


 Cite this: *RSC Adv.*, 2020, 10, 9663

One-pot multicomponent synthesis of thieno[2,3-*b*]indoles catalyzed by a magnetic nanoparticle-supported [Urea]₄[ZnCl₂] deep eutectic solvent†

 The Thai Nguyen^{ab} and Phuong Hoang Tran ^{*ab}

In this study, we have developed the synthesis of thieno[2,3-*b*]indole dyes *via* a multicomponent reaction of cheap and available reagents such as sulfur, acetophenones, and indoles using a magnetic nanoparticle-supported [Urea]₄[ZnCl₂] deep eutectic solvent as a green catalyst. The synthesis of a series of diversely functionalized thieno[2,3-*b*]indole has been successfully performed in a one-pot reaction. Among a total of 25 compounds synthesized, there are 21 new compounds with full characterization such as FT-IR, ¹H and ¹³C NMR, HRMS (ESI). Due to the deep eutectic solvent coated surface of the magnetic nanoparticles, the catalyst could be recovered by an external magnet and reused in five consecutive runs without a considerable decrease in catalytic activity.

 Received 24th January 2020
 Accepted 28th February 2020

DOI: 10.1039/d0ra00773k

rsc.li/rsc-advances

1. Introduction

Thieno[2,3-*b*]indoles are common structural motifs in a variety of biological activities and therapeutically useful compounds leading to potential applications in treating epilepsy, neurological diseases,¹ plant-growth regulatory properties,² battery devices,³⁻⁷ and polymer conductors.⁴ Due to the potential applications of thieno[2,3-*b*]indoles, the development of new and efficient synthetic protocols for these compounds has received considerable attention. Traditionally, thieno[2,3-*b*]indoles are synthesized using functionalized 2-alkynylphenylisothiocyanates, 1-alkylisatins, and indole-2-thiones as starting materials, and require several steps to afford the desired products (Scheme 1).⁸⁻¹⁰ Although these classical approaches have been extensively used, there is ongoing research for new methods that have simpler work-ups, are atom-economical with good to excellent yields, use recyclable catalysts and use readily inexpensive starting materials. To date, only two reports have described the synthesis of thieno[2,3-*b*]indoles *via* a one-pot multicomponent reaction which provides a greener approach than the classical methods. Li and coworkers reported the Brønsted acid-catalyzed three-component synthesis of thieno[2,3-*b*]indoles starting from indoles, alkenes or alkynes, and sulfur powder.¹¹ Ni and coworkers demonstrated the synthesis of 3-substituted thieno[2,3-*b*]indoles *via* cascade cyclization by Brønsted acid-promoted annulation of indoles, ketones, and sulfur powder.¹² However, the low yields, narrow substrate

scope, and the inability to recycle catalysts remain a crucial challenge to industrial application.

The development of new catalysts is necessary for the advancement of organic synthesis.¹³⁻¹⁵ Over the last decade, several synthetic approaches have been reported in the field by employing deep eutectic solvents as catalysts.¹⁶⁻²⁰ However, significant drawbacks including difficult catalyst recovery, tedious product separation, and the requirement of greater stoichiometric amounts of DES as dual catalyst/solvent limited their widespread application.^{15,21-23} Hence, the idea for using supported DES always exists in our mind, which combines the prominent feature of DES and the benefits of heterogeneous catalysts, including ease of handling, separation, and recycling. Magnetic nanoparticles (MNPs) have recently been shown to be catalyst supports due to their easy preparation, low cost, high stability, high surface area, and simple recovery by magnetic forces.²⁴ Due to these unique properties, many MNPs supported catalysts have been employed in magnetic catalysts in organic syntheses,²³⁻³¹ biomass conversion,³² and biodiesel production.³³ Thus, we focused our attention on the combination of the attractive properties of MNPs and DESs to produce a novel catalyst which can be highly dispersed in the reaction mixture (like a homogenous catalyst) and easily separated by a magnet and reused (like a heterogeneous catalyst).³⁴

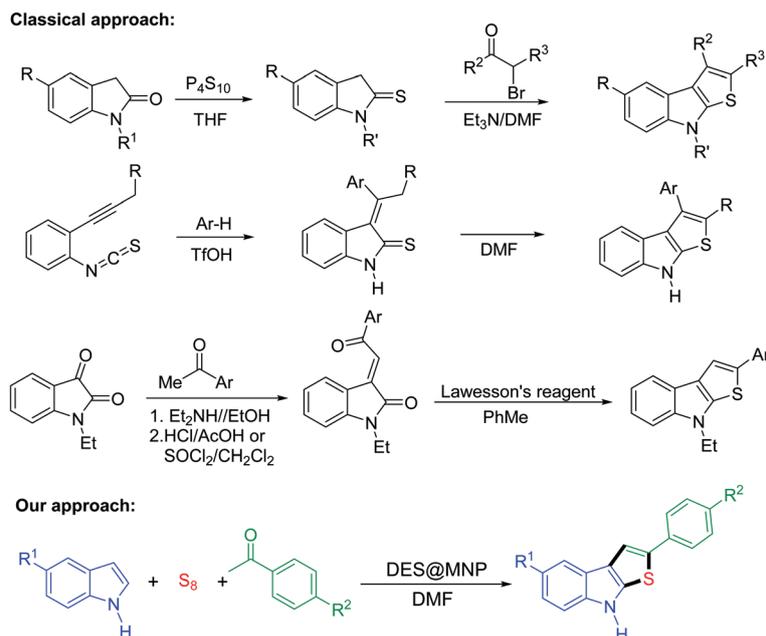
Because of the concerns about the development of a novel catalyst in green and sustainable chemistry, we report here the one-pot three-component synthesis of thieno[2,3-*b*]indoles from readily available indoles, acetophenones, and sulfur powder using a magnetically recyclable DES@MNP catalyst. The outstanding features of the current work are good to excellent yield, high regioselectivity, a broad substrate scope, and a recyclable catalyst.

