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REVIEW

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Recent advances in the synthesis of thienoindole analogs and their diverse applications

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Thiophene-fused heterocyclic organosulfur systems, especially the thieno[3,2-*b*]indole moiety have attracted significant attention because they show a wide spectrum of biological activities such as antituberculosis, antitumor, antifungal, antibacterial, and human 5-HT5A receptor binding inhibition. Moreover, they also find applications in material chemistry and chemical engineering. Thus, due to their intriguing properties and applications, researchers are continually attempting to create more effective and environment-friendly methods for their preparation. In this review, we present a complete assessment of the current advances in the field of thieno[3,2-*b*]indole synthesis.

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1 Introduction

Thiophene-fused heterocyclic organosulfur systems have piqued the interest of chemists around the world as they exhibit a diverse set of biological properties and are considered safe compounds for agricultural and pharmaceutical applications.¹

The thieno[3,2-*b*]indole moiety is specifically useful in the development of antituberculosis,² human 5-HT5A receptor binding inhibition, antitumor,³ anti-infective, anti-osteoar-thritis,⁴ antibacterial,⁵ and antifungal⁶ drugs and also potent in

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Moreover, it is an important type of π -extended electron-rich system, which can be used for designing molecules for photosensitive and photovoltaic devices. In the past few years, it has also been widely used in designing and engineering fused molecules for organic electronic application, which basically works *via* the electron push–pull mechanism. This moiety is present in several functionalized organic dyes^{7–9} such as MKZ-39 and DPP-r-TI, which is an effective photosensitizer for photothermal and photodynamic therapies and polymers^{10,11} such as PTITBT, PTTICN, PTTIF, and PTTI (Fig. 2).

According to the fusion of the thiophene ring on the indole ring, thienoindole can be categorized into different types. It is a tricyclic heterocyclic compound in which a thiophene ring is fused to an indole ring having one nitrogen atom. The fusion



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may occur in four different ways, *i.e.*, 8*H*-thieno[2,3-*b*]indole 5, 4*H*-thieno[3,2-*b*]indole 6, 4*H*-thieno[3,4-*b*]indole 7 and 6*H*-thieno[3,2-*e*]indole 8, resulting in four important types of thienoindoles (Fig. 3).

2 Synthesis of thienoindoles

In the past few decades, thiophene-fused indoles have gradually been established as a novel class of valuable compounds having intriguing chemical activities and distinct biological activities. Plant growth regulators are critically important for crop production in high yield and enhanced quality, and therefore many plant growth-regulating chemicals have been synthesized



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to yield good seedlings by promoting root elongation. Nowadays, synthetic chemicals are used at various stages of rice plant development. A streptomycete strain identified as Streptomyces albogriseolus MJ286-76F7 produces a novel active chemical named thienodolin, an alkaloid having a thienoindole skeleton, which exhibits growth-promoting and growth-inhibiting activities in rice seedlings. In 1950, P. A. S. Smith and co-workers reported the first synthesis of 4H-thieno[3,2-b]indole 6 from the diazotization of o-nitroaniline 9 and thiophene 10 via the formation of 2-(2-nitrophenyl)thiophene 11 (Scheme 1a). Later, in 1960, Kobayashi et al. synthesized thieno [2,3-b] indoles 5, starting from 3-(2-oxo-2-phenylethyl)indolin-2-one 12 and phosphorus pentasulfide 13 (Scheme 1b). In the early 90 s, Nakamura et al. structurally elucidated and Kanbe et al. isolated and characterized thienodolin by actively extracting thienodolin from a Streptomyces albogriseolus culture broth using ethyl acetate as the solvent followed by purification *via* preparative HPLC and silica gel column chromatography. It was found that when rice seedlings were treated with 1.2 \times 10 $^{-6}$ to 1.2 \times 10^{-5} M thienodolin, it exhibited growth-promoting activity, whereas 4.0×10^{-5} M thie nodolin showed inhibitory activity in rice seedlings.

2.1 Synthesis of thieno[2,3-*b*]indole by radical cyclization

Singh *et al.*¹² synthesized substituted thieno[2,3-*b*]indole **15** using the radical cyclization approach in 2011 (Scheme 2). However, although this is an effective approach for the synthesis of thieno[2,3-*b*]indole in high yield, the substrate, (*o*-bromoindolyl)acrylonitrile **14**, used is synthesized in many steps *via* the base-induced condensation of (*o*-bromoindolyl) acrylonitrile with various aryl/heteroaryldithioesters. Also, the ¹H NMR spectrum of the substrate showed that the substrate is



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produced as an inseparable mixture of (E)/(Z) isomers. Moreover, the push-pull nature of the double bond forced the substrate to undergo thermal (Z)/(E) isomerization.

Thieno[2,3-*b*]indole and its derivatives were also prepared *via* the nitrene-mediated Cadogan cyclization of 3-(*o*-nitrophenyl)thiophene, $AlCl_3$ -induced electrophilic recyclization of 2-(2-furyl)aryl isothiocyanates and oxidative cyclization of indolin-2-thiones. However, most of these methods are afflicted by drawbacks of multistep precursor synthesis, limited scope and generality. Accordingly, considering their applications in pharmaceuticals and materials science, more versatile and effective strategies for the synthesis of thiophene-fused heterocycles are needed.

2.2 Synthesis of 2-substituted thieno[2,3-*b*]indoles by Lawesson's reagent (LR)

Igrashev and co-workers reported the convenient, short and reliable synthesis of 2-substituted thieno[2,3-*b*]indoles from



Fig. 3 Types of thienoindoles.

readily available reagents involving the two-step reaction-aldolcrotonic condensation of the starting materials and treatment of the intermediate with Lawesson's reagent. The reaction of the intermediate with **LR** takes place in two steps. Initially, the ethylidene double bond of indolin-2-ones undergoes reduction, and then Paal–Knorr cyclization occurs to give the tricyclic product.

