

Elimination Reactions of Aryl Furylacetates Promoted by $R_2NH-R_2NH_2^+$ in 70 mol % MeCN(aq). Effects of β -Aryl on the Ketene-Forming Transition-State

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Ketene-forming elimination from 2-X-4-nitrophenyl furylacetates (**1a-d**) promoted by $R_2NH-R_2NH_2^+$ in 70 mol % MeCN(aq) has been studied kinetically. When X = Cl and NO₂, the reactions exhibited second-order kinetics as well as Brønsted $\beta = 0.37-0.54$ and $|\beta_{lg}| = 0.31-0.45$. The Brønsted β decreased with a poorer leaving group and $|\beta_{lg}|$ increased with a weaker base. The results are consistent with an E2 mechanism. When the leaving group was changed to a poorer one [X = H (**1a**) and OCH₃ (**1b**)], the reaction mechanism changed to the competing E2 and E1cb mechanisms. A further change to the E1cb mechanism was realized for the reaction of **1a** with *i*-Pr₂NH/*i*-Pr₂NH₂⁺ in 70 mol % MeCN-30 mol % D₂O. By comparing the kinetic results in this study with the existing data for ArCH₂C(O)OC₆H₃-2-X-4-NO₂, the effect of the β -aryl group on the ketene-forming elimination was assessed.

Key Words : Elimination, E2 and E1cb Mechanism, β -Aryl Group effect

Introduction

Extensive studies of the structure-reactivity relationships in the ketene-forming elimination reactions have led to the qualitative understanding of the relationship between the reactant structure and the mechanism.¹⁻¹¹ Earlier, we reported that the ketene-forming elimination reactions from *O*-aryl phenylacetates (**3**) promoted by $R_2NH-R_2NH_2^+$ in 70 mol % MeCN(aq) proceeded by an E2 mechanism,^{11a} while the reactions of *O*-aryl thienylacetates (**2a-c**) proceeded by a competing E2 and E1cb mechanism under the same condition.^{11g} The results have been attributed to the higher acidity of the C β -H bond in the latter,¹² which would stabilize the carbanion intermediate such that the E1cb mechanism can compete with the E2. Alternatively, the smaller aromatic resonance energy of the thiophene than that of benzene may also have contributed to the stability of the carbanion.¹³ To assess the relative importance of C β -H bond acidity and the aromatic resonance energy of the β -aryl group on the ketene-forming eliminations, we have now studied the reactions of aryl furylacetates (**1**) with $R_2NH-R_2NH_2^+$ in 70 mol % MeCN(aq) (eq 1). Since the pK_a values of 2-methylfuran and toluene are identical,¹² whereas the aromatic resonance energy of furan is much lower than that of benzene,¹³ this seemed to be an ideal system for such a purpose. We have investigated the reaction mechanism under various conditions. By comparing with the existing data for eliminations of ArCH₂C(O)OC₆H₃-2-X-4-NO₂ [Ar = thiophene (**2**), phen-

yl (**3**)] with $R_2NH-R_2NH_2^+$ in 70 mol % MeCN(aq), the effect of β -aryl group was assessed.

Results

Aryl furylacetates **1a-d** were synthesized by the reaction between 2-furylacetic acid, substituted phenols, 2-chloro-1-methylpyridinium iodide, and Et₃N in CH₂Cl₂ as reported previously.^{11a,14} The spectral and analytical data for the new compounds were consistent with the proposed structures.

For the reactions of **1a-d** with $R_2NH-R_2NH_2^+$ in 70 mol % MeCN(aq), the yields of aryloxides determined by comparing the UV absorption of the infinity samples of the kinetic runs with those of the authentic aryloxides were in the range

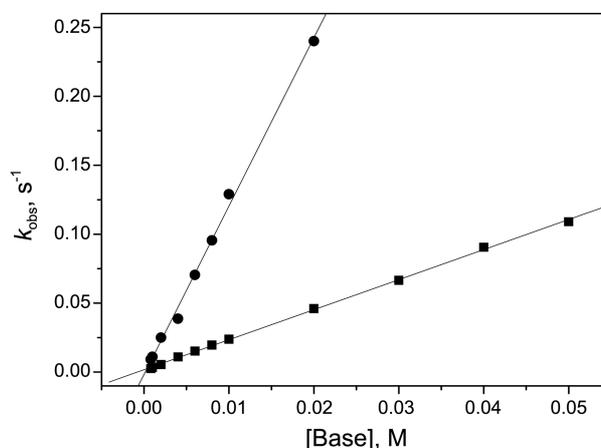


Figure 1. Plots of k_{obs} versus base concentration for eliminations from 2-chloro-4-nitrophenyl furylacetate (**1c**, ■) and 2,4-dinitrophenyl furylacetate (**1d**, ●) promoted by $R_2NH-R_2NH_2^+$ in 70 mol % MeCN(aq) at 25.0 °C, [*i*-Pr₂NH]/[*i*-Pr₂NH₂⁺] = 1.0, $\mu = 0.10$ M (Bu₄N⁺Br⁻).