^aDepartment of Organic Chemistry, Faculty of Chemistry, University of Science, Ho Chi Minh City 721337, Vietnam. E-mail: thphuong@hcmus.edu.vn; Tel: +84 903706762

^bVietnam National University, Ho Chi Minh City 721337, Vietnam

† Electronic supplementary information (ESI) available. See DOI: 10.1039/d0ra00773k



Scheme 1 Approaches to thieno[2,3-*b*]indoles.

2. Experimental

2.1. Preparation of the magnetic nanoparticle-supported Lewis acidic deep eutectic solvent (DES@MNP)

Synthesis of silica-coated Fe₃O₄ nanoparticles (Fe₃O₄@SiO₂@(CH₂)₃Cl). The magnetic nanoparticle-supported deep eutectic solvent catalyst was prepared following the procedure shown in Scheme 1. Firstly, the magnetic nanoparticle Fe₃O₄ was synthesized by the coprecipitation method reported in the previous literature. The mixture of FeCl₂ and FeCl₃ was dissolved in 200 mL water under ultrasonic irradiation. The potassium hydroxide solution (1.0 M) was added slowly until pH 12.0. After 120 min sonication, Fe₃O₄ was washed with water and ethanol to pH 7.5, separated by a magnetic and drying at 80 °C for 12 h. Fe₃O₄ was coated with a layer of silica by refluxing mixture Fe₃O₄ (1.0 g), 3.0 mL ammonia solution, and 3.0 mL of tetraethylorthosilicate (TEOS) in 100 mL ethanol for 8 h. Next, Fe₃O₄@SiO₂ (5.0 g) and (3-chloropropyl)triethoxysilane (4.820 g, 20.0 mmol) in 100 mL toluene were refluxed for 12 h. The prepared Fe₃O₄@SiO₂@(CH₂)₃Cl was separated using a permanent magnet, followed by washing with ethanol and dried at 80 °C for 6 h.

Synthesis of DES. The [Urea]₄[ZnCl₂] DES was synthesized by adding urea (1.2 g, 20.0 mmol) to zinc chloride (0.680 g, 5.0 mmol) and stirring to afford a transparent liquid at 100 °C for 30 min.

Synthesis of DES@MNP. Fe₃O₄@SiO₂@(CH₂)₃Cl (3.0 g) and DES [Urea]₄[ZnCl₂] (1.880 g, 5.0 mmol) were added round-bottomed flask and stirred vigorously at 100 °C for 18 h. After completion of the reaction, the DES@MNP was collected by an external magnet and washed with ethanol (3 × 20.0 mL) and dried under reduced pressure at 60 °C for 6 h.

2.2. General experimental procedure for the synthesis of 2-phenyl-8*H*-thieno[2,3-*b*]indole

Indole (1.0 mmol, 117 mg), acetophenone (2.0 mmol, 240 mg), and elemental sulfur power (5.0 mmol, 160 mg) were added into a round-bottom flask precharged with DES@MNP (10 mol%, 30 mg) in DMF (1.5 mL). The mixture reaction was then stirred at 140 °C for 12 h under the opening atmosphere. After completion of the reaction as checked by TLC, and the reaction mixture dissolved in 5.0 mL ethyl acetate. The DES@MNP catalyst was separated out by magnet and washed further with anhydrous acetone (3 × 5.0 mL) and ethanol (3 × 5.0 mL) for the next experiments. The reaction mixture was diluted with water (15.0 mL) and extracted with EtOAc (3 × 20.0 mL). The organic layer was dried over sodium sulfate, and the solvent was recovered under reduced pressure. Next, the residue was purified by column chromatography on silica gel using dichloromethane/petroleum ether = 2 : 3 as the eluent to afford the pure product. The product identity was confirmed by melting point, FT-IR, ¹H NMR, ¹³C NMR, and MS.

