

## Novel aziridination of $\alpha$ -halo ketones: an efficient nucleophile-induced cyclization of phosphoramidates to functionalized aziridines

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### Abstract

A novel and efficient aziridination of  $\alpha$ -halo ketones is reported. The reaction of  $\alpha$ -halo ketones with diethyl *N*-arylphosphoramidates affords diethyl *N*-aryl-*N*-(2-oxoalkyl)phosphoramidates which undergo reductive ( $\text{H}^-$ -induced) cyclization with sodium borohydride followed by sodium hydride to give 1,2-disubstituted and 1,2,3-trisubstituted aziridines. The cyclization induced by  $\text{NCS}^-$  or  $\text{PhS}^-$  affords substituted aziridines functionalized at C-2. The reactions give excellent yields and are highly diastereoselective in favour of *cis* aziridines. © 2007 Elsevier Ltd. All rights reserved.

**Keywords:** Aziridines;  $\alpha$ -Halo ketones; Phosphoramidates; Nucleophile-induced; Reduction; Cyclization reactions

Owing to the inherent strain in small ring heterocycles, they are useful as feedstocks in organic synthesis to provide functionalized carbon chains. Aziridines have been synthetic targets as well as building blocks in synthesis since Gabriel's 1888 discovery of the smallest nitrogen-containing heterocycle.<sup>1–3</sup> In terms of synthetic transformations, the utility of aziridines derives from their selective ring-opening reactions,<sup>4,5</sup> which often form the basis for more complex target syntheses.<sup>6,7</sup> Yudin and co-workers have demonstrated a number of connections between aziridines and the products of their ring-opening.<sup>8</sup>

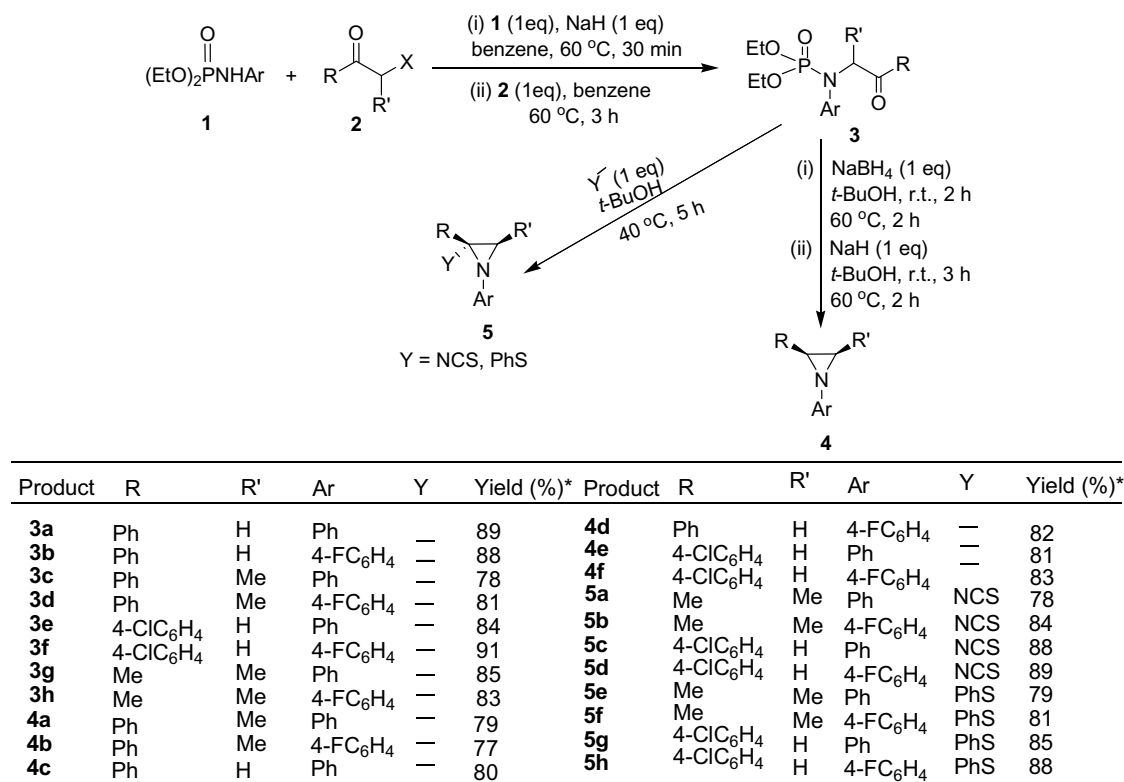
Antibiotic, anticancer and antitumour activities of aziridine-containing natural products, such as azinomycins,<sup>9–11</sup> mitomycins,<sup>12,13</sup> FR-900482, FR-66979 and related compounds,<sup>14</sup> are of significant interest. Of these, azinomycins possess a broad spectrum of activity against cancers, including tumours, and mitomycin C has been in clinical use since the 1960s for the treatment of a wide range of tumours. FR-900482 and FR-66979 are structurally related

