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Brønsted acid-catalysed desilylative heterocyclisation to form substituted furans†

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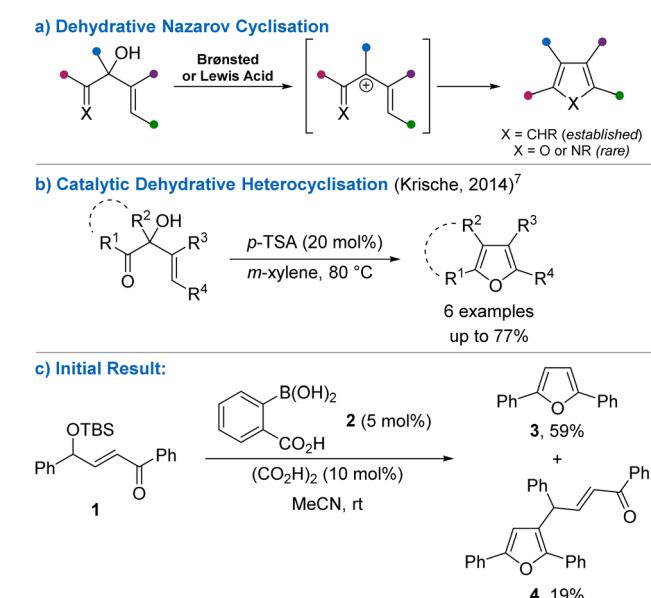
Heterocyclisation of *tert*-butyldimethylsilyl (TBS) protected γ -hydroxy- α,β -unsaturated ketones catalysed by *para*-toluenesulfonic acid (*p*-TSA) to form substituted furans is reported. The reaction proceeds under mild conditions at room temperature in methanol to give a range of furan products (21 examples, up to 98% yield). Mechanistic experiments suggest the reaction proceeds *via* *in situ* deprotection followed by catalytic dehydrative heterocyclisation.

Introduction

Small heterocyclic ring systems are ubiquitous molecular scaffolds within synthetic chemistry. For example, substituted furans are found within natural products and synthetic biologically active compounds, and they also have many uses as reaction substrates and/or intermediates. Consequently, various protocols have been developed for their preparation from a wide array of starting materials.¹ Advances in the practical synthesis of heterocycles, including substituted furans, under mild and sustainable conditions from readily available substrates is essential for continued development in a range of fields, including the pharmaceutical and agrochemical industries.²

The Nazarov cyclisation is an established strategy for the synthesis of cyclopentenones from divinyl ketones. Activation with either a Lewis or Brønsted acid forms an intermediate pentadienyl cation, which undergoes 4π -electrocyclisation to construct the cyclopentyl ring. Modern variations of this process have found different ways of accessing the key pentadienyl cation intermediate,³ including through dehydration of divinyl alcohols to form substituted cyclopentadienes (Scheme 1a).⁴ However, this strategy has rarely been extended to the synthesis of five-membered heterocycles. Würthwein and co-workers reported that α -hydroxy- β,γ -unsaturated oximes undergo dehydration in the presence of stoichiometric triflic acid to form 1-azapentadienyl cations, with subsequent electrocyclisation giving trisubstituted pyrroles.⁵ Das and co-workers found that highly functionalised Baylis–Hillman adducts also undergo dehydrative heterocyclisation promoted

by stoichiometric methanesulfonic acid to form functionalised benzofuran derivatives.⁶ Subsequently, Krische and co-workers showed that β,γ -unsaturated α -hydroxyketones, formed from ruthenium-catalysed hydrohydroxyalkylation of alkynes with diols, undergo dehydrative heterocyclisation in the presence of catalytic *para*-toluenesulfonic acid (*p*-TSA) to form a range of tetrasubstituted furans (Scheme 1b).⁷ There are also a few sporadic reports of isomeric γ -hydroxy- α,β -unsaturated ketones and their derivatives undergoing Brønsted acid-promoted dehydrative cyclisation.⁸ Similarly, γ -hydroxy- α,β -unsaturated ketones formed through *in situ* Knoevenagel condensation of 1,3-diketones and α -hydroxyaldehydes undergo facile cyclisation and elimination to form highly substituted furans, a



Scheme 1 Catalytic heterocyclisation reactions.

