



Cite this: *RSC Sustainability*, 2024, **2**, 2249

Received 29th April 2024
Accepted 27th May 2024

DOI: 10.1039/d4su00206g
rsc.li/rscsus

Green synthesis of thioamide derivatives in an environmentally benign deep eutectic solvent (DES)†

Susmita Mandal, ^a Archana Jain ^{*b} and Tarun K. Panda ^{*a}

A mild, direct, simple, and highly efficient green protocol is described for synthesizing thioamides. A wide variety of thioamides are obtained in good-to-excellent yields by the reaction of readily available starting materials, substituted aldehydes/ketones, secondary amines, and elemental sulfur in a choline chloride–urea (1 : 2)-based deep eutectic solvent (DES). This protocol, which can be applied without an additional catalyst support, contributes significantly to enhancing sustainability by reducing energy consumption and waste reduction, involving biodegradability, improved reactivity, and versatility. The DES can be recycled and used several times without any significant loss in terms of its activity.

Sustainability spotlight

With the growing need for sustainable chemistry, deep eutectic solvents (DESs) drive the field of synthetic chemistry towards more green practices. In most conventional methods, organic solvents are used which have adverse effects on the environment. In this paper, we report the synthesis of a wide variety of thioamides in good to excellent yield under mild conditions using a mixture of choline chloride and urea (1 : 2) (DES) as an environmentally benign solvent as well as a catalyst without using any toxic organic solvent and additional catalyst. Our work aligns with the 12 principles of green chemistry and the following UN sustainable development goals: SDG 9 (industry), SDG 12 (responsible consumption and production), and SDG 13 (climate action).

Introduction

The development of environmentally benign and sustainable synthetic methodologies is undeniably necessary in the modern era of organic synthesis, owing to ever-increasing concerns over environmental contamination and its impact. In addition to our scientific obligations, it is now a moral and ethical responsibility to protect our environment for future generations. In this context, the 12 principles of green chemistry, first proposed by Anastas and Warner in 1998,¹ have transformed the way we approach chemical synthesis. The principles of green chemistry promote the design and development of chemical processes that minimize environmental impact, reduce waste, and maximize efficiency.^{2,3} Recently, deep eutectic solvents (DESs) have emerged as a potential new class of solvents that not only follow the principles of green chemistry, but also have a wide range of applications in various fields.^{4,5} DESs, first reported by Abbott and co-workers,⁶ exhibit numerous unique properties such as

low volatility, high thermal stability, and variable polarity.^{7,8} Additionally, they are easily accessible, relatively nontoxic, low-cost, and biodegradable.⁹ DESs have been employed as media/catalysts in a variety of chemical reactions, such as biomass processing,^{10,11} polymerizations,¹² material synthesis,^{13,14} electrochemistry,^{15,16} nanotechnology,¹⁷ and multicomponent reactions (MCRs).^{18,19} Literature reports show that DESs have the potential to play an important part in enhancing the sustainability of MCRs.²⁰ Their green solvent properties and versatility render them valuable for researchers striving to develop more sustainable chemical synthetic methods.²¹

Thioamides, characterized by the presence of the $-\text{C}(=\text{S})\text{NH-}$ functional group, have been employed as indispensable building blocks in the synthesis of bioactive compounds,^{22,23} pharmaceutical agents,^{24–26} agrochemicals,²⁷ and functional materials with tailored properties.²⁸ Among the myriad of various traditional synthetic methods reported in the literature,^{29–39} one of the most efficient methods for thioamide synthesis is the Willgerodt–Kindler (WK) reaction, comprising the three-component condensation of a carbonyl compound, a sulfur reagent, and an amine (primary or secondary) in one pot.^{40,41} The reported protocols produce good results in many cases. However, most of them involve the use of expensive chemicals, such as sulfur reagents, metal catalysts, and toxic organic solvents, as well as long reaction times, excess reactants, harsh reaction conditions, and strenuous work-up

^aDepartment of Chemistry, Indian Institute of Technology Hyderabad, Kandi, Sangareddy, Telangana, 502284, India. E-mail: tpanda@chy.iith.ac.in; Tel: +91-40-2301-6254

^bDepartment of Physics and Chemistry, Mahatma Gandhi Institute of Technology, Gandipet, Hyderabad, Telangana, 500 075, India. E-mail: archanajain_chem@mgiit.ac.in

† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d4su00206g>

methods. In some instances, the yield of products is also low. To avoid such limitations, there is a significant trend toward substituting most traditional organic solvents or harsh reaction conditions with ones that have a reduced impact on the environment while still achieving the same outcomes. Ionic liquids and DESs have recently emerged as potential substitutes for toxic organic solvents.⁴² Hitherto, only a few examples of thioamide synthesis using ionic liquids and DESs have been reported (Scheme 1).^{43–45} The protocols reported involved the use of relatively toxic ionic liquid/metal-based DESs, high temperature conditions, presence of toxic organic solvents, an excess of reactants, and limited substrate scope.

Given this context, we were interested in showcasing the pivotal role of metal-free DESs in the synthesis of a series of biologically and pharmaceutically important thioamide derivatives *via* a multicomponent approach (*i.e.*, Willgerodt–Kindler reaction), while aligning with the principles of green chemistry. Recently, we reported the application of a choline chloride (ChCl)–urea (1 : 2)-based DES in the synthesis of α -amino-phosphorus derivatives.⁴⁶ This DES is composed of the naturally occurring hydrogen-bond acceptor (HBA) choline chloride and hydrogen-bond donor (HBD) urea in a 1 : 2 molar ratio.

