>2</td>
<td>CH₃</td>
<td>Br</td>
<td>11:89</td>
</tr>
<tr>
<td>3</td>
<td>CH₃</td>
<td>OMe</td>
<td>10:90</td>
</tr>
<tr>
<td>4</td>
<td>CH₂OMe</td>
<td>Br</td>
<td>17:83</td>
</tr>
<tr>
<td>5(^c)</td>
<td>O(t-Bu)</td>
<td>H</td>
<td>3:97</td>
</tr>
<tr>
<td>6(^c)</td>
<td>O(t-Bu)</td>
<td>Me</td>
<td>3:97</td>
</tr>
<tr>
<td>7(^c)</td>
<td>O(t-Bu)</td>
<td>Br</td>
<td>20:80</td>
</tr>
<tr>
<td>8(^c)</td>
<td>O(t-Bu)</td>
<td>OMe</td>
<td>19:81</td>
</tr>
</tbody>
</table>

\(^a\) Z/E ratio determined by \(^1\)H NMR analysis of the crude product obtained after aqueous work-up of the reaction mixture. See ESI for full details.

\(^b\) Pre-formed ylides were used unless otherwise indicated.

\(^c\) Ylide 18 was generated in situ from the (commercially available) parent phosphonium salt (tert-butoxycarbonyl)methyltriphenylphosphonium chloride 19. Reactions of this ylide were carried out at 20 °C.

\(^d\) Unless otherwise indicated, reactions were stirred at -78 °C for 20 minutes and then allowed to warm to 20 °C. The Z/E ratios were determined by \(^1\)H NMR analysis of the crude product obtained after aqueous work-up of the reaction mixture. See Supporting Information for full details.

This result was reported in reference 22.

The reaction of acetonylendienetriphenylphosphorane (16) with benzaldehyde (2a) yields the expected very high E-selectivity in reaction with benzaldehyde (Table 2 entry 1), while its reactions with ortho-heteroatom substituted benzaldehydes (2d-e) are somewhat more Z-selective (Table 2 entries 2 & 3). The reaction of 3-methoxy-2-oxopropylidene triphenylphosphorane (17) with 2-bromobenzaldehyde (2d) shows a further increase in Z-selectivity (Table 2 entry 4). Thus the trends observed for the reactions of 2-oxoalkylidemethyldiphenylphosphoranes detailed in Table 1 are reproduced here – including the operation of the cooperative effect – albeit with far less dramatic shifts in Z-selectivity.

Also included in Table 2 are some results from reactions of an ester-stabilised ylide (18), two of which have been reported in a previous publication.\(^2\) This ylide reacts with each of benzaldehyde (2a) and 2-methylbenzaldehyde (2f) in THF at 20 °C with, again, very high E-selectivity (see Table 2 entries 5 & 6) while its reactions with ortho-heteroatom substituted benzaldehydes are somewhat more Z-selective (Table 2 entries 7 & 8), again clearly demonstrating the operation of the ortho-heteroatom effect. The absence of the effect from the reaction of 2-methylbenzaldehyde (Table 2 entry 6) indicates that the
operation of the effect, as in our previous study, is dependent on the ortho-substituent of the aldehyde bearing a lone pair of electrons.

3. Discussion

It has previously been established that reactions of ester-stabilised methylidiphosphonium ylides are under kinetic control. Although irreversibility of OPA formation has not been explicitly demonstrated in reactions of keto-stabilised ylides, it seems reasonable by analogy to assume that the reactions detailed here are irreversible, especially in light of the increased Z-selectivity observed here, which occurred in polar-aprotic THF solvent where high E-selectivity is the norm. Since the ortho-heteroatom effect observed in our previous study is also present in reactions of 2-oxoalkylidenemethylidiphosphonanes, and since the ortho-heteroatom-induced increase in Z-selectivity is augmented by greater steric bulk at carbon-3 of the ylide oxoalkylidene moiety (cooperative effect), we conclude that the anomalous Z-selectivity in these reactions is a consequence of the existence of a cis-selective TS (shown in Fig. 1(b) above) similar to that proposed by us previously.

In the proposed TS, the new factor is a bonding interaction between the phosphorus and the ortho-heteroatom, causing the TS to be of sufficiently low energy that it can become competitive with the normally favoured trans-selective cycloaddition TS. The crowded environment about phosphorus in the proposed TS, results in a significant reorganisation of the phosphorus-substituents in comparison with other Wittig TSs. The position of the aldehyde aryl group is dictated by the phosphorus-heteroatom bond, and, as a result, 1-3 steric interactions (between the aryl group and the ylide phosphorus substituents, Fig. 1(b)) are relatively unimportant in this TS. However, there is a concomitant increase in the potential for 2-3 steric interactions, which are minimized if the ylide α-carbon substituent occupies a pseudo-apical position in the forming ring. Furthermore, such a TS would benefit from the favourable anti-parallel orientation of the dipoles along the aldehyde C=O and ylide C(=O)O bonds, respectively, as is proposed to occur in E-selective reactions of stabilised ylides. As a result of the above factors, this TS is cis-selective.