3. Results and discussion

3.1. Catalyst characterization

Initially, the DES@MNP was synthesized following the procedure presented in Scheme S1 (see ESI†).³⁵ First, Fe₃O₄ was prepared by a coprecipitation method similar to a previously reported literature, followed by a SiO₂ coating procedure afforded Fe₃O₄@SiO₂. Subsequently, Fe₃O₄@SiO₂ was functionalized using (3-chloropropyl)triethoxysilane by surface capping method to provide Fe₃O₄@SiO₂@(CH₂)₃Cl. The precursor [Urea]₄[ZnCl₂] deep eutectic solvent was prepared



by mixing of ZnCl_2 with urea (1 : 4 molar ratio) at $100\text{ }^\circ\text{C}$ until a clear solution was obtained. Finally, the treatment of DES with $\text{Fe}_3\text{O}_4@\text{SiO}_2@(\text{CH}_2)_3\text{Cl}$ in acetone at $100\text{ }^\circ\text{C}$ for 18 h provides the DES@MNP catalyst. The structure of DES@MNP was characterized by various techniques including FT-IR, Raman, TGA, SEM, TEM, XRD, VSM, EDX and ICP-MS (see the ESI†).

FT-IR spectra of $\text{Fe}_3\text{O}_4@\text{SiO}_2$ magnetic nanoparticles showed peaks at around 580, 1095, 1645, and 3419 cm^{-1} , which are attributed to the Fe–O–Fe, Si–O–Si and O–H groups (Fig. S1, ESI†). For DES-supported $\text{Fe}_3\text{O}_4@\text{SiO}_2$, additional peaks were observed for C=C, C–N, N–H, which confirms successful immobilization (Fig. S1, ESI†). The characteristic peaks of DES@MNP were at the same wavenumbers with the precursors, with a small shift due to the interaction between DES and the support. The Raman peak at 485 cm^{-1} can be interpreted as stretching vibration of the Zn–Cl bond (Fig. S2, ESI†). The stability of the DES@MNP was proved by the thermogravimetric (TG) analysis. DES@MNP showed a sharp decrease in weight (40%) was observed at $250\text{--}650\text{ }^\circ\text{C}$, which is attributed to the organic composition grafting. Thus, the DES@MNP is stable at around $250\text{ }^\circ\text{C}$ (Fig. S3, ESI†).

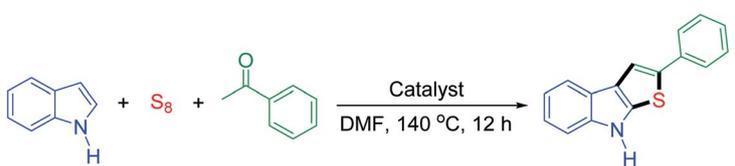
The magnetic properties of the DES@MNP and the neat Fe_3O_4 were recorded by vibrating sample magnetometry (Fig. S4, ESI†). The saturation magnetization of these samples was found to decrease from 69.2 to 30.8 emu g^{-1} due to the silica coating and the organic grafting on the surface of Fe_3O_4 . However, it was sufficient for the recovery of the DES@MNP from the reaction mixture using an external magnet. In

addition, the XRD pattern of the coated nanoparticles functionalized DES was carried out to confirm the crystalline nature. The XRD pattern (Fig. S5, ESI†) of the DES@MNP showed the diffraction peak positions and the relative intensities of the peaks, which matched well with the standard Fe_3O_4 sample (JCPDS file no. 19-0629). The SEM image of DES@MNP revealed that the catalyst was obtained in uniform nanometer-sized particles (Fig. S6a, ESI†). TEM image showed a well-defined core-shell structure with relatively nanosize distribution, a dark Fe_3O_4 core surrounded by a grey silica shell, and the average size of the obtained particles are 15–25 nm (Fig. S6b, ESI†). The chemical composition of DES@MNP was recorded by energy-dispersive X-ray spectroscopy (EDX), confirming the presence of C, O, N, Fe, Zn, Si, and Cl elements in the structure of the catalyst (Fig. S7, ESI†). The EDX revealed that the mass percent of Zn was 26.69%. This calculation was in good agreement with ICP-MS data for Zn analysis reporting the content of elemental Zn as 277 600 ppm, corresponding to Zn^{2+} loading of 4.10 mmol g^{-1} .

3.2. The activity of DES@MNP for the synthesis of 2-phenyl-8H-thieno[2,3-b]indole

In a preliminary survey of reactions, DES@MNP was found to be catalytically active in a one-pot multi-component reaction of indole, acetophenone, and sulfur as a model reaction (Table 1). Previously, this reaction can also be accomplished with acetic acid as a homogeneous catalyst.¹² However, the method suffers from difficulties in the recovery of the catalyst, moderate yield, high temperature for longer reaction time. These drawbacks

Table 1 Screening of different catalysts on the synthesis of 2-phenyl-8H-thieno[2,3-b]indole



Entry ^a	Type of catalysts	Catalyst (mol%)	Yield ^b (%)
1	Brønsted acids	H_2SO_4	72
2		HCl	70
3		H_3PO_4	67
4		CH_3COOH	75
5		CF_3COOH	80
6	Lewis acids	AlCl_3	73
7		FeCl_3	57
8		ZnCl_2	80
9		SnCl_4	65
10		CuCl_2	42
11	Control experiment	Fe_3O_4	—
12		$\text{Fe}_3\text{O}_4@\text{SiO}_2$	—
13		$\text{Fe}_3\text{O}_4@\text{SiO}_2@(\text{CH}_2)_3\text{Cl}$	—
14		$[\text{Urea}]_4[\text{ZnCl}_2]$	82
15		$\text{Fe}_3\text{O}_4@\text{SiO}_2@(\text{CH}_2)_3\text{--[Urea]}_4[\text{ZnCl}_2]$	87

^a Reaction conditions: acetophenone (2.0 mmol), indole (1.0 mmol), sulfur (5.0 mmol) and catalysis (10 mol%) were heated in DMF (1.5 mL) at $140\text{ }^\circ\text{C}$ for 12 h. ^b Yield of 2-phenyl-8H-thieno[2,3-b]indole was isolated by column chromatography (dichloromethane/petroleum ether).