When isatins **16** are treated with methyl ketones **17** in mild base, *e.g.*, secondary and tertiary amines, aldol-type adduct **18** is formed, which undergoes dehydration under acidic conditions to form the crotonic condensation product 3-(2-oxo-2-(hetero) arylethylidene)indolin-2-one **19**. Compound **19** is more stable than compound **18** given that compound **19** is an unsaturated 1,4-diketone. The carbon–carbon double bond of **19** undergoes reduction in the presence of Na₂S₂O₄,¹³ H₂/Pd(C)¹⁴ or Me₃P–H₂O (ref. 15) (Scheme 3) to give indolin-2-one **20**. Compound **20**, which bears a 4-oxobutyramide fragment (1,4-dicarbonyl derivatives), undergoes Paal–Knorr reaction in the presence of thionation agents such as P₄S₁₀ or Lawesson's reagent and gets cyclized into thieno[2,3-*b*]indole **21**.

Depending on the reaction conditions, the Paal–Knorr reaction produces pyrroles, furans or thiophenes from 1,4diketones. Thiophene is obtained by using sulfurization agents such as phosphorus pentasulfide and Lawesson's reagent. **LR** is used as a thiation agent and is a powerful dehydrator, driving the reaction towards completion. This reagent has a fourmembered ring of alternating phosphorus and sulfur atoms.

Although this synthetic strategy appears to be appropriate, it has little preparative interest given that thienoindoles are obtained in low yields. For example, 2-methyl-8*H*-thieno[2,3-*b*] indole was obtained in 15% yield *via* this four-step pathway using unsubstituted isatin and acetone.

This procedure was further modified using path A to path D to enhance the overall yield of the desired product. Thieno[2,3-*b*]indole **21a** was prepared using 1-ethyl-isatin **16a** and acetophenone **17a** in the presence of base and ethanol *via* path **A** to path **D** (Scheme 4).



Scheme 1 (a) First synthesis of thieno[3,2-b]indole via diazotization of o-nitroaniline and thiophene. (b) First synthesis of thieno[2,3-b]indole from 3-phenacyloxindole and phosphorous pentasulfide.



2.2.1 Path A. Compound **18a** was refluxed with **LR** in toluene for 1 h and the target compound **21a** was obtained in very poor yield (10%).

2.2.2 Path B. This is the conventional path to get 19a *via* the dehydration of aldol adduct 18a. Further, the reduction of 19a generates indolin-2-ones 20a, which undergoes cyclization using LR to give substituted thienoindole 21a (25%).

2.2.3 Path C. This is a one-pot synthetic route, which involves the reaction of isatin 16a with (phenacylidene)triphenylphosphorane 22 to give intermediate 19a, which gets cyclized to give 54% of 21a. However, the limitation of this method is the use of phosphorane derivative 22 (formed by prefunctionalization of acetophenone 17a), which is very expensive.

2.2.4 Path D. This pathway involves the refluxing of intermediate 19a with LR in toluene for 1 h, giving 21a in 57% yield.

Lawesson's reagent first acts as the source of H_2S to reduce C=C in **19a**, and then acts as the thiation agent to give **21a** *via* Paal-Knorr reaction. Thus, the four-step procedure is reduced to a two-step procedure, leading to an overall good yield of the product. Hence, path D is the most effective, shortest and most

robust method for the synthesis of thienoindoles. In some specific cases, path C is also used as an alternative synthetic route.

Thieno[2,3-*b*]indoles containing electron-withdrawing groups such as 4-cyano or 2-nitro-phenyl substituent at the C-2 position were prepared in high yields of 92% and 82%, respectively (Fig. 4).

2.3 Synthesis of thienoindoles *via* Pd-catalyzed cross coupling reaction

2.3.1 Pd-Catalyzed cross-coupling reactions. Palladium can transfer two electrons and form complexes in the 0 and +2 oxidation state. According to Pauling's scale, Pd has an electronegativity of 2.2, which leads to the formation of relatively stable and non-polar Pd–C bond. Thus, Pd is extensively used in synthesis. Due to the capacity of Pd to interact with non-polar bonds, a heteroatom a lone pair of electrons can easily undergo oxidative addition, transmetalation and reductive elimination. Heck, Negishi and Suzuki pioneered the work on Pd-catalyzed cross-coupling reactions and were awarded the Nobel prize in 2010.

Scheme 2



Scheme 3 Synthesis of 2-substituted thieno[2,3-b]indole 21 via thionation of indolin-2-ones 19.

2.3.2 Suzuki–Miyaura cross-coupling reaction. The Suzuki–Miyaura reaction is a coupling reaction between aryl halides and organoborane reagents, including boranes, boronic acids and boronic esters. Organoboranes are non-toxic, air and moisture resistant and can be easily handled. The non-polar nature of the C–B bond (because of the relatively lower electronegativity of boron) makes it more stable than other metal-carbon bonds such as Li, Mg, Si, Al, Zr, Cu and Sn.

2.3.2.1 Role of base and solvent. In most organoboron compounds, the C–B bond is extremely covalent and the complex does not undergo transmetalation easily. Thus, the notable and significant role of the base such as K_2CO_3 , K_3PO_4 , Na_2CO_3 , NaOH and NaHCO₃ in the Suzuki–Miyaura reaction is to activate the organoboron derivative by making a hypervalent, anionic boron-"ate" complex, which promptly undergoes

transmetalation. In an alternative process, the base displaces the halide in the $[PdXR^2]$ complex to form the $[Pd(OtBu)R^2]$ complex (Scheme 5).

The activity and selectivity of the Suzuki–Miyaura reaction are influenced by the solvent such as PhMe, DMF, 1,4-dioxane, benzene, THF and CH₃CN. Moreover, a mixture of organic solvents and water can be utilized to increase the rate, selectivity and yield of the coupled product. In the current scenario of synthesis, a mixture of toluene and 1,4-dioxane is used as the organic solvent and water as the co-solvent.