Here, we report a green methodology for the synthesis of thioamide derivatives *via* a Willgerodt–Kindler (W–K) reaction between aldehydes/ketones, different secondary amines, and sulfur powder in a ChCl–urea (1 : 2)-based DES under mild conditions. This methodology positively contributes to enhancing sustainability by: (i) reducing the need for additional toxic solvents/catalysts, thereby improving the overall efficiency of the reaction in terms of high yield and selectivity, (ii) cutting energy consumption through low-temperature reactions due to its relatively low melting point, (iii) lowering waste generation by using reagents in stoichiometric ratio and since the DES is easily recyclable, it allows the recovery of reactants and products, and (iv) reducing the environmental burden due to its biodegradable nature. We also describe mechanistic insights

and versatility of the DES with various substrates (up to 40 examples). Furthermore, we evaluate the reusability of the DES.

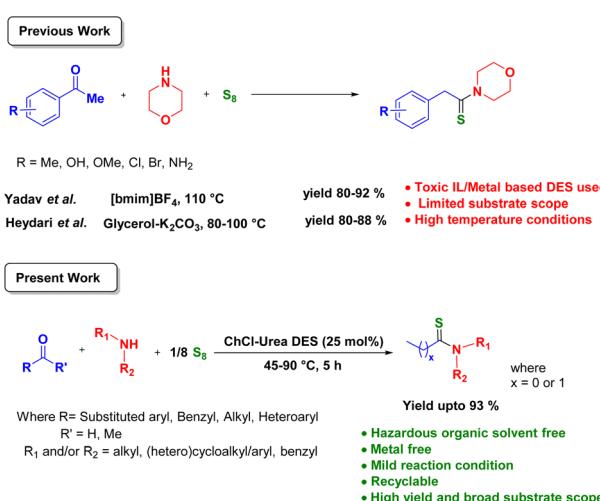
Results and discussion

In this investigation, we present a simple and green methodology for synthesizing a series of thioamide derivatives *via* a multicomponent approach (*i.e.*, Willgerodt–Kindler reaction) using a DES [ChCl–urea (1 : 2)] as the reaction medium as well as a catalyst, without adding any other organic solvent or catalyst, under mild reaction conditions.

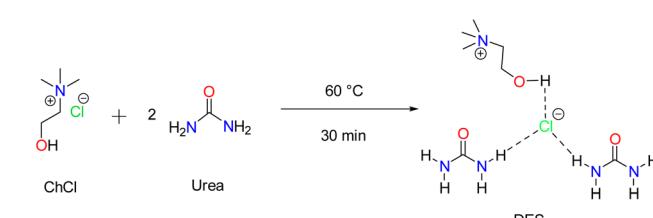
The DES was synthesized by following the previously reported method of heating a mixture of choline chloride (1 mol) and urea (2 mol) at 60 °C for 30 minutes (Scheme 2).⁴⁷ The colorless liquid formed was used for thioamidation reaction as such without any purification.

To begin our investigation, we selected the thioamidation reaction between *p*-tolualdehyde (**1a**), diethylamine (**2a**), and sulfur powder as the model reaction. The three reactants, **1a**, **2a**, and sulfur powder, in a 1 : 1 : 0.125 molar ratio, were reacted in 25 mol% of DES [ChCl–urea (1 : 2)] in one pot with constant stirring at 45 °C for five hours. Progress of the reaction was monitored by thin-layer chromatography (TLC). After completion, the mixture was cooled to room temperature and then diluted with water and ethyl acetate. To our delight, the corresponding thioamide derivative was isolated in a yield of 93% (Table 1, entry 3). The DES could easily be recycled by evaporating the water under vacuum.

The thioamidation reaction can be optimized by altering several parameters, *viz.*, reaction temperature, reaction time, DES dosage, and subtraction of DES (Table 1). Initially, when the reaction between **1a**, **2a**, and sulfur powder in a 1 : 1 : 0.125 molar ratio was performed in the absence of DES at 45 °C for 24 hours under neat conditions, no product was formed (Table 1, entry 1). This explains the role of the DES as an effective catalyst as well as medium. The high polarity and unique hydrogen-bonding properties of the DES influence the reactivity of reactants. It also accelerates the reaction speed and improves selectivity by solvating and stabilizing reactive intermediates or transition states in the catalytic reactions.⁴⁸ Even in the DES, this reaction at room temperature did not afford any product (Table 1, entry 2). This may be because at 45 °C, mass transfer of reactants in the DES increases, facilitating more effective reactant contact. Other DESs, such as ChCl–oxalic acid (1 : 1) and ChCl–PTSA (1 : 1) were also tested, but we were unsuccessful in obtaining the products (Table 1, entries 6, 7). However, using

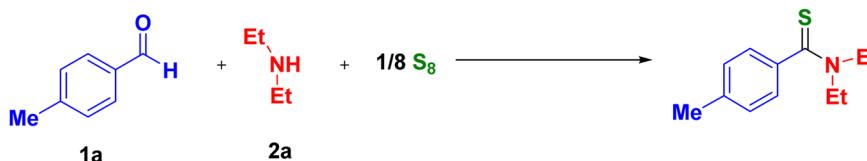


Scheme 1 Schematic representation of previous methods and present work using ionic liquid/DES.



Scheme 2 Synthesis of the DES from choline chloride and urea.