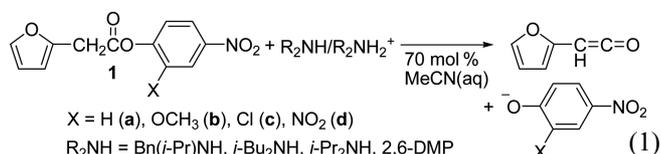


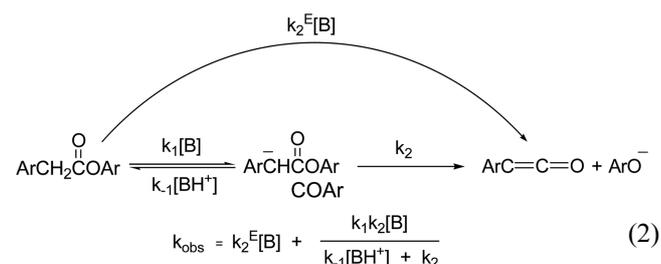
Table 1. Rate constants for ketene-forming elimination from ArCH₂C(O)OC₆H₃-2-X-4-NO₂ (Ar = furyl)^a promoted by R₂NH-R₂NH₂⁺ in 70 mol % MeCN(aq) at 25.0 °C

R ₂ NH ^b	pK _a ^c	10 ² k ₂ ^E , M ⁻¹ s ^{-1d,e}			
		X = H (1a)	X = OMe (1b)	X = Cl (1c)	X = NO ₂ (1d)
Bn(<i>i</i> -Pr)NH	16.8	0.217	0.254	36.0	309
<i>i</i> -Bu ₂ NH	18.2	0.766	0.686	120	1054
<i>i</i> -Pr ₂ NH	18.5	1.62	1.26	218	1220
2,6-DMP ^f	18.9	3.52	3.83	443	1935

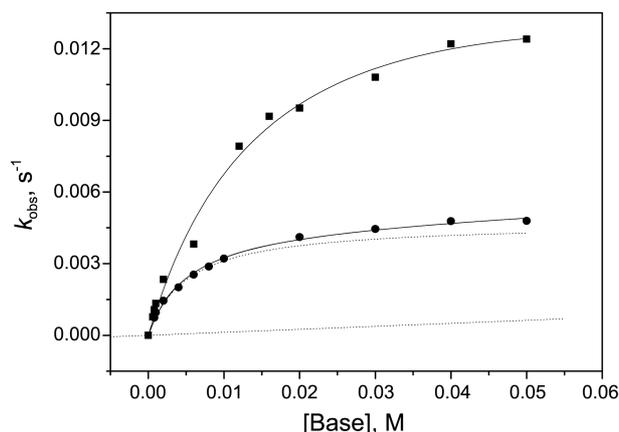
^a[Substrate] = 5.0 × 10⁻⁵ M. ^b[R₂NH] = 8.0 × 10⁻⁴ - 5.0 × 10⁻² M. ^cReference 15. ^dAverage of three or more rate constants. ^eEstimated uncertainty, ± 3%. ^f*cis*-2,6-Dimethylpiperidine.

of 94-98%. The rates of elimination reactions were followed by monitoring the increase in the absorption at the λ_{max} for the aryloxides in the range of 400-434 nm. Excellent pseudo-first order kinetics plots, which covered at least three half-lives were obtained. The rate constants are summarized in Tables S1-S4 in Supporting Information. For the reactions of 1c and 1d with R₂NH-R₂NH₂⁺ in 70 mol % MeCN(aq), the plots of k_{obs} versus base concentration were straight lines passing through the origin, indicating that the reactions are second-order, first order to the substrate and first order to the base (Figures 1, S8 and S9). The slopes are the overall second-order rate constants k₂^E. Values k₂^E for eliminations from 1c and 1d were summarized in Table 1.