2.3.3 Buchwald–Hartwig amination reaction. A Pd catalyzed cross-coupling reaction between amines and aryl halides forms the C–N bond. In the case of $P(t-Bu)_3$, a monodentate ligand, the active $Pd[P(t-Bu)_3]$ is formed. Imine is obtained as



Scheme 4 Various paths undertaken for enhancing the yield of thienoindoles.



Fig. 4 Thieno[2,3-b]indole containing electron-withdrawing groups.

a by-product due to the β -hydride elimination reaction, which occurs when the H-atom is at the α -position to the N-atom.

Toluene and 1,4-dioxane are frequently used as solvents given that they have a high boiling point and can solubilize most organic compounds. Generally, strong bases such as NaOtBu and KOtBu in toluene are used to increase the reaction rate and product yield.

2.3.4 Synthesis of thieno[3,2-*b*]**indoles and thieno**[3,4-*b*]**indoles.** Thieno[3,2-*b*]**indoles 6 and thieno**[3,4-*b*]**indoles 7a**

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Scheme 5 (a) Mechanism of Suzuki-Miyaura cross-coupling reaction; (b) Buchwald-Hartwig coupling reaction.

were firstly synthesized in 1982 *via* two methods (Scheme 6), as follows:

(a) Reaction of substituted indole with thioamide.¹⁶

(b) Reaction of 3-hydroxy thiophene with hydrazine.²

However, these methods were tedious, inflexible and no improvement was reported until 2000. Later, the development

of Pd cross-coupling chemistry facilitated the synthesis of thienoindoles and a palladium-catalyzed two step procedure has been developed, which involves a Suzuki reaction to form a C–C bond between benzene and thiophene, followed by ring closure. Ring closure reaction can be of three types, as follows:



Classical synthetic approaches for synthesizing 6 and 7a Scheme 6



7 Site-selective 2,3-Scheme Suzuki cross-coupling of dibromothiophene.

(a) nitrene insertion,¹⁷ (b) oxidative C-N coupling¹⁸ and (c) Cadogan cyclization.19

Thieno[3,2-b]indoles were synthesized efficiently via the siteselective Suzuki-Miyaura coupling of 2,3-dibromothiophene with 2-bromophenylboronic acid,20 and subsequent two-fold palladium catalyzed C-N coupling (Buchwald-Hartwig reaction). In the first step, 2,3-dibromothiophene 23 is converted to 3bromo-2-(2-bromophenyl)thiophene 24 in 82% yield (Scheme 7).

3-bromo-2-(2-bromophenyl)thiophene Further, 24 undergoes Pd-catalyzed two-fold C-N coupling with p-methyl aniline 28 to give 25a (Scheme 8). Four ligands were screened using NaOtBu in toluene and Pd₂(dba)₃ to enhance the yield of 25a and it was observed that the use of bidentate ligands dppf and (S)-BINAP gave 25a in 97% and 67% yield, respectively, whereas the bulky monodentate ligands SPhos and

Table 1 Results of ligand screening

S. No.	Ligand [mol%]	Yield (%)
1	$P(tBu)_3 \cdot HBF_4[10\%]$	86
2	dppf [5%]	97
3	SPhos [10%]	65
4	(S)-BINAP[5%]	67
3 4	SPhos [10%] (S)-BINAP[5%]	65 67

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 $P(tBu)_3 \cdot HBF_4$ gave 25a in 65% and 86% yield, respectively. However, the bidentate ligand DPPF was found to give the highest yield,²¹ i.e., 97% (Table 1).

Using these optimized conditions, compound 24 was treated with electron-rich amines, namely, 4-(methylthio)aniline and panisidine, which produced the corresponding thienoindoles 25b and 25c in excellent yields. When electron-poor, benzylic and aliphatic amines were used, 1,1'-bis(diphenylphosphino) ferrocene (dppf) resulted in poor yield of the products. Thus, dppf was replaced by (S)-BINAP, which gave the corresponding thienoindoles 25d and 25e in good yields. Treatment of 1-



Fia. 5 Yield of products having different substituents.



Scheme 8 Pd-Catalyzed C-N coupling of 3-bromo-2-(2-bromophenyl)thiophene 24 to give 25a.

naphthylamine gave the product 25f in a comparatively low yield (78%), whereas sterically hindered amine gave the product 25g in very good yield. Hence, the maximum yield was observed for electron-rich amines. In total, 20 substituted thieno[3,2-*b*] indoles were synthesized using this method, some of which are shown in Fig. 5.

Also, thieno[3,4-*b*]indoles were synthesized *via* the siteselective Suzuki cross-coupling reaction of 3,4-dibromothiophene **26** to form 3-bromo-4-(2-bromophenyl)thiophene **24** in 78% yield (Scheme 9).

Further, 3-bromo-4-(2-bromophenyl)thiophene 24 undergoes Pd-catalyzed two-fold C–N coupling with *p*-methyl aniline 28 to give 29a (Scheme 10).

Upon screening of various ligands, the monodentate ligand $P(tBu)_3$. HBF₄ was found to be the most effective, giving the corresponding thieno[3,4-*b*]indole **29a** in 95% yield (Table 2).



Scheme 10 Two-fold C–N coupling of compound 24.

Table 2	Table 2 Results of ligand screening			
S. No.	Ligand [mol%]	Yield (%)		
1	$P(tBu)_3 \cdot HBF_4[10\%]$	95		
2	dppf [5%]	43		
3	SPhos [10%]	65		
4	(S)-BINAP[5%]	36		

Table 3 Percentage yield of 31 and 32 using various R groups

R	31 (yield%)	32 (yield%)
(b) Cl	37	30
(c) F	43	33
(d) <i>t</i> -Bu	43	26

At last, the practicality of the strategy was examined for the preparation of 5-substituted thieno[3,2-*b*]indoles. The synthesis began with two successive site-selective Suzuki cross-coupling reactions of 2,3,5-tribromothiophene **30** utilising the earlier reported conditions. 2,3,5-Tribromothiophene **31a–d**, and later to 3-bromo-2-(2-bromophenyl)-5-arylthiophene **32a–d** (Scheme 11), giving product **33** in low yield. This is presumably because of the extra aryl group at position-5, which may affect the overall electronic nature²² of the molecule (Table 3).