Table 1 Optimization of reaction conditions for the reaction between *p*-tolualdehyde, diethylamine, and elemental sulfur^a

Entry	1a (equiv.)	2a (equiv.)	S (equiv.)	DES	Loading of DES (mol%)	Time (h)	Temp. (°C)	Yield ^a (%)
1	1	1	1	—	—	24	45	—
2	1	1	1	ChCl-urea (1 : 2)	25	24	rt	—
3	1	1	1	ChCl-urea (1 : 2)	25	5	45	93
4	1	1	2	ChCl-urea (1 : 2)	25	5	45	93
5	1	2	1	ChCl-urea (1 : 2)	25	5	45	93
6	1	1	1	ChCl-OA (1 : 1)	25	5	45	—
7	1	1	1	ChCl-PTSA (1 : 1)	25	5	45	—
8	1	1	1	ChCl-urea (1 : 1)	25	5	45	74
9	1	1	1	ChCl-urea (1 : 2)	30	5	45	93
10	1	1	1	ChCl-urea (1 : 2)	20	5	45	83
11	1	1	1	ChCl-urea (1 : 2)	25	5	60	93
12	1	1	1	ChCl-urea (1 : 2)	25	4	45	86
13	1	1	1	ChCl-urea (1 : 2)	25	8	45	93

^a Isolated yield. OA-oxalic acid, PTSA-*p*-toluenesulfonic acid.

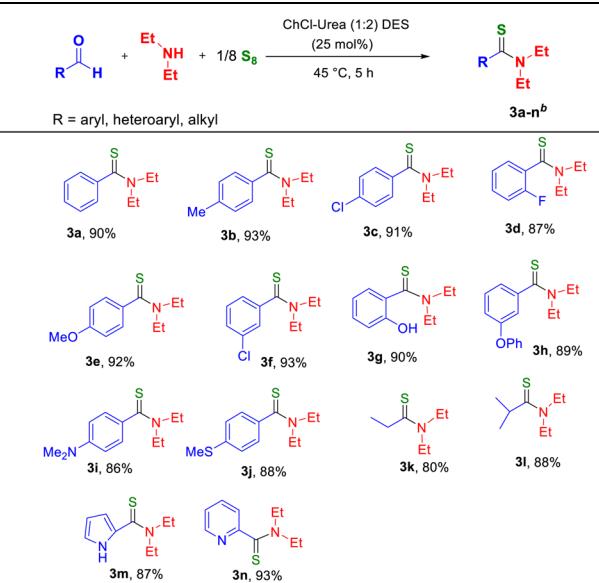
the DES in lower proportion ChCl:urea (1 : 1) led to the formation of the corresponding product in lower yield (74%) under similar conditions (Table 1, entry 8). For further screening, we changed the quantities of the starting materials, and the best result was obtained when **1a**, **2a**, and sulfur were used in a 1 : 1 : 0.125 molar ratio (Table 1, entries 3–5). Next, we optimized the amount of DES, and the highest yield was obtained with 25 mol% of DES (Table 1, entries 3, 9, and 10). Other parameters, such as reaction temperature (Table 1, entries 3, 10, 11) and time (Table 1, entries 11–13) were also analyzed. The ideal reaction temperature and time were 45 °C and five hours, respectively.

With the established optimized conditions, an extensive study was undertaken to examine the scope of this protocol, using a variety of aldehydes/ketones and amines in the DES [ChCl : urea (1 : 2)]. At first, the reactions of various aldehydes with diethylamine and sulfur powder were performed in the DES ChCl-urea (1 : 2) in 1 : 1 : 1 : 0.125 molar ratio at 45 °C for five hours. After the completion of each reaction, the corresponding thioamide (**3a**–**3n**) was isolated and characterized by ¹H and ¹³C NMR spectroscopy (see the ESI†). The results are shown in Table 2. We observed that aromatic aldehydes containing electron-releasing and electron-withdrawing groups reacted smoothly under mild conditions, and the corresponding thioamides were obtained in good-to-excellent yields (Table 2, **3a**–**3j**). Aliphatic aldehydes also reacted well and gave good yields (Table 2, **3k** and **3l**). Interestingly, heteroaromatic aldehydes such as pyrrole 2-carbaldehyde and pyridine 2-carbaldehyde also performed well when our method was used and afforded good-to-excellent yields (Table 2, **3m** and **3n**).

Next, we expanded our protocol to demonstrate the scope of the DES with the different types of amines. Under the given

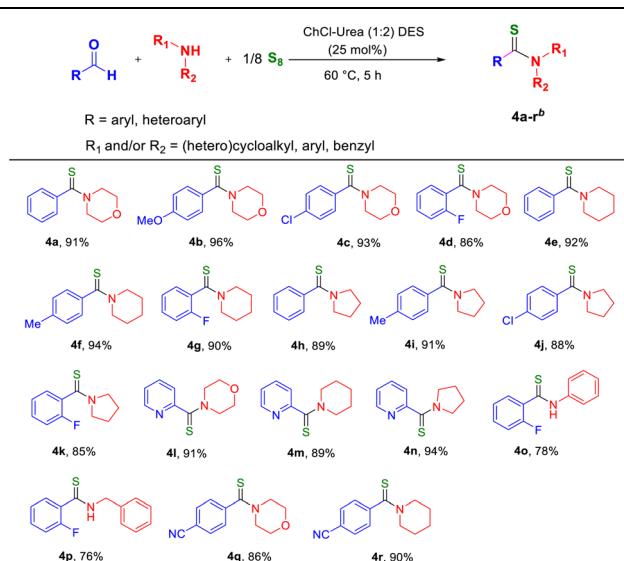
optimized conditions, the cyclic secondary amines (*viz.*, morpholine, piperidine, and pyrrolidine) reacted well with aromatic aldehydes containing electron-releasing as well as electron-withdrawing groups.