Table 2 Substrate scope for the synthesis of 2-phenyl-8*H*-thieno[2,3-*b*]indole derivatives

$\text{Indole } 1 + \text{S}_8 + \text{Aromatic ketone } 2 \xrightarrow[\text{DMF, 140 }^\circ\text{C, 12 h}]{\text{DES@MNP}} \text{Product } 3$

Entry ^a	Indoles 1	Aromatic ketones 2	Products 3	Yield ^b (%)
1			3a	87
2			3b	81
3			3c	83
4			3d	77
5			3e	65
6			3f	59
7			3g	80
8			3h	77
9			3i	78
10			3j	80
11			3k	71
12			3l	75
13			3m	75
14			3n	78
15			3o	80
16			3p	79
17			3q	75



Table 2 (Contd.)

Entry ^a	Indoles 1	Aromatic ketones 2	Products 3	Yield ^b (%)
18			3r	78
19			3s	81
20			3t	89
21			3u	80
22			3v	81
23			3w	81
24			3x	87
25			3y	54

^a Reaction conditions: aromatic ketones (2.0 mmol), indoles (1.0 mmol), sulfur (5.0 mmol) and DES@MNP (10 mol%) were heated in DMF (1.5 mL) at 140 °C for 12 h. ^b Yield of 2-phenyl-8*H*-thieno[2,3-*b*]indole was isolated by column chromatography.

can be overcome by using DES@MNP as a magnetically recoverable catalyst. To our delight, the expected product was obtained in excellent yield (87%) at 140 °C for 12 h when DES@MNP was employed. The reactions were also tested using Brønsted acids as the catalysts at 140 °C for 12 h in DMF; the desired product was obtained in 67–72% yield (Table 1, entries 1–5). Next, several Lewis acids were screened in the model reaction (Table 1, entries 6–10). Among these Lewis acids, ZnCl₂ was shown to be a suitable catalyst that provided the desired product in 80% yield. At this point, we revealed that ZnCl₂ could play an essential role in promoting the condensation of indole, sulfur, and acetophenone and this is the reason we developed the new catalyst with ZnCl₂ active sites. Additionally, the control experiment in the absence of catalyst did not generate the expected product. The support Fe₃O₄, Fe₃O₄@SiO₂, and Fe₃O₄@SiO₂-(CH₂)₃Cl were also not reactive under these conditions (Table 1, entries 11–13). The typical [Urea]₄[ZnCl₂] DES provided the desired product in 82% yield, but the homogeneous DES was difficult to recover and reuse with the loss of the catalytic activity. The combination of nanoparticle and DES increased

the reaction yield as well as selectivity and made the recovery of the catalyst much more comfortable by using an external magnet. The reaction proceeds effectively because the nanocatalyst possesses a high surface area increasing the adsorption of reactants on the surface of the active Zn²⁺ sites on the DES@MNP. Interestingly, DES@MNP showed a higher yield in comparison with HCOOH catalyst (75% yield) which was reported by Ni and coworkers.¹²

The additive DMF plays a vital role in the yield and regioselectivity control for the thieno[2,3-*b*]indole synthesis.¹² Next, we turned our attention to the optimization of the additives to obtain the major corresponding regioselective product where the phenyl substituent was attached at the C-2 position in the thieno[2,3-*b*]indole. The results are presented in Table S1 (ESI[†]). Among these additives, DMF was the best additive under the current method. The reaction proceeds efficiently with DABCO or *N*-ethylmorpholine to provide the desired product in 75 and 81%, respectively.

When solid catalysts promote organic synthesis, the solvent plays a vital role in the reaction rate, depending on the nature of



these catalysts. The solvent effect on the catalytic activity of DES@MNP was further investigated, and the results are shown in Table S2 (ESI†). The three-component cyclization reaction was carried out at 140 °C, using acetophenone, indole, and sulfur molar ratio of 1 : 2 : 5, in the presence of 10 mol% DES@MNP catalyst and 0.3 mL DMF as an additive, in 1.5 mL organic solvent. A significant drop in reaction yield was observed for the other solvents, whereas a higher reaction activity was observed in DMF as a double role of additive and solvent. It was proposed that the DMF solvent could form the active precursor when interacted with sulfur and acetophenone, affording reaction rate enhancement (please see the proposed mechanism in Scheme S2†).