to mitomycins and show similar anticancer activity. The activity of all these compounds lies in the role of aziridines as powerful alkylating agents.<sup>15</sup> Furthermore, several synthetic aziridines have also been reported to exhibit useful biological properties such as irreversible inhibition of glutamate racemase,<sup>16</sup> and diaminopimelic acid epimerase (DAP),<sup>17</sup> and high level cytotoxicity against melanoma cell lines.<sup>18</sup>

Numerous methods are available for differently substituted aziridines, which include aziridination of olefins,<sup>19–21</sup> carbene and ylide addition to imines,<sup>22–24</sup> and cyclization of  $\beta$ -amino alcohols,<sup>25,26</sup>  $\alpha$ -halo imines,<sup>27</sup>  $\beta$ -amino halides,<sup>25,26</sup> and  $\beta$ -azido alcohols.<sup>25,26,28</sup> Very recently, an excellent review by De Kimpe and co-workers covered the synthesis and reactivity of C-heteroatom-substituted aziridines,<sup>29</sup> which also includes C-sulfur-substituted aziridines similar to those reported in the present Letter.  $\alpha$ -Halo imines are important intermediates in aziridine synthesis, which involves a two-step process starting from  $\alpha$ -halo ketones or aldehydes.

In continuation of our efforts to develop new one-pot cyclization processes,<sup>30</sup> we report herein a two-step synthetic protocol for the synthesis of aziridines starting from

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\* Yields of isolated and purified products.

Scheme 1. Synthesis of aziridines **4** and **5** from  $\alpha$ -halo ketones **2**.

$\alpha$ -halo ketones. The synthesis involves a novel one-pot nucleophile-induced intramolecular cyclization of  $\alpha$ -halo ketone-derived phosphoramidates **3** to yield aziridines **4** and **5** (Scheme 1). Thus, treatment of diethyl *N*-aryl-phosphoramidates **1** with sodium hydride in benzene followed by  $\alpha$ -halo ketones **2** afforded diethyl *N*-aryl-*N*-(2-oxoalkyl)phosphoramidates **3**<sup>31</sup> in 78–91% yields. Conventional reduction of phosphoramidates **3** with sodium borohydride afforded the corresponding alcohols in addition to a small amount of aziridines **4**. Thus, phosphoramidates **3** were reduced with sodium borohydride in *t*-butanol followed by treatment with sodium hydride in the same vessel to produce the corresponding aziridines **4**<sup>32</sup> in 77–83% yields (Scheme 1). Furthermore, phosphoramidates **3**, on reaction with potassium thiocyanate, or sodium thiophenolate in *t*-butanol afforded functionalized aziridines **5**<sup>33</sup> in 78–89% yields (Scheme 1). The synthesis of a few aziridines similar to **5** has been previously reported from 1,3-diphenyl-2-chloroaziridines via displacement of the chloride ion by hydride or thiolates,<sup>29,34,35</sup> however, the present method provides a direct access to this class of aziridines.

