FULL PAPER

HBF_4 Catalysed Nucleophilic Substitutions of Propargylic Alcohols

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Abstract: The activity of HBF_4 (aqueous solution) as a catalyst in propargylation reactions is presented. Diverse types of nucleophiles were employed in order to form new C–O, C–N and C–C bonds in technical acetone, and in air. Good to excellent yields were obtained using low acid loading (typically 1 mol %) under simple reaction conditions and good chemoselectivity.

Introduction

The direct nucleophilic substitution of alcohols is of high interest as it provides access to a wide variety of derivatives, with the formation of water as the only by-product. Indeed, the ACS Green Chemistry Institute Pharmaceutical Roundtable identified OH activation for nucleophilic substitutions as a priority area currently used in the preparation of pharmaceutical intermediates that would greatly benefit from the development of better methods.[1] Unarguably, propargylic substitutions have progressed substantially since the pioneering work of Nicholas on dicobalt octacarbonyl-stabilised propargylic cations.[2] The versatility of the propargylic moiety as a synthon in organic chemistry as well as its occurrence in natural products and synthetic pharmaceuticals have been the main driving forces for these advances. Furthermore, propargylic alcohols are easily prepared from the corresponding aldehyde or ketone *via* addition of an alkynyl anion. However, propargylic substitution reactions remain underdeveloped when compared to allylic substitutions. Diverse transition metals,[3] such as ruthenium, palladium, gold or silver,[4] have been successfully used in this context. However, the cost of the catalyst, together with its selectivity (metal-allenylidene *vs* metal-propargylic intermediates) remain important issues to solve.

The direct displacement of 'activated' alcohols -such as benzylic, allylic, and propargylic alcohols- can also be achieved using Brønsted or Lewis acids through simple S_N1 reactions.[5] Important advantages of Brønsted acids over Lewis acids often include lower catalytic loadings and easier handling as they are generally more stable towards oxygen and water.

Sulfonic acids are the most commonly used Brønsted acids for the nucleophilic substitution of propargylic alcohols as described in extensive work by Sanz and co-workers with *p*-toluenesulfonic acid.[6],[7] Inorganic acids, such as phosphomolybdic acid on silica, have also been studied with *C*-, *N*- and *O*-nucleophiles.[8] Depending on the substrates, the reactions required either 10 mol % of acid at room temperature, or 1 mol % in refluxing toluene. An additional asset of these inorganic acids is their straightforward separation from the organic products through a simple basic workup. A common feature for all these catalytic systems is their compatibility with air and reagent-grade solvents, although they are mostly undesirable ones (toxic, costly to dispose of such as $MeNO_2$).

HBF_4 is a common acid in academic and industrial laboratories that has found diverse applications in synthesis, either as a reagent (nucleophilic fluorination,[9] synthesis of vinylidene metal complexes[10]), or catalyst (amidation of olefins,[11] Biginelli reaction,[12] acylation of aldehydes[13]). In particular, the Friedel-Crafts alkylation of benzylic alcohols has been reported using an excess of $HBF_4{\cdot}OEt_2$ solution at -78 °C.[14] Even though high diastereoselectivities could be achieved with this methodology, the excess of acid and low reaction temperatures represent important drawbacks. Herein, we report the use of HBF_4 as a highly efficient catalyst for S_N1 reactions of propargylic alcohols with different nucleophiles under mild, simple reaction conditions.

Results and Discussion

In a first step, the effect of different solvents was tested on the reaction of propargylic alcohol **1a** with MeOH to give **2a** with 1 mol % of HBF_4 (Table 1). These reactions were run in air with 2 equivalents of MeOH and using a commercially available 48 wt% solution of HBF_4 in water as catalyst. No attempts were made to optimise the reaction times. Whereas sluggish reactions were observed in THF or in water (Table 1, entries 1 and 2), high conversions were obtained in DCM, acetonitrile, and acetone (Table 1, entries 4-6). Overall, acetone was chosen as our preferred solvent because of its greener profile.[15] It is important to note that all tested solvents were technical grade, and in particular, the acetone employed in these reactions was standard laboratory washing acetone. Furthermore, the formation of fluoro derivatives or α,β-unsaturated compounds derived from a Meyer-Schuster rearrangements[16] was not observed either in the model reaction or during the study of the scope of the reaction.