Buchwald's biaryl ligand SPhos gave the final product **33a** in the highest yield, *i.e.*, 96% (Scheme 12), whereas ligands such as $(t-Bu)_3P \cdot HBF_4$, dppf and xantphos gave compound **33a** in 84%, 36% and 43% yield, respectively.

In conclusion, thieno[3,2-*b*]indoles and thieno[3,4-*b*]indoles have been synthesized *via* a new, more efficient and convenient synthetic methodology, namely, Buchwald–Hartwig cross-coupling. Also, the role of the ligand was found to be more crucial in the second step.

2.4 Synthesis of thieno[3,2-*b*]indole *via* Cadogan reductive cyclization

Dehaen *et al.*¹⁹ synthesized thieno[3,2-*b*]indole in two steps. The first step is the Suzuki–Miyaura coupling reaction between *o*-nitrophenyl boronic acid 34 and 2-bromothiophene 35. The recent literature revealed that arylboronic acids substituted with electron-withdrawing groups (here, nitro at the ortho position) undergo extensive deboronation under standard Suzuki–Miyaura conditions, which employ aq. Na₂CO₃ as the base. This premature destruction of the C–B bond causes low yields. Hence, to reduce or protect from proto-deboronation, Suzuki–Miyaura coupling reaction has been performed under microwave-enhanced conditions. The second step is the nitrene-mediated reductive cyclization of 2-(2-nitrophenyl) thiophene **36** under MW irradiation, which leads to a dramatic rate enhancement given that the usual method demands drastic conditions and long reaction time (Scheme 13).



Scheme 11 Coupling reaction of 2,3,5-tribromothiophene 30 to give compound 32.



Scheme 12 Synthesis of 5-substituted thieno[3,2-b]indole 33.



Scheme 13 Synthesis of 2-(2-nitrophenyl)thiophene 36 via Suzuki– Miyaura coupling reaction.

Further, a mixture of compound **36** and triethyl phosphite was suspended in a 10 mL sealed glass vial and irradiated with 300 W power at 210 °C. Bunyan and Cadogan demonstrated that aromatic *C*-nitroso-compounds are promptly deoxygenated by triethyl phosphite. Hence, deoxygenation of **36** led to the formation of a nitroso-compound, which readily underwent deoxygenation, resulting in the formation of an indole ring *via* a nitrene intermediate.²³ The reaction took 15 min to go to completion and acidic work-up removed the phosphate by-products and product **37** was obtained in good yield (Scheme 14).

Dehaen *et al.* used 2-nitro-phenylboronic acid given that it gives access to the biaryl compounds needed for the Cadogan

cyclization and also heterocyclic boronic acids are expensive and difficult to synthesize. Hence, it is a more versatile, efficient and economic method for synthesizing thieno[3,2-*b*]indoles.

2.5 Metal-free approach for synthesizing regioselective thieno[2,3-*b*]indole

Penghui *et al.*²⁴ described an effective metal-free approach for synthesizing substituted thieno[2,3-*b*]indole **40** and **41** with high regioselectivity and great functional group tolerance. In this approach, the cascade cyclization occurs *via* the acid-promoted annulation of ketone **38**, indole **39** and sulfur powder, where the solvent DMF and the additive control the regioselectivity of the reaction (Scheme 15).

2.5.1 Multi-component reactions. Multicomponent reactions are very efficient given that they are an easy and atomeconomic approach, which is highly advantageous compared to the conventional methods of synthesis. In this process, more than two starting materials combine to form a product, which contains almost all the employed atoms. MCRs can be divided into three types, *i.e.*, domino or cascade, sequential and consecutive MCR. Domino reactions take place without the



Scheme 14 Cadogan reductive cyclization to thieno[3,2-b]indole 37.



requirement of additional reagents or without the need for changing the reaction conditions, *i.e.*, everything needed for the reaction is there at the beginning. In the case of sequential MCR, the functionality necessary for the second step is created in the first step but an additional reagent must be added for the second reaction to occur. In consecutive MCR, the subsequent addition of reagent is done together with changing the reaction conditions from one step to another. Each type of MCR provides high structural and functional diversity.^{25,26}

Elemental sulfur was used as the source of sulfur given that it is abundant in nature, non-toxic and stable under normal conditions. Its low price and high purity make it a good choice. In recent years, elemental sulfur had found great applications in C–S bond-forming reactions. Most of the reported C–S bond formation reactions using elemental sulfur are catalyzed by transition metals. However, a few reactions are also available that do not need any transition metal.²⁷ Our reaction is among these types of reactions (Scheme 16).

1-Methyl-1*H*-indole **39**, acetophenone **38** and sulfur powder were reacted using different additives and solvents. The desired

product, *i.e.*, 3-phenylthieno[2,3-b]indole 40 was obtained when chlorobenzene was used as the solvent and 50 mol% HI as an additive at 130 °C. Further, the yield of product 40 was improved using PhCF₃ and anisole as additives and 1,4-dioxane as the solvent. The yield of 40 was further improved using PhCF₃/1,4dioxane, which was further greatly enhanced when 1 equivalent of L-phenylalanine was used together with PhCF3 and 1,4dioxane. 2-Phenylthieno[2,3-b]indole 41 was obtained when DMF was chosen as the solvent, whose yield was increased when acetic acid was used as the acid instead of HI. This is because when DMF was used as the solvent, the regioselectivity of the cyclization process switches as a result of the change in the polarity of solvent. This is a direct cyclization reaction. The reaction yield of 41 and the ratio of 41:40 further increased when the temperature was increased to 150 °C and the ratio of 39:38 changed to 1:2.3.