In contrast to the reactions of 1c and 1d, the plots of k_{obs} versus base concentration for the reactions of 1a and 1b were curves at low base concentration and straight lines at higher base concentration (Figures 2 and S1-S7). The data were analyzed with a nonlinear regression analysis program by assuming that the reaction proceeds by concurrent E2 and E1cb mechanisms (eq. 2).^{11a,11d} By utilizing the computer program, the k₂^E, k₁, and k₋₁/k₂ values that best fit with Eq. (2) were calculated, and the plots were dissected into the E2 and E1cb components. In all cases, the correlations between the calculated and the experimental data were excellent. The calculated k₂^E values are included in Table 1 and k₁ and k₋₁/k₂ values are summarized in Table 2. The k₁ increased and k₋₁/k₂ decreased with a stronger base. For a given base, k₁ decreased and k₋₁/k₂ ratio increased as the leaving group was changed to a poorer one.



The k₂^E values for 1a-d showed excellent correlations with the pK_a values of the promoting bases on the Brønsted plots (Figure 3). The β values are in the range of 0.37-0.54 and

**Figure 2.** Plots of k_{obs} versus base concentration for eliminations from 4-nitrophenyl furylacetate (1a) promoted by R₂NH-R₂NH₂⁺ in 70 mol % MeCN(aq) (●) and 70 mol % MeCN-30 mol % D₂O (■) at 25.0 °C, [*i*-Pr₂NH]/[*i*-Pr₂NH₂⁺] = 1.0, μ = 0.10 M (Bu₄N⁺Br⁻). The closed circles (●) and squares (■) are the experimental data, and the solid line shows the fitted curve by using eq. 2 (see text). The curve for the reaction in 70 mol % MeCN(aq) is dissected into the E2 and E1cb reaction components (dashed lines). Calculated values of k₂^E, k₁, and k₋₁/k₂ for 1a and 1b are summarized in Tables 1 and 2.**Table 2.** The k₁ and k₋₁/k₂ values for elimination from ArCH₂C(O)OC₆H₃-2-X-4-NO₂ (Ar = furyl, 1a-d)^a promoted by R₂NH-R₂NH₂⁺ in 70 mol % MeCN(aq) at 25.0 °C

R ₂ NH ^b	k ₁ , M ⁻¹ s ^{-1c,e}		k ₋₁ /k ₂ , M ^{-1c,e}	
	X = H (1a)	X = OMe (1b)	X = H (1a)	X = OMe (1b)
Bn(<i>i</i> -Pr)NH	0.260	0.238	967	1920
<i>i</i> -Bu ₂ NH	0.728	0.319	880	636
<i>i</i> -Pr ₂ NH	0.922(1.13) ^f	0.602	196(67.9) ^f	300
2,6-DMP ^g	1.77	1.19	381	594

^a[Substrate] = 5.0 × 10⁻⁵ M. ^b[R₂NH] = 8.0 × 10⁻⁴-5.0 × 10⁻² M. ^cCalculated from k_{obs} by using eq. 2. ^dThe slopes of the plots of log k₁ versus pK_a of the base for 1a and 1b are 0.38 ± 0.04 and 0.29 ± 0.12, respectively. ^eThe slopes of the plots of log k₋₁/k₂ versus pK_a of the base for 1a and 1b are -0.17 ± 0.12 and -0.27 ± 0.07, respectively. ^fSolvent was 70 mol % MeCN-30 mol % D₂O. ^g*cis*-2,6-Dimethylpiperidine.

Table 3. Brønsted β values for elimination from ArCH₂C(O)OC₆H₃-2-X-4-NO₂ (Ar = furyl) promoted by R₂NH-R₂NH₂⁺ in 70 mol % MeCN(aq) at 25.0 °C

	X = H (1a)	X = OMe (1b)	X = Cl (1c)	X = NO ₂ (1d)
pK _{lg} ^a	20.7	20.6	18.1	16.0
β	0.54 ± 0.12	0.51 ± 0.1	0.49 ± 0.07	0.37 ± 0.02

^aReference 16.

decrease with a better leaving group (Table 3).

When the reaction of 1a with *i*-Pr₂NH/*i*-Pr₂NH₂⁺ was conducted in 70 mol % MeCN-30 mol % D₂O, the k_{obs} values were larger than those measured in 70 mol % MeCN(aq) (Figure 2 and Table S2), and the rate data could be fitted with the E1cb mechanism, that is, the 2nd term in Eq. (2). Calculated values of k₁ and k₋₁/k₂ for this reaction are