The use of different *O*-nucleophiles was first explored (Scheme 1). Propargylic alcohol **1a** was reacted with different primary and secondary alcohols to form the expected ethers **2a**-**h** in high yields under our standard conditions. When chiral alcohols were

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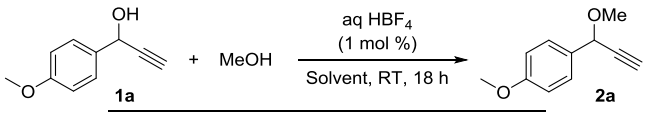
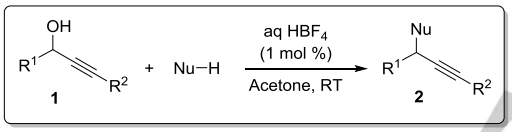
used, the corresponding ethers **2g** and **2h** were formed as a mixture of inseparable diastereoisomers. A tertiary alcohol, *t*-BuOH, only led to low yields of the ether **2i**, and the major product of that reaction (51% conversion) formed from dimerization of the starting propargylic alcohol (**3a**, *vide infra* for further details). On the other hand, an *ortho*-disubstituted aryl group was not detrimental to the reactivity of the propargylic alcohol, as

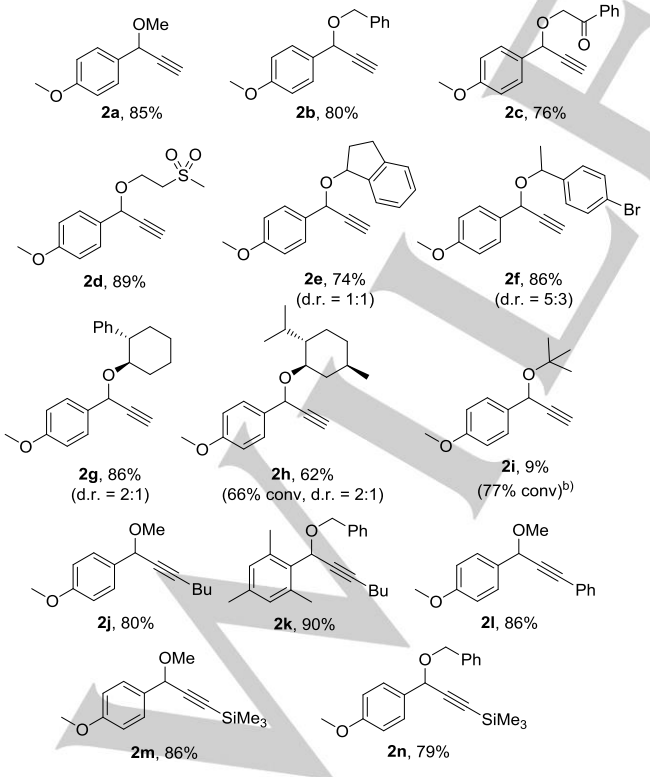
Table 1. Solvent screening for propargylation reactions.



Entry	Solvent	Conv (%)[a)]
1	THF	26
2	Water	47
3	Toluene	77
4	MeCN	93
5	DCM	>95
6	Acetone	>95

[a)] ^{1}H NMR Conversions

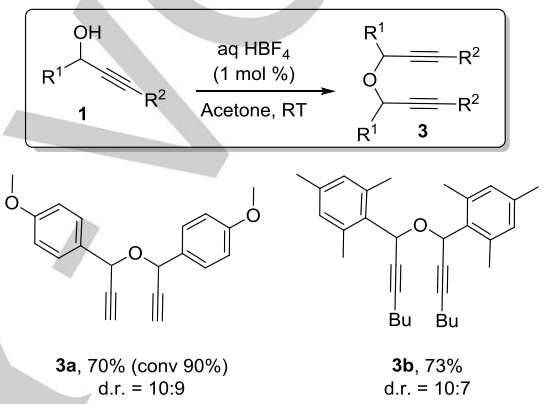




Scheme 1. C–O bond formation with HBF_4 at room temperature.[a)] d.r. = diastereoisomeric ratio.

