

UC Berkeley

UC Berkeley Previously Published Works

Title

α -Phenylthioaldehydes for the effective generation of acyl azolium and azolium enolate intermediates.

Permalink

<https://escholarship.org/uc/item/5kr078sr>

Journal

Chemical Science, 15(24)

ISSN

2041-6520

Authors

Ewing, Paul

Majhi, Pankaj

Prentice, Callum

et al.

Publication Date

2024-06-19

DOI

10.1039/d3sc06879j

Peer reviewed

EDGE ARTICLE

Cite this: *Chem. Sci.*, 2024, 15, 9369

All publication charges for this article have been paid for by the Royal Society of Chemistry

α -Phenylthioaldehydes for the effective generation of acyl azolium and azolium enolate intermediates†

Paul M. D. A. Ewing,^{ab} Pankaj Kumar Majhi,^a Callum Prentice,^a Claire M. Young,^a Karlotta van Rees,^b Polly L. Arnold,^b Eli Zysman-Colman^b*^{cd} and Andrew D. Smith^b*^{ea}

α -Phenylthioaldehydes are readily prepared using a simple multi-step procedure and herein are introduced as a new precursor for the NHC-catalysed generation of acyl azolium and azolium enolate intermediates that are of widespread synthetic interest and utility. Treatment of α -phenylthioaldehydes with an NHC precatalyst and base produces an efficient redox rearrangement *via* a Breslow intermediate, elimination of thiophenolate, and subsequent rebound addition to the generated acyl azolium to give the corresponding thiol ester. In the presence of an external alcohol, competition between redox rearrangement and redox esterification can be controlled through judicious choice of the *N*-aryl substituent within the NHC precatalyst and the base used in the reaction. With NEt_3 as base, NHCs bearing electron-withdrawing (*N*- C_6F_5 or *N*- $\text{C}_6\text{H}_2\text{Cl}_3$) substituents favour redox rearrangement, while triazolium precatalysts with electron-rich *N*-aryl substituents (*N*-Ph, *N*-Mes) result in preferential redox esterification. Using DBU, redox esterification is preferred due to transesterification of the initially formed thiol ester product. Additionally, α -phenylthioaldehyde-derived azolium enolates have been used in enantioselective formal [4 + 2]-cycloaddition reactions to access dihydropyridinone heterocycles with high enantioselectivity (up to >95 : 5 dr, 98 : 2 er).

Received 21st December 2023

Accepted 9th May 2024

DOI: 10.1039/d3sc06879j

rsc.li/chemical-science

1. Introduction

Acyl azolium and azolium enolate intermediates have been harnessed as key species within a range of enantioselective reactions using *N*-heterocyclic carbenes (NHCs).¹ For example, a variety of effective kinetic resolution and dynamic kinetic resolution processes have been developed that use chiral NHCs as catalysts and exploit *in situ* acyl azolium generation.¹ Similarly, azolium enolate intermediates have been widely utilised to access important heterocyclic scaffolds *via* formal [2 + 2], [3 + 2], [4 + 2] and higher cycloadditions in an enantioselective manner.¹ A variety of strategies have been developed to access these reactive species, with an overview of the current methods, as well as their interconnectivity, given in Fig. 1. Acyl azolium intermediates I can be generated directly from the nucleophilic addition–elimination reaction between an NHC and

a carboxylic acid derivative such as an ester, anhydride or acid chloride.^{2,3} Alternatively, they can be prepared from the stoichiometric oxidation of an *in situ* generated Breslow intermediate from an aldehyde using so called “oxidative” NHC

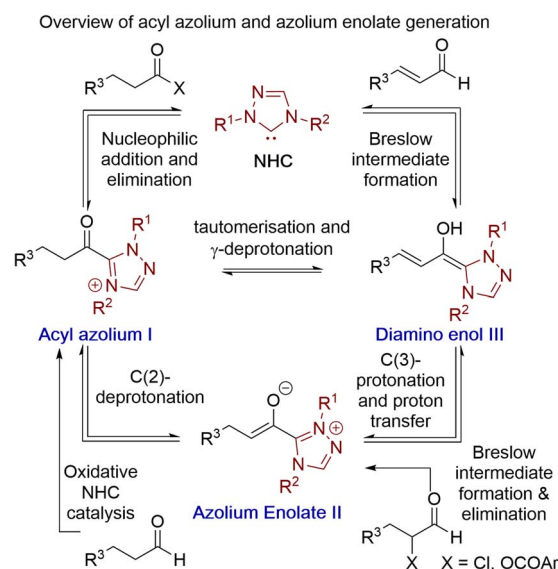


Fig. 1 Established access to acyl azolium I and azolium enolate II intermediates.

^aEastCHEM, School of Chemistry, University of St Andrews, St Andrews, Fife, KY16 9ST, UK. E-mail: ads10@st-andrews.ac.uk; eli.zysman-colman@st-andrews.ac.uk

^bEastCHEM, School of Chemistry, University of Edinburgh, EH9 3JF, UK

^cDept of Chemistry, University of California, Berkeley, CA, 94720, USA. E-mail: pla@berkeley.edu

^dChemical Sciences Division, Lawrence Berkeley National Laboratory, Berkeley, CA 94720, USA