The increased Z-selectivity in reactions of ortho-heteroatom substituted benzyldenylides with ylides that have greater steric bulk at carbon-3 of the oxoalkylidene moiety (cooperative effect) can be rationalised using the same TS arguments. Thus the increased 2-3-interactions that would be engendered by the larger substituent on the ylide α-carbon result in a greater bias towards the cis-selective TS since it is better able to accommodate steric bulk in this position. That the cooperative effect exists in reactions of keto-stabilised ylides but not in reactions of ester-stabilised ylides is consistent with our initial working hypothesis: that the alkyl group of the ester moiety in ester-stabilised ylides is too far removed from the ylide α-carbon for its steric bulk to have a bearing on the shape of the TS, while for keto-stabilised ylides the closer proximity of the variable alkyl group to the ylide α-carbon means that differences in this group result in different stereoselectivity in the cycloaddition step.

The existence of the proposed TS is also consistent with selectivity being dependent on the through-space phosphorus-heteroatom bond, and thus essentially independent of the electrophilicity of the carbonyl group – i.e. the effect is not a result of through-bond electronic effects exerted by the ortho-heteroatom.

In the reactions of β-heteroatoms substituted aldehydes that we have studied, the steric effects of the phosphorus substituents, and the influence of these effects on the observed diastereoselectivity, are consistent across all ylide types, including keto-stabilised ylides. We had previously noted that the magnitude of the shift in selectivity for Z-alkene or cis-OPA in reactions of β-heteroatom substituted aldehydes (compared with analogous unsubstituted aldehydes) decreases in line with the steric bulk of the phosphonium moiety. In the context of our proposed TS model, it is easy to envisage how sterically bulky substituents on phosphorus would reduce the efficacy of phosphorus-heteroatom bonding. In reactions of non-stabilised ylides, the magnitude of this shift decreases in the order PhDBP > Ph3P > EtPhP; for semi-stabilised ylides the trend is PhDBP ≈ MePh3P > Ph3P; and for both ester-stabilised and keto-stabilised ylides we are now in a position to conclude that the trend is MePh3P > Ph3P. The consistency and generality of these results supports the existence of the proposed TS.

4. Conclusion

Reactions of keto-stabilised ylides and triphenylphosphine-derived ester-stabilised ylides with ortho-heteroatom substituted benzaldehydes show an anomalous increase in Z-selectivity compared with the reactions of the same ylides with benzaldehyde itself or with 2-methylbenzaldehyde. These results are entirely consistent with those previously observed in reactions of non-stabilised, semi-stabilised and ester-stabilised ylides with β-heteroatom substituted ylides, although the magnitudes of the shifts towards Z-selectivity are not as great in the present study. Most significantly, we have demonstrated that the cooperative effect that operates in reactions of benzylides with ortho-heteroatoms substituted benzaldehydes is replicated in reactions of keto-stabilised ylides with the same aldehydes. We conclude that these reactions are irreversible, and all occur through a similar cis-selective [2+2] cycloaddition transition state in which a bond exists between the phosphorus and the β-heteroatom substituent on the aldehyde. The consistency of the results further strengthens the argument that all Li salt-free Wittig reactions occur by the [2+2] cycloaddition mechanism proposed by Vedejs and co-workers, and modified for reactions of stabilised ylides by Aggarwal, Harvey and co-workers.

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Supplementary Data

A Supplementary Data document is available containing details of experimental procedures, assignments of alkene Z/E ratios, and full characterization of all new phosphonium salts and alkenes.
References

18. For further details of the experimental evidence that has been amassed in support of the [2+2] cycloaddition mechanism (including our own contribution to this body of evidence), as well as details of historical significance to its development, we refer the interested reader to reference 7.
19. As an aside, we note that betaines are not involved in the mechanism of the Li-salt free Wittig but, if generated through a non-Wittig route, will collapse to OPA and hence give phosphine oxide and alkene as shown as Scheme 2.
20. As a further aside, we also note that almost all Li-salt free Wittig reactions are under kinetic control. As also shown in Scheme 2, a very small number of reactions of non-stabilised ylides are known in which the formation of cis-OPA is reversible, resulting in enhanced production of trans-OPA (and hence E-alkene) relative to the initial amount formed of this intermediate. Most of these involve ethylides and benzaldehydes - see reference 7 for a full discussion of the phenomenon and an exhaustive list of the examples for which it is known to operate.
21. Reaction vessels and NMR tubes were protected from light where possible by being wrapped in aluminium foil. NMR samples of the crude reaction mixtures were obtained as soon as the work-up of the reaction was complete.