2.5.2 Substrate scope of 3-substituted thieno[2,3-*b*]indole synthesis. Product 43 was obtained in moderate to excellent yield (31–83%) using various substituted indoles (Scheme 17). 43a was formed in 83% when the substrate had a methoxy group as the substituent and 66% of 43b was formed when the substituent was the cyano group. Bulky substrates such as 1-(3-bromophenyl)ethanone and 1-(*o*-tolyl)ethanone gave 47% of 43c and 37% of 43d, respectively, due to steric hindrance (Fig. 6).

Further, the yield of product 44 was affected by the position of the functional group on the indole ring. When it was present on C-5, C-6 and C-7, then the yield of the product was good, whereas when it was present at the C-4 position, the yield decreased dramatically (Scheme 18).

2.5.3 Substrate scope of 2-substituted thieno[2,3-*b*]indole synthesis. **41a** was synthesized using different ketones in yield of up to 85%. The product was obtained in good yield when aromatic acetophenones were used, irrespective of the position of the functional group, whereas aliphatic ketones such as ⁱPrCOCH₃ **38a** gave 2-isopropyl-8-methyl-8*H*-thieno[2,3-*b*]indole **41a** in moderate yield (Scheme 19).



Scheme 16 Schematic representation of the synthesis of 3-phenylthieno[2,3-b]indole 40 and 2-phenylthieno[2,3-b]indole 41.



(+19 other examples)

Scheme 17 Synthesis of 3-substituted thieno[2,3-b]indole analogs 43 using different substituted indoles 42.





Scheme 18 Synthesis of 3-substituted thieno[2,3-*b*]indole analogs by varying the position of the functional group on the indole ring.



Scheme 19 Synthesis of 2-substituted thieno[2,3-*b*]indole analogs using different substituted ketones.



Scheme 20 Synthesis of 2-substituted thieno[2,3-*b*]indole analogs using different indoles.

Further, indoles bearing various substituents gave the product in a yield of up to 83%. When Me was at the C-6 or C-7 position of the indole moiety, the yield of product **45** decreased slightly to 67% and 73%, respectively, as shown in Scheme 20.

2.5.4 Scheme for preparing 3-phenylthieno[2,3-*b*]indole. When 1-methyl-indole reacted with acetophenone in the

absence of elemental sulfur, it gave 1-methyl-3-(1-phenylvinyl)-1*H*-indole **46** as an intermediate, which on reaction for 4 h with HI and sulfur powder in air at 130 °C, gave 3-phenylthieno[2,3-*b*] indole **40** in 95% yield. This reaction proceeded *via* a [4 + 1]-type synthetic route to give our product²⁸ **40** (Scheme 21).

2.5.5 Mechanism for 2-phenylthieno[2,3-b]indole synthesis. When acetophenone 38 was treated with sulfur powder in dry DMF under acidic conditions, *N*,*N*-dimethyl-2phenylethanethioamide 47 was obtained (Willgerodt–Kindler reaction),^{29,30} which resonates to form the intermediate *N*,*N*dimethyl-2-phenylethanethioamide 48. Moreover, 47 reacted with 1-methyl-1*H*-indole to give intermediate 49, which was converted to 2-phenylthieno[2,3-b]indole 41 under acidic conditions (Scheme 22).

2.6 Three-component metal free synthesis of thieno[2,3-*b*] indole in sulfur powder

Li *et al.*³¹ synthesized thieno[2,3-*b*]indole from indoles and alkenes or alkynes in the presence of sulfur powder and in the absence of metal. This is a simple and efficient method in which a Brønsted acid promotes the formation of substituted thieno [2,3-*b*]indole, where DMF is essential for converting the reactants into the fused products. Substituted 1-methylindole **39a**, substituted alkyne **50** and sulfur powder were treated in acid at 150 °C and in metal-free conditions using DMF as a solvent to obtain product **51**. However, in the absence of DMF, no product was obtained (Scheme 23).

Further, product **41** was obtained in 70% yield when AcOH was used as an acid. Moreover, inorganic acids such as hydrochloric acid acted as the most efficient acid and gave the product in 86% yield. Reducing either the reaction temperature or HCl concentration reduced the yield of the product (Scheme 24).

2.6.1 Substrate scope for the formation of substituted thieno[2,3-*b*]indole. The yield of the product depends on the substrate used (Scheme 25).

53 was obtained in 31% to 82% when substituted alkynes were used. *Para*-substituted phenylacetylene such as *p*-ethoxyphenylacetylene reacted with 1-methylindole **39** in the presence of sulfur powder to give the product 2-(4-ethoxyphenyl)-8-methyl-8*H*-thieno[2,3-*b*]indole **53a** in 82% yield. When 1-chloro-2-ethylbenzene was used as the substrate, 31% of 2-(2-chlorophenyl)-8-methyl-8*H*-thieno[2,3-*b*]indole **53b** was formed due to the steric hindrance caused by the substrate and when aliphatic alkynes such as 1-octyne were used as the substrate, 31% of 2-hexyl-8-methyl-8*H*-thieno[2,3-*b*]indole **53c** was formed because aliphatic alkynes do not favour annulation (Fig. 7).

Further, different substituted indoles **39a** were reacted with phenylacetylene **52** and sulfur powder and the product was



Scheme 21 Schematic representation of 3-phenylthieno[2,3-b]indole synthesis.



Scheme 22 Mechanism for the formation of 2-phenylthieno[2,3-*b*]indole.



Scheme 23 Strategy for the synthesis of indole-fused heterocycle.





Scheme 25 Synthesis of thieno[2,3-*b*]indole analogs by varying the substituents on the alkyne.

obtained in yields of up to 81%. When methyl was substituted at the C-5 position of 1-methyl-1*H*-indole, 61% of 5,8-dimethyl-2-phenyl-8*H*-thieno[2,3-*b*]indole was obtained, and when it is substituted at the C-7 position, 56% of 7,8-dimethyl-2-phenyl-8*H*-thieno[2,3-*b*]indole was formed (Scheme 26).

