Stereoselective synthesis of 2,3-disubstituted dihydrobenzofuran using alkyne Prins type cyclization to vinylogous carbonates

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Abstract. An intramolecular, alkyne Prins type cyclization of vinylogous carbonates derived from *o*-alkynyl phenols is developed for the stereoselective construction of *trans*-2,3-disubstituted dihydrobenzofuran derivatives. Strong Lewis acids like TMSOTf catalyse this reaction efficiently. The presence of mildly electron donating groups on aryl rings increases the efficiency of the reaction.

Keywords. Vinylogous carbonates; Prins cyclization; Lewis acids; Sonogashira coupling.

1. Introduction

2,3-Disubstituted dihydrobenzofuran moiety is ubiquitous in nature and molecules possessing this skeleton display remarkably diverse biological activity. Many of these natural products are constituents of traditional medicines and have been implicated with diverse biological activity such as antimicrobial,¹ antioxidant,² antimitotic,³ antiangiogenic,⁴ neuritogenic⁵ and even HIV integrase inhibition activity.⁶ As a result, a variety of methods have been developed over the years for the stereoselective synthesis of this motif.⁷ Acid catalysed [3 + 2] cycloaddition of styrene derivatives with quinones,⁸ cyclodehydration of hydroxyphenols,⁹ anion or radical induced cyclization of substituted iodophenols,¹⁰ Rh(I) catalysed catalytic C–H insertion,¹¹ reactions of 2-hydroxyaryl- α,β -unsaturated ketones with dimethylsulfonium carbonylmethylides,¹² Pd-catalysed cyclizations¹³ and reduction of benzofurans¹⁴ are some of the prominent methods used for the synthesis of 2,3-disubstituted dihydrobenzofurans. Chiral Rh(II)-catalysed intramolecular C–H insertion reaction has also been reported for enantioselective synthesis of this motif.¹⁵ Though majority of these methods give access to *cis* isomer of 2,3-disubstituted dihydrobenzofuran, it is the *trans* isomer which is more prevalent in many of the natural products. All these methods have their advantages but many suffer from moderate diastereoselectivity. Thus, there is still need

for developing highly diastereoselective strategies for this important class of molecules. Over the years, Prins cyclization has emerged as a useful method for the synthesis of tetrahydropyrans (THPs) and tetrahydrofurans (THFs).¹⁶ Vinylogous carbonate, which has proved to be excellent functional group for the synthesis of cyclic ethers under radical¹⁷ and anionic conditions,¹⁸ has also been found to be useful in the Lewis/Bronsted acid mediated Prins cyclization leading to THP derivatives.¹⁹ Even though most of the efforts on Prins cyclization used olefins; alkynes too have been shown to participate efficiently in the synthesis of THFs and THPs.²⁰ In continuation of our interest on using vinylogous carbonates in the synthesis of cyclic ethers,²¹ particularly under non-radical conditions,²² we describe here an efficient and a highly diastereoselective synthesis of *trans*-2,3-disubstituted dihydrobenzofuran employing alkyne Prins cyclization to vinylogous carbonates.

2. Results and discussion

Recently, Cho *et al.* reported TMSOTf mediated synthesis of 5- and 6-exocyclic products, *cis*-2,3,5-trisubstituted THFs, and *cis*-2,3,6-trisubstituted THPs by Prins type reaction between terminally substituted alkynyl alcohols and aldehydes.²⁰ When this type of reaction was attempted by them with 2-alkynylphenols **1** with aldehydes **2** in the presence of TMSOTf, it generated the chalcones **3**, chroman-4-ones **4** and hydrated products **5** depending on the substituent on the alkyne in moderate yields (scheme 1).²³ These products are

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Scheme 1. TMSOTf-Promoted addition of alkynes to aldehydes.

Scheme 2. Retrosynthesis for 2,3-disubstituted dihydrobenzofurans.

Scheme 3. Synthesis of the vinylogous carbonate **7a**.

formed by the reaction of alkyne directly with the aldehyde rather than by formation of oxonium ion from phenol and aldehyde followed by subsequent cyclization.

We envisioned that if the phenolic OH of 2-alkynylphenol is engaged in the reaction for generating oxonium ion in the presence of Lewis acid, the reaction could lead to 2,3-disubstituted dihydrobenzofuran via a

Prins type cyclization. However, this oxonium ion generation by intermolecular reaction of a phenol and an aldehyde is non-trivial. To circumvent this problem, we argued that the 2,3-disubstituted dihydrobenzofurans **6** could be readily assembled by carrying out intramolecular alkyne Prins cyclization of the alkynyl vinylogous carbonates **7** using an appropriate Lewis acid

Table 1. Optimization of the alkyne Prins cyclization for the synthesis of 2,3-disubstituted dihydrobenzofurans.