In all cases, the corresponding thioamides were obtained in good-to-excellent yields (Table 3, **4a**–**4k**). We also tested the

Table 2 Substrate scope of the DES with different aldehydes for the synthesis of thioamide derivatives^{a,b}

^a Reaction conditions: aldehyde (1.56 mmol), diethylamine (1.56 mmol), and sulfur powder (0.195 mmol) in 25 mol% DES ChCl-urea (1 : 2) at 45 °C for 5 h. ^b Isolated yields.

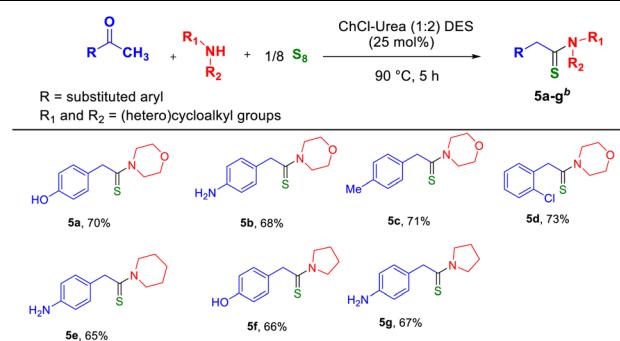


Table 3 Substrate scope of the DES with different amines for the synthesis of thioamide derivatives^{a,b}

^a Reaction conditions: aldehyde (1.56 mmol), amine (1.56 mmol) and sulfur powder (0.195 mmol) in 25 mol% DES ChCl-urea (1:2) at 60 °C for 5 h. ^b Isolated yields.

reactivity of cyclic secondary amines with a heterocyclic aldehyde, pyridine 2-carboxyaldehyde, under similar conditions and it afforded the pyridine-based thioamide motifs in excellent yields (Table 3, **4l–4n**). Primary amines, such as aniline and benzyl amine, also gave thioamides in good yields (Table 3, **4o** and **4p**). The thioamidation reaction was also performed with benzaldehyde bearing a reducible functional group such as nitro or cyano to show the limitations of this methodology. It was observed that 4-nitro benzaldehyde did not give any product when reacted with diethyl amine as well as heterocyclic amines such as morpholine and piperidine under the optimized conditions. However, 4-cyano benzaldehyde gave the corresponding products in good yield with morpholine and piperidine (Table 3, **4q** and **4r**) but did not afford any product with diethylamine. The corresponding thioamide compounds (**4a–4r**) were isolated and characterized using ¹H and ¹³C NMR spectroscopy (see the ESI†). This study demonstrated that our methodology is compatible with a broad range of aldehydes and amines bearing a variety of functional groups including methyl, methoxy, chloro, and fluoro, under mild conditions (Tables 2 and 3).

Upon achieving these results, we were keen to extend our methodology to various ketones. The W-K reaction with alkyl aryl ketones is quite interesting and challenging as it involves the reversal of polarity where the carbonyl –C(=O)– group is reduced and the terminal methyl group is oxidized, and new C=S and C–N bonds are formed. For the thioamidation reaction, different cyclic secondary amines, such as morpholine, piperidine, and pyrrolidine, were employed with the ketones bearing electron-donating and withdrawing groups, which afforded the corresponding thioamides (**5a–5g**) in moderate-to-

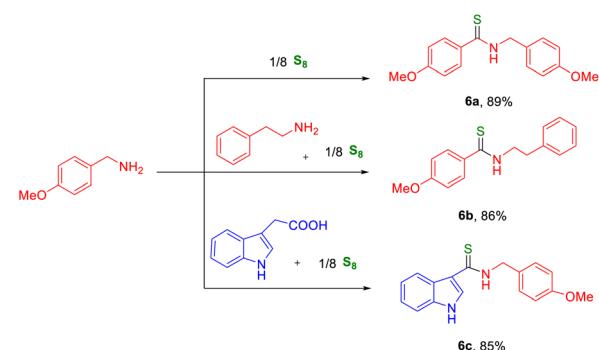
Table 4 Substrate scope of the DES with ketones and secondary amines for synthesis of thioamide derivatives^{a,b}

^a Reaction conditions: ketone (1.56 mmol), amine (1.56 mmol) and sulfur powder (0.195 mmol) in 25 mol% DES ChCl-urea (1:2) at 90 °C for 5 h. ^b Isolated yields.

good yields. For the ketones, the reaction conditions were similar to those used for aldehydes, except that the reaction was performed at a slightly higher temperature of 90 °C (Table 4). All the compounds (**5a–5g**) were isolated and characterized using ¹H and ¹³C NMR spectroscopy (see the ESI†).

We continued our exploration of the thioamidation reaction using different types of primary amines and acids. In one case, we used two same and different types of amines and sulfur powder in a 1:1:0.125 molar ratio and obtained the corresponding thioamide products in good yields (Table 5, **6a** and **6b**). In another case we used acid, primary amine, and sulfur powder in a 1:1:0.125 molar ratio and obtained the corresponding thioamide in good yield (Table 5, **6c**). Here, we tried the reaction using aliphatic and aromatic secondary amines as well as with aliphatic acid but failed to obtain any product.

