

Enantioselective Acylation of Silyl Ketene Acetals through Fluoride Anion-Binding Catalysis

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Supporting Information

ABSTRACT: A highly enantioselective acylation of silyl ketene acetals with acyl fluorides has been developed to generate useful α , α -disubstituted butyrolactone products. This transformation is promoted by a new thiourea catalyst and 4-pyrrolidinopyridine and represents the first example of enantioselective thiourea anion-binding catalysis with fluoride.

Pyridine derivatives induce significant rate accelerations in acyl transfer reactions through the generation of electrophilic *N*-acylpyridinium ion intermediates.¹ While pyridine itself is a competent nucleophilic catalyst, analogues bearing strongly electron-donating substituents such as 4-(dimethylamino)pyridine $(DMAP)^2$ and 4-pyrrolidinopyridine $(PPY)^3$ are up to 4 orders of magnitude more reactive in representative acyl transfer reactions.⁴ The amplified reactivity of aminopyridine derivatives is ascribable both to a greater equilibrium concentration of the acylpyridinium ion intermediate and to increased electrophilicity of that intermediate as a result of looser ion pairing.⁵ Both effects can be enhanced, in principle, through stabilization of the counteranion by a specific hydrogen-bond donor (Scheme 1).⁶ We have shown that chiral urea and thiourea derivatives can catalyze enantioselective reactions via mechanisms involving anion binding,⁷ and Seidel has demonstrated recently the successful application of this concept in the context of chiral thiourea-DMAP cocatalyzed acylative kinetic resolutions of primary amines.⁸ This strategy is fundamentally different from classical approaches to catalytic asymmetric acyl transfer reactions that rely on construction of chiral nucleophilic catalysts.9

We sought to apply this reactivity principle to the *C*-acylation of enolate equivalents, a transformation that affords versatile β -dicarbonyl derivatives with generation of a quaternary stereocenter. We report here the highly enantioselective acylation of silyl ketene acetals with acyl fluorides promoted by the new thiourea catalyst 3d and PPY.¹⁰ This method provides a preparatively useful route to α, α -disubstituted butyrolactones¹¹ and represents the first example of enantioselective thiourea anion-binding catalysis with fluoride.¹²

Benzoylation of silyl ketene acetal 1 was selected as a model reaction. A lead result afforded lactone 2a in 19% yield and 41% ee by acylation of 1 with benzoic anhydride in the presence of phenylpyrrolidine-derived thiourea 3a and DMAP (Table 1, entry 1). Slightly improved enantioselectivity and yield were obtained by replacing DMAP with PPY (entry 2). However, a more striking dependence on the identity of the acylating agent

Scheme 1. Anion-Binding/Nucleophilic Cocatalysis of Acyl Transfer

Mechanism of catalysis of acyl transfer by pyridine derivatives:



was observed, with benzoyl chloride being completely unreactive (entry 3) and benzoyl fluoride affording **2a** in 84% yield and 81% ee (entry 4). Accordingly, chiral catalyst optimization studies were carried out on the PPY-catalyzed acylation of **1** with benzoyl fluoride.

Catalysts with the general structure of 3 have been demonstrated recently to be broadly effective in a wide range of enantioselective transformations⁷ and also proved optimal in the present study. Notably, Schreiner's thiourea 13 (4) and Seidel's optimal catalyst for the acylative kinetic resolution of primary amines^{8b,c} (5) were considerably less reactive than 3. The right-hand tert-leucinethiourea-3,5-bis(trifluoromethyl)anilide fragment of 3 is a common feature in most of these systems, and it was found to be strictly superior to other amino acid-H-bond donor-anilide combinations in the model acylation reaction. Enantioselectivity also proved strongly responsive to the structure of the left-hand 2-arylpyrrolidino fragment. Systematic variation of the aryl group of the pyrrolidine led to the identification of 3d as the most enantioselective catalyst. Although elucidation of the mechanism of stereoinduction must await detailed mechanistic investigation, the important influence of the arylpyrrolidino component of the thiourea catalyst on both reaction rate and enantioselectivity¹⁴ is suggestive of differential stabilizing noncovalent interactions with the cationic electrophile in the transition states leading to the major and minor enantiomers of product.¹⁵

Under the optimized reaction conditions, silyl ketene acetal 1 underwent acylation with a variety of substituted benzoyl fluorides (Table 2).¹⁶ Substrates containing both electron-donating

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Table 1. Catalyst and Reaction Optimization



		1	/ 00	/	
entry ^a	catalyst	catalyst	(X =)	(%)	(%)
1	3a	DMAP	OBz	19	41
2	3a	РРҮ	OBz	21	52
3	3a	PPY	Cl	0	_
4	3a	PPY	F	84	81
5	3b	PPY	F	88	87
6	3c	PPY	F	29	75
7	3d	PPY	F	80	92
8	4	PPY	F	9	—
9	5	PPY	F	20	<5
10^d	3d	PPY	F	86	93