† Electronic supplementary information (ESI) available: Full characterisation and HPLC spectra. CCDC 2310270. For ESI and crystallographic data in CIF or other electronic format see DOI: <https://doi.org/10.1039/d3sc06879j>



catalysis, with a range of either transition metal or organic oxidants compatible with this process.⁴ C(2)-deprotonation of an acyl azolium species **I** leads to the azolium enolate **II**, while in the reverse direction the selective protonation of an azolium enolate **II** leads to the generation of an acyl azolium species **I**.⁵ First demonstrated independently by Scheidt⁶ and Bode,⁷ the generation of acyl azolium species **I** from enals has also been developed. C(3)-protonation of an enal-derived Breslow intermediate diamino enol **III** and proton transfer furnishes the azolium enolate **II**, which undergoes tautomerisation to form the acyl azolium species **I**.^{8,9} An alternative and direct method to generate the azolium enolate *via* addition of an NHC to a disubstituted ketene was first demonstrated by Ye¹⁰ and Smith¹¹ through formal [2 + 2]-cycloaddition reactions. However, due to limited synthetic diversity of reactive ketene substrates alternative processes to allow the generation of azolium enolates have been developed. For example, Chi and co-workers showed that 4-nitrophenyl esters could be used as azolium enolate precursors,¹² while the most common process uses α -functionalised aldehydes as precursors.¹³ In this area, elimination of an α -leaving group from an *in situ* generated Breslow intermediate provides the desired enolate. Despite being prone to decomposition, the most common starting materials are α -chloroaldehydes, that have been used to access enantioenriched heterocycles.^{14,15} Furthermore, while α -alkyl- α -chloro-substituted aldehydes are generally tolerated, only limited precedent for the use of α -unsubstituted or α -aryl substituted precursors have been developed.¹⁶ In previous work, we introduced bench stable α -aroyloxyaldehydes as alternative acyl azolium and azolium enolate precursors.^{13a} They have since been applied in [2 + 2], [3 + 2] and [4 + 2] cycloadditions,¹⁷ redox α -aminations,¹⁸ as well as kinetic resolutions and desymmetrisations,¹⁹ but again α -unsubstituted or α -aryl substituted derivatives were not tolerated.^{13a} Building upon these prior demonstrations, in this manuscript the synthesis and reactivity of α -phenylthioaldehydes as acyl azolium and azolium enolate intermediate precursors is described (Fig. 2). Notably, in this work thiophenolate eliminated during azolium enolate formation is harnessed as an *in situ* nucleophile to turnover the NHC catalyst from an acyl azolium intermediate, leading to thiol ester products resulting from redox rearrangement. This process is tolerant of both α -unsubstituted and α -aryl substituted derivatives. In the

presence of an external alcohol, competition between redox rearrangement and redox esterification can be controlled through judicious choice of the *N*-aryl substituents within the NHC precatalyst and the base used in the reaction. In addition, α -phenylthioaldehyde-derived azolium enolates have been used in enantioselective formal [4 + 2]-cycloaddition reactions to access dihydropyridinone heterocyclic products with high enantioselectivity (up to >95 : 5 dr, 98 : 2 er).

2. Results and discussion

2.1 Synthesis of α -phenylthioaldehydes

Initial studies commenced by demonstrating a viable route to a model starting material. α -Phenylthioaldehyde **3** bearing an α -phenyl substituent was synthesised by modification of a two-step procedure first reported by Nozaki and co-workers (Scheme 1).²⁰ Starting from commercially available methoxymethyl(phenylsulfide) **1**, deprotonation and subsequent aldol reaction gave **2** in 85% yield (85 : 15 dr), with mesylation promoting rearrangement to give **3** in 85% yield. Modification of the reported procedure for the second step, including the use of dichloromethane as solvent and dry loading on silica, resulted in reproducible product yields and simple purification (see ESI† for details). Using this general approach, a range of differently substituted α -phenylthioaldehydes **5–11** were prepared, incorporating α -alkyl as well as an α,α -disubstituted aldehyde. The unsubstituted aldehyde **4** was prepared by an alternative two-step procedure from the corresponding acid involving methyl ester formation and selective reduction with DIBAL (see ESI† for further details).

2.2 NHC-catalysed redox rearrangement

Studies subsequently investigated the proposed NHC-catalysed internal redox rearrangement of α -phenyl- α -phenylthioaldehyde **3** to give the corresponding thiol ester **12** as a model substrate for reaction development. Aldehyde **3** was treated with a variety of triazolium NHC precatalysts with different *N*-aryl substituents (**13–16**) and bases.²¹ In the presence of *N*-mesityl triazolium precatalyst **13** (10 mol%) and

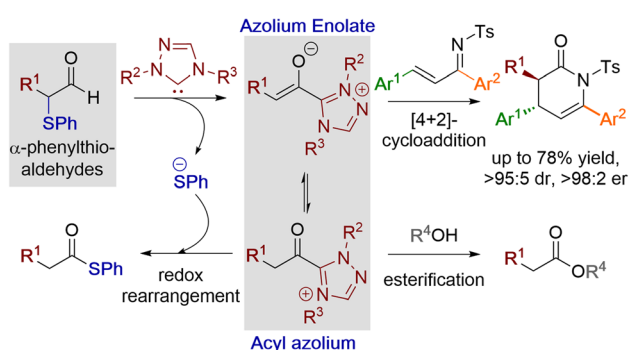
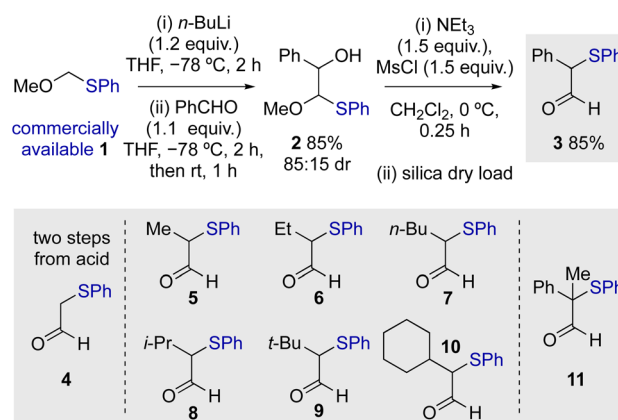