The nucleophile-induced cyclization of (2-oxoalkyl)-phosphoramidates **3** to aziridines **4** and **5** was highly diastereoselective in favour of the *cis* isomers. Diastereomeric ratios in the crude isolates were checked by <sup>1</sup>H NMR to note any alteration of these ratios during subsequent purification. The crude isolates of **4a**, **4b**, **5a**, **5b**, **5e** and **5f** were

found to be diastereomeric mixtures containing 94–97% of the *cis* isomer. Strong NOEs were observed between 2-H/3-H and 2-Me/3-Me of these aziridines, which conclusively demonstrate their *cis* stereochemistry (Fig. 1). The *cis* stereochemistry of aziridines **4** was also supported by comparison of the *J* values of 2-H and 3-H with those reported in the literature.<sup>27g</sup>

The formation of aziridines **4** is best explained through intramolecular attack of the alkoxide ion **7** on the phosphorus atom (Scheme 2). This assumption is supported by the exclusive formation of aziridines **4** on addition of a base, such as sodium hydride, to facilitate alkoxide ion formation. On the other hand, aziridines **5** are formed by

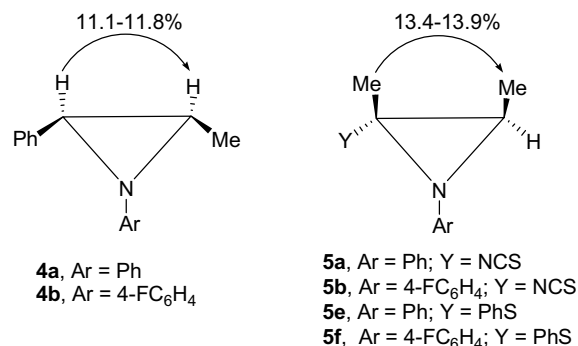
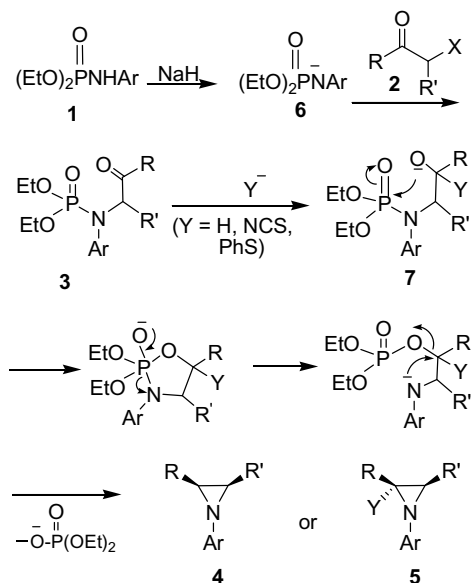


Fig. 1. NOE experiments on aziridines **4** and **5**.



Scheme 2. Tentative mechanism for the nucleophile-induced cyclization of phosphoramidates **3** to aziridines **4** and **5**.

attack of a nucleophile ( $NCS^-$  or  $PhS^-$ ) on the carbonyl carbon of **3** followed by intramolecular attack of the alkoxide ion on the phosphorus atom (Scheme 2).

In summary, we have developed a general and efficient method for the synthesis of substituted and functionalized aziridines by nucleophile-induced cyclization of readily available  $\alpha$ -halo ketone-derived phosphoramidates in a one-pot procedure, which may find application in organic synthesis.

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- General procedure for the synthesis of diethyl N-aryl-N-(2-oxoalkyl)phosphoramidates 3*: To a solution of diethyl N-arylphosphoramidate **1** (5 mmol) in dry benzene (5 mL) was added dropwise a solution of sodium hydride (120 mg, 5 mmol) in dry benzene (10 mL) with stirring at rt. After the addition was complete and the evolution of hydrogen gas (effervescence) had ceased, the reaction mixture was stirred at 60 °C for 30 min and then cooled to rt. Next, a solution  $\alpha$ -halo ketone **2** (5 mmol) in dry benzene (5 mL) was added and the reaction mixture was stirred at 60 °C for 3 h. The solvent was evaporated under reduced pressure, the residue washed with water and crystallized from *n*-hexane to afford an analytically pure sample of **3**. Physical data of representative compounds: Compound **3a**: White crystals, Yield 89%, mp 161–163 °C. IR (KBr)  $\nu_{max}$  3046, 2951, 1745, 1599, 1508, 1463, 1383, 758, 704  $cm^{-1}$ .  $^1H$  NMR (400 MHz;  $CDCl_3/TMS$ )  $\delta$ : 1.19 (t, 6H,  $J = 7.4$  Hz,  $2 \times Me$ ), 4.12 (q, 4H,  $J = 7.4$  Hz,  $2 \times OCH_2$ ), 4.57 (s, 2H,  $CH_2$ ), 7.13–7.92 (m, 10H<sub>arom</sub>).  $^{13}C$  NMR (100 MHz;  $CDCl_3/TMS$ )  $\delta$ : 15.3, 52.7, 61.2, 125.9, 126.7, 128.5, 130.7, 131.9, 132.8, 135.7, 138.3, 193.7. EIMS ( $m/z$ ): 347 ( $M^+$ ). Anal. Calcd for  $C_{18}H_{22}NO_4P$ : C, 62.24; H, 6.38; N, 4.03. Found: C, 62.58; H, 6.27; N, 4.22. Compound **3g**: White crystals, Yield 85%, mp 143–145 °C. IR (KBr)  $\nu_{max}$  3042, 2949, 1748, 1603, 1511, 1460, 1378, 1035, 761, 706  $cm^{-1}$ .  $^1H$  NMR (400 MHz;  $CDCl_3/TMS$ )  $\delta$ : 1.21 (t,

