

## Concise asymmetric syntheses of (*S*)-ethyl 4-methyloctanoate and its acid: aggregation pheromones of rhinoceros beetles of the genus *Oryctes*

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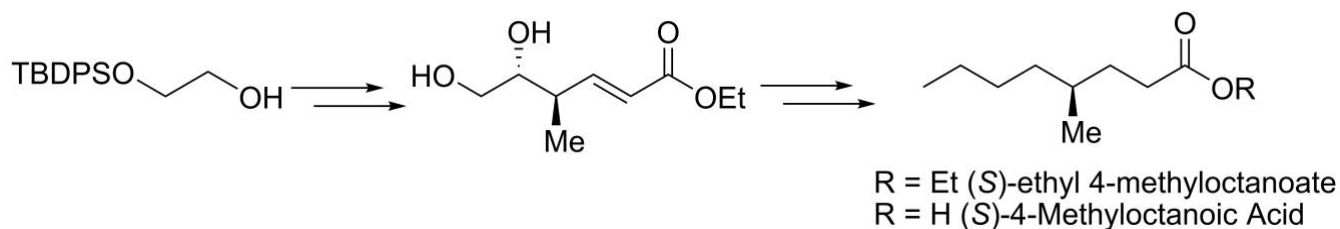
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### Abstract

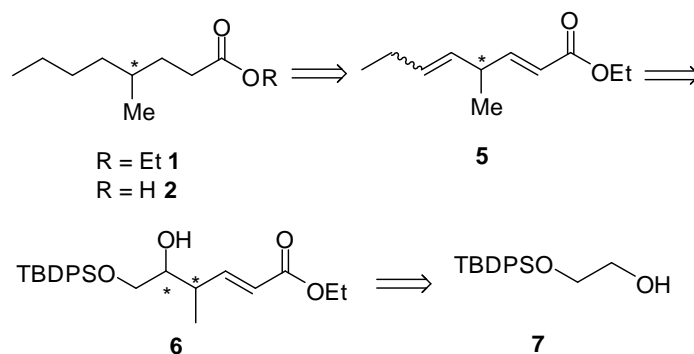
A concise stereoselective approach to the asymmetric syntheses of (*S*)-ethyl 4-methyloctanoate and its acid, main aggregation pheromones of the genus *Oryctes* and an important fragrance compounds, is described. The synthesis utilizes the organocatalyzed MacMillan's cross aldol reaction as key step. The synthetic approach outlined permits synthesis of a variety of enantiopure branched methyl unsaturated and saturated fatty acids.



**Keywords:** Rhinoceros Beetle, Aggregation Pheromone, MacMillan's cross Aldol reaction, Wittig olefination, Fragrance



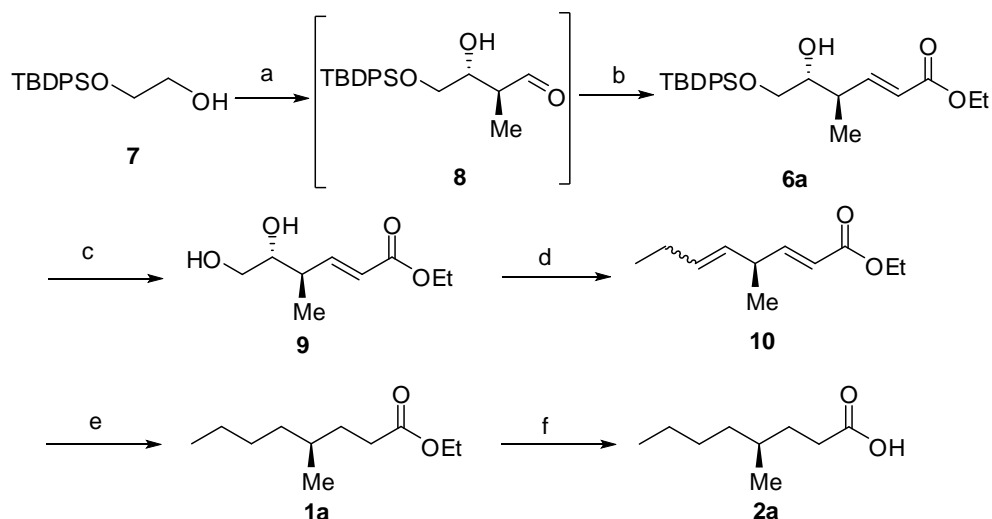
organocatalyzed MacMillan's cross aldol reaction of aldehyde derived from **7** with propionaldehyde and Wittig homologation reaction. The (*R*)- and (*S*)- configuration of the target compounds ethyl 4-methyloctanoate (**1**) and its acid **2** can be manipulated by simply changing organocatalyst under MacMillan's cross aldol conditions.