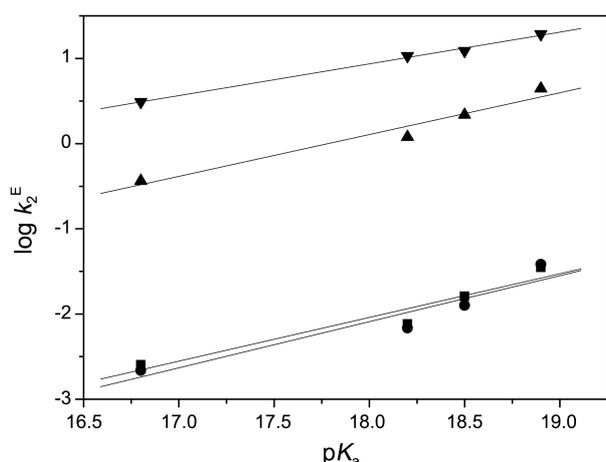


Figure 3. Brønsted plots for the elimination from $\text{ArCH}_2\text{C}(\text{O})\text{-OC}_6\text{H}_3\text{-2-X-4-NO}_2$ ($\text{Ar} = \text{furyl}$) promoted by $\text{R}_2\text{NH-R}_2\text{NH}_2^+$ in 70 mol % $\text{MeCN}(\text{aq})$ at $25.0\text{ }^\circ\text{C}$ [$\text{X} = \text{H}$ (**1a**, ●), OMe (**1b**, ■), Cl (**1c**, ▲), NO_2 (**1d**, ▼)].

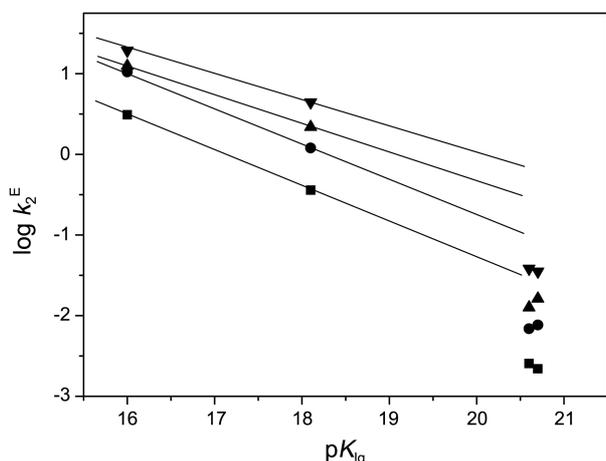


Figure 4. Plots $\log k_2^{\text{E}}$ versus $\text{p}K_{\text{lg}}$ values of the leaving group for the elimination from $\text{ArCH}_2\text{C}(\text{O})\text{OC}_6\text{H}_3\text{-2-X-4-NO}_2$ ($\text{Ar} = \text{furyl}$, **1a-d**) promoted by $\text{R}_2\text{NH-R}_2\text{NH}_2^+$ in 70 mol % $\text{MeCN}(\text{aq})$ at $25.0\text{ }^\circ\text{C}$. [$\text{R}_2\text{NH} = \text{Bn}(i\text{-Pr})\text{NH}$ (■), $i\text{-Bu}_2\text{NH}$ (▲), $i\text{-Pr}_2\text{NH}$ (●), 2,6-DMP (▼)].

summarized in Table 2.

The plots of the $\log k_2^{\text{E}}$ versus the leaving group $\text{p}K_{\text{lg}}$ values are depicted in Figure 4. As reported for **2a** and **2b**,^{11g} the data for **1a** and **1b** showed large negative deviation from the straight lines. Therefore, the β_{lg} values were calculated without the data for **1a** and **1b**. The β_{lg} values are in the range 0.31–0.45 and decrease as the $\text{p}K_{\text{a}}$ value of the base increase (Table 4).

Table 4. Brønsted β_{lg} values for Elimination from $\text{ArCH}_2\text{C}(\text{O})\text{-OC}_6\text{H}_3\text{-2-X-4-NO}_2$ ($\text{Ar} = \text{furyl}$) promoted by $\text{R}_2\text{NH-R}_2\text{NH}_2^+$ in 70 mol % $\text{MeCN}(\text{aq})$ at $25.0\text{ }^\circ\text{C}$

R_2NH	$\text{Bn}(i\text{-Pr})\text{NH}$	$i\text{-Bu}_2\text{NH}$	$i\text{-Pr}_2\text{NH}$	2,6-DMP ^b
$\text{p}K_{\text{a}}^{\text{a}}$	16.8	18.2	18.5	18.9
β_{lg}	-0.45	-0.45	-0.36	-0.31

^aReference 15. ^bcis-2,6-Dimethylpiperidine.

To provide additional evidence for the reaction mechanism, the H-D exchange experiment was conducted by mixing **1a** with $i\text{-Pr}_2\text{NH}/i\text{-Pr}_2\text{NH}_2^+$ in 70 mol % MeCN -30 mol % D_2O at $-10\text{ }^\circ\text{C}$. The reactant was recovered immediately after mixing. The NMR spectrum indicated that approximately 35% of furfuryl C-H bond of **1a** was converted to the C-D bond.