- 6H,  $J = 7.4$  Hz,  $2 \times \text{Me}$ ), 1.45 (d, 3H,  $J = 7.8$  Hz, Me–C), 2.12 (s, 3H, Me–CO), 3.72 (q, 1H,  $J = 7.8$  Hz, CH), 4.10 (q, 4H,  $J = 7.4$  Hz,  $2 \times \text{OCH}_2$ ), 6.93–7.21 (m, 5H<sub>arom</sub>).  $^{13}\text{C}$  NMR (100 MHz;  $\text{CDCl}_3/\text{TMS}$ )  $\delta$ : 15.5, 16.3, 27.2, 57.6, 61.5, 122.9, 126.4, 129.9, 137.5, 193.6. EIMS ( $m/z$ ): 299 ( $\text{M}^+$ ). Anal. Calcd for  $\text{C}_{14}\text{H}_{22}\text{NO}_4\text{P}$ : C, 56.18; H, 7.41; N, 4.68. Found: C, 55.91; H, 7.19; N, 4.82.
32. **General procedure for the synthesis of substituted aziridines 4:** A mixture of phosphoramidate **3** (5 mmol) and sodium borohydride (190 mg, 5 mmol) in dry *t*-butanol (25 mL) was stirred at rt for 2 h, then at 60 °C for 2 h. Sodium hydride (120 mg, 5 mmol) was added to the above reaction mixture at rt. Stirring was continued for 3 h at rt, then at 60 °C for 2 h. Water (30 mL) was added, the mixture was extracted with ether ( $3 \times 50$  mL), the combined organics dried over anhydrous sodium sulfate, filtered and evaporated to dryness. The crude product thus obtained was crystallized from *n*-hexane twice to afford an analytically pure sample of **4**. Physical data of representative compounds: Compound **4a**: White crystals, Yield 79%, mp 30–31 °C. IR (KBr)  $\nu_{\text{max}}$  3040, 2943, 1605, 1508, 1462, 1380, 1032, 762, 703  $\text{cm}^{-1}$ .  $^1\text{H}$  NMR (400 MHz;  $\text{CDCl}_3/\text{TMS}$ )  $\delta$ : 1.40 (d, 3H,  $J = 5.8$  Hz, Me), 3.02–3.12 (m, 1H, 3-H), 3.41 (d, 1H,  $J = 7.3$  Hz, 2-H), 6.89–7.18 (m, 10H<sub>arom</sub>).  $^{13}\text{C}$  NMR (100 MHz;  $\text{CDCl}_3/\text{TMS}$ )  $\delta$ : 21.3, 41.7, 55.1, 122.8, 123.9, 126.4, 132.5, 133.1, 135.5, 137.8, 138.9. EIMS ( $m/z$ ): 209 ( $\text{M}^+$ ). Anal. Calcd for  $\text{C}_{15}\text{H}_{15}\text{N}$ : C, 86.08; H, 7.22; N, 6.69. Found: C, 86.29; H, 7.08; N, 6.35. Compound **4e**: White crystals, Yield 81%, mp 41–43 °C. IR (KBr)  $\nu_{\text{max}}$  3043, 2942, 1601, 1511, 1465, 1381, 761, 706  $\text{cm}^{-1}$ .  $^1\text{H}$  NMR (400 MHz;  $\text{CDCl}_3/\text{TMS}$ )  $\delta$ : 1.73 (dd, 1H,  $J = 10.9, 7.5$  Hz, 3Ha), 2.13 (dd, 1H,  $J = 10.9, 5.9$  Hz, 3Hb), 2.87 (dd, 1H,  $J = 7.5, 5.9$  Hz, 2-H), 6.73–7.11 (m, 7H<sub>arom</sub>), 7.31–7.36 (m, 2H<sub>arom</sub>).  $^{13}\text{C}$  NMR (100 MHz;  $\text{CDCl}_3/\text{TMS}$ )  $\delta$ : 37.1, 45.4, 125.7, 126.7, 132.2, 133.5, 134.9, 136.2, 137.2, 139.5. EIMS ( $m/z$ ): 229, 231 ( $\text{M}^+$ ,  $\text{M}^+ + 2$ ). Anal. Calcd for  $\text{C}_{14}\text{H}_{12}\text{ClN}$ : C, 73.20; H, 5.27; N, 6.10. Found: C, 73.03; H, 5.39; N, 5.89.