[a)] Isolated yields, ^{1}H NMR conversions are provided in parentheses when lower than 95%. [b)] 51% ^{1}H NMR conversion into dimer **3a**.

exemplified with the formation of **2k**. Also, unlike most transition-metal-based methodologies,[3] the reaction is not limited to terminal alkynes, and alkyl (**2j**, **2k**), aryl (**2l**) or silyl groups (**2m**, **2n**) at the acetylenic position did not have any major effect on the outcome of the reaction (Scheme 1). Also, several functional groups (ketone, halogen or sulfone) were shown to be compatible with the reaction conditions. In the absence of any other nucleophile, the starting propargylic alcohol dimerised to form the symmetrical ether as a mixture of diastereoisomers (Scheme 2).



Scheme 2. HBF_4-catalysed dimerization of propargylic alcohols.[a)] d.r. = diastereoisomeric ratio.

a) Isolated yields, ^{1}H NMR conversions are provided in parentheses when lower than 95%.

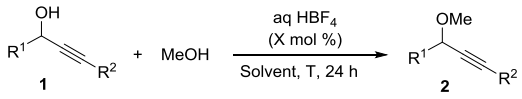
On the other hand, when R^1 on the starting propargylic alcohol was not an electron-rich aryl group, no reaction was observed at room temperature. In most cases however, the formation of the desired ethers was possible by increasing the reaction temperature and/or the acid loading (Table 2). Good yields could be then obtained, except for a nitro-substituted substrate (Table 2, entry 4). It is important to note that as no decomposition, or undesired reactions were observed at room temperature, this difference in reactivity could be used to selectively functionalise a more complex molecule with two electronically dissimilar propargylic alcohols (*vide infra*).

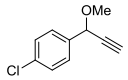
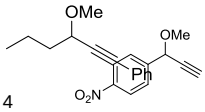
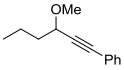
We next explored the use of nitrogen nucleophiles in this substitution reaction. The inherent basicity of most amines is an obvious potential limitation of any Brønsted acid-catalysed reaction as they might simply neutralise the catalyst. Our conditions, however, could be successfully applied to different carbamates and sulphonamides, as well as weakly basic anilines (Scheme 3), and the expected products **4** were prepared in good yields at room temperature, with the exception of **4e**.

Carbon nucleophiles were also investigated and diketones as well as electron-rich arenes reacted to form the expected products **5** in good to excellent yields (Scheme 4). Very similar results were obtained with pentane-2,4-dione and a variety of substituted propargylic alcohols. For the formation of **5d**, with an electron neutral aryl group at the propargylic position, a higher

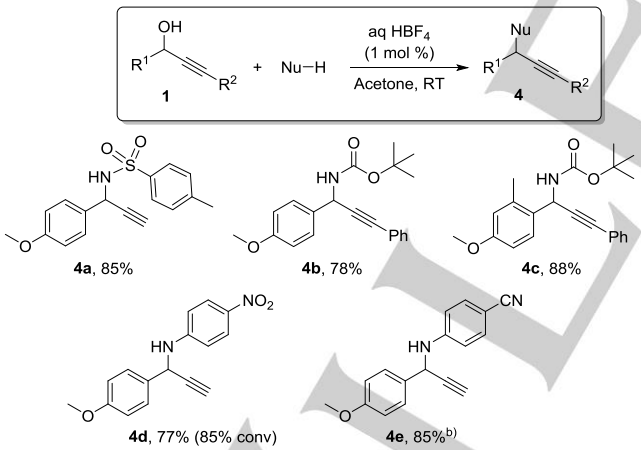
catalyst loading and an elevated temperature were required in order to obtain high conversions.

Table 2. HBF_4-catalysed reaction of electron-poor/neutral propargylic alcohols.