Next, the effect of reaction time and temperature were investigated (ESI, Table S3 and S4†). The three-component reaction was conducted in DMF for 12 h, using acetophenone : indole : sulfur molar ratio of 1 : 2 : 5 in the presence of 10 mol% of DES@MNP catalyst. It was shown that no expected product was detected for the reaction at 80 °C. The low yield of the product was obtained at 100 °C. Increasing the reaction

temperature to 140 °C led to a significantly improved reaction yield of 2-phenyl-8*H*-thieno[2,3-*b*]indole. The reaction time was also screened and the best yield was observed at 12 h (please see in ESI, Table S4†). The reactant molar ratio was investigated in the model reaction at 140 °C for 12 h. It was found that the reaction yield was significantly affected by the reactant molar ratio. The results revealed that the use of fewer than 4 equivalents of sulfur gave rise to the low yields of the desired product. However, when 4 or 5 equivalents of sulfur were employed, the yields dramatically increased.

To further define the scope and limitations of the present method, various indoles and acetophenones were examined. As shown in Table 2, the process was highly useful for the preparation of thieno[2,3-*b*]indole derivatives (Fig. 1). Acetophenone bearing various substituents at the *para* position could smoothly proceed under the present method provided the desired product in good to excellent yields. When acetophenone and acetophenone bearing electron-donating groups were employed to couple with indole, the desired product was obtained in higher yields than 4-fluoroacetophenone and 2-

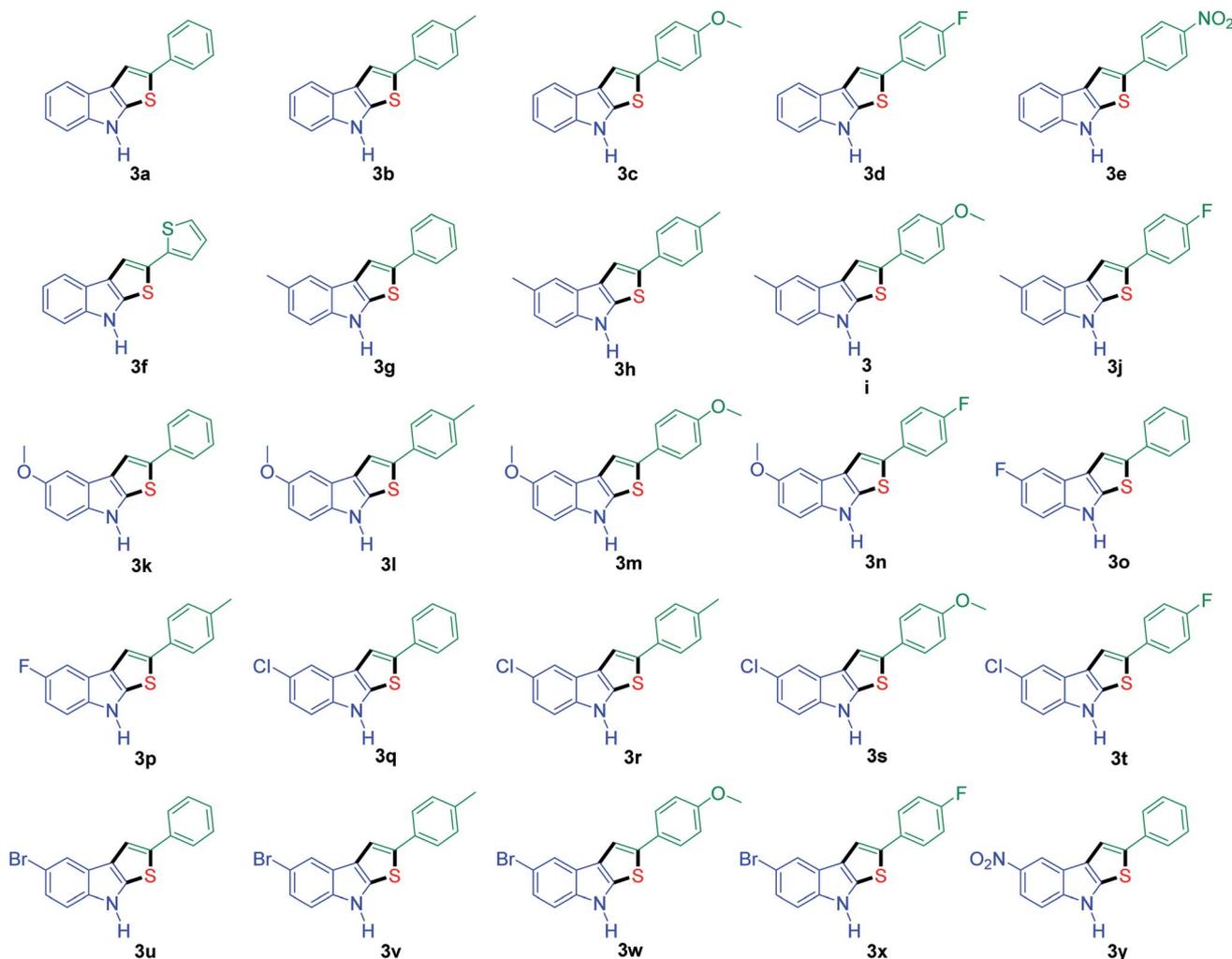


Fig. 1 The structure of various 2-phenyl-8*H*-thieno[2,3-*b*]indole compounds.



