Tributylphosphine as an Efficient Catalyst in C-Vinylation of Naphthol Derivatives

ALIREZA HASSANABADI*a, MOHAMMAD H. MOSSLEMINb and ZAHRA ALSADAT RASOLHOSSEINIb

*aDepartment of Chemistry, Islamic Azad University, Zahedan Branch, P.O. Box 98135-978, Zahedan, Iran
bDepartment of Chemistry, Islamic Azad University, Yazd Branch, P.O. Box 89195-155, Yazd, Iran
ar_hasanabadi@yahoo.com

Received 20 May 2012 / Accepted 4 June 2012

Abstract: A diastereospecific c-vinylation of naphthol derivatives is described from a reaction of dialkyl acetylenedicarboxylate and naphthol derivatives, in the presence of tributylphosphine as an efficient catalyst, in excellent yield. The presented one-pot method is simple and effective with mild conditions.

Keywords: C-Vinylation, Dialkyl acetylenedicarboxylates, Tributylphosphine, Naphthol derivatives

Introduction

Vinylation of aromatic compounds is an important reaction in organic synthesis and much attention has been paid to the synthesis of vinyl compounds and their functionalized derivatives. However, there are few reports on direct vinylation of aromatic compounds1. In 1908, pure O-vinyl phenol was synthesized for the first time by decarboxylation of o-hydroxycinnamic acid2. Since then, a number of methods were devoted to the synthesis of vinylphenols. Electrophilic acylation of phenol followed by reduction and dehydration was employed in the commercial production of p-vinylphenol by Maruzen Petrochemicals Co3. Another method that utilized benzylic oxidation of ethylphenol was reported4. Halophenol derivatives could be vinylated by the Heck reaction5. Pyrolysis of heterocyclic compounds was reported6. The palladium-catalyzed arylation of olefins with aryl halides is a well-established method of carbon–carbon bond formation. Because this versatile reaction, discovered by Heck7 and Mizoroki8, is tolerant of a wide variety of functional groups on either coupling partner, it has gained much interest for the preparation of elaborate arenes from relatively simple starting materials9. Ortho-vinylation reaction of anilines was reported in the literature10. In recent years there has been increasing interest in the synthesis of organophosphorus compounds, particularly those bearing a carbon atom bound directly to a
phosphorus atom. This interest has resulted from the recognition of the value of such compounds for a variety of industrial and chemical synthetic uses. Organophosphorus compounds also have been extensively used in organic synthesis as useful reagents as well as ligands of a number of transition metal catalysts. However, there are few reactions in which organophosphorus(III) species work as catalysts.

Development of a new approach for a highly stereoselective and efficient C-vinylation of naphthol derivatives is still in demand. Previously, in our investigations of stable phosphorus ylides, many condensation reactions between triphenylphosphine and dialkyl acetylenedicarboxylate in the presence of C-H, N-H, O-H or S-H acidic organic compounds were reported. Herein, we introduce a new method for the diastereospecific synthesis of (E)-C-vinyl naphthol derivatives from a reaction between dialkyl acetylenedicarboxylate and naphthol derivatives in the presence of tributylphosphine.

**Experimental**

Melting points were determined with an Electrothermal 9100 apparatus. IR spectra were recorded on a Shimadzu IR-470 spectrometer. H, C spectra were recorded on BRUKER DRX-500 AVANCE spectrometer in CDCl₃ using TMS as internal standard. The chemicals used in this work were purchased from Fluka (Buchs, Switzerland) and were used without further purification.

**General procedure for preparation of compounds (4a-g)**

To a magnetically stirred solution of naphthol derivative (2 mmol) and tributylphosphine (2 mmol) in 10 mL acetone, a mixture of dialkyl acetylenedicarboxylate (2 mmol) in 2 mL acetone was added at −4 °C over 5 min. The reaction mixture was then allowed to stir for 12 h. The solvent was evaporated at reduced pressure. The residue was precipitated in a solution of diethyl ether-hexane. The solid was filtered and washed with diethyl ether to give the pure product.

**Results and Discussion**

The reaction of dialkyl acetylenedicarboxylate (2) with naphthol derivatives (3), in the presence of tributylphosphine (1) as an efficient catalyst, leads to diastereospecific C-vinylation of those in high yields (Figure 1).

![Figure 1. Synthesis of C-vinylation of naphthol derivatives](image)

These reactions were carried out in acetone at −4 °C. The compounds 4a-g (Table 1) result from the initial addition of tributylphosphine (1) to dialkyl acetylenedicarboxylate (2) and subsequent protonation of the 1:1 adduct by naphthol derivative 3. Then, the positively charged ion 5 is attacked by the anion 6 to form ylide 7. Phosphonium ylide undergoes 1,2-H shift to give compound 8. Next, tributylphosphine is going out for generating alkenes 4. The proposed mechanism for C-vinylation of naphthol derivatives is shown in Figure 2.
Table 1. C-Vinylation of naphthol derivatives

<table>
<thead>
<tr>
<th>Entry</th>
<th>X</th>
<th>R</th>
<th>Product</th>
<th>Time/h</th>
<th>Yield %a</th>
<th>m.p./°C</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>Found</td>
</tr>
<tr>
<td>1</td>
<td></td>
<td>Me</td>
<td>4a</td>
<td>12</td>
<td>90</td>
<td>179-181</td>
</tr>
<tr>
<td>2</td>
<td></td>
<td>Et</td>
<td>4b</td>
<td>12</td>
<td>84</td>
<td>62-64</td>
</tr>
<tr>
<td>3</td>
<td></td>
<td>i-Pr</td>
<td>4c</td>
<td>12</td>
<td>88</td>
<td>84-86</td>
</tr>
<tr>
<td>4</td>
<td></td>
<td>Me</td>
<td>4d</td>
<td>12</td>
<td>95</td>
<td>178-180</td>
</tr>
<tr>
<td>5</td>
<td></td>
<td>Et</td>
<td>4e</td>
<td>12</td>
<td>93</td>
<td>166-168</td>
</tr>
<tr>
<td>6</td>
<td></td>
<td>Me</td>
<td>4f</td>
<td>12</td>
<td>92</td>
<td>165-167</td>
</tr>
<tr>
<td>7</td>
<td></td>
<td>Et</td>
<td>4g</td>
<td>12</td>
<td>90</td>
<td>153-155</td>
</tr>
</tbody>
</table>

aYield refer to the pure isolated products

Figure 2. Suggested mechanism for formation of compound 4

The compounds 4a-g were characterised by their ¹H, ¹³C NMR and IR spectroscopy and elemental analyses. The structures of the those were proved by comparing the ¹³C{¹H} NMR spectral data of related compounds reported elsewhere. The chemical shift of the olefinic proton in the ¹H NMR spectrum of compounds are consistent with the E-geometry of the carbon-carbon double bond.

Conclusion

We have shown that tributylphosphine as an efficient catalyst has advantages in the C-vinylation of naphthol derivatives such as shorter reaction times, simple work-up and affords excellent yield. The present method carries the advantage that, not only is the reaction performed under neutral conditions, but the educts can be mixed without any activation or modification.

Acknowledgement

We gratefully acknowledge financial support from the Research Council of Islamic Azad University of Zahedan and The Islamic Azad University of Yazd of Iran.
References