33. **General procedure for the synthesis of functionalized aziridines 5:** A mixture of phosphoramidate **3** (5 mmol) and potassium thiocyanate, or sodium thiophenolate (5 mmol) in *t*-butanol (25 mL) was heated at 40 °C for 5 h. Water (30 mL) was added and the mixture was extracted with dichloromethane ( $3 \times 50$  mL), the combined organics dried over anhydrous sodium sulfate, filtered and evaporated to dryness. The crude product thus obtained was recrystallized from *n*-hexane twice to afford an analytically pure sample **5**. Physical data of representative compounds: Compound **5a**: White crystals, Yield 78%, mp 58–59 °C. IR (KBr)  $\nu_{\text{max}}$  3042, 2947, 1602, 1507, 1461, 1377, 1035, 759, 705  $\text{cm}^{-1}$ .  $^1\text{H}$  NMR (400 MHz;  $\text{CDCl}_3/\text{TMS}$ )  $\delta$ : 1.32 (d, 3H,  $J = 5.6$  Hz, 3-Me), 1.81 (s, 3H, 2-Me), 2.47 (q, 1H,  $J = 5.6$  Hz, 3-H), 6.81–7.10 (m, 5H<sub>arom</sub>).  $^{13}\text{C}$  NMR (100 MHz;  $\text{CDCl}_3/\text{TMS}$ )  $\delta$ : 19.3, 29.5, 48.7, 112.5, 123.1, 132.3, 133.7, 138.7. EIMS ( $m/z$ ): 204 ( $\text{M}^+$ ). Anal. Calcd for  $\text{C}_{11}\text{H}_{12}\text{N}_2\text{S}$ : C, 64.67; H, 5.92; N, 13.71. Found: C, 64.35; H, 5.97; N, 13.81. Compound **5g**: White crystals, Yield 85%, mp 79–81 °C. IR (KBr)  $\nu_{\text{max}}$  3041, 2948, 1601, 1510, 1460, 1383, 765, 703  $\text{cm}^{-1}$ .  $^1\text{H}$  NMR (400 MHz;  $\text{CDCl}_3/\text{TMS}$ )  $\delta$ : 1.99 (d, 1H,  $J = 11.1$  Hz, 3-Ha), 2.56 (d, 1H,  $J = 11.1$  Hz, 3-Hb), 6.93–7.21 (m, 12H<sub>arom</sub>), 7.39–7.42 (m, 2H<sub>arom</sub>).  $^{13}\text{C}$  NMR (100 MHz;  $\text{CDCl}_3/\text{TMS}$ )  $\delta$ : 44.3, 60.5, 122.6, 124.6, 126.7, 130.5, 131.7, 132.5, 133.7, 134.6, 136.8, 138.1, 138.8, 139.7. EIMS ( $m/z$ ): 337, 339 ( $\text{M}^+$ ,  $\text{M}^+ + 2$ ). Anal. Calcd for  $\text{C}_{20}\text{H}_{16}\text{ClNS}$ : C, 71.10; H, 4.77; N, 4.15. Found: C, 71.37; H, 4.61; N, 4.29.
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