## An Efficient One Pot Synthesis of Bicyclic Dienones

## Farhat Rezgui and Mohamed Moncef El Gaïed\*

Laboratoire de Chimie Organique, Faculté des Sciences de Tunis Campus Universitaire, 1060 Tunis, Tunisia

Bicyclic dienones **4** are prepared in a one pot process from the reaction of 2-(acetoxymethyl)cyclohex-2-enone **1** with 1,3-dicarbonyl compounds **2** in the presence of  $K_2CO_3$  in refluxing absolute ethanol.

Synthetic methods for dienones **4** are fairly sparse.<sup>1–3</sup> We have recently reported<sup>4</sup> that cyclization of intermediates **3**, obtained from the reaction of 2-(acetoxymethyl)cyclohex-2-enone  $1^{5,6}$  with 1,3-dicarbonyl compounds **2** in the presence of Et<sub>3</sub>N, afforded compounds **4** (Scheme 1). The overall yield of this Robinson annulation,<sup>7</sup> effected in a two-step sequence, is *ca.* 20–40%.

As an improvement to this method, we report herein a direct access to dienones 4 in satisfactory yields.

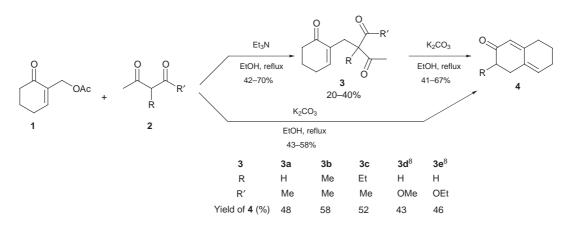
2-(Acetoxymethyl)cyclohex-2-enone 1 reacts with 1,3dicarbonyl compounds  $2^9$  in the presence of a large excess (4 equiv.) of anhydrous  $K_2CO_3$  in refluxing absolute ethanol by an  $S_N2$  type reaction to give products 3. These intermediates generate, *in situ*, *via* cleavage followed by cyclization, the target bicyclic dienones 4 in 43–58% overall yield (Scheme 1).

In summary, we have developed a simple one-step method for the preparation of bicyclic dienones **4** in overall yield higher than those obtained in our previous paper.<sup>4</sup> 2 h. The solvent was evaporated *in vacuo* and the reaction mixture was partitioned between water (25 ml) and diethyl ether (50 ml). Aqueous 2M HCl (5 ml) was added to dissolve salts and the mixture was extracted with diethyl ether ( $3 \times 20$  ml). The ether extracts were dried over MgSO<sub>4</sub> and concentrated *in vacuo*. The crude product was purified by column chromatography [1:4 then 4:1 diethyl ether–light petroleum (bp 40–60 °C).

Dienone **4a** yellow oil; IR (CHCl<sub>3</sub>): 1659, 1630, 1589 cm<sup>-1</sup>;  $\delta_{\rm H}(300 \,{\rm MHz}, {\rm CDCl}_3)$ : 6.10–5.93 (m, 1H), 5.75 (s, 1H), 2.65 (t, 2H,  $J = 7 \,{\rm Hz}$ ), 2.50–2.45 (m, 4H) 2.29–2.27 (m, 2H), 1.85–1.76 (m, 2H);  $\delta_{\rm C}$  (75 MHz, CDCl<sub>3</sub>): 199.3, 155.9, 131.8, 131.7, 123.1, 37.3, 30.8, 29.4, 25.8, 22.0; MS: m/z 148 (M<sup>+</sup>, 69), 133 (12), 120 (43), 105 (37), 91 (100), 77 (22), 65 (20), 51 (21) (Found: C, 81.2; H, 8.1. C<sub>10</sub>H<sub>12</sub>O requires C, 81.08; H, 8.11%).