S.No	Lewis acid	Equiv.	Time (h)	Yield (%) ^a	d.r. ^b
1	BF ₃ ·OEt ₂	1.1	24	74	(≥19:1)
2	TMSOTf	1.1	6	79	(≥19:1)
3	TMSOTf	1.8	5	65	(≥19:1)
4	TiCl ₄	1.1	15	30	(≥19:1)
5	FeCl ₃	1.1	24	0 ^c	-
6	CF ₃ SO ₃ H	1.1	15	44	(≥19:1)
7	CF ₃ CO ₂ H	1.1	24	0 ^c	-

^aIsolated yield. ^bDetermined on crude reaction mixture by ¹H NMR. ^cUnreacted starting material recovered

Table 2. Synthesis of the iodo vinylogous carbonates **8**.

Sr. No.	R ¹	R ²	Product	Yield (%) ^a
1.	Me	H	8b	89
2.	^t Bu	H	8c	87
3.	Cl	H	8d	90
4.	Ph	H	8e	64 ^b
5.	C ₄ H ₄	C ₄ H ₄	8f	90

^aIsolated yield. ^b24% of *cis* isomer of vinylogous carbonate was also obtained

(scheme 2). It was rationalized that the vinylogous carbonate moiety derived from phenol will serve as a surrogate for oxonium ion. The alkynyl vinylogous carbonates **7** in turn can be synthesized from the corresponding iodides **8** by Sonogashira coupling reaction. The *o*-iodo vinylogous carbonates **8** can be obtained from substituted *o*-iodophenol derivatives **9**.

To test the hypothesis, synthesis of the alkyne **7a** was initiated. Thus, reaction of *o*-iodophenol (**9a**) with ethyl propiolate in the presence of N-methylmorpholine (NMM) gave the *o*-iodo vinylogous carbonate **8a** in good yield. The iodo vinylogous carbonate **8a** was then subjected to Pd catalysed Sonogashira coupling reaction to furnish the enyne **7a** in excellent yield (scheme 3).

The feasibility of the envisaged intramolecular alkyne Prins cyclization of enyne **7a** was studied next using various Lewis acids. Reaction of the enyne **7a** with BF₃·OEt₂ in CH₂Cl₂ at 0°C gratifyingly gave the 2,3-disubstituted dihydrobenzofuran **6a** in good yield and excellent diastereoselectivity (table 1, entry 1). The reaction time was substantially reduced when TMSOTf was used as the Lewis acid for effecting this transformation with slight increase in yield (table 1, entry 2). Increasing the amount of TMSOTf did not show further improvements in the reaction time or the yield (table 1, entry 3). When TiCl₄ was used to promote the reaction, it resulted in the formation of dihydrobenzofuran **6a** in only 30% yield; whereas FeCl₃ was found to be ineffective in this reaction (table 1, entries 4–5). While a strong Bronsted acid like CF₃SO₃H gave moderate yield albeit with good diastereoselectivity, the milder trifluoroacetic acid failed to promote the alkyne Prins cyclization (table 1, entries 6–7). Based on these results it was clear that mild Lewis or Bronsted acids

are not good for effecting this alkyne Prins cyclization and hence TMSOTf was chosen as the catalyst of choice for further study. The structure of the dihydrobenzofuran derivative **6a** rests secured from its spectral data.¹ The stereochemistry of dihydrobenzofuran **6a** was ascertained as *trans*, based on coupling constants of H₂ and H₃ protons (*J* = 6.3 Hz) which was comparable with that reported earlier for similar compounds. Further support was provided by the NOE studies in which irradiation of H₃ proton resulted in enhancement in the CH₂CO₂Et intensity and vice versa (see supporting information).

To study the scope of this alkyne Prins cyclization for the synthesis of *trans*-2,3-disubstituted dihydrobenzofurans, synthesis of various vinylogous carbonates **8** was undertaken. Various iodophenols **9** were subjected to Michael addition with ethyl propiolate to give the corresponding *o*-iodo vinylogous carbonates **8** (table 2). In general, the formation of iodo vinylogous carbonates **8** was uneventful and in all the cases excellent yields were obtained. Only in the case of the iodophenol **9e**, the reaction led to formation of a mixture of *cis* and *trans* vinylogous carbonates **8e** which could be separated by column chromatography on silica gel.

Table 3 describes the scope of this intramolecular alkyne Prins cyclization to vinylogous carbonates. In all the cases, the requisite alkynyl vinylogous carbonates **7** were prepared by Sonogashira coupling of the iodide **8** with aryl acetylene **10** using Pd(OAc)₂/PPh₃/CuI as the catalyst. In general, the reaction proceeded smoothly furnishing the coupling products **7**.

