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Convenient Synthesis of  $\alpha$ -Trifluoromethylated Acylolins From  $\alpha$ -Hydroxy or  $\alpha$ -Amino AcidsMasami Kawase (河瀬雅美)<sup>1</sup>, Setsuo Saito<sup>1</sup>, and Teruo Kurihara<sup>2</sup><sup>1</sup>Faculty of Pharmaceutical Sciences, Josai University, Saitama, Japan;<sup>2</sup>Faculty of Science, Josai University, Saitama, Japan.

**Abstract.**  $\alpha$ -Trifluoromethylated acylolins (2 and 6) have been prepared from  $\alpha$ -hydroxy acids (1), *N*-acylprolines (5) or *N*-acyl-*N*-alkyl  $\alpha$ -amino acids (8) by novel transformation reactions with trifluoroacetic anhydride (TFAA) in the presence of pyridine. The former reaction of 1 could proceed through mesoionic 1,3-dioxolium-4-olates, whereas the latter two reactions of  $\alpha$ -amino acids (5 and 8) could involve mesoionic 1,3-oxazolium-5-olates. The reaction of 1 with TFAA shows more potential for practical applications because of the ready availability of the starting materials and ease of manipulation.