*Dienone* **4b** yellow oil; IR (CHCl<sub>3</sub>): 1659, 1630, 1591 cm<sup>-1</sup>;  $\delta_{\rm H}(60 \,{\rm MHz}, {\rm CDCl}_3)$ : 6.17–5.90 (m, 1H), 5.67 (s, 1H), 2.77–2.10 (m, 7H), 2.01–1.54 (m, 2H), 1.23–1.04 (m, 3H);  $\delta_{\rm C}$  (75 MHz, CDCl<sub>3</sub>): 202.0, 155.0, 131.8, 131.7, 122.2, 41.0, 37.6, 30.4, 25.7, 22.0, 15.4; MS: *m/z* 162 (M<sup>+</sup>, 100), 147 (21), 134 (74), 119 (51), 105 (22), 91 (53) (Found: C, 81.4; H, 8.6. C<sub>10</sub>H<sub>14</sub>O requires C, 81.48; H, 8.64%).

*Dienone* **4c** yellow oil;  $\delta_{\rm H}$ (300 MHz, CDCl<sub>3</sub>): 6.00–5.98 (m, 1H), 5.57 (s, 1H), 2.64–2.59 (m, 1H), 2.38–2.33 (m, 3H), 2.20–2.15 (m, 3H), 1.72–1.66 (m, 3H), 1.33-1.29 (m, 1H), 0.84 (t, 3H, J = 6 Hz);  $\delta_{\rm C}$ 





## Experimental

<sup>1</sup>H NMR spectra were recorded in CDCl<sub>3</sub> solutions at 60 or 300 MHz with tetramethylsilane as an internal reference. <sup>13</sup>C NMR were recorded at 75 MHz with CDCl<sub>3</sub> as the internal reference. Chemical shifts are given in ppm ( $\delta$ ) and coupling constants *J* are reported in Hz. IR spectra were obtained on a Perkin Elmer Paragon 1000 PC IR spectrometer. Mass spectra were measured on a Hewlett-Packard 5890 spectrometer at 70 eV (EI). Column chromatography was performed using silica gel 60 (70–230 mesh).

*Preparation of Dienones* **4**.—A representative experimental procedure for the preparation of **4a** is described. A mixture of 2-(acetoxymethyl)cyclohex-2-enone **1** (1.00 g, 6 mmol), pentane-2,4dione (0.684 g, 6 mmol), commercial anhydrous potassium carbonate (3.32 g, 24 mmol) and 20 ml of absolute ethanol, was refluxed for (75 MHz, CDCl<sub>3</sub>): 201.9, 154.8, 132.1, 131.7, 122.4, 47.8, 34.5, 30.5, 26.0, 23.2, 22.2, 11.5; MS: m/z 176 (M<sup>+</sup>, 16), 148 (100), 120 (76), 91 (63), 77 (24) (Found: C, 81.9; H, 9.0.  $C_{12}H_{16}O$  requires C, 81.82; H, 9.09%).

We thank Professor Labassi, T. (FST) for interesting and valuable discussions.

Received, 15th March 1999; Accepted, 22nd April 1999 Paper E/9/02012H

## References

- 1 R. C. Klix and R. D. Bach, J. Org. Chem., 1987, 52, 580.
- 2 W. von E. Doering, L. Birladeanu, X. Cheng, T. Kitagawa and
- K. Sarma, J. Am. Chem. Soc., 1991, 113, 4558.
- 3 J. Colonge and P. Brison, Bull. Soc. Chim. Fr., 1961, 98.

<sup>\*</sup> To receive any correspondence.

- 4 F. Rezgui and M. M. El Gaïed, Tetrahedron, 1997, 53, 15711 and references cited therein.
- A. B. Smith, III, B. D. Dorsey, M. Ohba, A. T. Lupo, Jr. and
  M. S. Malamas, J. Org. Chem., 1988, 53, 4314; F. Rezgui
  and M. M. El Gaïed, *Tetrahedron Lett.*, 1998, 39, 5965.
  I. Takao, K. Kunimoto, S. Masanobu, N. Kiyohino and N. 5
- 6 Takemitsu, Chem. Abstr., 1987, 106, 49660m.
- 7 R. E. Gawley, Synthesis, 1976, 777.
- Et<sub>3</sub>N (1 equiv.) was used to obtain 3d and 3e which were con-8 verted in situ by addition of K2CO3 (2 equiv.) into 4d and 4e (both equivalent to 4a), respectively; spectroscopic data for compounds 3d and 3e are consistent with those reported in ref. 4.
- 9 A. Chamekh and H. Amri, Tetrahedron Lett., 1998, 39, 375.