Moreover, 1-methylindole **39** was reacted with substituted styrene **55** and sulfur powder as the source of sulfur to get up to 76% of 2-substituted thieno[2,3-*b*]indole. Styrene derivatives





with either an electron-donating group such as methoxy and methyl, or halogen (F, Cl or Br) gave the product in good yield upon reacting with 1-methylindole. Overall, the reaction yield was influenced by the placement of the substituent on the phenyl motif of styrene. The use of 1-chloro-2-vinyl benzene as the substrate gave product **56** in 30% yield (Scheme 27).

The reactions of substituted indoles **39a** such as methyl, methoxy, bromo, fluoro and chloro gave the product in good yield independent of the position of the substituent at the C-5, C-6 or C-7 position. Even good yield of the product was obtained when unprotected indole was used as the substrate (54–79%, Scheme 28).

Plausible mechanism involves the reaction of phenylacetylene **52** with DMF and sulfur powder to form intermediate **47**, which resonates to form **48**. Further, **47** reacts with 1-methylindole **39** to form **49**, which ultimately gets converted to 8methyl-2-phenyl-8*H*-thieno[2,3-*b*]indole **41**, the desired product^{32,33} in moderate yield (Scheme 29).

2.7 Preparation of thieno[3,2-*b*]indole by halogen dance and sequential coupling reaction

Hayashi *et al.*³⁴ reported the synthesis of thieno[3,2-*b*]indole from 2,5 dibromothiophene 57 and lithium diisopropylamide (LDA), which resulted in the formation of transient thienyl anion species 58 *via* the mediated halogen dance reaction,³⁵ which led to the development of two chemical bonds in one pot, followed by Negishi coupling³⁶ and Suzuki–Miyaura coupling or Buchwald–Hartwig amination *via* tandem catalysis in the



Scheme 26 Extension of the substrate scope with respect to indoles.



Scheme 27 Synthesis of thieno[2,3-b]indoles using different substituted styrenes.



(+ 11 other examples)

Scheme 28 Reaction of different substituted indoles with styrene under the optimized reaction conditions.



Scheme 29 Proposed mechanism for the formation of thieno[2,3-b] indole derivative 41.

presence of 1,1'-bis(diphenylphosphino)ferrocene (dppf) and tri-*tert*-butylphosphine (tBu_3P) ligand to give thieno[3,2-b] indole.

The favourable environment for halogen dance with 2,5dibromothiophene 57 occurred by its deprotonation at the 4position with 1.3 eq. LDA at -78 °C for 5 min.³⁷ Moreover, the lithiated species gets rearranged to 5-lithio-4-brominated thiophene **58** at -78 °C *via* halogen dance, which on treatment with ZnCl₂ in tetramethylethylenediamine³⁸ (TMEDA 1.4 equiv.) at 0 °C formed thienylzinc species **58a**, which was further subjected to coupling conditions using the transition metal catalyst Pd(PPh₃)₄ and protected iodoaniline at 60 °C for 24 h, forming the coupled product **59** in 69% yield (Scheme 30).

Hayashi *et al.*³⁴ reported the synthetic approach of **59** towards ligand-controlled Suzuki–Miyaura coupling and Buchwald–Hartwig amination *via* assisted tandem catalytic transformation. Here, 2-(3,5-dibromothiophen-2-yl)-*N*-phenylaniline **59** gave π -conjugated thienoindole *via* Suzuki–Miyaura coupling^{39,40} and C–N bond formed by Pd-catalyzed reaction in the presence of aryl halide and amine groups in a stoichiometric amount of base (intramolecular Buchwald–Hartwig amination).^{41–43} Arylation at the α -position of thiophene resulted in the formation of 2-(3-bromo-5-(*p*-tolyl)thiophen-2-yl)-*N*-phenylaniline **60** and amination of the remaining β -bromo group with 2-aminophenyl group led to the formation of 4-phenyl-2-(*p*-tolyl)-4*H*-thieno[3,2-*b*]indole **61** (Scheme 31).

The above-mentioned reaction failed in the absence of additive, as reported by Okano and coworkers,⁴⁴ demonstrating that the additive plays an important role in significant amination for getting the desired product. Despite the longer response time of 24 h at 125 °C, additives such as NaOtBu were detected to be insufficient. The addition of $tBu_3P \cdot HBF_4$ (20 mol%) notably promoted amination to form 4-phenyl-2-(*p*-tolyl)-4*H*-thieno[3,2-*b*]indole **61** in 69% yield. A reduced amount of the phosphorus ligand in the additive led to a lower yield of **61**. This outcome suggests that tBu_3P , a monodentate ligand, needs to coordinate Pd with dppf, a bidentate ligand to form the efficient catalyst *in situ* for intramolecular amination (Scheme 32).

The one-pot reaction was also reported at 125 $^{\circ}$ C for 5 h by one-shot addition, which included all the required reagents for forming aryl substituted thieno[3,2-*b*]indole (Scheme 33).

A few of the limitations of the above-mentioned one-pot reaction were justified based on the variation in the Ar group at the second position of thieno[3,2-*b*]indole **61**. The above-











mentioned reaction conditions (Scheme 20) were not applicable for Ar = 4-nitrophenylboronic acid given that it has low solubility in 1,4-dioxane, and thus water as the co-solvent in a ratio



Scheme 33 Synthetic route for one-pot single-shot addition.

of 4 : 1 mainly helps to maximize the yield of 2-(2-nitrophenyl)-4-phenyl-4*H*-thieno[3,2-*b*]indole (Ar = 2-nitrophenyl) to 43%.