acetylthiophene. To our delight, good to excellent yields were observed when the methyl, methoxyl, halo, and nitro groups were located at the C-5 in the NH-unprotected indole ring. Indoles with halogens were all reactive, but 5-bromoindole produced the product in slightly higher yields than 5-fluoroindole and 5-chloroindole. 5-Nitroindole, which is known to deactivate many organic reactions in which it is used, was found to be a feasible reagent in our method producing 5-nitro-2-phenyl-8*H*-thieno[2,3-*b*]indole in 54% yield. It is noteworthy that these unprotected 1*H*-indoles worked well in this protocol to provide the desired product in high yields without the need for protection of the NH position when compared with previous literature.¹² Additionally, significant regioselectivities of products were observed in over 95% making the isolation of products more efficient. Thus, the novel DES@MNP is a highly active catalyst for the synthesis of thieno[2,3-*b*]indole derivatives.

Although several homogeneous catalysts in our research such as CF₃COOH, ZnCl₂, [Urea]₄[ZnCl₂] also exhibited high activity in the reaction (please see in Table 1). However, the main drawback is the difficulty in the recovery and reuse of these homogeneous catalysts. The prominent feature of the current method is the ease of separation of DES@MNP by an external magnet. The DES@MNP was recovered and reused over five consecutive runs, by repeatedly separating the DES@MNP catalyst from the reaction mixture by a magnet, washed with an organic solvent, dried under vacuum in 30 min, and then reused for next cycle. Interestingly, the reaction still afforded 83% yield of the desired product in the fifth run (Fig. 2). Remarkably, the structure of the recovered DES@MNP catalyst could be maintained under the current method, as confirmed by FT-IR (Fig. S8, ESI[†]).

The active sites on the heterogeneous catalyst can dissolve into the liquid phase during a reaction and act as a homogeneous catalyst. To confirm whether active zinc species dissolved from the DES@MNP promoted to the formation of the desired product (A), the leaching test was investigated. The model reaction was stopped after 8 h, separation of DES@MNP from the reaction mixture by a magnet. The

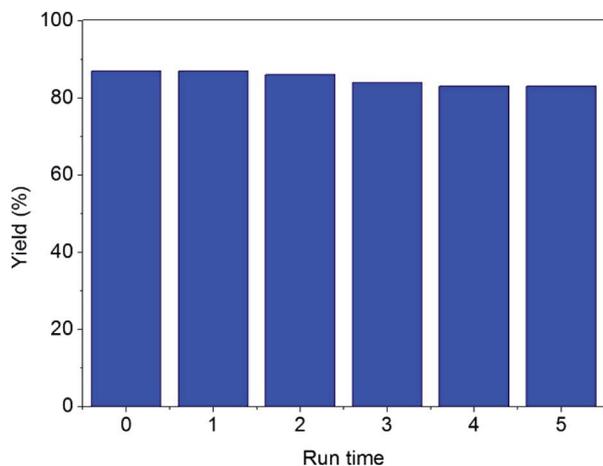


Fig. 2 Recycling experiments.