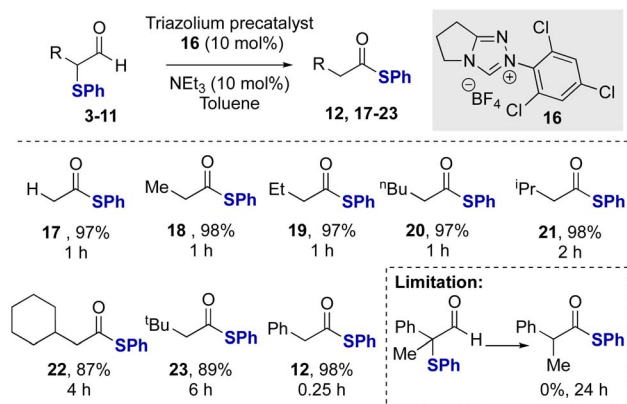
Fig. 2 α -Phenylthioaldehydes as acyl azolium and azolium enolate precursors.



Scheme 1 Synthesis of α -phenylthioaldehydes.

Cs_2CO_3 as a base, thiol ester **12** was formed in 97% yield in 16 hours (Table 1, entry 1). Lowering the catalyst loading to 5 mol% resulted in full conversion of the aldehyde but required an extended reaction time of 24 hours (Table 1, entry 2). Changing the base from Cs_2CO_3 to the organic bases NEt_3 or DBU gave comparable reaction yields (Table 1, entry 3 and 4). The effect of alteration of the *N*-aryl substituent within the precatalyst was next explored using NEt_3 as the base. *N*-Phenyl substituted NHC precatalyst **14** (10 mol%) gave 95% conversion to the thiol ester **12**, while both *N*-pentafluorophenyl precatalyst **15** and *N*-2,4,6-trichlorophenyl precatalyst **16** (10 mol%) gave high product yield with a significantly shorter reaction time of only 15 minutes (Table 1, entries 5 and 6). Using NHC precatalyst **16** at 5 mol% loading led to good product conversion within 2 hours (Table 1, entry 8). Control reactions in the absence of NHC precatalyst gave no rearrangement product or degradation of the aldehyde (Table 1, entry 9).

After demonstrating the feasibility of this redox rearrangement process, its scope and limitations were established using *N*-2,4,6-trichlorophenyl precatalyst **16** and NEt_3 as base. The unsubstituted aldehyde **4** gave the corresponding thiol ester **17** in 97% isolated yield (Scheme 2). α -Phenylthioaldehydes bearing α -alkyl substituents were tolerated in this redox rearrangement although the rate of rearrangement, and thus required reaction time, is significantly affected by the steric bulk of the α -substituent. For example, using linear Me, Et and *n*-Bu α -substituents, the rearrangement was complete within 1



Scheme 2 Redox rearrangement of α -phenylthioaldehydes.^a All reactions were conducted in a flame-dried Schlenk flask under a nitrogen atmosphere using 0.166 mmol of aldehyde **3–11**, 10 mol% NEt_3 .

hour. The incorporation of branched α -substituents required longer reaction times for effective rearrangement, with α -*i*-Pr aldehyde **8** requiring two hours to give **21** at full conversion. Similarly, the α -cyclohexyl and α -*t*-Bu aldehydes **9** and **10** required 4 and 6 hours, respectively, for full conversion to the corresponding products **22** and **23**.²² These results contrast the efficient rearrangement of the α -aryl substituted derivative **3**, where full conversion to the product **12** was observed within just 15 minutes. This is hypothesized to reflect the ability of the conjugating α -phenyl substituent to stabilize the azolium enol/enolate intermediate within this process.²³ A limitation of this methodology showed that α -phenylthioaldehyde **11** bearing a tertiary centre was unsuccessful in this catalytic rearrangement, presumably due to steric hindrance inhibiting NHC addition to the aldehyde.

Table 1 Optimisation of NHC-catalysed redox rearrangement^a

Entry	Conditions	Yield ^{b,c} (%)	Time ^d (h)
1	13 , Cs_2CO_3	>95 (97)	16
2 ^e	13 , Cs_2CO_3	>95	24
3	13 , NEt_3	>95 (97)	16
4	13 , DBU	>95	16
5	14 , NEt_3	>95	16
6	15 , NEt_3	>95 (97)	0.25
7	16 , NEt_3	>95 (98)	0.25
8 ^e	16 , NEt_3	>95 (95)	2
9	NEt_3	0	24

^a All reactions conducted in a flame-dried Schlenk flask under a nitrogen atmosphere using aldehyde **3** (0.166 mmol), base (10 mol%), NHC precatalyst **13–16** (10 mol%). ^b Conversion was determined by ^1H NMR spectroscopy of the crude reaction mixtures. ^c Yields in parentheses correspond to isolated yields after chromatographic purification. ^d Reaction times refer to aliquots taken from the reaction mixture at various time points followed by ^1H NMR spectroscopic analysis. ^e 5 mol% catalyst was used.