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process that is used in the valorisation of aldose sugars (known as the Garcia-Gonzalez reaction).⁹

Building upon our interest in the dehydrative substitution of alcohols using arylboronic acids as Brønsted acid catalysts,^{10,11} we were interested in investigating such systems as mild, catalytic methods of generating 1-oxapentadienyl cations that would undergo 4π -electrocyclisation to give substituted furans. To investigate this, *tert*-butyldimethylsilyl (TBS) protected γ -hydroxy- α,β -unsaturated ketone **1** was prepared as a precursor to the desired alcohol substrate. However, silyl ether **1** did not undergo smooth deprotection using tetrabutylammonium fluoride (TBAF), and resulted in either unexpected oxidation of the enone when performed under air, or deprotection followed by tautomerisation into the corresponding 1,4-diketone under an inert atmosphere.¹² As TBS ethers are also known to undergo acidic deprotection, it was reasoned that a one-pot deprotection-heterocyclisation process may be possible to form the furan directly from **1**. For example, Sammond and Sammakia reported that TBS protected γ -hydroxy- α,β -unsaturated ketones similar to **1** undergo deprotection/cyclisation in the presence of stoichiometric triethylamine trihydrofluoride.^{13a} A furan side product was observed in the synthesis of sacrolide A when a more complex unsaturated system was reacted with HF pyridine complex,^{13b} while single examples of similar furan formations have been reported using *p*-TSA^{13c} and H_2SO_4 .^{13d} To investigate whether this process could be rendered catalytic under mild reaction conditions, **1** was treated with 2-carboxybenzeneboronic acid **2** (5 mol%) and oxalic acid (10 mol%) in acetonitrile at room temperature (Scheme 1c). Pleasingly, furan **3** was obtained in 59% yield, alongside a side product that was isolated and characterised as substituted furan **4**. Arylboronic acids are known to catalyse dehydrative Friedel–Crafts alkylation reactions of electron-rich (hetero)arenes, accounting for the formation of **4**.¹⁴ Given the promising initial result for the one-pot deprotection-heterocyclisation, we decided to optimise this protocol with the aim of developing milder conditions compared to the previous reports and fully exploring the reaction scope and limitations.

Results and discussion

First, the reaction of **1** was further optimised through variation of key reaction parameters (Table 1).¹² An initial solvent screen revealed that the reaction also occurred in methanol forming furan **3** in a modest 17% yield by NMR analysis but, importantly, in the absence of side product **4** and the remainder of the material was unreacted **1** (entry 1). Next, a range of alternative Brønsted acid catalysts (5 mol%) were screened in methanol at room temperature. Acetic acid, oxalic acid, and trifluoroacetic acid (entry 2) showed little reactivity but strongly acidic methanolic HCl gave a promising 45% conversion to furan **3** with no formation of side product **4** (entry 3). The use of *para*-toluenesulfonic acid (*p*-TSA·H₂O) resulted in a further improvement, providing a clean 63% conversion into furan **3**

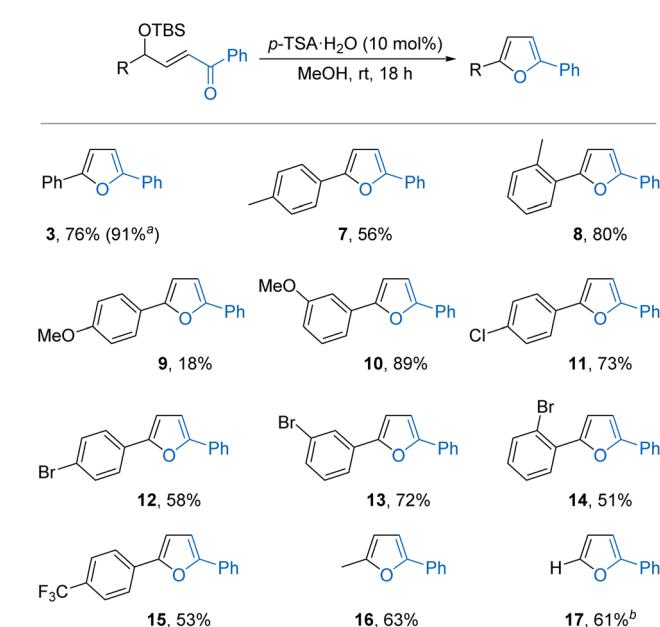
Table 1 Reaction optimisation

Entry	SiR ₃	Catalyst (mol%)	Yield ^a (%)
1	SiMe ₂ <i>t</i> -Bu 1	2 (5) + (CO ₂ H) ₂ (10)	17
2	SiMe ₂ <i>t</i> -Bu 1	TFA (5)	<5
3	SiMe ₂ <i>t</i> -Bu 1	HCl (5)	45
4	SiMe ₂ <i>t</i> -Bu 1	<i>p</i> -TSA·H ₂ O (5)	63
5 ^b	SiMe ₂ <i>t</i> -Bu 1	<i>p</i> -TSA·H ₂ O (10)	99 (76) ^c
6	Si(<i>i</i> -Pr) ₃ 5	<i>p</i> -TSA·H ₂ O (5)	50
7	SiPh ₂ <i>t</i> -Bu 6	<i>p</i> -TSA·H ₂ O (5)	10
8	SiMe ₂ <i>t</i> -Bu 1	None	0