The Pd-dppf catalyst plays an important role in Suzuki-Miyaura coupling, which led to C–C bond formation at the α position of thiophene by releasing the bromo group. Further, ligands were exchanged from Pd-dppf to Pd- $(tBu_3P)_2$ and the reaction moves towards C–N bond formation in which the β bromo group undergoes oxidative addition of the Pd- $(tBu_3P)_2$ catalyst and reductive elimination to yield 4-phenyl-2-(aryl)-4*H*thieno[3,2-*b*]indole **61**. The maximum yield was obtained when tBu_3P was used as an additive for amination. The following sequential coupling reactions (Scheme 34) were reported as tandem catalytic pathways.³⁴

2.8 Synthesis of 2-(hetero)aryl-substituted thieno[3,2-*b*] indole *via* Fisher indolization

Irgashev *et al.*⁴⁵ reported the Fischer indolization^{46,47} methodology for the development of 2-(hetero)aryl-substituted thieno [3,2-b]indole **69** from 5-(hetero)arylthiophen-3(2*H*)-ones **68** and phenyl hydrazine **62**. The intermediate 5-(hetero)arylthiophen-



Scheme 34 Tandem catalytic pathway for sequential coupling reaction.

3(2H)-one was acquired in two steps, *i.e.*, the reaction of α bromocinnamates or their other hetero derivatives with methyl thioglycolate, followed by the base treatment of 5-(hetero)aryl-3hydroxysubstituted 2-thenoates to yield 2-(hetero)arylsubstituted thieno[3,2-*b*]indole.

Convenient and cheap synthetic approaches are urgent to meet the growing demand of hetero-arylated thieno[3,2-b] indoles, which are widely used in optoelectronic material engineering. In this context, the retro-synthetic approach towards the synthesis of thienoindoles is shown in the following scheme (Scheme 35).

An appropriate route for the synthesis of alkyl 3hydroxythiophene-2-carboxylates (2-thenoates) *via* Fiesselmann thiophene synthesis⁴⁸ was reported. It is a condensation reaction of 1,3-*C*,*C*-dielectrophilic substrates and alkyl thioglycolates upon treatment with base. 5-(Hetero)aryl-3-hydroxysubstituted 2-thenoates were synthesized in two steps starting from substrate 2-(hetero)aroylacetates⁴⁹ **64** or 3-(hetero)arylpropiolates⁵⁰ **65** or 2-bromo-3-(hetero)arylacrylates **66**. Moreover, 2-bromo-3-(hetero)arylacrylates **66** were easily available and also act as a 1,3-dielectrophilic three-carbon center for the Fiesselmann method. Further, **68** was formed by ester hydrolysis and *in situ* decarboxylation of alkyl 3-hydroxythiophene-2carboxylates **67** (Scheme 36).

The maximum yield of the product was reported when 2bromo-3-(hetero)arylacrylate **66** and methylthioglycolate were slowly added to NaOMe base (4 equiv.) in dry MeOH and refluxing the reaction mixture for 5 h resulted in the formation of methyl-5-aryl-3-hydroxythiophene-2-carboxylates **70** (Scheme 37).



Scheme 35 Retrosynthetic route of thieno[3,2-*b*]indole *via* Fisher indole synthesis.



Scheme 36 Synthetic route of 2-(hetero)aryl-substituted thieno[3,2-b]indoles via Fiesselmann and Fischer method.



Scheme 38 Synthetic route to methyl 4-hydroxy-5'-phenyl-[2,2'-bithiophene]-5-carboxylate 71.



Scheme 39 Synthetic approach towards the formation of 2-(5-phenylthiophen-2-yl)-4H-thieno[3,2-b]indole 73.

Methyl 5'-bromo-4-hydroxy-[2,2'-bithiophene]-5-carboxylate **70a** was synthesized from methyl-2-bromo-3-(5bromothiophen-2-yl)acrylate **66a**, which was in turn prepared *via* the esterification of 2-bromo-3-(5-bromothiophen-2-yl) acrylic acid⁵¹ (Scheme 38). The ester having a Br group was changed to 5-phenylthien-2-yl-connected derivative **71** *via* Suzuki–Miyaura reaction.

Further, saponification⁴⁵ of methyl-4-hydroxy-5'-phenyl-[2,2'bithiophene]-5-carboxylate **71** in the presence of NaOH and DMSO-H₂O for 1.5 h at 140 °C under an argon atmosphere yielded 5'-phenyl-[2,2'-bithiophen]-4(5*H*)-one **72** in 94–95% yield, which on further treatment with phenylhydrazine **62** in glacial CH₃COOH for 1 h at 120 °C, yielded 2-(5-phenylthiophen-2-yl)-4*H*-thieno[3,2-*b*]indole **73** in 58% yield (Scheme 39).

2.9 One-pot approach for synthesizing thieno[2,3-*b*]indole starting from indoline-2-thiones and nitroalkene-based adducts

Mane *et al.*⁵² reported a novel approach for the synthesis of thieno [2,3-*b*]indoles *via* the base-assisted [3 + 2]annulation of indoline-2thione involving nitroalkene adducts, *i.e.*, Morita–Baylis–Hillman (MBH)^{53,54} **76** and Rauhut–Currier (RC)⁵⁵ **78** adduct. The α -hydrazino- α , β -unsaturated nitroalkene intermediate was synthesized in excellent yields from imidazole or DMAP directed Morita–Baylis– Hillman reaction of nitroalkene and azodicarboxylates.⁵⁶ Nitroalkene-based Rauhut–Currier adducts⁵⁷ are effective synthons for synthesizing various functionalized heterocycles. Also, these synthons are great Michael acceptors, and also employed in Cascade Michael addition–cyclization reactions. The MBH adduct and RC adduct were synthesized, as shown in Scheme 40.



Scheme 40 Synthesis of nitroalkene-based MBH and RC adducts.



The reaction of 1-methylindoline-2-thione **79a** and α -hydrazinonitroalkenes **76** was performed by using potassium acetate/ acetic acid as an additive and CH₃CN as the solvent to obtain diisopropyl-1-(8-methyl-3-(*p*-tolyl)-8*H*-thieno[2,3-*b*]indol-2-yl) hydrazine-1,2-dicarboxylate **80** in 78% yield.