2.3 NHC-catalysed redox esterification

Further investigation explored α -phenylthioaldehydes in NHC-catalysed redox esterification reactions using benzyl alcohol. Using α -phenyl- α -phenylthioaldehyde **3** as a model, at the onset the potential for mixtures of benzyl ester **24** and thiol ester **12** products, arising either directly from competition between nucleophiles, or from transesterification, was considered (Table 2). The use of NEt_3 as base was initially probed, with *N*-mesityl substituted triazolium precatalyst **13** leading exclusively to benzyl ester product **24** (entry 1). Using *N*-phenyl precatalyst **14** gave a 75 : 25 mixture of benzyl ester **24** : thiol ester **12** products, with benzyl ester **24** isolated in 70% yield (entry 2). The use of *N*-pentafluorophenyl precatalyst **15** gave a 50 : 50 ratio of benzyl ester **24** to thiol ester **12** (entry 3), while *N*-2,4,6-trichlorophenyl precatalyst **16** led to a 10 : 90 ratio of benzyl ester **24** to thiol ester **12** (entry 4). Further studies considered the effect of the base on product selectivity. Using DBU, *N*-mesityl precatalyst **13** again gave the benzyl ester **24** as the major product (entry 5). However, using *N*-2,4,6-trichlorophenyl triazolium precatalyst **16** and DBU, the benzyl ester product **24** was generated exclusively, indicating a switch in product

Table 2 A Redox esterification optimisation^a

Reaction scheme showing the redox esterification of aldehyde **3** (Ph-CH(SPh)-CHO) using NHC precatalysts **13-16** (10 mol%), base (1.5 equiv.), and BnOH (2.0 equiv.) in toluene to yield benzyl ester **24** and thiol ester **12**.

Chemical structures of NHC precatalysts **13**, **14**, **15**, and **16** are shown in a dashed box below the reaction scheme.

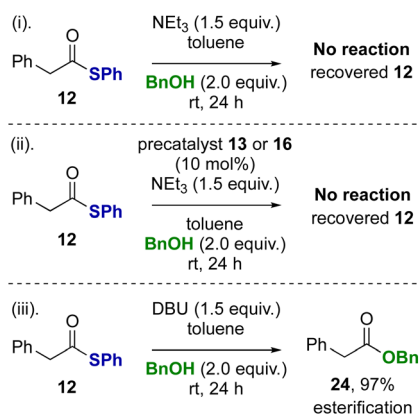
Entry ^a	Precat, base	Conv. ^b	24 ^{b,c} (%)	12 ^{b,c} (%)
1	13 , NEt ₃	>95	>95 (97)	<5
2	14 , NEt ₃	>95	76 (70)	23 (21)
3	15 , NEt ₃	>95	50 (46)	50 (47)
4	16 , NEt ₃	>95	10 (5)	90 (87)
5	13 , DBU	>95	>95 (95)	<5
6	16 , DBU	>95	>95 (98)	<5

^a All reactions conducted using aldehyde **3** (0.166 mmol), base (1.5 equiv.), NHC precatalyst **13-16** (10 mol%), BnOH (2 equiv.).

^b Conversion, as well as ratio of ester and thiol ester products, determined by ¹H NMR spectroscopic analysis of the crude reaction product. ^c Parentheses indicate isolated yields after chromatographic purification.

selectivity with base for precatalyst **16** (entry 6). Similarly, the use of benzylamine in this redox transformation using precatalyst **13** and DBU led to productive amide bond formation in 91% yield (see ESI† for further details).

Control reactions were subsequently performed to rationalise the product distributions observed (Scheme 3). Potential transesterification to account for the interconversion between thiol ester **12** and benzyl ester **24** during the reaction was tested (reactions i–iii). Treatment of thiol ester **12** with BnOH (2 equiv.) and NEt₃ (reaction i) or NEt₃ in the presence NHC precatalysts **13** or **16** (reaction ii) returned exclusively starting material, with no transesterification observed. However, treatment of thiol ester **12** with DBU and benzyl alcohol (2 equiv.) showed complete transesterification to benzyl ester **24** within 24 hours (reaction iii).



Scheme 3 Control studies: esterification of thiol ester.

To probe if these results were unique to α -phenyl- α -phenylthioaldehyde, α -methyl substituted substrate **5** was tested in the redox esterification reaction (Table 3). Using NHC precatalysts **13** or **16** and either NEt₃ or DBU, similar trends in product distributions were observed; using NEt₃ a reduced rate of product formation was observed, with *N*-mesityl precatalyst **13** favouring benzyl ester **25** (Table 3, entry 1) and *N*-2,4,6-trichlorophenyl precatalyst **16** favouring thiol ester **26** (Table 3, entry 2). The use of DBU promoted full conversion selectively to the benzyl ester **25** irrespective of NHC precatalyst (Table 3, entries 3 and 4). Changing from benzyl alcohol to methanol with DBU as the base led to selective formation of the methyl ester **27** (entries 5 and 6).

2.5 Interpretation of experimental results and mechanistic construct

These experimental results are consistent with the electronic nature of the *N*-aryl substituent within the NHC and the base as the main factors influencing selectivity in these processes. Firstly, with NEt₃, NHCs derived from triazolium precatalysts **15** and **16** bearing electron-withdrawing *N*-aryl substituents favour redox rearrangement, consistent with the observation of enhanced conversion rate to the thiol ester in comparison to triazolium precatalysts **13** and **14**. Switching to triazolium precatalysts with more electron-rich *N*-aryl substituents (triazoliums **13** and **14**), results in preferential redox esterification in the presence of BnOH. The choice of base also influences product selectivity as the presence of DBU overrides the intrinsic selectivity of the triazolium precatalyst due to

Table 3 Redox esterification: probing generality with alcohol and substrate^a

Reaction scheme showing the redox esterification of aldehyde **5** (Me-CH(SPh)-CHO) using NHC precatalysts **13** or **16** (10 mol%), base (1.5 equiv.), and ROH (2.0 equiv.) in toluene to yield benzyl ester **25** (R = Bn) and thiol ester **26** (R = Me).

Chemical structures of NHC precatalysts **13** and **16** are shown in a dashed box below the reaction scheme.