Entry	Product	2	Conditions	Yield (%)[a]
1		**2o**	1 mol % HBF_4 Toluene, 80°C	70
2		**2p**	5 mol % HBF_4 Acetone, 30°C	59 (70)
3		**2q**	5 mol % HBF_4 Toluene, 80°C	88 (94)
4		**2r**	5 mol % HBF_4 Toluene, 80°C	N.R.
5		**2s**	5 mol % HBF_4 Toluene, 80°C	70 (90)

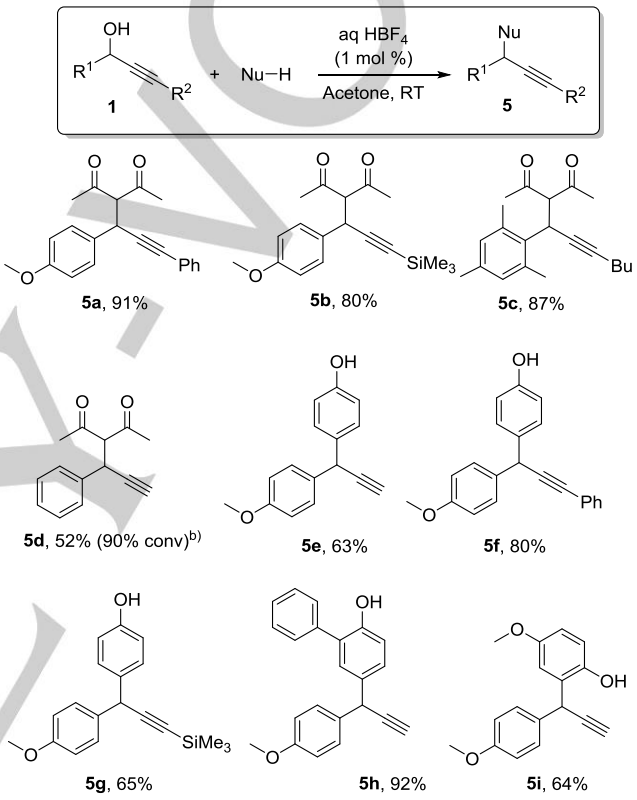
[a] Isolated yields, ^{1}H NMR conversions are provided in parentheses when lower than 95%. N.R. = No reaction



Scheme 3. C–N bond formation with HBF_4 at room temperature.[a]
a) Isolated yields, ^{1}H NMR conversions are provided in parentheses when lower than 95%. [b] Reaction carried out with 5 mol % of HBF_4 at 60°C.

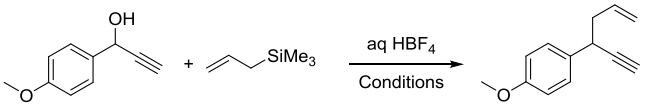
Phenol also reacted efficiently in a Friedel-Crafts type reaction,[17] to give *para*-substituted derivatives **5e**-**g** exclusively. When using 2-phenylphenol as the nucleophile, the hydroxyl group had a stronger directing power than the arene, as expected (Scheme 4, **5h**). We then tested a phenol with a second strongly activating group at the *para* position (4-methoxyphenol, for the formation of **5i**). In this case, only the product derived from reaction at the *ortho* position to the phenol was isolated.

Surprisingly, allyltrimethylsilane proved to be a very poor reaction partner for the substitution of propargylic alcohols with HBF_4 as the catalyst. Low conversions were obtained in either acetone or hot toluene, even when higher catalyst or nucleophile loadings were used (Table 3). Overall, the best results were obtained in MeCN at 80°C, and still product **5j** could only be isolated in 50% yield. It is important to note that alcohol **1a** was stable under the studied conditions and besides the expected product **5j**, only **1a** and ether **3a** were evidenced in the ^{1}H NMR spectra. Hence, no amide formation, which could potentially take place *via* a Ritter reaction,[18], [19] was observed under these conditions.



Scheme 4. C–C bond formation with HBF_4 at room temperature.[a]
[a] Isolated yields, ^{1}H NMR conversions are provided in parentheses when lower than 95%; [b] Reaction carried out with 5 mol % of HBF_4 in MeCN at 80°C.