**Scheme 1.** Retrosynthetic approach to asymmetric syntheses of ethyl 4-methyloctanoate (**1**) and its corresponding ethyl ester **2**.

Our approach for the asymmetric synthesis of (*S*)-ethyl 4-methyloctanoate (**1a**) and its acid **2a** is depicted in Scheme 2. Syntheses of target compounds (*S*)-**1a** and (*S*)-**2a** began with readily available monosilylated ethylene glycol **7** which was subjected to a two-step sequence including Swern oxidation followed by L-proline catalyzed MacMillan's cross aldol reaction to afford the diastereomeric mixture of aldehyde **5** with >99% ee (*anti/syn* 9:1)<sup>26</sup> and subsequent 2C-Wittig olefination without separation with (ethoxycarbonylmethyl-ene)triphenylphosphorane furnished unsaturated ester **6a** following a literature procedure.<sup>27-28</sup>

With olefin derivative **6a** in hand, we then subjected it to silyl ether deprotection to get the diol intermediate **9**. Pleasingly, the minor diastereomer was separated out during column chromatography purification. The NaIO<sub>4</sub> mediated oxidative cleavage of terminal diol **9** followed by treatment of aldehyde intermediate with phosphonium salt (propyltriphenyl-phosphonium bromide) in presence of *n*-BuLi as a base furnished the olefin derivative **10**. In view of the fact that the incipient olefin would be ultimately hydrogenated to get the target compound **1a** we did not examine the olefin geometry of derivative **10**. Accordingly, the olefin derivative **10** on hydrogenation at one atmospheric pressure in the presence of 10% Pd-C furnished the target compound (*S*)-ethyl 4-methyloctanoate (**1a**) in 76% yield (over three steps); [ $\alpha$ ]<sub>D</sub><sup>25</sup> +1.61 (*c* 1.0, CHCl<sub>3</sub>); {lit.<sup>4</sup> [ $\alpha$ ]<sub>D</sub><sup>20</sup> +1.67 (*c* 1.35, CHCl<sub>3</sub>)}. Finally, saponification reaction of the (*S*)-ethyl 4-methyloctanoate (**1a**) with LiOH in aqueous THF solvent at room temperature afforded the target compound (*S*)-4-methyloctanoic acid (**2a**) in 92% Yield; [ $\alpha$ ]<sub>D</sub><sup>25</sup> +1.46 (*c* 1.0, CHCl<sub>3</sub>); {lit.<sup>23</sup> [ $\alpha$ ]<sub>D</sub><sup>20</sup> +1.5 (*c* 1.4, CHCl<sub>3</sub>)}. The physical and spectroscopic data were in full agreement with those reported literature.



**Scheme 2. Reagents and conditions:** (a) i) (COCl)<sub>2</sub>, DMSO, Et<sub>3</sub>N, dry CH<sub>2</sub>Cl<sub>2</sub>, -78 °C to rt, 3 h; ii) C<sub>3</sub>H<sub>6</sub>O, 10 mol % L-Proline, dioxane, 4 °C, 48 h, 94% (over two steps); (b) Ph<sub>3</sub>PCHCOOEt, dry THF, rt, 24 h, 92%; (c) TBAF, THF, rt, 4 h, 95%; (d) i) NaIO<sub>4</sub>, THF:H<sub>2</sub>O (1:1), rt, 20 min.; ii) propyltriphenylphosphonium bromide, *n*-BuLi, THF, -78 °C to rt, 3 h; (e) H<sub>2</sub>, 10% Pd-C, MeOH, rt, 4 h, 76% (over three steps); (f) LiOH, THF:H<sub>2</sub>O (4:1), rt, 6h, 92%.