Recycling the reaction media and catalysts is crucial for waste reduction – it also aligns with green chemistry principles and contributes to sustainability.⁴⁹ To assess reusability, we conducted the MCR using *p*-tolualdehyde, diethylamine, and

Table 5 Substrate scope of the DES with primary amines and primary amine/acid for synthesis of thioamide derivatives^{a,b}

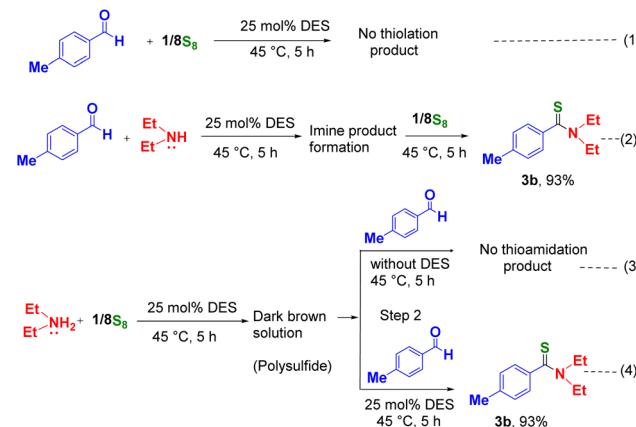
^a Reaction conditions: amine/acid (1.56 mmol), amine (1.56 mmol) and sulfur powder (0.195 mmol) in 25 mol% DES ChCl-urea (1:2) at 90 °C for 5 h. ^b Isolated yields.



sulfur powder in the DES under optimized conditions (at 45 °C for five hours). After the reaction was completed, the DES was recovered from the water phase by evaporation at 85 °C under reduced pressure. The recovered DES was further dried at 70 °C for three hours under reduced pressure to remove any traces of water and then subjected to the next run of the same reaction without adding more DES. We were able to reuse the DES medium four times for the same reaction with negligible loss of product yields (Fig. 1).

We also tested this methodology for industrial applications by performing a gram-scale reaction between 3.7 g of *p*-tolualdehyde, 2.3 g of aniline, and 1 g of sulfur powder in DES ChCl-urea (1 : 2) (25 mol%) under solvent-free conditions with constant stirring at 45 °C for five hours. The corresponding thioamides generated had a yield of 93%. These results highlight the efficacy of DES ChCl-urea (1 : 2) for the large-scale, ecofriendly synthesis of a series of biologically and pharmaceutically important thioamide derivatives. Additionally, we evaluated green chemistry metrics to demonstrate the environment-friendly nature of our methodology.⁵⁰ Ideally, high atom economy and a low *E*-factor value, together with high process mass intensity is crucial for green reactions. In this study, we found that the reaction has a small *E*-factor value (0.17), a high reaction mass efficiency value (RME = 85.5%), high atom economy (AE = 92%), and high process mass intensity (PMI = 1.17) (see the ESI† for green metrics calculation). This reaction avoids toxic solvents, expensive reagents, and harsh conditions.

To clarify the mechanism of this reaction, we performed some control reactions under the optimized conditions. At first, the *p*-tolualdehyde and sulfur powder were heated at 45 °C for five hours in the DES, we observed that no product was generated by the thiolation reaction (Scheme 3, eqn (1)). Later when aldehydes and diethylamine were reacted together at 45 °C for five hours in the DES, we observed the formation of the imine product which yielded thioamide on reaction with sulfur under the similar reaction conditions (Scheme 3, eqn (2)). Then, diethylamine was reacted with elemental sulfur powder at both room temperature and 45 °C for one hour. In both cases, a dark, brown-colored solution was obtained due to the formation of polysulfides. Subsequently, when we added aldehyde to the



Scheme 3 Control experiments.

brown-colored solution obtained in each case (in the absence of the DES) (Scheme 3, eqn (3)), we observed that the expected thioamide product was not formed. Only when the aldehyde was added to the reaction mixture of diethylamine and sulfur and heated at 45 °C in the presence of the DES (Scheme 3, eqn (4)), the expected thioamide product was formed. Results from our control experiments suggest that the formation of polysulfide (I) may take place even at room temperature under neat conditions and thioamide formation may be feasible in the DES medium at a slightly elevated temperature (45 °C) only. In this catalytic reaction, the DES may play a crucial role in inducing the reactivity of reactants by solvating and stabilizing reactive intermediates or transition states due to its high polarity and unique hydrogen-bonding properties. Therefore, the presence of the DES may be essential for the thioamidation reaction.

Based on results from our control experiments and the previous literature, we propose a most plausible mechanism of thioamidation reaction in the DES (Scheme 4).^{41,51–53} The first