Entry ^a	R	Precat, base	Conv. ^b	Ester ^{b,c} (%)	26 ^{b,c} (%)
1	Bn	13 , NEt ₃	40	80	20
2	Bn	16 , NEt ₃	>95	43 (40)	57 (55)
3	Bn	13 , DBU	>95	>95 (97)	<5
4	Bn	16 , DBU	>95	>95 (98)	<5
5	Me	13 , DBU	45	>95	<5
6	Me	16 , DBU	>95	>95 (95)	<5

^a All reactions conducted using aldehyde **5** (0.166 mmol), base (1.5 equiv.), NHC precatalyst **13** or **16** (10 mol%), ROH (2 equiv.).

^b Conversion, as well as ratio of ester and thiol ester products, determined by ¹H NMR spectroscopic analysis of the crude reaction product. ^c Parentheses indicate isolated yields after chromatographic purification.

transesterification of any thiol ester product. These observations can be explained by considering mechanistic observations reported by Bode²⁴ and Berkessel.^{5a} Bode showed that triazolylidenes bearing electron-donating *N*-aryl substituents (such as *N*-mesityl) favour azolium enolate formation, while *N*-aryl groups bearing electron-withdrawing substituents (C₆F₅ or C₆H₂Cl₃) favour acyl azolium intermediates.²⁵ A detailed mechanistic study by Berkessel and co-workers has recently shown that NHC-catalysed redox esterification of aldehydes with imidazolium based NHC catalysts proceeds through an azolium enolate intermediate.^{5a}

With these principles in mind, a plausible mechanism for the processes reported here is outlined in Fig. 3. Initially, the triazolium precatalyst is deprotonated by base to generate the corresponding NHC. Addition to the α -phenylthioaldehyde generates the corresponding tetrahedral adduct **I**, which after proton transfer gives the corresponding Breslow intermediate **II**. Elimination of thiophenolate gives the azolium enol intermediate ion pair **III**. For triazolium precatalysts with electron-withdrawing *N*-aryl substituents (**15** and **16**) tautomerisation to the acyl azolium **IV** is preferred, and reaction of **IV** with the thiophenolate counterion generates the thiol ester. For triazolium precatalysts bearing electron donating *N*-aryl substituents such as **13** and **14** (when NEt₃ is used as base), the azolium enolate intermediate **V** is preferred. Reaction of **V** with alcohols (R'OH) *via* a concerted deprotonation-nucleophilic attack as outlined by Berkessel forms the ester product. In the presence of DBU, the thiol ester can be converted to the corresponding ester product.

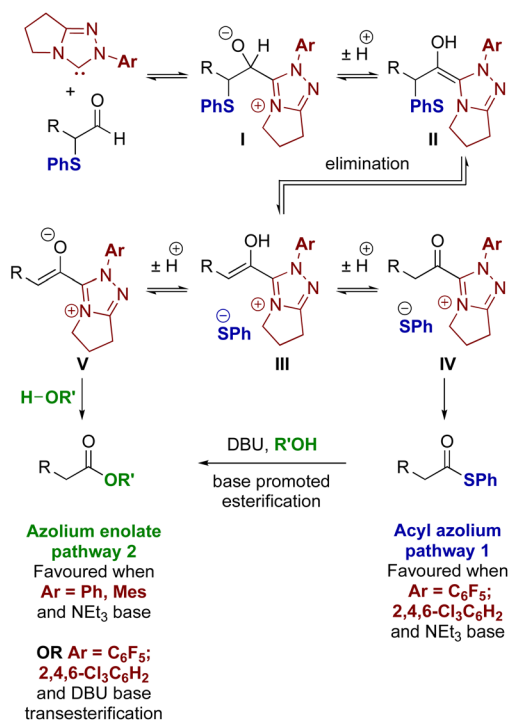
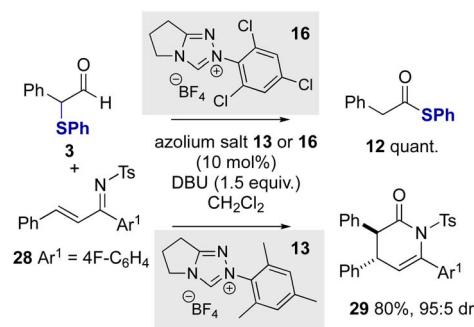


Fig. 3 Proposed mechanism for acyl azolium and azolium enolate generation.

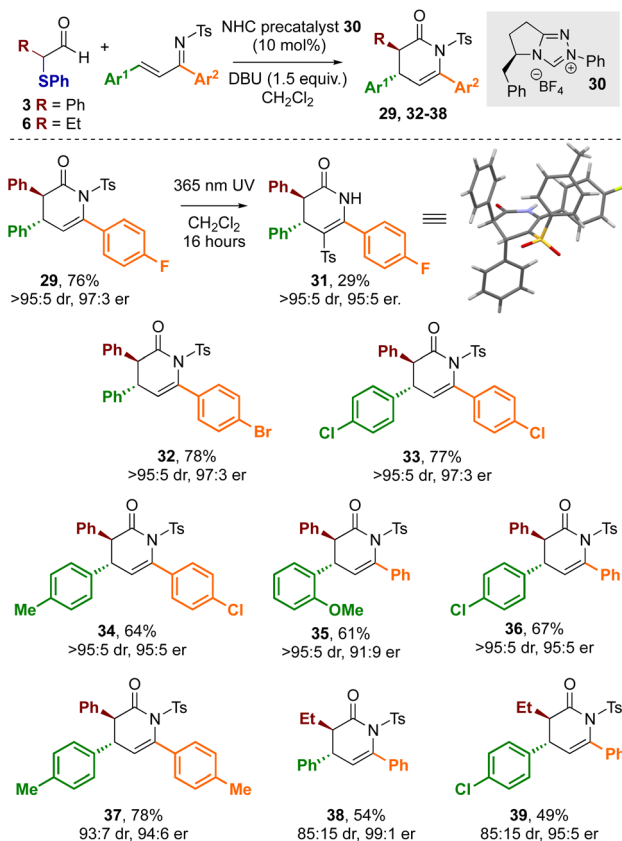
2.6 NHC-catalysed formal [4 + 2] cycloadditions