Table 3. HBF_4-mediated reactions with an allylsilane.

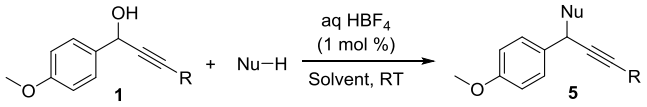


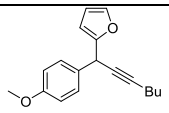
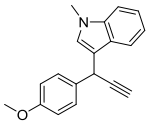
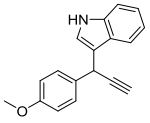
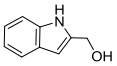
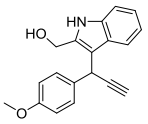
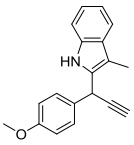
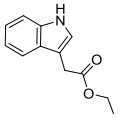
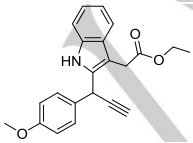
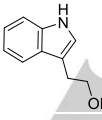
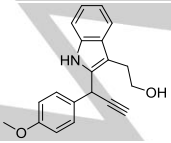
Entry	Conditions	Conv (%)[a]
1	2 equiv NuH, HBF_4 (1 mol %) Acetone, RT	18
2	2 equiv NuH, HBF_4 (5 mol %) Toluene, 80°C	30
3	3 equiv NuH, HBF_4 (5 mol %) Toluene, 80°C	35
4	2 equiv NuH, HBF_4 (5 mol %) MeCN, 80°C	72 (50)

[a] ^{1}H NMR conversions, isolated yield is provided in parentheses.

We next moved to electron rich heterocycles (Table 4). When furan was used as nucleophile, the ^{1}H NMR of the reaction crude showed the formation of a complex mixture of products. Prompt purification allowed the isolation of **5k** in moderate yield but it is important to note that **5k** still decomposed rapidly after purification. These observations were perhaps not surprising as many furan derivatives are well-known to be acid-sensitive. This is also the case for some indoles, but as it can be seen in Table 4, better yields were obtained with this important family of heterocyclic nucleophiles.[20]

Table 4. HBF_4-catalysed propargylation reactions of heterocycles.

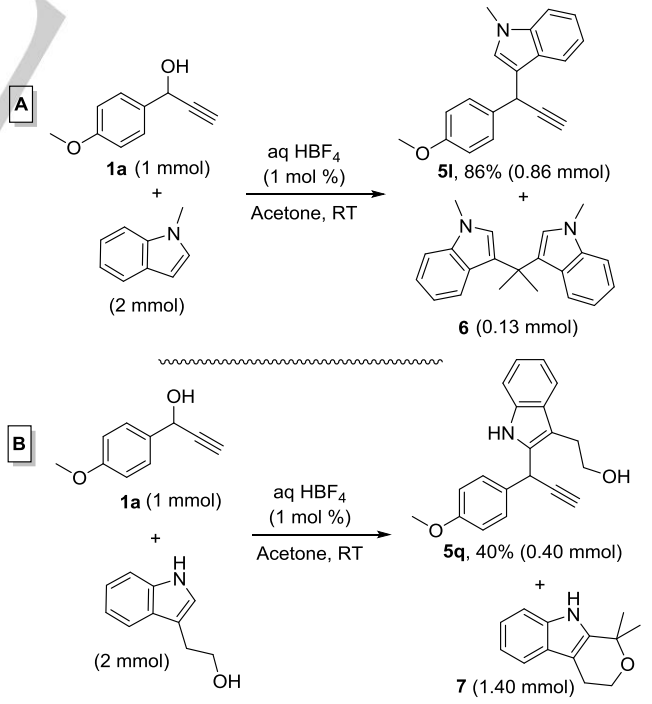


Nu—H	Product	5	Solvent	Yield (%)[a]
		**5k**	Acetone	41
		**5l**	DCM	85
		**5m**	DCM	70
		**5n**	Toluene	12 (20)
		**5o**	Acetone	58 (80)
		**5p**	DCM	30 (70)
		**5q**	DCM	60

[a] Isolated yields, ^{1}H NMR conversions are provided in parentheses when lower than >95%; [b] Reaction carried out with 5 mol % of HBF_4 in toluene at 80°C.