^a Determined by ¹H NMR using 1,4-dinitrobenzene as an internal standard. ^b 18 hours. ^c Isolated yield.

(entry 4). Increasing the catalyst loading to 10 mol% and the time to 18 hours gave complete conversion of **1**, allowing furan **3** to be isolated in 76% yield (entry 5). The use of substrates bearing bulkier silyl protecting groups such as triisopropylsilyl (TIPS) **5** and *tert*-butyldiphenylsilyl (TBDPS) **6** gave reduced yields (entries 6 and 7),¹⁵ while in the absence of a Brønsted acid catalyst no conversion of starting material **1** was observed (entry 8). The scope and limitations of the catalytic heterocyclisation were then explored under the optimised conditions (Scheme 2).

The synthetic potential was initially demonstrated by performing the reaction on gram scale (5.7 mmol of **1**) to give 1.13 g of furan **3** in 91% yield. Next, a series of TBS protected γ -hydroxy- α,β -unsaturated ketones was conveniently prepared

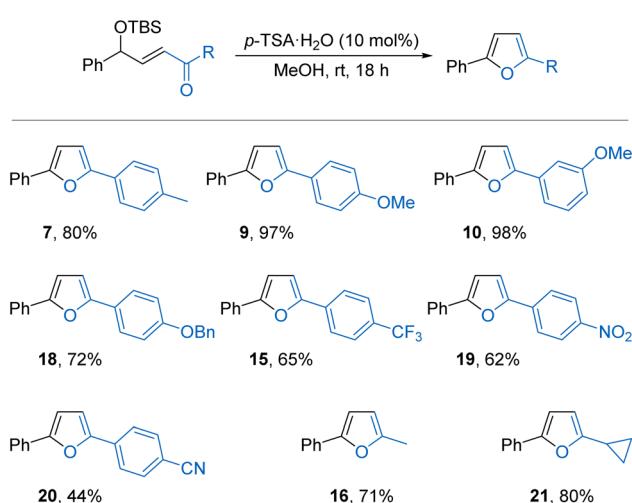


Scheme 2 Scope and limitations. Reactions performed on a 0.2 mmol scale in MeOH (1 M). ^a Performed on 5.7 mmol scale. ^b Performed on a 0.4 mmol scale in MeOH (0.1 M).



through a Wittig reaction between the corresponding TBS protected α -hydroxy aldehyde and an acyl phosphorane.^{12,16} First, various aryl substituents on the aldehyde component were trialled. Mildly electron-donating 2- and 4-methyl substituents were tolerated, forming products **7** and **8** in good yields. However, a strongly electron-donating 4-methoxy group was highly reactive, giving only 18% of furan **9** alongside a complex mixture of side-products. It may be that the increased stability of a possible cationic intermediate formed in this case facilitates unwanted reactivity, including solvolysis and Friedel-Crafts alkylations.^{10a,14} Moving the methoxy group to the 3-position where it is formally electron-withdrawing to the benzylic position was more successful,¹⁷ forming furan **10** in 89% yield. Aryl rings bearing halogen substituents in either the 2-, 3-, or 4-positions were well tolerated, forming products **11–14** in good yields. A strongly electron-withdrawing 4-trifluoromethyl group was also tolerated to give **15** in 53% yield. Alkyl substitution on this portion of the starting material was also possible, with 2-methyl-5-phenylfuran **16** formed in a good 63% yield. Singly substituted 2-phenylfuran **17** underwent side reactions at the standard 1 M substrate concentration in methanol but could be effectively isolated in 61% yield from a more dilute 0.1 M reaction.