Further work sought to employ α -phenylthioaldehydes in an enantioselective C–C bond forming reaction between azolium enolates and α,β -unsaturated tosyl imines.^{12a} Given the trends in reactivity observed with variation of the *N*-aryl substituent of the NHC, initial studies probed the effect of the catalyst *N*-aryl substituent on product distribution in the racemic series using α -phenylthioaldehyde **3** and α,β -unsaturated tosyl imine **28** as substrates (Scheme 4). Consistent with the previous observations, *N*-2,4,6-trichlorophenyl triazolium precatalyst **16** gave exclusively thiophenyl ester **12** in quantitative yield with no dihydropyridone product observed. However, when using *N*-mesityl precatalyst **13** in the presence of DBU as base the product from preferential azolium enolate formation was observed, giving the desired (\pm)-cycloadduct **29** in 80% yield as a single diastereoisomer (>95 : 5 dr).

Building on these observations, the use of enantiopure NHC precatalyst **30** bearing an electron rich *N*-Ph substituent was investigated for enantioselective cycloaddition reactions. Using **30** with DBU as the base and α -phenylthioaldehyde **3** with α,β -unsaturated tosyl imine **28** gave the desired dihydropyridinone product **29** in 76% yield in excellent stereoselectivity (>95 : 5 dr, 97 : 3 er, Scheme 5). The relative and absolute configuration of **29** was unambiguously determined by X-ray crystallographic analysis of derivative **31**,²⁶ derived from facile *N*- to *C*-sulfonyl transfer of the product,²⁷ and is consistent with the selectivity observed in previous NHC-catalysed [4 + 2] cycloadditions (Scheme 4).^{12a} Using aldehyde **3** a variety of substrates were explored in this protocol through variation of the aromatic substituents within the α,β -unsaturated tosyl imine reactant. In all cases, the corresponding dihydropyridinones **32**–**37** were isolated in good yields (61–78%) and in excellent stereoselectivity (typically >95 : 5 dr and \sim 95 : 5 er), demonstrating the use of α -phenylthioaldehyde **3** as an effective azolium enolate precursor. Furthermore, the demonstration of this process on a preparative laboratory scale was investigated, allowing the preparation of **36** using the corresponding imine (1.5 mmol scale rather than 0.2 mmol), giving **36** in 57% yield (438 mg, 95 : 5 dr, 95 : 5 er). To further develop the scope of this process, extension to aliphatic aldehyde **6** as the azolium enolate precursor was applied, giving dihydropyridinones **38** and **39** in 54 and 49% yield respectively and in excellent stereoselectivity (85 : 15 dr, 99 : 1 er and 85 : 15, 95 : 5 er respectively).



Scheme 4 Evaluation of azolium enolate generation with NHC catalysts.



Scheme 5 Scope of formal [4 + 2] cycloadditions.^a All reactions were conducted in a flame dried Schlenk flask under a nitrogen atmosphere using 0.4 mmol of **3**, 0.2 mmol of the corresponding imine.^b Yields are isolated yields ^c dr was determined by ¹H NMR spectroscopy of the crude reaction mixtures, er of purified product was determined by HPLC analysis on a chiral stationary phase. X-ray representation; blue = N; red = O; dark grey = C; light grey = H; yellow = S; green = F.

3. Conclusions

In summary, α -phenylthioaldehydes were prepared by a simple two-step synthesis from commercially available reagents. They can act as acyl azolium and azolium enolate precursors for a range of NHC-mediated catalytic processes. Due to the efficient leaving group ability and nucleophilicity of thio-phenolate,²⁸ a variety of thiol esters can be prepared from these species *via* redox rearrangement. In the presence of an external alcohol, competition between redox rearrangement and redox esterification can be controlled through judicious choice of the *N*-aryl substituent within the NHC pre-catalyst and the base used in the reaction. With NEt₃ as base, NHCs bearing electron withdrawing (*N*-C₆F₅ or *N*-C₆H₂Cl₃) substituents favour redox rearrangement, while triazolium pre-catalysts with electron-rich *N*-aryl substituents (*N*-Mes) result in preferential redox esterification. Using DBU, redox esterification is preferred due to transesterification. α -Phenylthioaldehyde-derived azolium enolates are also used for enantio- and diastereoselective [4 + 2]-cycloaddition reactions with α,β -unsaturated tosyl imines, giving dihydropyridinone derivatives with excellent stereo-control (up to >95 : 5 dr, >98 : 2 er).

Data availability

All data (experimental procedures and characterization) that support the findings of this study are available within the article and its ESI.† Crystallographic data for compound **31** has been deposited with the Cambridge Crystallographic Data Centre under deposition numbers 2310270. The research data supporting this publication can be accessed from “ α -phenylthioaldehydes for the effective generation of acyl azolium and azolium enolate intermediates”. University of St Andrews Research Portal. DOI: <https://doi.org/10.17630/b0dfa038-03b4-4054-a2a3-72a4e1593b16>. PURE ID: 302087534.

Author contributions

ADS conceived the project; PE and PKM carried out experimental studies in consultation with CMY. CP carried out preliminary studies. ADS and PKM wrote the manuscript. KvR carried out single crystal X-ray analysis. ADS and PLA raised the funding. All authors reviewed, edited, and agreed on the finalised version of the manuscript.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

The research leading to these results has received funding from the EPSRC Centre for Doctoral Training in Critical Resource Catalysis (CRICAT) (PE, EP/L016419/1), the University of St Andrews (CP), the EPSRC (CMY, EP/S019359/1) and the EU (Marie-Curie Fellowship to PKM, Advanced Research Grant to PLA, 740311). Part of the work was supported by the U.S. Department of Energy (DOE), Office of Science, Office of Basic Energy Sciences, Chemical Sciences, Geosciences, and Biosciences Division, in the Catalysis Program at the Lawrence Berkeley National Laboratory under contract no. DE-AC02-05CH11231 (PLA). ADS and EZ-C thank the EPSRC Programme Grant “Boron: Beyond the Reagent” (EP/W007517/1) for support.