^{*a*} Reactions run on a 0.08 mmol scale. ^{*b*} Yields determined by ¹H NMR analysis relative to *p*-xylene as an internal standard. ^{*c*} Enantiomeric excess determined by HPLC analysis on commercial chiral columns. ^{*d*} Reaction run using 5 mol % thiourea catalyst and 6 mol % PPY at -60 °C in TBME (0.01 M) for 8 h.

and electron-withdrawing groups, as well as 2-naphthoyl fluoride, provided acylation products in high yields and enantioselectivities. Substitution at the meta and para positions was well tolerated; however, substitution at the ortho position resulted in a complete loss of reactivity. This observation supports the notion of an acylpyridinium intermediate, as the rates of reactions that proceed through such species are known to be severely affected by ortho substitution on the acylating agent.¹⁷ A number of silvl ketene acetals were also viable substrates, with derivatives bearing electron-rich and electron-poor arene substituents, as well as heteroaromatic functionality, participating in acylation with benzoyl fluoride in good yields and enantioselectivities.¹⁸ The efficiency of this protocol is illustrated in the acylation of 1 with 2-naphthoyl fluoride on a preparative scale using only 0.5 mol % thiourea catalyst (Scheme 2). The product of this reaction was purified by crystallization from the crude reaction mixture without the need for chromatography. A single recrystallization provided pure acylation product 2b in 70% overall yield and >99% ee.



entry ^a	Product	time	yield ^b (%)	ee^{c} (%)
1		6 h	88	92
2^d		6 h	85	95
3		24 h	91	95
4		6 h	90	91
5		4 h	95	86
6		16 h	77	92
7		6 h	95	92
8		6 h	78	87
9		6 h	75	93
10		6 h	80	88
11		24 h	76	89
12		6 h	76	89
13		6 h	77	89

^{*a*} Reactions run on a 0.20 mmol scale. ^{*b*} Isolated yield of purified product. ^{*c*} Enantiomeric excess determined by HPLC analysis on commercial chiral columns. ^{*d*} The structure and absolute configuration of **2b** was established by X-ray crystallography, and the stereochemistry of all other products was assigned by analogy.

No reaction between silyl ketene acetal 1 and benzoyl fluoride is observed in the absence of either thiourea catalyst or nucleophilic catalyst. This observation indicates the thiourea is playing a role not only in the enantioselectivity-determining acylation event but also in the generation of the key acylpyridinium ion intermediate. It is likely that the outstanding hydrogen-bondaccepting ability of the fluoride anion is important in this

Scheme 2. Preparative-Scale Reaction



Scheme 3. Proposed Catalytic Cycle



regard.¹⁹ Furthermore, activation of the silyl ketene acetal by fluoride or benzoate seems necessary for acylation to occur, particularly given the complete absence of reactivity observed with benzoyl chloride under the standard reaction conditions.²⁰ Consistent with this hypothesis, variation of the silyl group of the silyl ketene acetal has a measurable influence on the rate of the reaction, with larger silyl groups leading to diminished reaction rates. However, the identity of the silyl group has a negligible effect on reaction enantioselectivity, indicating that it plays no significant role in the organization of the stereoselectivitydetermining step.²¹ This result raises the possibility of a thioureabound enolate as a key intermediate.²²

A proposed catalytic cycle that is consistent with these observations is presented in Scheme 3. As noted, the thiourea catalyst activates benzoyl fluoride for reaction with PPY, presumably via initial complexation to the carbonyl group of the acyl fluoride (A). A thiourea-bound acylpyridinium/fluoride intermediate (B) is then proposed, in which the thiourea is associated to the fluoride anion and the catalyst arene substituent is engaged in a stabilizing interaction with the acylpyridinium cation.¹⁵ Reaction of **B** with the silyl ketene acetal likely proceeds via a pentavalent silicate intermediate²³ and is proposed to be rate-determining on the basis of the observed dependence of the overall rate on the identity of the silyl group. However, the independence of reaction enantioselectivity on the identity of the silyl group points to a thioureabound enolate such as C as the intermediate involved in enantiodetermining acylation.

In conclusion, a highly enantioselective acylation of silyl ketene acetals with acyl fluorides has been developed to generate useful α, α -disubstituted butyrolactone products. The remarkable hydrogen-bond acceptor properties and silaphilicity of the fluoride anion facilitate an efficient reaction protocol with low catalyst loadings and high yields and selectivities. A more complete mechanistic elucidation of this acylation reaction that includes analysis of the basis for enantioinduction is the focus of ongoing studies.

ASSOCIATED CONTENT

Supporting Information. Complete experimental procedures and characterization data for acylation products and all isolated intermediates, ¹H and ¹³C NMR spectra of acylation products, HPLC traces of racemic and enantioenriched acylation products, catalyst and silyl ketene acetal optimization data, and crystallographic information for compound **2b**. This material is available free of charge via the Internet at http://pubs.acs.org.

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