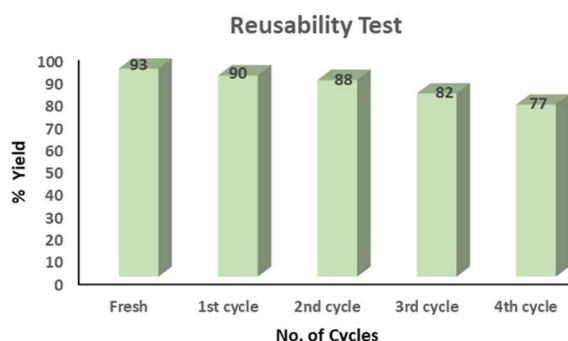
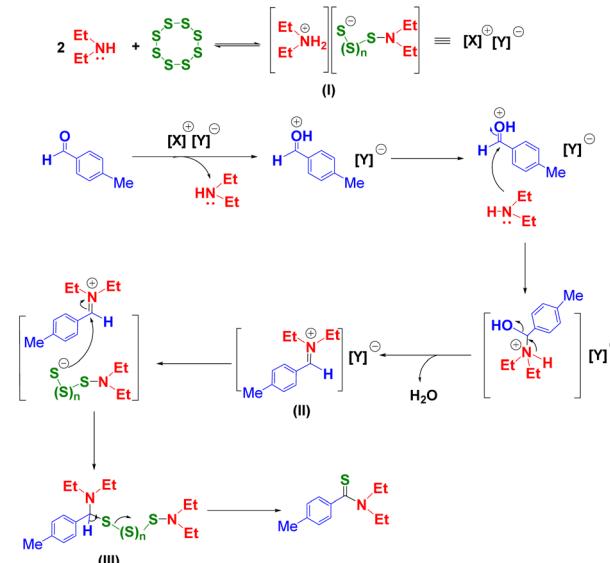
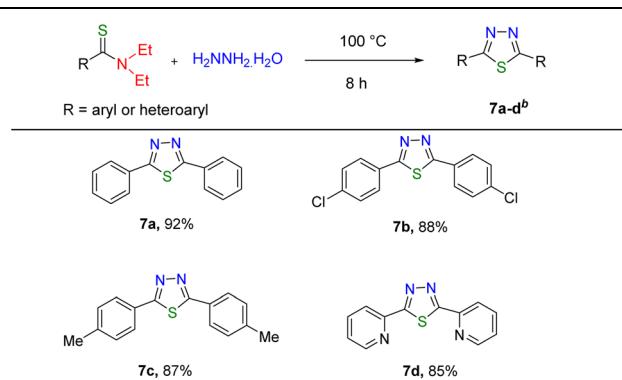


Fig. 1 Recyclability test of the DES. Reaction conditions: *p*-tolualdehyde (1.56 mmol), diethylamine (1.56 mmol) and sulfur powder (0.195 mmol) in 25 mol% DES ChCl-urea (1 : 2) at 45 °C for 5 h. Isolated yields.



Scheme 4 Plausible mechanism of thioamidation reaction.



Table 6 Further reaction of thioamides with hydrazine hydrate^{a,b}

^a Thioamide (1 equiv.) and hydrazine hydrate (10 equiv.) at 100 °C for 8 h. ^b Isolated yields.

step in this reaction involves a cleavage of the S-S bond of elemental sulfur by the nucleophilic attack of a lone nitrogen on the corresponding secondary amine to form the polysulfide anion (**I**) in a reversible way. Next, upon addition of the aldehyde, the protonation of aldehyde molecule enhances the electrophilicity of carbonyl carbon of the aldehyde and it thus reacts easily with secondary amine to form the imine intermediate (**II**). We anticipate that the driving force in the formation of the imine compound *in situ* might be the hydrogen bond interaction in the OH group of the intermediate formed with the DES.⁴⁷ Thereafter, the nucleophilic attack of polysulfide anion on the imine intermediate (**II**) led to intermediate **III** in which the hydrogen on the methylene group oxidizes and gives the final product.

We also present here the application of thioamide derivatives in the synthesis of thiadiazoles. Thiadiazoles are five-membered heterocyclic compounds which are extensively applied in medicinal, agricultural, industrial, and polymer chemistry. In particular, 1,3,4-thiadiazole compounds are important owing to their characteristics – antiviral,⁵⁴ antitubercular,⁵⁵ anti-depressant,⁵⁶ anti-microbial,⁵⁷ and anticancer.⁵⁸ 1,3,4-Thiadiazole derivatives have been used in the synthesis of different drugs like acetazolamide,⁵⁹ furidazine, megazol,⁶⁰ etc. There are various reports on thiadiazole derivative synthesis.^{61–64} Here, we have demonstrated the synthesis of 2,5-disubstituted 1,3,4-thiadiazole compounds from the reaction between thioamides we prepared with hydrazine hydrate at 100 °C for eight hours under neat conditions (Table 6). The corresponding thiadiazole compounds (**7a–7d**) were isolated and characterized using ¹H and ¹³C NMR spectroscopy (see the ESI†).

Conclusion

In summary, we have reported here a simple, novel, efficient, and scalable methodology for thioamide synthesis in a ChCl-urea DES without any other additive or organic solvent. Thioamides were obtained at the 45–60 °C temperature range within five hours with yields of 76–93% under optimized conditions

using aldehyde, secondary amine, and sulfur powder in a 1 : 1 : 0.125 molar ratio. We also used this protocol for ketones and obtained the corresponding thioamides in ~65–90% yield. Furthermore, we demonstrated the synthesis of thiadiazole derivatives using the thioamides prepared. The main benefits of this protocol are the use of the DES as a green solvent as well as a non-toxic catalyst, lower reaction temperature, and most importantly, easy recovery and reusability of the DES. We also performed a gram-scale reaction, which yielded good results. To the best of our knowledge, this is the first report on thioamidation of aldehydes using a metal-free basic DES.