Discussion

Mechanism of Eliminations from 1. Results of kinetic investigations and product studies reveal that the reaction of **1c** and **1d** with $\text{R}_2\text{NH-R}_2\text{NH}_2^+$ in 70 mol % $\text{MeCN}(\text{aq})$ proceed by an E2 mechanism. The reactions produced elimination product and exhibited second-order kinetics, thereby ruling out all but bimolecular pathway. The $(\text{E1cb})_{\text{R}}$, $(\text{E1cb})_{\text{ip}}$, and internal return mechanisms, which would exhibit either a specific base catalysis or Brønsted β values near unity,^{16,17} were negated by the observed general base catalysis with the Brønsted β ranging from 0.37 to 0.54. Moreover, the β values increased with a poorer leaving group. This effect corresponds to a positive interaction coefficient, *i.e.*, $p_{\text{xy}} = \partial\beta/\partial\text{p}K_{\text{lg}} = \partial\beta_{\text{lg}}/\partial\text{p}K_{\text{BH}} > 0$, which describes the interaction between the base catalysis and the leaving group.^{17,18} The positive interaction coefficient provides additional support for an E2 mechanism in which the $\text{C}_{\beta}\text{-H}$ and $\text{C}_{\alpha}\text{-OAr}$ bonds are partially cleaved in the transition state.

When the leaving group was changed to a poorer one (**1a** and **1b**), the plots of k_{obs} versus the base concentration were curves (Figures 2 and S1-S7). The result could be explained with a competing E2 and E1cb mechanism, as explained below. First, the rate data fitted well with Eq. (2) and the shapes of dissected lines were typical for an E2 and E1cb mechanisms. Second, the calculated values of k_2^{E} , k_1 , and k_{-1}/k_2 are consistent with the competing mechanism; (i) The calculated k_2^{E} values for **1a** and **1b** and the experimentally measured k_2^{E} values for **1c** and **1d** show excellent correlation on the Brønsted plots (Figure 3), as expected for the E2 mechanism. (ii) The increase in the k_1 value with a stronger base is in good agreement with the deprotonation step of the E1cb mechanism. (iii) The decrease in the k_{-1}/k_2 ratio with a stronger base can be attributed to the decreased acidity of the R_2NH_2^+ which would reduce k_{-1} .

It is interesting to note that the k_2^{E} values for **1a** and **1b** showed large negative deviation in Figure 4. This outcome could be explained, if the extent of the $\text{C}_{\alpha}\text{-OAr}$ bond cleavage is smaller than those for **1c** and **1d**. Since the β values for **1a** and **1b** are larger than those for **1c** and **1d**, the transition state of the E2 pathway would be skewed toward the E1cb-like with greater extent of proton transfer and smaller degree of cleavage of the $\text{C}_{\alpha}\text{-OAr}$ bond. This would destabilize the transition state as a result of the smaller extent of double bond formation due to the inefficient charge transfer from the β - to α -carbon and retard the rate. This result underlines the unusual sensitivity of the ketene-forming transition state to the reactant structure in the borderline between the E2 and E1cb.

The mechanistic diversity of the ketene-forming elimination was further demonstrated in the reaction of **1a** with *i*-Pr₂NH/*i*-Pr₂NH₂⁺ in 70 mol % MeCN-30 mol % D₂O. There is convincing evidence in support of the mechanism change from the competing E2 and E1cb to E1cb under this condition. First, the *k*_{obs} values measured in the deuterated solvent could be fitted to the 2nd term of Eq. (2), indicating an exclusive E1cb mechanism (Figure 2). Second, the larger *k*₁ value in the deuterated solvent is also consistent because the basicity of R₂NH would increase in the poorer solvating D₂O.¹⁹ Third, since the *k*₂ value should be almost the same in the two solvents, the 3-fold smaller *k*₋₁/*k*₂ value in the deuterated solvent predicts that *k*₋₁^{H₂O}/*k*₋₁^{D₂O} ≈ 3.0 (Table 2). Since all of the R₂NH₂⁺ should be converted to R₂ND₂⁺ in the deuterated solvent, the large primary isotope effect is not unexpected. Fourth, the smaller *k*₋₁^{D₂O} would increase the 2nd term of Eq. (2) more in the deuterated than in undeuterated solvent to make the *k*_{obs} values larger. If it becomes the predominant term, a change of the mechanism would occur. Fifth, the partial H-D exchange of the furfuryl C-H bond in the recovered starting material from the reaction of **1a** with *i*-Pr₂NH/*i*-Pr₂NH₂⁺ in 70 mol % MeCN-30 mol % D₂O provides additional support for the E1cb mechanism.

