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Title: Highly efficient Cu(I)-catalyzed trifluoromethylation of aryl(heteroaryl) enol acetates with  $CF_3$  radicals derived from  $CF_3SO_2Na$  and TBHP at room temperature

Author: Yang Lu Yaming Li Rong Zhang Kun Jin Chunying

Duan

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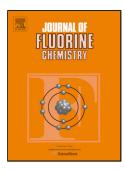
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## ACCEPTED MANUSCRIPT

Highly efficient Cu(I)-catalyzed trifluoromethylation of aryl(heteroaryl) enol acetates with  $CF_3$  radicals derived from  $CF_3SO_2Na$  and TBHP at room temperature

Yang Lu, Yaming Li\*, Rong Zhang, Kun Jin, Chunying Duan\*

State Key Laboratory of Fine Chemicals, School of Chemical Engineering, Dalian

University of Technology, Dalian 116024, China

Tel./fax: +86 411 84986295; E-mail address: ymli@dlut.edu.cn (Y. Li)

OAc
$$R^{1} = \text{aryl, heteroaryl}$$

$$R^{2} = \text{H. Me. Ph}$$

$$R^{1} = \text{Aryl, heteroaryl}$$

$$R^{2} = \text{H. Me. Ph}$$

$$R^{2} = \text{H. Me. Ph}$$

$$R^{3} = \text{If mol% Cul}$$

$$5 = \text{equiv TBHP}$$

$$CH_{3}CN, RT, 2 = \text{H. Me. Ph}$$

$$14 = \text{examples}$$

$$61-96\% \text{ yields}$$

**ABSTRACT:** An efficient method for the Cu(I)-catalyzed synthesis of α-trifluoromethyl ketones via the addition of CF<sub>3</sub> to aryl(heteroaryl) enol acetates by using the readily available CF<sub>3</sub>SO<sub>2</sub>Na (Langlois reagent) has been developed. The reaction is experimentally simple and carried out at room temperature, providing good to excellent yields with wide functional group tolerance.

*Keywords:* Trifluoromethylation; Enol acetates; Synthetic methods; α-CF<sub>3</sub> ketone; Radical addition

#### 1. Introduction

Trifluoromethylated compounds have been widely utilized as the important building blocks for the synthesis of pharmaceuticals, agrochemicals and specialty materials due to the stereoelectronic property of the CF<sub>3</sub> moiety and the increased bioavailability [1]. Therefore, it has been of great interest to develop more efficient, operationally simple and

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