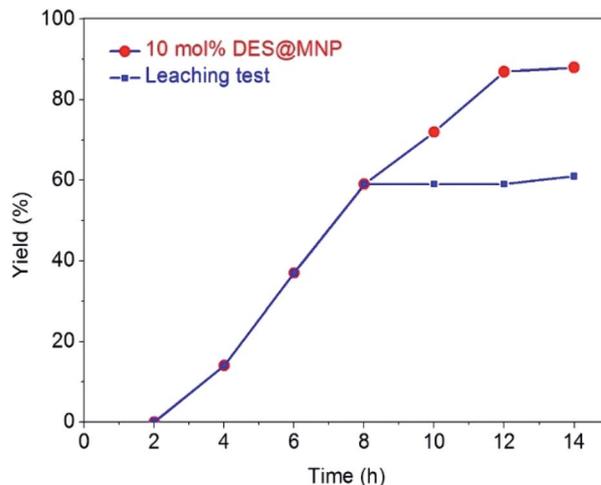


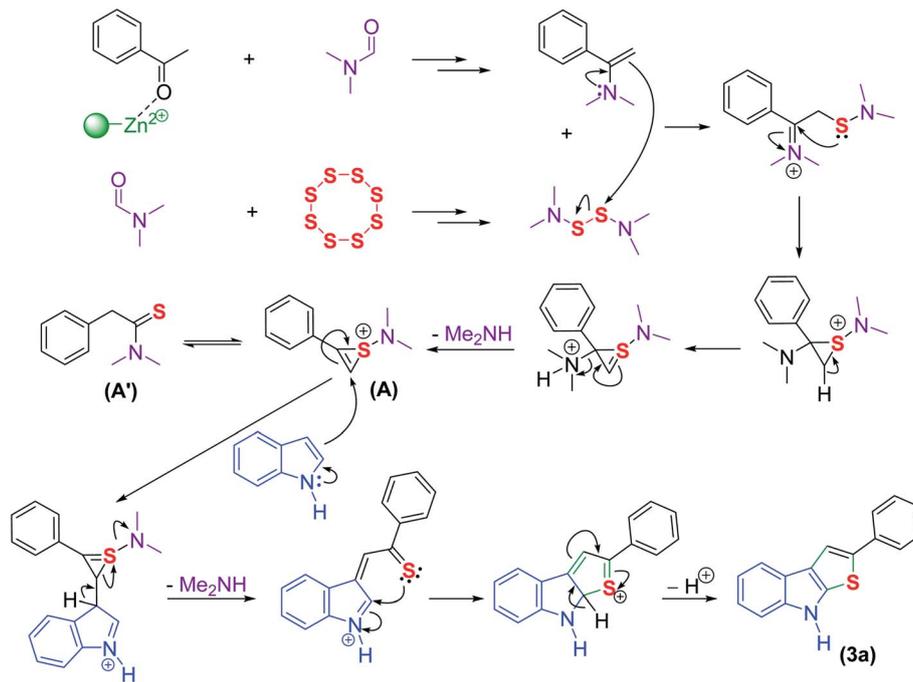
Fig. 3 Leaching test.

reaction mixture without catalyst was then stirred for further 8 h and analyzed by GC-MS. It was found that no additional desired product was obtained in the absence of DES@MNP catalyst (Fig. 3). Thus, the leaching of zinc from the catalyst to the solution was negligible. Additionally, the content of Zn species in reaction filtrate was recorded by ICPMS, and the result was measured less than 1 ppm of Zn. Obviously, there was no contribution from the leaching of active Zn species in the reaction solution, and the synthesis of thieno[2,3-*b*]indole could only occur in the presence of the DES@MNP.

3.3. Reaction mechanism research

To determine the mechanism insights for the synthesis of thieno[2,3-*b*]indole *via* a multicomponent reaction, some mechanistic experiments were carried out. According to the above control experiments and previous literature, a proposed mechanism was depicted in Scheme 2. Assisted by DES@MNP, the reaction proceeds *via* the initial formation of 1-(dimethylamino)-2-phenyl-1*H*-thiiren-1-ium (A), which is a resonance to form A' (Willgerodt–Kindler reaction). The intermediate (A) reacts with indole *via* a ring-opening addition, followed by ring closure and then eliminated one molecule of dimethylamine to afford the target product. To check the conformability of the plausible mechanism for the same reaction with the absence of indole catalyzed by DES@MNP, we also carried out the reaction of acetophenone, elementary sulfur, and DMF under optimized conditions. The (A') product was obtained in a yield of 65%. Next, the indole was then subjected to the reaction mixture and the desired product was achieved in 87% yield (ESI[†]). Thus, it is not doubtful that the multicomponent synthesis of thieno[2,3-*b*]indole reported herein must also undergo formation the precursor (A) and the DMF solvent plays an important role in the reaction mechanism. Although the reaction mechanism is not clear now, the protocol possesses attractive merits including simply and efficiently recyclable catalyst, wide scope of substrates, high yield, and high regioselectivity.





Scheme 2 Proposed reaction mechanism.

4. Conclusion

In conclusion, we have developed a catalyst for the synthesis of thieno[2,3-*b*]indole *via* the one-pot multicomponent reaction of indoles, acetophenones, and sulfur. The corresponding products are smoothly formed in high yields and regioselectivity under mild conditions. More importantly, we have described the first example of using DES@MNP-catalyzed the thieno[2,3-*b*]indole synthesis *via* multicomponent reaction with cheap and readily available reactants. DES@MNP was shown to be the extremely powerful catalyst to activate the reaction leading to total conversions even various substrates bearing electron-rich or electron-poor substituents. Further development and applications of multicomponent reactions of direct construction of thienoindole from elemental sulfur and indoles are currently under investigation in our laboratory. We hope that DES@MNP will find their applications in many organic syntheses, especially in green and sustainable syntheses of biologically active compounds.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This research is funded by Vietnam National Foundation for Science and Technology Development (NAFOSTED) under grant number 104.01-2019.26. We thank MSc Duy-Khiem Nguyen Chau (North Carolina State University, USA) and MSc Mai Hoang Ngoc Do (IPH-HCM) for their help.