## Conclusions

In conclusion, we have developed a simple and flexible approach to the total syntheses of (*S*)-ethyl 4-methyloctanoate (**1a**) and its acid **2a** from readily available starting materials. The synthetic sequence demonstrates the application of organocatalyzed MacMillan's cross aldol reaction and Wittig olefination as the key steps. The synthetic approach described has significant potential for stereochemical variations and further extension to other stereoisomers and their analogues.

## Experimental Section

**Ethyl (4*R*,5*R*,*E*)-5,6-dihydroxy-4-methylhex-2-enoate (9).** To a stirred solution of unsaturated ester **6a** (1.4 g, 3.28 mmol) in dry THF (10 mL) TBAF (4.91 mL, 4.92 mmol) was added dropwise at room temperature. After stirring the reaction mixture at room temperature for 4 h, the reaction mixture was diluted with water and ethyl acetate. The organic layer was washed with brine, dried over anhydrous Na<sub>2</sub>SO<sub>4</sub> and the solvent was evaporated under reduced pressure. Silica gel column chromatography (hexane/EtOAc, 4:1) furnished olefin diol **9** (988 mg, 95%) as a colorless liquid. [α]<sub>D</sub><sup>25</sup> +9.56 (*c* 1.0, CHCl<sub>3</sub>); IR (CH<sub>2</sub>Cl<sub>2</sub>) *v*: 3425, 2957, 2926, 1720, 1478, 1268, 1220 cm<sup>-1</sup>; <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>) δ: 6.97 (dd, *J* 8.28, 15.6 Hz, 1H), 5.87 (d, *J* 14.68 Hz, 1H), 4.19 (q, *J* 7.32, 14.2 Hz, 2H), 3.71-3.59 (m, 2H), 3.56-3.49 (m, 1H), 3.08 (brs, 2H), 2.54-2.45 (m, 1H), 1.29 (t, *J* 7.32 Hz, 3H), 1.1 (d, *J* 6.84 Hz, 3H); <sup>13</sup>C NMR (100 MHz, CDCl<sub>3</sub>) δ: 166.8, 150.3, 121.9, 74.8, 64.5, 60.4, 39.5, 15.8, 14.2. HRMS (ESI), *m/z* calcd for C<sub>9</sub>H<sub>16</sub>O<sub>4</sub>Na [M + Na]<sup>+</sup> 211.0941; found 211.0952.

**Ethyl (*S*)-4-methyloctanoate (1a).** To a stirred solution of NaIO<sub>4</sub> (226 g, 1.06 mmol) in H<sub>2</sub>O:THF (1:1, 10 mL) was added a suspension of olefin diol **9** (100 mg, 0.53 mmol) in THF (3 mL) and stirred for 20 min at room

temperature. To the reaction mixture EtOAc and water were added, the aqueous layer was extracted with EtOAc (3 x 15 mL). The combined organic layer was washed with brine, dried over anhydrous Na<sub>2</sub>SO<sub>4</sub>, and concentrated in vacuo. The aldehyde intermediate obtained was used as such for next reaction without further purification.

To a suspension of propyltriphenylphosphonium bromide (264 mg, 0.69 mmol) in dry THF (5 mL) dropwise *n*-BuLi (2.5 M in hexane, 0.3 mL, 0.79 mmol) at -78 °C was added. The reaction mixture was warmed to rt over a period of 1 h before being cooled to -78 °C. To the obtained orange solution above crude aldehyde dissolved in THF (3 mL) was added and the reaction mixture was left to warm to rt over 2 h before the addition of H<sub>2</sub>O (10 mL). The resulting suspension was extracted with EtOAc (3 x 10 mL). The combined organic layers were washed with brine, dried over anhydrous Na<sub>2</sub>SO<sub>4</sub> and concentrated under reduced pressure to afford olefin intermediate **10** as a colorless liquid which was used for the next reaction.

