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Development of a flow process for an easy and fast access to 2-pyrone derivatives

Grazia Isa C. Righetti, Francesca Tentori, M. Elisabetta Brenna, Cristian Gambarotti

Department of Chemistry, Materials and Chemical Engineering "Giulio Natta", Politecnico di Milano, piazza Leonardo da Vinci 32, I-20133 Milano, Italy.

Electronic Supplementary Information

Continuous flow reaction's details

The continuous flow reactions were performed using an E-Series Integrated Flow Chemistry system from Vapourtec (Alfatech s.p.a., Genoa, Italy) equipped with a 10 mL PFA coiled tubular reactor (1.0 mm i.d. \times 12.75 m length). The coil reactor was kept under controlled pressure using back pressure regulators (BPR) from IDEX Health & Science L.L.C. (Microcolumn s.r.l., Desio, Italy).



The reactor used in the research

Spectra of isolated compounds

The ¹H-NMR was recorded on a Bruker AV 400 MHz instrument, equipped with a 5 mm multinuclear probe, and 32 scans were acquired with an acquiring time of 3 seconds for each spectrum. The chemical shifts are referred to the signal of the solvent residual peak (¹H: CDCl₃ = 7.26 ppm, DMSO-d6 = 2.5 ppm, D₂O = 4.79 ppm; ¹³C: CDCl₃ = 77.36 ppm, DMSO-d6 = 40.45 ppm; *J. Org. Chem.* **1997**, *62*, 7512-7515)

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- **Figure S4:** ¹³C-NMR (100.6 MHz, CDCl₃) of 3-hydroxy-2-pyrone (**3**)
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Figure S1: ¹H-NMR (400 MHz, DMSO-d6) of pyrone 2 sodium salt



Figure S2: ¹³C-NMR (100.6 MHz, DMSO-d6) of pyrone 2 sodium salt



Figure S3: ¹H-NMR (400 MHz, CDCl₃) of 3-hydroxy-2-pyrone (3)



Figure S4: ¹³C-NMR (100.6 MHz, CDCl₃) of 3-hydroxy-2-pyrone (3)



Figure S5: ¹H-NMR (400 MHz, D₂O) of 3-hydroxy-2-pyrone (**3**)