References

- For selected reviews, see: (a) D. M. Flanigan, F. Romanov-Michailidis, N. A. White and T. Rovis, *Chem. Rev.*, 2015, **15**, 9307–9387; (b) R. S. Menon, A. T. Biju and V. Nair, *Chem. Soc. Rev.*, 2015, **44**, 5040–5052; (c) *N-Heterocyclic carbenes in organocatalysis*, ed. A. T. Biju, Wiley VCH, Weinheim, 2019; (d) M. N. Hopkinson, C. Richter, M. Schedler and F. Glorius, *Nature*, 2014, **510**, 485–496; (e) K. J. R. Murauski, A. A. Jaworski and K. A. Scheidt, *Chem. Soc. Rev.*, 2018, **47**, 1773–1782; (f) S. J. Ryan, L. Candish and D. W. Lupton, *Chem. Soc. Rev.*, 2013, **42**, 4906–4917; (g) A. Mondal, A. Ghosh and A. T. Biju, *Chem. Rec.*, 2022, **22**, e202200054; (h) Y. Liu, Y. Wang, X. Wu and Y. R. Chi, *Chem. Rec.*, 2023, **23**, e202200219.

- 2 G. A. Grasa, R. M. Kissling and S. P. Nolan, *Org. Lett.*, 2002, **4**, 3583–3586.
- 3 G. W. Nyce, J. A. Lamboy, E. F. Connor, R. M. Waymouth and J. L. Hedrick, *Org. Lett.*, 2002, **4**, 3587–3590.
- 4 C. D. Risi, A. Brandolese, G. D. Carmine, D. Ragno, A. Massi and O. Bortolini, *Chem.–Eur. J.*, 2023, **29**, e2022024.
- 5 For selected examples, see: (a) A. Biswas, J.-M. Neudörfl, N. E. Schlörer and A. Berkessel, *Angew. Chem., Int. Ed.*, 2021, **60**, 4507–4511; (b) Y. Suzuki, K. Yamauchi, K. Muramatsu and M. Sato, *Chem. Commun.*, 2004, **23**, 2770–2771; (c) T. Kano, K. Sasaki and K. Maruoka, *Org. Lett.*, 2005, **7**, 1347–1349; (d) J. E. Thomson, K. Rix and A. D. Smith, *Org. Lett.*, 2006, **8**, 3785–3788; (e) B. E. Maki, A. Chan, E. M. Phillips and K. A. Scheidt, *Org. Lett.*, 2007, **9**, 371–374; (f) L. Candish and D. W. Lupton, *Org. Lett.*, 2010, **12**, 4836–4839; (g) B. Wanner, J. Mahatthananchai and J. W. Bode, *Org. Lett.*, 2011, **13**, 5378–5381; (h) C. Zhang, J. F. Hooper and D. W. Lupton, *ACS Catal.*, 2017, **7**, 2583–2596; (i) Y. Liu, Q. Chen, C. Mou, L. Pan, X. Duan, X. Chen, H. Chen, Y. Zhao, Y. Lu, Z. Jin and Y. R. Chi, *Nat. Commun.*, 2019, **10**, 1675; (j) Y. Liu, P. K. Majhi, R. Song, C. Mou, L. Hao, H. Chai, Z. Jin and Y. R. Chi, *Angew. Chem., Int. Ed.*, 2020, **59**, 3859–3863; (k) X. Yang, P. K. Majhi, H. Chai, B. Liu, J. Sun, T. Liu, Y. Liu, L. Zhou, J. Xu, J. Liu, D. Wang, Y. Zhao, Z. Jin and Y. R. Chi, *Angew. Chem., Int. Ed.*, 2021, **60**, 159–165.
- 6 (a) A. Chan and K. A. Scheidt, *Org. Lett.*, 2005, **7**, 905–908; (b) B. E. Maki, E. V. Patterson, C. J. Cramer and K. A. Scheidt, *Org. Lett.*, 2009, **11**, 3942–3945.
- 7 S. S. Sohn and J. W. Bode, *Org. Lett.*, 2005, **7**, 3873–3876.
- 8 For selected reviews, see: (a) X.-Y. Chen and S. Ye, *Synlett*, 2013, **24**, 1614–1622; (b) H. U. Vora, P. Wheeler and T. Rovis, *Adv. Synth. Catal.*, 2012, **354**, 1617–1639; (c) J. Douglas, G. Churchill and A. D. Smith, *Synthesis*, 2012, **44**, 2295–2309; (d) A. Ghosh and A. T. Biju, *Angew. Chem., Int. Ed.*, 2021, **60**, 13712–13724; (e) S. Mondal, S. R. Yetra, S. Mukherjee and A. T. Biju, *Acc. Chem. Res.*, 2019, **52**, 425–436.
- 9 M. He, G. J. Uc and J. W. Bode, *J. Am. Chem. Soc.*, 2006, **128**, 15088.
- 10 Y.-R. Zhang, L. He, X. Wu, P.-L. Shao and S. Ye, *Org. Lett.*, 2008, **10**, 277–280.
- 11 N. Duguet, C. D. Campbell, A. M. Slawin and A. D. Smith, *Org. Biomol. Chem.*, 2008, **6**, 1108–1113.
- 12 (a) L. Hao, Y. Du, H. Lv, X. Chen, H. Jiang, Y. Shao and Y. R. Chi, *Org. Lett.*, 2012, **14**, 2154–2157; (b) L. Hao, C. W. Chuen, R. Ganguly and Y. R. Chi, *Synlett*, 2013, **24**, 1197–1200; (c) B. Liu, R. Song, J. Xu, P. K. Majhi, X. Yang, S. Yang, Z. Jin and Y. R. Chi, *Org. Lett.*, 2020, **22**, 3335–3338.