Effect of β-Aryl Group on the Ketene-Forming Transition State. Table 5 compares the rates and transition state parameters for the eliminations from ArCH₂C(O)OC₆H₃-2-X-4-NO₂. The relative rate of E2 reaction changed from 1.0 to 10 to 1.2 as the β-aryl group was varied from furyl (**1a**) to thienyl (**2**) to phenyl (**3**). Since the p*K*_a values of 2-methylfuran, 2-methylthiophene, and toluene in DMSO are 43, 42, and 43, respectively,¹² the faster rate observed for **2** can be attributed to the higher acidity of the C_β-H bond. However, *b* and |β_{lg}| values of **1a**, **2**, and **3** are almost the same within experimental errors, indicating that structure of the E2 transition states are not significantly influenced by the β-aryl group. In addition, the *k*₁ value is larger for **2** than that for **1a** is consistent with the higher acidity of the C_β-H bond. The larger *k*₋₁/*k*₂ value for **2** is as expected for the E1cb mechanism. Although the lower basicity of the conjugate base of **2** would reduce both *k*₋₁ and *k*₂, the effect should be more pronounced for the slower step (*k*₂). This would predict a larger *k*₋₁/*k*₂. The most interesting result from this study is that **1a** and **2**, but not **3**, showed mechanistic changes from E2 to a competing E2 and E1cb. While the change of the mechanism for **2** can be attributed to the higher acidity of the C_β-H bond of **2** (see above), the difference between **1a** and **3** cannot be explained similarly because the acidities of their C_β-H bonds are the same. Alternatively, the negative charge density developed at the β-carbon can better be stabilized by the furyl than phenyl group because of the smaller aromatic resonance energy.¹³ This would stabilize the carbanion intermediate for **1a** such that the E1cb mechanism can compete with E2. A similar explanation can be applied to the competing E2 and E1cb mechanisms for **2**, although the higher acidity of the C_β-H bonds would undoubtedly contribute to the carbanion stability.

In conclusion, we have studied the elimination reactions of

Table 5. Transition state Parameters for ketene-forming eliminations from ArCH₂C(O)OC₆H₄-4-NO₂ Promoted by R₂NH-R₂NH₂⁺ buffers in 70 mol % MeCN(aq) at 25.0 °C

	Ar = furyl (1a)	Ar = thienyl (2) ^a	Ar = phenyl (3) ^b
rel. rate (<i>k</i> ₂ ^E) ^c	1.0	10	1.2
β	0.54 ± 0.12	0.64 ± 0.09	0.77 ± 0.03
β _{lg}	-0.45	-0.49 ± 0.01	-0.43
<i>k</i> ₁	0.26	0.53	
<i>k</i> ₋₁ / <i>k</i> ₂	967	2100	

^aReference 11g. ^bReference 11a. ^cR₂NH = Bn(*i*-PrNH).

aryl furylacetates (**1a-d**) promoted by R₂NH-R₂NH₂⁺ in 70 mol % MeCN(aq). The reactions of **1c** and **1d** proceeded by a concerted E2 mechanism. The mechanism changed to the competing E2 and E1cb mechanisms as the leaving group was changed to a poorer one [X = H (**1a**) and OCH₃ (**1b**)]. For elimination from **1a** with *i*-Pr₂NH/*i*-Pr₂NH₂⁺, a further change to the E1cb mechanism was realized in 70 mol % MeCN-30 mol % D₂O. When the β-aryl groups was changed from phenyl to thienyl to furyl, the structure of the E2 transition state remained almost the same although *k*₂^E value changed according to the acidity of the C_β-H bond. Noteworthy is the mechanistic diversity observed in the ketene-forming eliminations from **1** by the changes in the leaving group and the base-solvent system.

Experimental Section

Materials. Aryl furylacetates **1a-d** were prepared from 2-furanacetic acid and substituted phenols in the presence of Et₃N and 2-chloro-1-methylpyridinium iodide in CH₂Cl₂ as previously reported.^{11a,14} The yield (%), IR (KBr, C=O, cm⁻¹), ¹H NMR (400 MHz, CDCl₃, *J* values are in Hz), and ¹³C NMR (100 MHz), and mass spectral data for the new compounds are as follows.

2-(4-Nitrophenyl)furylacetate (1a): Yield 75%; IR 1760 cm⁻¹; ¹H NMR δ 3.98 (s, 2H), 6.37 (m, 2H), 7.29 (d, *J* = 9.24, 2H), 7.41 (s, 1H), 8.25 (d, *J* = 9.24, 2H); ¹³C NMR δ 34.1, 108.8, 110.8, 122.4, 125.2, 142.6, 145.5, 146.2, 155.2, 167.0; LRMS (EI); *m/z* 247 [M⁺] (64), 109 (12), 108 (79), 81 (100), 82 (34), 63 (6), 53 (46), 50 (5).

