Molbank 2006, M506

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## 3β-Acetoxy-elemo-lanost-8-en-24-one

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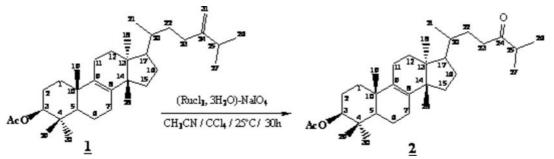
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Received: 1 August 2005 / Accepted: 17 October 2006 / Published: 1 December 2006

Keywords: Euphorbia resinifera, latex, catalytic oxidation.



The sodium periodate is prepared in situ with a equimolecular quantity of soda NaOH (0.5g; 12.50 mmol) and periodic acid H<sub>5</sub>IO<sub>6</sub> (2.85g; 12.50 mmol), the mixture is stirred at 0°C. After 15 min, 5 ml of CCl4, 5 ml of H<sub>3</sub>CCN and 93.03mg (0.12mmol) of ruthenium trichloride<sup>1,2</sup> were added. The mixture was stirred during 15min, and then 1.50g (3.12 mmol) of  $\underline{1}^3$  was added. The reaction was left under stirring at 25°C for 30h, then 20 ml of distilled water was added and the fractional mixture was extracted with the dichloromethane. After filtration on a silica gel column to eliminate RuO4, the organic layer was recovered, dried by Na<sub>2</sub>SO<sub>4</sub> and evaporated under reduce pressure. The residue was purified on silica gel column using hexane: ethyl acetate (97:3) as eluent to give  $\underline{2}$  (1.14g, 2.35 mmol) in 76 % yield.

Melting point: 110-111 °C (Hexane)

MS (EI, 70eV): 484 (M<sup>+.</sup>).

<sup>1</sup>H NMR (300 MHz, CDCl<sub>3</sub>) d(ppm): 4.50 (3H-3, dd:  $J_1 = 12$  Hz,  $J_2 = 4$  Hz); 2.13 (COO<u>CH3</u>, s); 0.76 (3H-18, s); 0.85 (3H-19, s); 0.93 (3H-21, d, J = 6 Hz); 2.60 (2H-23, m); 1.45 (3H-26, d: J = 2 Hz); 0.80 (3H-28, s); 0.95 (3H-29, s); 1.05 (3H-30, s).

<sup>13</sup>C NMR (75 MHz, CDCl<sub>3</sub>) δ (ppm): 35.03 (C-1); 27.60 (C-2); 81.03 (C-3); 36.18 (C-4); 50.03 (C-5); 37.95 (C-6); 28.07 (C-7); 133.71 (C-8); 134.04 (C-9); 36.3 (C-10); 20.25 (C-11); 25.37 (C-12); 44.22 (C-13); 50.22 (C-14); 30.84 (C-15); 30.26 (C-16); 51.15 (C-17); 18.60 (C-18); 18.75 (C-19); 37.23 (C-20); 21.53 (C-21); 37.23 (C-22); 51.32 (C-23); 215.50 (C-24); 31.68 (C-25); 22.74 (C-26); 18.80 (C-27); 15.55 (C-28); 16.45 (C-29); 171.09 (<u>CO</u>OCH3); 21.2 (COO<u>CH3</u>).

MS (m/z): 484 (25%), 426 (54%), 298 (65%).

## Acknowledgments

We would like to thank Pr. A. Idmessaaoud for helpful discussions.

## References

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