- 13 (a) K. B. Ling and A. D. Smith, *Chem. Commun.*, 2011, **47**, 373–375; (b) J. E. Taylor, A. T. Davies, J. J. Douglas, G. Churchill and A. D. Smith, *Tetrahedron: Asymmetry*, 2017, **28**, 355–366; (c) N. T. Reynolds and T. Rovis, *J. Am. Chem. Soc.*, 2005, **127**, 16406–16407; (d) Y. Kawanaka, E. M. Phillips and K. A. Scheidt, *J. Am. Chem. Soc.*, 2009, **131**, 18028–18029.
- 14 H. U. Vora and T. Rovis, *J. Am. Chem. Soc.*, 2010, **132**, 2860–2861.
- 15 M. He, G. J. Uc and J. W. Bode, *J. Am. Chem. Soc.*, 2006, **128**, 15088–15089.
- 16 S. Chen, L. Hao, Y. Zhang, B. Tiwari and Y. R. Chi, *Org. Lett.*, 2013, **15**, 5822–5825.
- 17 (a) A. T. Davies, A. M. Z. Slawin and A. D. Smith, *Chem.–Eur. J.*, 2015, **21**, 18944–18948; (b) A. T. Davies, M. D. Greenhalgh, A. M. Z. Slawin and A. D. Smith, *Beilstein J. Org. Chem.*, 2020, **16**, 1572–1578; (c) R. W. F. Kerr, M. D. Greenhalgh, A. M. Z. Slawin, P. L. Arnold and A. D. Smith, *Tetrahedron: Asymmetry*, 2017, **28**, 125–134; (d) A. T. Davies, J. E. Taylor, J. Douglas, C. J. Collett, L. C. Morrill, C. Fallan, A. M. Z. Slawin, G. Churchill and A. D. Smith, *J. Org. Chem.*, 2013, **78**, 9243–9257; (e) A. T. Davies, P. M. Pickett, A. M. Z. Slawin and A. D. Smith, *ACS Catal.*, 2014, **4**, 2696–2700; (f) N. Attaba, J. E. Taylor, A. M. Z. Slawin and A. D. Smith, *J. Org. Chem.*, 2015, **80**, 9728–9739.
- 18 J. E. Taylor, D. S. B. Daniels and A. D. Smith, *Org. Lett.*, 2013, **15**, 6058–6061.
- 19 (a) S. Lu, S. B. Poh and Y. Zhao, *Angew. Chem. Int. Ed.*, 2014, **53**, 11041–11045; (b) S. Lu, X. Song, S. B. Poh, H. Yang, M. W. Wong and Y. Zhao, *Chem.–Eur. J.*, 2017, **23**, 2275–2281.
- 20 (a) T. Sato, M. Inoue, S. Kobara, J. Otera and H. Nozaki, *Tetrahedron Lett.*, 1989, **30**, 91–94; (b) T. Sato, J. Otera and H. Nozaki, *J. Org. Chem.*, 1990, **55**, 6116–6121.
- 21 M. S. Kerr, J. R. de Alaniz and T. Rovis, *J. Org. Chem.*, 2005, **70**, 5725–5728.
- 22 As a measure of increasing steric hindrance in this series we noted that the “A” value of a substituent increases from Me to *t*-Bu (Me, A value 1.7 kcal/mol; Et, A value 1.75, *i*-Pr A value 2.15, *t*-Bu A value 4.9 kcal/mol). This trend for increased steric hindrance is consistent with the increased time taken to generate full conversion to the product, J. A. Hirsch, in *Topics in Stereochemistry*, 1967, pp. 199–222.
- 23 R. A. L. Peerboom, L. J. de Koning and N. M. M. Nibbering, *J. Am. Soc. Mass Spectrom.*, 1994, **5**, 159–168.
- 24 (a) J. Mahatthananchai and J. W. Bode, *Acc. Chem. Res.*, 2014, **47**, 696–707; (b) C. J. Collett, R. S. Massey, O. R. Maguire, A. S. Batsanov, A. C. O'Donoghue and A. D. Smith, *Chem. Sci.*, 2013, **4**, 1514–1522; (c) S. K. Giri, R. Gour and K. P. R. Kartha, *RSC Adv.*, 2017, **7**, 13653–13667.
- 25 J. Mahatthananchai and J. W. Bode, *Chem. Sci.*, 2012, **3**, 192–197.
- 26 CCDC 2310270 contains the supplementary crystallographic data for **31**. These data can be obtained free of charge from The Cambridge Crystallographic Data Centre.
- 27 (a) P.-P. Yeh, J. E. Taylor, D. G. Stark, D. S. B. Daniels, C. Fallan, J. C. Walton and A. D. Smith, *Org. Biomol. Chem.*, 2017, **15**, 8914–8922; (b) C. Simal, T. Lebl, A. M. Z. Slawin and A. D. Smith, *Angew. Chem., Int. Ed.*, 2012, **51**, 3653–3657.
- 28 For an NHC-catalyzed process involving the addition of thiophenols to the C=C of an enal see Q. Wang, S. Wu, J. Zou, X. Liang, C. Mou, P. Zheng and Y. R. Chi, *Nat. Commun.*, 2023, **14**, 4878.