In order to investigate the influence of substitution on the nucleophile, different indole derivatives were reacted with the same propargylic alcohol **1a**. 1-Methyl and 1-H-indoles reacted regioselectively through C-3, as expected, leading to the formation of **5l** and **5m**, respectively, in good yields. Related indole-2-methanol, in contrast, only gave very low conversions. This might be due to a higher instability under acidic conditions or its low solubility either in acetone, or DCM. Even under more forcing conditions (5 mol % of acid at 80°C), no evidence for reaction through the hydroxyl group could be detected. We next screened different indoles with a substituent on C-3. Gratifyingly, products **5o**-**q**, derived from a Friedel-Crafts reaction on C-2 could be prepared under very simple reaction conditions. To the best of our knowledge, this is the first example of the synthesis of 2,3-disubstituted indoles via Brønsted acid-catalysed propargylation reactions.[21]

Some of the reactions in Table 4 were carried out in DCM, instead of acetone, to avoid the formation of undesired by-products. It has previously been reported that indoles can react with ketones or aldehydes as electrophiles under acidic conditions.[22] Indeed, when 1-methylindole was reacted with propargylic alcohol **1a** under our standard conditions in acetone, the expected product **5l** was formed preferentially, but it was contaminated with bis-indole **6** (Scheme 5, A). The high yield obtained of **5l** indicates that the indole reacts preferentially with the propargylic cation formed from **1a** and that the reaction between the excess of indole and acetone is quite sluggish.



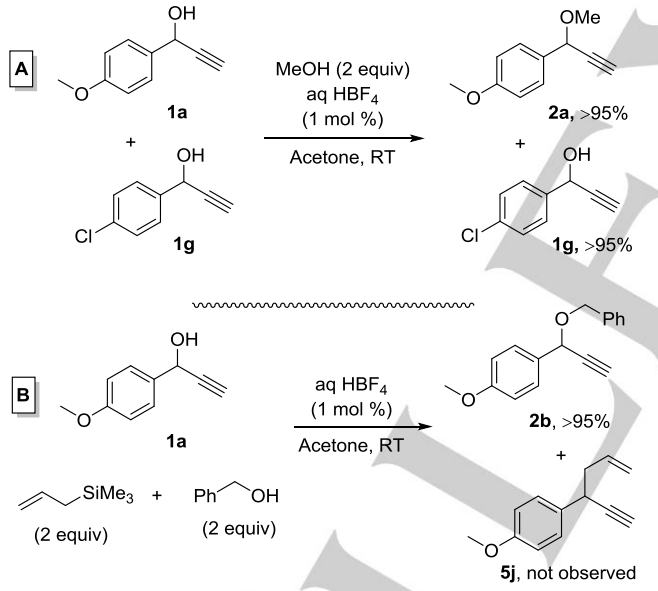
Scheme 5. Undesired reactions of indoles in acetone. Isolated yields are provided.

Even if both compounds can be separated by column chromatography the formation of **6** remains undesirable and hence acetone was avoided as a solvent for these reactions.

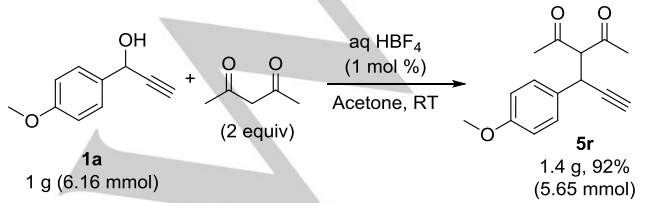
More problematic was the reaction of **1a** with tryptophol as nucleophile (Scheme 5, B). In this case the expected product could only be isolated in 40% yield because of a competitive oxa-Pictet-Spengler acid-catalysed cyclocondensation of tryptophol with the solvent,[23] which consumed 70% of the available nucleophile.