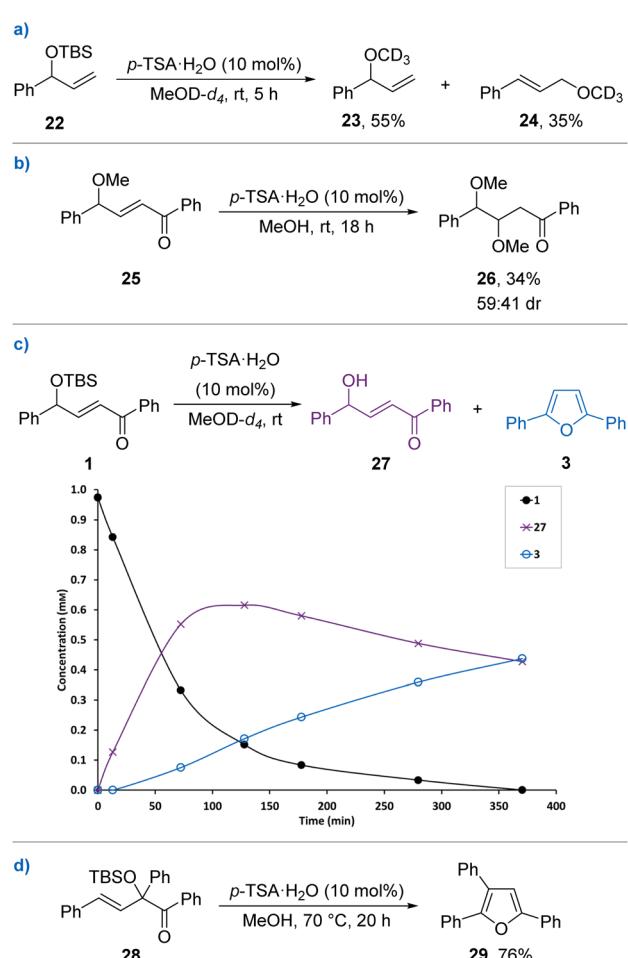
The acyl phosphorane component was then varied using 2-silyloxy-2-phenylacetaldehyde as the standard aldehyde (Scheme 3). In contrast to aryl substitution in the γ -position, electron-rich aryl rings were much better tolerated in the α' -position with 4-methyl and 4-methoxy substituted furans **7** and **9** isolated in 80% and 97% yield, respectively. The use of 3-methoxy and 4-benzyloxy substituted rings was also successful, giving products **10** and **18** in excellent yields. In these cases, the electron-donating aryl substituents are not conjugated with the allylic cation intermediate and therefore have less of an influence on the outcome of the cyclisation compared with substitution in the γ -position. Strongly electron-withdrawing aryl substituents including 4-trifluoromethyl,



Scheme 3 Scope and limitations. Reactions performed on a 0.2 mmol scale in MeOH (1 M).

4-nitro, and 4-cyano, were also better tolerated in the α' -position compared with the γ -position, with furans **15**, **19**, and **20** formed in reasonable yields. Alkyl substitution was again possible, with 2-methyl and 2-cyclopropyl substituted furans **16** and **21** isolated in 71% and 80% yields, respectively. Attempts to form tri-substituted furans through α -substitution of the enone were unsuccessful as the substrates could not be prepared.

Next, a series of control experiments were performed to gain further mechanistic understanding of the cyclisation process (Scheme 4). The likely silyl deprotection of the starting material was initially studied using model TBS protected allylic alcohol **22** that is incapable of undergoing heterocyclisation. Reacting **22** under the standard conditions in MeOD-*d*₄ and monitoring over time by ¹H NMR using 1,4-dinitrobenzene as an internal standard showed rapid silyl deprotection into the corresponding allylic alcohol, which underwent solvolysis over time to form a mixture of branched and linear methyl ethers **23** and **24** (Scheme 4a). To investigate the possibility of solvolysis occurring in the heterocyclisation process, methyl ether **25** was used as a substrate (Scheme 4b). No furan formation occurred after 18 hours; however, a small amount of conjugate addition product **26** was observed as a 59 : 41 mixture of dia-



Scheme 4 Control experiments.