Author contributions

Susmita Mandal: methodology, validation, formal analysis, data curation. Archana Jain: writing (original draft), data curation, project administration. Tarun K. Panda: funding acquisition, supervision, project administration, reviewing, and editing.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work was supported by the Science and Engineering Research Board (SERB), Government of India, under the Teaching Associateship for Research Excellence (TARE) scheme (project no. TAR/2020/000003). Instrumental facilities were provided by the Indian Institute of Technology, Hyderabad (IITH).

References

- 1 P. T. Anastas and J. C. Warner, *Green Chemistry: Theory and Practice*, Oxford University Press, New York, 1998, p. 30, By permission of Oxford University.
- 2 A. DeVierno Kreuder, T. House-Knight, J. Whitford, E. Ponnusamy, P. Miller, N. Jesse, R. Rodenborn, S. Sayag, M. Gebel, I. Aped, I. Sharfstein, E. Manaster, I. Ergaz, A. Harris and L. Nelowet Grice, *ACS Sustain. Chem. Eng.*, 2017, **5**, 2927–2935.
- 3 O. V. Kharissova, B. I. Kharisov, C. M. O. González, Y. P. Méndez and I. López, *R. Soc. Open Sci.*, 2019, **6**, 191378.
- 4 E. L. Smith, A. P. Abbott and K. S. Ryder, *Chem. Rev.*, 2014, **114**, 11060–11082.
- 5 L. Lomba, C. B. García, M. P. Ribate, B. Giner and E. Zuriaga, *Appl. Sci.*, 2021, **11**, 10156.
- 6 A. P. Abbott, G. Capper, D. L. Davies, H. L. Munro, R. K. Rasheed and V. Tambyrajah, *Chem. Commun.*, 2001, **1**, 2010–2011.
- 7 A. P. Abbott, D. Boothby, G. Capper, D. L. Davies and R. K. Rasheed, *J. Am. Chem. Soc.*, 2004, **126**, 9142–9147.
- 8 C. Ruß and B. König, *Green Chem.*, 2012, **14**, 2969–2982.
- 9 T. El Achkar, H. Greige-Gerges and S. Fourmentin, *Environ. Chem. Lett.*, 2021, **19**, 3397–3408.

10 M. Francisco, A. Van Den Bruinhorst and M. C. Kroon, *Green Chem.*, 2012, **14**, 2153–2157.

11 S. Xia, G. A. Baker, H. Li, S. Ravula and H. Zhao, *RSC Adv.*, 2014, **4**, 10586–10596.

12 F. Del Monte, D. Carriazo, M. C. Serrano, M. C. Gutiérrez and M. L. Ferrer, *ChemSusChem*, 2014, **7**, 999–1009.

13 D. V. Wagle, H. Zhao and G. A. Baker, *Acc. Chem. Res.*, 2014, **47**, 2299–2308.

14 D. Carriazo, M. C. Serrano, M. C. Gutiérrez, M. L. Ferrer and F. del Monte, *Chem. Soc. Rev.*, 2012, **41**, 4996–5014.

15 A. R. Hillman, K. S. Ryder, C. J. Zaleski, V. Ferreira, C. A. Beasley and E. Vieil, *Electrochim. Acta*, 2014, **135**, 42–51.

16 P. Sebastián, E. Vallés and E. Gómez, *Electrochim. Acta*, 2014, **123**, 285–295.

17 A. Abo-Hamad, M. Hayyan, M. A. H. AlSaadi and M. A. Hashim, *Chem. Eng. J.*, 2015, **273**, 551–567.

18 F. G. Calvo-Flores and C. Mingorance-Sánchez, *ChemistryOpen*, 2021, **10**, 815–829.

19 R. C. Cioc, E. Ruijter and R. V. A. Orru, *Green Chem.*, 2014, **16**, 2958–2975.

20 L. S. Longo and M. V. Craveiro, *J. Braz. Chem. Soc.*, 2018, **29**, 1999–2025.

21 P. Liu, J. W. Hao, L. P. Mo and Z. H. Zhang, *RSC Adv.*, 2015, **5**, 48675–48704.

22 Z. Moussa, Z. M. A. Judeh, M. A. M. S. El-Sharief and A. M. S. El-Sharief, *ChemistrySelect*, 2020, **5**, 764–798.

23 H. Verma, B. Khatri, S. Chakraborti and J. Chatterjee, *Chem. Sci.*, 2018, **9**, 2443–2451.

24 P. Zoumpoulakis, C. Camoutsis, G. Pairas, M. Soković, J. Glamočlija, C. Potamitis and A. Pitsas, *Bioorg. Med. Chem.*, 2012, **20**, 1569–1583.

25 F. Wang, R. Langley, G. Gulten, L. G. Dover, G. S. Besra, W. R. Jacobs and J. C. Sacchettini, *J. Exp. Med.*, 2007, **204**, 73–78.

26 J. R. Kirshner, S. He, V. Balasubramanyam, J. Kepros, C. Y. Yang, M. Zhang, Z. Du, J. Barsoum and J. Bertin, *Mol. Cancer Ther.*, 2008, **7**, 2319–2327.

27 J. Yu and X. Jiang, *Adv. Agrochem.*, 2023, **2**, 3–14.

28 Y. Huang, R. Hu and B. Z. Tang, *Sulfur-Containing Polymers: from Synthesis to Functional Materials*, WILEY-VCH GmbH, 2021, pp. 1–37.