2-(2-Methoxy-4-nitrophenyl)furylacetate (1b): Yield 78 %; IR 1774 cm⁻¹; ¹H NMR δ 3.90 (s, 3H), 3.99 (s, 2H), 6.38 (m, 2H), 7.20 (d, *J* = 8.56, 1H), 7.41 (d, *J* = 1.72, 1H), 7.83 (d, *J* = 2.4, 1H), 7.88 (dd, *J* = 2.4, 8.56, 1H); ¹³C NMR δ 33.7, 56.5, 107.7, 108.7, 110.7, 116.4, 123.1, 142.5, 144.9, 146.4, 146.5, 151.6, 166.5; LRMS (EI); *m/z* 277 [M⁺] (14), 108 (99), 81 (100), 53 (27).

2-(2-Chloro-4-nitrophenyl)furylacetate (1c): Yield 58 %; IR 1780 cm⁻¹; ¹H NMR δ 4.03 (s, 2H), 6.37 (m, 2H), 7.35 (d, *J* = 8.88, 1H), 7.42 (d, 1.72, 1H), 8.15 (dd, *J* = 2.72, 8.88, 1H), 8.33 (d, *J* = 2.72, 1H); ¹³C NMR δ 33.7, 109.0, 110.8, 123.2, 124.3, 125.9, 128.2, 142.7, 145.9, 151.9, 166.1; LRMS (EI); *m/z* 281 [M⁺] (42), 109 (14), 108 (29), 82 (64), 81 (100), 63 (11), 53 (53).

2-(2,4-Dinitrophenyl)furylacetate (1d): Yield 49%; IR

1745 cm^{-1} ; ^1H NMR δ 4.08 (s, 2H), 6.39 (m, 1H), 7.42 (m, 1H), 7.50 (d, $J = 9.06$, 1H), 8.51 (dd, $J = 2.72$, 9.06, 1H), 8.96 (d, $J = 2.72$, 1H); ^{13}C NMR δ 33.6, 109.3, 110.8, 121.3, 121.8, 126.7, 129.1, 131.6, 142.8, 145.4, 148.4, 166.3; LRMS (EI); m/z 292 [M^+] (23), 125 (9), 108 (27), 82 (40), 81 (100), 53 (38).

Reagent grade acetonitrile and secondary amine were fractionally distilled from CaH_2 . The base-solvent solutions were prepared by dissolving equivalent amount of R_2NH and R_2NH_2^+ in 70 mol % MeCN(aq). In all cases, the ionic strength was maintained to 0.1 M with $\text{Bu}_4\text{N}^+\text{Br}^-$.

Kinetic Studies. Reactions of **1a-d** with $\text{R}_2\text{NH}/\text{R}_2\text{NH}_2^+$ in 70 mol % MeCN(aq) were followed by monitoring the increase in the absorbance of aryloxides at 400-434 nm with a UV-vis spectrophotometer as described.^{11a-11c}

Product Studies. The products of eliminations from **1a-d** promoted by $\text{R}_2\text{NH}/\text{R}_2\text{NH}_2^+$ in 70 mol % MeCN(aq) were identified as reported.^{11a} The yields of aryloxides determined by comparing the UV absorptions of the infinity samples with those for the authentic aryloxides were in the range of 94-98%.

H-D Exchange Experiment. To determine whether the β -furfuryl protons of **1a** undergo H-D exchange, **1a** (0.047 g, 0.19 mmol) was reacted with $i\text{-Pr}_2\text{NH}/i\text{-Pr}_2\text{NH}_2^+$ (0.05 M, 5 mL) in 70 mol % MeCN-30 mol % D_2O at -10°C . The reaction was quenched by adding dilute HCl(aq) immediately after mixing. The recovered reactant was isolated by extracting the products with ethyl acetate followed by the silica gel column chromatography using ethyl acetate/hexane = 1/4 as eluent. The proton NMR spectrum of the recovered reactant from **1a** was identical to that of the substrate except that the integration at δ 3.98 decreased by approximately 35%.

Control Experiments. The stabilities of **1a-d** were determined as reported.^{11b-11d} The solutions of **1** in MeCN were stable for at least two weeks when stored in the refrigerator.

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Support Information Available. Observed rate constants for elimination from **1a-d** promoted by $\text{R}_2\text{NH}/\text{R}_2\text{NH}_2^+$ in 70 mol % MeCN(aq), plots of k_{obs} versus base concentration, and spectroscopic and analytical data for the new compounds

are available on request from the correspondence author (20 pages). e-mail: sypyun@pknu.ac.kr.