References

- 1 J. Yang, Y. Cai, Y. Zhou, C. Zhang, P. Liang, B. Zhao, J. Shao, N. Fu, W. Huang and X. Dong, *Dyes Pigments*, 2017, **147**, 270–282.
- 2 K. Kanbe, M. Okamura, S. Hattori, H. Naganawa, M. Hamada, Y. Okami and T. Takeuchi, *Biosci., Biotechnol., Biochem.*, 2014, **57**, 632–635.
- 3 R. A. Irgashev, A. A. Karmatsky, S. A. Kozyukhin, V. K. Ivanov, A. Sadovnikov, V. V. Kozik, V. A. Grinberg, V. V. Emets, G. L. Rusinov and V. N. Charushin, *Synth. Met.*, 2015, **199**, 152–158.
- 4 F. Baert, C. Cabanetos, M. Allain, V. Silvestre, P. Leriche and P. Blanchard, *Org. Lett.*, 2016, **18**, 1582–1585.
- 5 H. Qi, Y. Zhang, Q. Tao, J. Chen, Z. Li, X. Xu, H. Tan, Q. Peng and W. Zhu, *Tetrahedron*, 2016, **72**, 7430–7437.
- 6 Y. Wen, W. Wu, Y. Li, Y. Li, T. Qin, Y. Tang, L. Wang and J. Zhang, *Org. Electron.*, 2016, **38**, 61–68.
- 7 Y. Wen, W. Wu, Y. Li, W. Zhang, Z. Zeng, L. Wang and J. Zhang, *J. Power Sources*, 2016, **326**, 193–202.
- 8 T. Otani, S. Kunimatsu, T. Takahashi, H. Nihei and T. Saito, *Tetrahedron Lett.*, 2009, **50**, 3853–3856.
- 9 R. A. Irgashev, A. A. Karmatsky, G. L. Rusinov and V. N. Charushin, *Beilstein J. Org. Chem.*, 2015, **11**, 1000–1007.
- 10 H. Z. Boeini, *Helv. Chim. Acta*, 2009, **92**, 1268–1272.
- 11 B. Li, P. Ni, H. Huang, F. Xiao and G.-J. Deng, *Adv. Synth. Catal.*, 2017, **359**, 4300–4304.
- 12 P. Ni, B. Li, H. Huang, F. Xiao and G.-J. Deng, *Green Chem.*, 2017, **19**, 5553–5558.
- 13 L. Wu, A. Mendoza-Garcia, Q. Li and S. Sun, *Chem. Rev.*, 2016, **116**, 10473–10512.



Paper

- 14 E. L. Smith, A. P. Abbott and K. S. Ryder, *Chem. Rev.*, 2014, **114**, 11060–11082.
- 15 Q. Zhang, K. De Oliveira Vigier, S. Royer and F. Jerome, *Chem. Soc. Rev.*, 2012, **41**, 7108–7146.
- 16 X.-d. Tang, Y.-f. Zhang and J.-j. Li, *Catal. Commun.*, 2015, **70**, 40–43.
- 17 Q. Wang, X. Yao, Y. Geng, Q. Zhou, X. Lu and S. Zhang, *Green Chem.*, 2015, **17**, 2473–2479.
- 18 C. Vidal, J. García-Álvarez, A. Hernán-Gómez, A. R. Kennedy and E. Hevia, *Angew. Chem., Int. Ed.*, 2016, **55**, 16145–16148.
- 19 S. B. Phadtare and G. S. Shankarling, *Green Chem.*, 2010, **12**, 458–462.
- 20 A. P. Abbott, D. Boothby, G. Capper, D. L. Davies and R. K. Rasheed, *J. Am. Chem. Soc.*, 2004, **126**, 9142–9147.
- 21 S. Khandelwal, Y. K. Tailor and M. Kumar, *J. Mol. Liq.*, 2016, **215**, 345–386.
- 22 F. Keshavarzipour and H. Tavakol, *J. Iran. Chem. Soc.*, 2015, **13**, 149–153.
- 23 D. Shahabi and H. Tavakol, *J. Mol. Liq.*, 2016, **220**, 324–328.
- 24 V. Polshettiwar, R. Luque, A. Fihri, H. Zhu, M. Bouhrara and J. M. Basset, *Chem. Rev.*, 2011, **111**, 3036–3075.
- 25 Y. Zhang and C. Xia, *Appl. Catal., A*, 2009, **366**, 141–147.
- 26 Q. Zhang, H. Su, J. Luo and Y. Wei, *Green Chem.*, 2012, **14**, 201–208.
- 27 Y. Kong, R. Tan, L. Zhao and D. Yin, *Green Chem.*, 2013, **15**, 2422–2433.
- 28 B. Karimi, F. Mansouri and H. Vali, *Green Chem.*, 2014, **16**, 2587–2596.
- 29 R. K. Sharma, S. Dutta, S. Sharma, R. Zboril, R. S. Varma and M. B. Gawande, *Green Chem.*, 2016, **18**, 3184–3209.
- 30 F. Keshavarzipour and H. Tavakol, *Appl. Organomet. Chem.*, 2017, **31**, e3682.
- 31 M. T. Jafari-Chermahini and H. Tavakol, *ChemistrySelect*, 2019, **4**, 1895–1902.
- 32 Q. Bao, K. Qiao, D. Tomida and C. Yokoyama, *Catal. Commun.*, 2008, **9**, 1383–1388.
- 33 Z. Wu, Z. Li, G. Wu, L. Wang, S. Lu, L. Wang, H. Wan and G. Guan, *Ind. Eng. Chem. Res.*, 2014, **53**, 3040–3046.
- 34 H. Tavakol and F. Keshavarzipour, *Appl. Organomet. Chem.*, 2017, **31**, e3811.
- 35 T. T. Nguyen, N.-P. Thi Le, T. T. Nguyen and P. H. Tran, *RSC Adv.*, 2019, **9**, 38148–38153.