To a solution of above olefin intermediate **10** in methanol (4 mL) catalytic amount of palladium on carbon (10%) was added and resulting reaction mixture was stirred under hydrogen atmosphere for 4 h at room temperature, at which point TLC indicated the completion of reaction. The reaction mixture was filtered through a pad of Celite, solvent was removed in vacuo and silica gel column chromatography (hexane/ EtOAc 10:1) furnished ethyl (*S*)-4-methyloctanoate (**1a**) (73 mg, 76%) as a colourless liquid. [ $\alpha$ ]<sub>D</sub><sup>25</sup> +1.61 (*c* 1.0, CHCl<sub>3</sub>) {lit.<sup>4</sup> [ $\alpha$ ]<sub>D</sub><sup>20</sup> +1.67 (*c* 1.35, CHCl<sub>3</sub>)}; IR (CH<sub>2</sub>Cl<sub>2</sub>)  $\nu$ : 2957, 2946, 1735, 1478, 1268, 1215 cm<sup>-1</sup>; <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>)  $\delta$ : 4.12 (d, *J* 7.32, 14.2 Hz, 2H), 2.33-2.26 (m, 2H), 1.70-1.60 (m, 1H), 1.47-1.41(m, 2H) 1.31-1.27 (m, 5H); 1.26 (t, *J* 6.88, 3H) 1.15-1.09 (m, 1H), 0.89 (t, *J* 6.88, 3H), 0.88 (d, *J* 6.40, 3H); <sup>13</sup>C NMR (100 MHz, CDCl<sub>3</sub>)  $\delta$ : 174.2, 60.2, 36.3, 32.4, 32.2, 31.9, 29.1, 22.9, 19.2, 14.2, 14.1. HRMS (ESI), *m/z* calcd for C<sub>11</sub>H<sub>22</sub>O<sub>2</sub>Na [M + Na]<sup>+</sup> 209.1512; found 209.1514.

**(S)-4-methyloctanoic acid (2a)**. A mixture of ethyl (*S*)-4-methyloctanoate (**1a**) (50 mg, 0.27 mmol), LiOH (34 mg, 0.81 mmol), THF (2 mL) and water (0.5 mL) was stirred at room temperature for 6 h. The organic solvent was removed under reduced pressure, and the residue obtained was then acidified to pH 5 with 1 N HCl, and the reaction mixture was extracted with EtOAc (3 x 10 mL). The combined organic layers were washed with brine, dried over anhydrous Na<sub>2</sub>SO<sub>4</sub> and concentrated under reduced pressure. Silica gel column chromatography (hexane /EtOAc 9:1) afforded the target compound (*S*)-4-methyloctanoic acid (**2a**) (39 mg, 92%) as a colorless liquid. [ $\alpha$ ]<sub>D</sub><sup>25</sup> +1.46 (*c* 1.0, CHCl<sub>3</sub>); {lit.<sup>23</sup> [ $\alpha$ ]<sub>D</sub><sup>20</sup> +1.5 (*c* 1.4, CHCl<sub>3</sub>)}. IR (CH<sub>2</sub>Cl<sub>2</sub>)  $\nu$ : 3450, 3071, 2957, 2946, 1715, 1468, 1268, 1215 cm<sup>-1</sup>; <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>)  $\delta$ : 2.42-2.31 (m, 2H); 1.73-1.63 (m, 1H); 1.50-1.40 (m, 2H); 1.33-1.10 (m, 6H), 0.92-0.80 (m, 6H); <sup>13</sup>C NMR (100 MHz, CDCl<sub>3</sub>)  $\delta$ : 180.3, 36.3, 32.3, 31.8, 31.6, 29.1, 22.9, 19.2, 14.1. HRMS (ESI), *m/z* calcd for C<sub>9</sub>H<sub>18</sub>O<sub>2</sub>Na [M + Na]<sup>+</sup> 181.1199; found 181.1219.

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## Supplementary Material

Copies of <sup>1</sup>H and <sup>13</sup>C NMR spectra of compounds **9**, **1a** and **2a** are available in the supplementary material file.

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26. The enantiomeric excess of intermediate **8** was determined by in situ reduction with NaBH<sub>4</sub> at 0 °C producing the corresponding diol. HPLC Chiralpak IB (250 x 4.6 mm), petroleum ether:EtOH (99:1) v/v, flow rate 1.0 ml/min, λ<sub>max</sub> = 220 nm, (2*R*, 3*S*) *anti* isomer t<sub>r</sub> = 14.29 min, (2*S*, 3*R*) *anti* isomer t<sub>r</sub> = 15.89 min, (2*R*, 3*R*) and (2*S*, 3*S*) *syn* isomers t<sub>r</sub> = 18.35, 19.88 min.
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