¹ All the compounds exhibited spectral data consistent with their structures, see supporting information.

Table 3. Scope of the alkyne Prins cyclization for the synthesis of 2,3-disubstituted dihydrobenzofurans.

S. No.	Iodide vinylogous carbonate (8)	Aryl acetylene (10)	Yield(%) ^a		Dihydrobenzofuran (6) ^b
			Step I	Step II	
1	8b	C ₆ H ₅ CCH	82	53	
2	8c	C ₆ H ₅ CCH	83	60	
3	8d	C ₆ H ₅ CCH	90	35	
4	8e	C ₆ H ₅ CCH	87	53	
5	8f	C ₆ H ₅ CCH	88	81	
6	8a	<i>p</i> -MeC ₆ H ₄ CCH	92	69	
7	8b	<i>p</i> -MeC ₆ H ₄ CCH	86	64	
8	8c	<i>p</i> -MeC ₆ H ₄ CCH	91	62	

Table 3. (continued).

S. No.	Iodide vinylogous carbonate (8)	Aryl acetylene (10)	Yield(%) ^a		Dihydrobenzofuran (6) ^b
			Step I	Step II	
9	8d	<i>p</i> -MeC ₆ H ₄ CCH	80	55
10	8e	<i>p</i> -MeC ₆ H ₄ CCH	85	36
11	8f	<i>p</i> -MeC ₆ H ₄ CCH	91	67
12	8a	<i>p</i> -(MeO)C ₆ H ₄ CCH	85	0 ^c
13	8a	<i>p</i> -(CN)C ₆ H ₄ CCH	83	0 ^c

^aIsolated yields. ^bd.r. was determined on crude reaction mixture by ¹H NMR and found to be ≥19:1 in all cases.^cDecomposition during the reaction

The alkyne Prins cyclization was studied on these alkynyl vinylogous carbonates **7**. The reaction was found to work very well with mildly electron donating alkyl substituents on any of the aryl rings (table 3, entries 1–2 and 6–8). The presence of electron withdrawing chlorine on one of the aryl ring led to the formation of the product in moderate yield (table 3, entry 4). Interestingly, when a methyl substituent was introduced in the other aryl ring, the yield of the reaction showed some improvement (table 3, entry 9). Phenyl substitution on the aryl ring in general resulted in the moderate efficiency (table 3, entries 4 and 10); whereas the naphthyl ring bearing substrates furnished the dihydrobenzofurans **6f** and **6l** in excellent yields. Interestingly, the presence of relatively stronger electron releasing (OMe) or electron withdrawing group (CN) in one of the aryl ring led to extensive decomposition and no dihydrobenzofuran formation could be detected (table 3, entries 12–13).

Formation of the 2,3-disubstituted dihydrobenzofuran can be explained with the help of the mechanism as shown in figure 1. Initially, the carbonyl group of the vinylogous carbonate moiety complexes with the Lewis acid leading to the formation of the oxonium ion **10**. This oxonium ion **10** is trapped by alkyne in a Prins type reaction generating a vinyl cation **11**. The vinyl cation **11** gets trapped with water during the work-up generating enol **12** which on tautomerization generates the 2,3-disubstituted dihydrobenzofuran **6**. Formation of *trans* isomer during tautomerization is perhaps the outcome of the protonation event which happens in such a way so as to minimize steric repulsion between the substituents on C2 and C3 of the dihydrobenzofuran moiety. It is pertinent to mention that changing the solvent from CH₂Cl₂ to ether did not result in formation of triflate derivative by trapping of vinyl cation **11** with the triflate ion as observed by Cho and co-workers.

Figure 1. Mechanism of formation for *trans*-2,3-disubstituted dihydrobenzofuran.

3. Conclusion

We have developed a highly stereoselective synthesis of *trans*-2,3-disubstituted dihydrobenzofurans using alkyne Prins cyclization to vinylogous carbonates. Generation of the oxonium ion by the intermolecular reaction of the phenolic hydroxyl group with an aldehyde is in general a non-trivial task. Notable feature of this study is that not only was this task made easy by engaging phenolic OH in the vinylogous carbonate moiety but also the reactivity of the generated oxonium ion was harnessed in the stereoselective transformation. The reaction is found to be sensitive to substituents on the aryl ring. In general, mildly electron donating groups are found to give better reactivity profile. Further efforts are underway to understand substituent effects and optimize these reactions for improving the substrate scope and applying them in target directed synthesis.

Supporting information

Synthetic procedures and characterization data for all the new compounds are given as electronic supporting information. See www.ias.ac.in/chemsci for details.

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