References

- (a) Holmquist, B.; Bruice, T. C. *J. Am. Chem. Soc. Res.* **1969**, *91*, 2993-3002. (b) Holmquist, B.; Bruice, T. C. *J. Am. Chem. Soc. Res.* **1969**, *91*, 3003-3006.
- Pratt, R. F.; Bruice, T. C. *J. Am. Chem. Soc.* **1970**, *92*, 5956-5964.
- Inoue, M.; Bruice, T. C. *J. Am. Chem. Soc.* **1982**, *104*, 1644-1653. (b) Inoue, M.; Bruice, T. C. *J. Org. Chem.* **1982**, *47*, 959-963.
- (a) William, A. *J. Chem. Soc., Perkin Trans. 2* **1972**, 808-812. (b) William, A.; Douglas, K. T. *Chem. Rev.* **1975**, 627-649.
- Tagaki, W.; Kobayashi, S.; Kurihara, K.; Kurashima, K.; Yoshida, Y.; Yano, J. *J. Chem. Soc. Chem. Commun.* **1976**, 843-845.
- Broxton, T. J.; Duddy, N. W. *J. Org. Chem.* **1981**, *46*, 1186-1191.
- Chandrasekar, R.; Venkatasubramanian, N. *J. Chem. Soc., Perkin Trans. 2* **1982**, 1625-1631.
- Douglas, K. T.; Alborz, M.; Rullo, G. R.; Yaggi, N. F. *J. Chem. Soc. Chem. Commun.* **1982**, 242-246.
- Isaac, N. S.; Najem, T. S. *J. Chem. Soc., Perkin Trans. 2* **1988**, 557-562.
- Chung, S. Y.; Yoh, S. D.; Choi, J. H.; Shim, K. T. *J. Korean Chem. Soc.* **1992**, *36*, 446-452.
- (a) Cho, B. R.; Kim, Y. K.; Maing Yoon, C. O. *J. Am. Chem. Soc.* **1997**, *119*, 691-697. (b) Cho, B. R.; Kim, Y. K.; Yoon, J. S.; Kim, J. C.; Pyun, S. Y. *J. Org. Chem.* **2000**, *65*, 1239-1241. (c) Cho, B. R.; Kim, N. S.; Kim, Y. K.; Son, K. H. *J. Chem. Soc., Perkin Trans. 2* **2000**, 1419-1422. (d) Cho, B. R.; Jeong, H. C.; Seung, Y. J.; Pyun, S. Y. *J. Org. Chem.* **2002**, *67*, 5232-5238. (e) Pyun, S. Y.; Lee, D. C.; Kim, J. C.; Cho, B. R. *Org. Biomol. Chem.* **2003**, *1*, 2734-2738. (f) Pyun, S. Y.; Cho, B. R. *Bull. Korean Chem. Soc.* **2005**, *26*, 1017-1024. (g) Cho, B. R.; Pyun, S. Y. *J. Org. Chem.* **2007**, *72*, 1098-1103. (h) Pyun, S. Y.; Cho, E. J.; Seok, H. J.; Kim, J. C.; Lee, S. H.; Cho, B. R. *Bull. Korean Chem. Soc.* **2007**, *28*, 917-920. (i) Pyun, S. Y.; Cho, E. J.; Seok, H. J.; Kim, J. C.; Lee, S. H.; Cho, B. R. *J. Phys. Org. Chem.* **2007**, *20*, 685-689.
- Bordwell, F. G. *Acc. Chem. Res.* **1988**, *21*, 456.
- Bernardi, F. *J. Mol. Struct.* **1988**, *163*, 173-177.
- Saigo, K.; Usui, M.; Kikuchi, K.; Shimada, E.; Mukaiyama, T. *Bull. Chem. Soc. Jpn.* **1977**, *50*, 1863-1866.
- Cho, B. R.; Lee, S. J.; Kim, Y. K. *J. Org. Chem.* **1995**, *60*, 2072-2076.
- Coetzee, J. F. *Prog. Phys. Org. Chem.* **1965**, *4*, 45-92.
- Gandler, J. R. *The Chemistry of Double Bonded Functional Groups*; Patai, S., Ed.; John Wiley and Sons: Chichester, 1989; vol. 2, part 1, pp 734-797.
- Lowry, T. H.; Richardson, K. S. *Mechanism and Theory in Organic Chemistry*; Harper and Row: New York, 1987. (a) pp 214-218. (b) pp 591-616. (c) pp 640-644.
- Gold, V. *Adv. Phys. Org. Chem.* **1969**, *7*, 259-267.