stereoisomers with the remainder unreacted starting material. This is surprising, as methyl ether **25** would likely have similar ionisation potential compared with the corresponding allylic alcohol yet is not a viable intermediate, but this experiment does show that heterocyclisation outcompetes solvolysis under the reaction conditions. However, the analogous methyl ether bearing a γ -4-methoxyphenyl substituent did form the expected furan **9** under the standard conditions, alongside the conjugate addition products.¹² These results therefore suggest that different reaction mechanisms may be in operation depending on the substituents present. The reaction of TBS γ -hydroxy-enone **1** was then studied by *in situ* ^1H NMR in MeOD-*d*₄ (Scheme 4c). As with model substrate **22**, TBS protected **1** was rapidly consumed, with deprotected alcohol **27** initially building-up in solution before its concentration decreased in line with formation of furan **3**. No other significant reaction intermediates were observed, while further time points could not be taken as furan **3** precipitates from solution at higher concentrations. Isolated γ -hydroxy-enone **27** was demonstrated to be a competent starting material, with furan **3** formed in 76% yield under the standard conditions.¹¹ The possibility of *in situ* isomerisation of alcohol **27** into the corresponding 1,4-diketone followed by Paal-Knorr furan formation was discounted, as the isolated 1,4-diketone does not react with *p*-TSA (10 mol%) in methanol. The reaction of **1** was also successful in the dark, forming furan **3** in 71% yield. This excludes possible photochemical alkene isomerisation and cyclisation as recently reported by Donohoe and co-workers for a set of related γ -hydroxy- α,β -unsaturated ketone substrates.¹⁸ Isomeric TBS α -hydroxy- β,γ -unsaturated ketone **28** was also trialled in the heterocyclisation reaction. The more sterically demanding tertiary substrate did not cyclise at room temperature, with starting material **28** returned alongside 15% of the γ -solvolysis product.¹² However, heating the reaction to 70 °C gave trisubstituted furan **29** in 76% yield (Scheme 4d). This provides support for the formation of an allylic cation inter-

mediate that can be accessed irrespective of the isomer of allylic alcohol used.

The available evidence cannot unambiguously determine the mechanism for heterocyclisation and suggests that more than one pathway may occur depending on the substrate (Scheme 5). The *in situ* NMR shows that acid-promoted deprotection of TBS γ -hydroxy- α,β -unsaturated ketone **30** is facile and forms alcohol **31**. The alcohol may then be protonated and ionised to form allylic cation (*E*)-**32**, which must be isomerised into (*Z*)-**32** before undergoing rapid 4 π -electrocyclisation followed by deprotonation to give the furan product **33**.¹⁹ Alternatively, alcohol **31** may undergo acid-catalysed conjugate addition of methanol to form **34**, which can cyclise to form hemiacetal **35**.²⁰ Successive acid-promoted eliminations of water and methanol would lead to furan **33**.

Conclusions

In conclusion, *p*-TSA (10 mol%) is an efficient catalyst for the heterocyclisation of TBS protected γ -hydroxy- α,β -unsaturated ketones to form substituted furans under mild conditions in methanol at room temperature. The process is thought to occur *via* *in situ* deprotection, followed by acid-promoted heterocyclisation with elimination of one equivalent of water. Further work on the catalytic activation of allylic alcohols for heterocycle formation is underway in our laboratory.²¹

Experimental

General procedure for catalytic heterocyclisation

The requisite TBS-protected γ -hydroxy- α,β -unsaturated ketone was dissolved in MeOH (1 M) and *p*-TSA·H₂O (10 mol%) was added. The reaction was stirred at rt for 18 h before being concentrated under reduced pressure. The crude material was purified by silica gel column chromatography (100% to 97/3 cyclohexane/Et₂O).

Author contributions

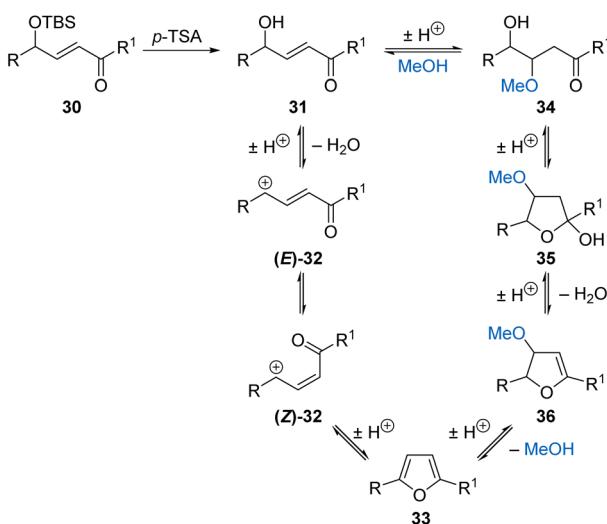
E. G. B. and M. S. R. performed all experimental work and collected the data. J. E. T. conceptualised the idea, supervised, and coordinated the project. E. G. B. and J. E. T. prepared the manuscript and ESI.†

Conflicts of interest

There are no conflicts to declare.

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Scheme 5 Plausible mechanistic pathways.



funding. We thank a reviewer for insightful comments regarding the alternative reaction mechanisms.

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