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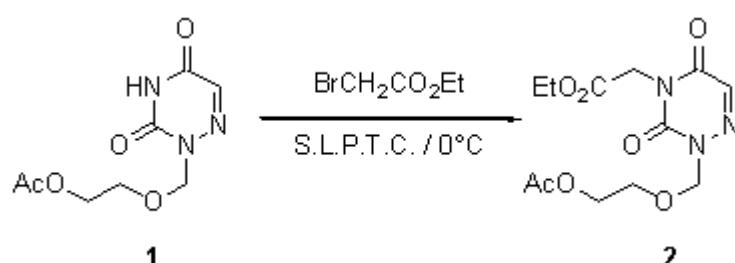
### 1-[(2-Acetoxyethoxy)methyl]-3-ethoxycarbonylmethyl-6-azauracil

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The product **2** was prepared via a direct condensation under solid-liquid phase transfer catalysis (S.L.P.T.C.) [1] conditions. To a solution of 0.02 mmole of tetraglyme in 4 ml of anhydrous THF, 0.11 mmole of potassium tert-butoxide is added. Then 0.1 mmole of the acyclonucleoside **1** [2] is added, the reaction mixture is stirred at room temperature for 15 min. The reaction mixture is cooled to 0°C and 0.11 mmole of alkylating agent in 2 ml of dry THF is added dropwise with stirring. When the addition is finished, the reaction mixture is stirred at 0°C for 30 min. The reaction mixture is then filtered and the filtrate is evaporated in vacuo to dryness. The residue is then chromatographed on a silica gel column and the expected acyclonucleoside **2** was isolated. Yield: 90 % (viscous and colourless).

Rf: 0.64 (CHCl<sub>3</sub> / MeOH, 9/1, V/V).

<sup>1</sup>H NMR (DMSO-d<sub>6</sub>): 1.20 (t, 3H, CH<sub>3</sub>CH<sub>2</sub>); 2.00 (s, 3H, COOCH<sub>3</sub>); 3.75 (m, 2H, OCH<sub>2</sub>CH<sub>2</sub>O); 4.10 (m, 2H, OCH<sub>2</sub>CH<sub>2</sub>O); 4.20 (q, 2H, CH<sub>3</sub>CH<sub>2</sub>); 4.75 (s, 2H, NCH<sub>2</sub>); 5.25 (s, 2H, OCH<sub>2</sub>N); 7.70 (s, 1H, H<sup>5</sup>).

UV (λ<sub>max</sub> (nm), H<sub>2</sub>O): 265.

MS (FAB, m/z): 316 [MH]<sup>+</sup>

Anal. calc. for C<sub>12</sub>H<sub>17</sub>N<sub>3</sub>O<sub>7</sub>: C 45.71, H 5.43, N 13.33; Found: C 45.69, H 5.40, N 13.40.

## References

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  2. Purkayastha, S.; Lazrek, H. B.; Panzica, R. P.; Naguib, F. N. M.; El-Kouni, M. H. *Nucleosides & Nucleotides* **1989**, *8*, 349-356.

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*Sample Availability:* Available from the authors.

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