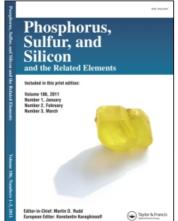
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Nucleophilic Epoxidation of α -Sulfonyl- α , β -Unsaturated Esters with m-CPBA

J. L. García Ruano^a; C. Fajardo^a; A. Fraile^a; M. R. Martićn^a

^a Departamento de Qu'i mica Orgánica, Universidad Autónoma de Madrid, Madrid, Spain

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J. L. García Ruano
C. Fajardo
A. Fraile
M. R. Martín
Departamento de Química Orgánica, Universidad Autónoma de Madrid,

INTRODUCTION

m-CPBA is usually described as an electrophilic oxidant that transfers oxygen to non electron-poor alkenes. There are a few reports about the epoxidation with *m*-CPBA of deactivated olefins. ^{1a-d} The reactions of 1-phenyl-(2-p-tolylsulfonyl)ethane^{1a,b} and 2-nitro-1-phenylpropene^{1c} using m-CPBA in basic aqueous media were described. We report herein the results obtained in the reaction of the title sulfones with m-CPBA.

RESULTS

The reactions of sulfinylfuranones **1a–c** with an excess of *m*-CPBA in CH₂Cl₂ at 0°C for 1.5–8 h afforded the corresponding sulfones **2a–c**, which were not isolated and in situ evolved into the sulfonyl oxiranes **3a-c** in good yields (75–89%). The reactions are stereoselective, affording only the products resulting from the anti approach of the reagent with respect to the OEt group. Under similar conditions, furanone 4 did not evolve into its epoxide derivative (Scheme 1).

R = H (1a, 2a, 3a); R = Me (1b, 2b, 3b); R = Et (1c, 2c, 3c)

SCHEME 1

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Address correspondence to C. Fajardo, Departamento de Química Orgánica, Universidad Autónoma de Madrid, Cantoblanco, 28049 Madrid, Spain. E-mail: cristina.fajardo@ uam.es

The above results suggest that m-CPBA is also able to act as a nucleophile on the proper substrates, which lead us to investigate the epoxidation of different α,β -unsaturated sulfones (5–6) with m-CPBA (Scheme 2 and Table I).

SCHEME 2

Eq. m -CPBA/eq. base	$T(^{\circ}C)$	Time	Ratio 5:7
2.10/0	$ \begin{array}{r} 20 \\ -20 \\ -20 \end{array} $	87 h	27:73
1.15/1.15 K ₂ CO ₃		40 min	0:100 (91)
1.15/1.15 KOH		10 min	0:100 (88)

The reaction with *m*-CPBA in dichloromethane was complete to give oxirane only in the case of the furanones **2**. Cyclopentenone **5** afforded only 73% of oxirane **7** after 87 h and the open-chain compounds did not undergo epoxidation under these conditions. The addition of base to solution of *m*-CPBA allowed us to obtain the sulfonyl-oxiranes **8** in good yields; an increase in the reaction rate of **2** and **5** was also detected.

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