29 W. Liu, C. Chen and H. Liu, *Beilstein J. Org. Chem.*, 2015, **11**, 1721–1726.

30 J. Li, X. Ren, G. Li, H. Liang, Y. Zhao, Z. Wang, H. Li and B. Yuan, *J. Sulfur Chem.*, 2020, **41**, 229–237.

31 T. B. Nguyen, L. Ermolenko and A. Al-Mourabit, *Org. Lett.*, 2012, **14**, 4274–4277.

32 T. B. Nguyen, M. Q. Tran, L. Ermolenko and A. Al-Mourabit, *Org. Lett.*, 2014, **16**, 310–313.

33 R. Liboska, D. Zylka and M. Bobek, *Synthesis*, 2002, 6–09.

34 B. Kaboudin, V. Yarahmadi, J. Y. Kato and T. Yokomatsu, *RSC Adv.*, 2013, **3**, 6435–6441.

35 A. D. Kale and D. S. Dalal, *ChemistrySelect*, 2022, **7**, e202203497.

36 J. Wei, Y. Li and X. Jiang, *Org. Lett.*, 2016, **18**, 340–343.

37 Y. Bian, X. Qu, Y. Chen, J. Li and L. Liu, *Molecules*, 2018, **23**, 2225.

38 N. Borthakur and A. Goswami, *Tetrahedron Lett.*, 1995, **36**, 6745–6746.

39 D. Brillon, *Synth. Commun.*, 1992, **22**, 1397–1401.

40 O. I. Zbruyev, N. Stiasni and C. O. Kappe, *J. Comb. Chem.*, 2003, **5**, 145–148.

41 K. Okamoto, T. Yamamoto and T. Kanbara, *Synlett*, 2007, 2687–2690.

42 D. A. Alonso, A. Baeza, R. Chinchilla, G. Guillena, I. M. Pastor and D. J. Ramón, *Eur. J. Org. Chem.*, 2016, **2016**, 612–632.

43 J. S. Yadav, B. V. S. Reddy, G. Kondaji, J. S. S. Reddy and K. Nagaiah, *J. Mol. Catal. A: Chem.*, 2007, **266**, 249–253.

44 I. Radfar, S. Abbasi, M. K. Miraki, E. Yazdani, M. Karimi and A. Heydari, *ChemistrySelect*, 2018, **3**, 3265–3267.

45 A. Gupta, J. K. Vankar, J. P. Jadav and G. N. Gururaja, *J. Org. Chem.*, 2022, **87**, 2410–2420.

46 S. Mandal, R. Narvariya, S. L. Sunar, I. Paul, A. Jain and T. K. Panda, *Green Chem.*, 2023, **25**, 8266–8272.

47 A. P. Abbott, G. Capper, D. L. Davies, R. K. Rasheed and V. Tambyrajah, *Chem. Commun.*, 2003, 70–71.

48 S. Khandelwal, Y. K. Tailor and M. Kumar, *J. Mol. Liq.*, 2016, **215**, 345–386.

49 V. Hessel, N. N. Tran, M. R. Asrami, Q. D. Tran, N. Van Duc Long, M. Escribà-Gelonch, J. O. Tejada, S. Linke and K. Sundmacher, *Green Chem.*, 2022, **24**, 410–437.

50 J. Martínez, J. F. Cortés and R. Miranda, *Processes*, 2022, **10**, 1274.

51 A. Gutiérrez-Hernández, A. Richaud, L. Chacón-García, C. J. Cortés-García, F. Méndez and C. A. Contreras-Celedón, *J. Org. Chem.*, 2021, **86**, 223–234.

52 X. Li, Q. Pan, R. Hu, X. Wang, Z. Yang and S. Han, *Asian J. Org. Chem.*, 2016, **5**, 1353–1358.

53 M. Carmack, *J. Heterocycl. Chem.*, 1989, **26**, 1319–1323.

54 D. Kumar, H. Kumar, V. Kumar, A. Deep, A. Sharma, M. G. Marwaha and R. K. Marwaha, *Med. Drug Discov.*, 2023, **17**, 100150.

55 S. G. Alegaon, K. R. Alagawadi, P. V. Sonkusare, S. M. Chaudhary, D. H. Dadwe and A. S. Shah, *Bioorg. Med. Chem. Lett.*, 2012, **22**, 1917–1921.

56 F. Clerici, D. Pocar, M. Guido, A. Loche, V. Perlini and M. Brufani, *J. Med. Chem.*, 2001, **44**, 931–936.

57 G. L. Almajan, S. F. Barbuceanu, G. Bancescu, I. Saramet, G. Saramet and C. Draghici, *Eur. J. Med. Chem.*, 2010, **45**, 6139–6146.

58 N. A. Al-Masoudi and Y. A. Al-Soud, *Nucleosides, Nucleotides Nucleic Acids*, 2008, **27**, 1034–1044.

59 J. R. A. Diaz, G. E. Camí, M. Liu-González, D. R. Vega, D. Vullo, A. Juárez, J. C. Pedregosa and C. T. Supuran, *J. Enzyme Inhib. Med. Chem.*, 2016, **31**, 1102–1110.

60 A. Abdel-Aziem, M. S. El-Gendy and A. O. Abdelhamid, *Eur. J. Chem.*, 2012, **3**, 455–460.

61 B. Stanovnik, *Compr. Org. Funct. Group Transform.*, 1995, **5**, 805–864.

62 A. R. Sayed, *Tetrahedron Lett.*, 2010, **51**, 4490–4493.

63 M. Zarei, *Tetrahedron*, 2017, **73**, 1867–1872.

64 V. Polshettiwar and R. S. Varma, *Tetrahedron Lett.*, 2008, **49**, 879–883.