Next, two competition experiments were performed to exploit the particular activity of HBF_4 in propargylation reactions (Scheme 6). Firstly, relatively electron-rich alcohol **1a** could be selectively reacted in the presence of **1g**, bearing a chloro substituent at the *para* position of the phenyl ring (Scheme 6, A). Also, the unexpected diminished reactivity of allylsilanes as nucleophiles was exploited when **1a** was reacted with 2 equivalents of benzyl alcohol and 2 equivalents of allyltrimethylsilane. Only **2b**, derived from the reaction with benzyl alcohol was formed under these conditions. Importantly, these chemoselectivities are not possible when using other Brønsted acids reported for this transformation such as *p*-toluenesulfonic acid,[6a] or phosphomolybdic acid.[8a]

Finally, a gram scale reaction was performed to further demonstrate the applicability of this reaction, and compound **5r** was isolated in high yield when using our optimised conditions (Scheme 7).



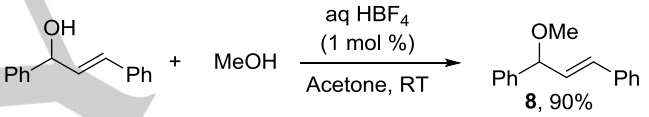
Scheme 6. Competition experiments with HBF_4. ^1H NMR conversions are provided.



Scheme 7. Gram scale reaction

Conclusions

The scope and limitations of HBF_4 as a practical catalyst for propargylation reactions have been explored. In general, good to excellent yields for the formation of C–O, C–N and C–C bonds were obtained under exceptionally simple reaction conditions. Challenging substrates such as electron-poor propargylic alcohols, or acid-sensitive indoles could be used with this methodology, even if slightly more forcing conditions were sometimes required. All reactions were carried out in air and in technical solvents, and the acid used was a commercially available aqueous solution. All the reactions were also completely regioselective, and no allene products were observed in any cases. Furthermore, many of the reactions were extremely clean and the desired products could be isolated analytically pure without the need for further purification after a simple aqueous work-up (*i.e.* **2a**, **2j**, **2l**-**o**, **5b**-**c**). Overall, this is a convenient and powerful methodology that does not employ a costly metal catalyst. The full potential of HBF_4 in this context remains to be uncovered. For instance, we were pleased to see that an allylic alcohol also reacted with MeOH at room temperature in very high yields (Scheme 8). Further applications of this synthetic protocol are currently being investigated in our laboratory.



Scheme 8. Reaction of an allylic alcohol with HBF_4.

Experimental Section

General procedure for the nucleophilic substitution of propargylic alcohols: In a vial fitted with a screw cap and a stirring bar, the propargylic alcohol **1** (1 mmol), nucleophile (2 mmol) and technical acetone (2 mL) were introduced. An aqueous solution 48 wt% in HBF_4 (1.2 µL,1 mol %) was then added and the reaction mixture was stirred at room temperature for 18 h. The resulting solution was quenched with a saturated aqueous solution of $NaHCO_3$ and extracted with ethyl acetate. The combined organic extracts were washed with brine, dried over anhydrous $MgSO_4$, filtered, and concentrated under reduced pressure to give the title compound. If needed, the crude product was then purified by column chromatography.

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Keywords: Propargylic Alcohols • Brønsted acid • Nucleophilic Substitution • Friedel-Crafts reaction • Homogeneous Catalysis

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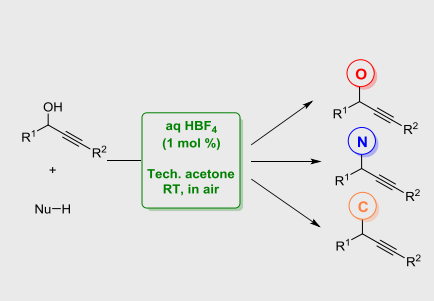
Entry for the Table of Contents

FULL PAPER

The activity of HBF_4 (aq solution) as a catalyst in propargylation reactions is presented. C–O, C–N and C–C bonds were formed in technical acetone, and in air. Good to excellent yields were obtained using low acid loading (typically 1 mol %) under mild reaction conditions.



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