Various organic and inorganic bases were tested as substitutes for KOAc to get the product in higher yield. Among them, K_2CO_3 , NaOH, Cs_2CO_3 (inorganic bases) and Et_3N (organic base) in the presence of CH_3CN resulted in the formation of dihydrothienoindole **80a** (Scheme 28), which was finally converted to thieno[2,3-*b*]indole derivative **80** (Scheme 41). In the synthesis of thieno-[2,3-*b*]indole *via* MBH reaction, firstly proton abstraction at the C-3 position of indoline-2-thione **79** takes place to form an anion, which attacks hydrazinonitroalkene **76** *via* Michael addition reaction to form an intermediate that is further activated by the H-bonding property of acetic acid to form **81a**. Subsequently, the removal of the nitro group was facilitated by the lone pair of the hydrazine moiety and an acyl iminium-type intermediate **81b** was generated. Following this, intramolecular 5-*exo*-trig cyclization of **80** occurred to form dihydrothienoindole **81c**, which underwent aerial oxidation to give aromatized thienoindole **80**.



Scheme 42 Mechanism for the synthesis of thieno[2,3-b]indole.



Fig. 8 Substrate scope for synthesizing functionalized thieno[2,3-b] indoles using several substituted RC-adducts of nitroalkenes.



Fig. 9 Reaction scope with distinctively substituted indoline-2-thiones.

The reaction can also occur in a different manner, where thio-enolization of **81a** takes place initially, and then intramolecular 5-*exo*-tet cyclization and aerial oxidation takes place to form thieno[2,3-*b*]indole **80** (Scheme 42).

Mane *et al.*⁵² also reported an alternative efficient approach for the synthesis of thieno[2,3-*b*]indole derivatives. They treated substituted RC adduct **78** with N-protected indoline-2-thione **79** in K₂CO₃ (1.0 eq.) and CH₃CN : H₂O solvent mixture to get 4-(8methyl-3-phenyl-8*H*-thieno[2,3-*b*]indol-2-yl)butan-2-one **82** in 76% yield (Scheme 43). In addition, it was found that LiCl and H₂O as additive enhanced the product yield.

The presence of electron-donating groups on the aryl ring of RC-adduct **78** significantly reduced the product (**82b** and **82c**) yield with respect to **82a** (Scheme 40, Fig. 8).

(1) Presence of numerous electron-donating groups at several positions resulted in good yield of products **82d** and **82e**.

(2) Presence of weak electron-withdrawing group such as 4chloro-substituted RC-adduct **82f** led to moderate yield (63%) of the product (Fig. 9).



Scheme 44 Mechanism for the formation of thieno[2,3-b]indole via the RC adduct.



Scheme 45 Nucleophilic attack on 3-aminothiophene.

Several N-protecting groups such as ethyl and benzyl resulted in excellent yield of thieno[2,3-*b*]indole derivatives **82g** and **82k**, whereas groups such as H and *n*-propyl led to a much lower yield of derivatives **82h** and **82m** (Fig. 9).

The base-directed Michael addition of indoline-2-thione **79** to RC-adduct **78**-formed intermediate **83a** and intramolecular thio-Mannich-type reaction in 5-*exo*-trig fashion led to intermediate **83b**, and further removal of HNO and H_2O gave the aromatized product. The overall mechanism involved in the synthesis of the thienoindole derivative is depicted in Scheme 44.

2.10 Synthesis of thieno[3,2-*b*]indole derivative *via in situ* generation of 3-aminothiophene

It is a convenient method to synthesize thieno[3,2-*b*]indole having a thien-2-yl, aromatic or styryl group at the C-2 position.

This method uses 5-substituted-3-aminothiophene-2carboxylate and proceeds *via* Fischer indolization. It involves two steps, where the first step is the saponification of 3-aminoester with NaOH, and in the second step, this sodium salt reacts with arylhydrazine in glacial CH_3COOH . In the latter step, decarboxylation of 3-aminothiophene-2-carboxylic acid takes place to give 3-aminothiophene, which further reacts with arylhydrazines under acidic conditions to form arylhydrazone, ultimately undergoing Fischer indolization to give the desired product⁵⁸ 73.

Thiophene derivatives with an amino group at the C-3 or C-3 and C-4 position have found numerous applications in the formation of thiophene-fused N-heterocycles such as thienoindoles.⁵⁹

2.10.1 Similarity in the behaviour of 3-aminothiophene and thiophene-3(2*H*)-one. 3-Aminothiophene can be considered the synthetic counterpart of thiophene-3(2H)-one 85a for annulation reactions. It has been observed that the 3-aminothiophene moiety shows an enamine nature and its protonation takes place on the C-2 position, thereby forming thiophene-3(2H)-iminium cation⁶⁰ 84, which can further react with the nucleophile at the C-3 position to give 85 (Scheme 45).

2.10.2 Strategy for preparing thieno[3,2-*b*]indole derivative. Igrashev *et al.* described an efficient method for



Scheme 46 Synthesis of thieno[3,2-*b*]indole from 5(hetero)arylthiophene-3(2*H*)-ones.



Scheme 47 Synthesis of thieno[3,2-b]indole from 5-substituted-methyl-3-aminothiophene-2-carboxylates.



Scheme 48 Formation of 5-substituted-3-aminothiophene-2-methylcarboxylate from methyl ketone.





Scheme 49 Synthesis of 2-phenylthieno[3,2-b]indole.



Scheme 50 Strategy for preparing 2-substituted-3-phenyl-8*H*-thieno[2,3-*b*]indole 99.

synthesizing 2-(hetero)aryl-substituted thieno[3,2-*b*]indole **73** by Fischer indolization of arylhydrazines and 5-(hetero)arylthiophene-3(2*H*)-ones **85**, which is obtained from 5-(hetero)aryl-3hydroxythiophene-2-carboxylates **70** (ref. 45) (Scheme 46).

Later, they described another method for the synthesis of 2substituted thieno[3,2-*b*]indoles starting from 5-substitutedmethyl-3-aminothiophene-2-carboxylates **87** *via* the *in situ* generation of 3-aminothiophene, which further participates in Fischer indolization with arylhydrazines (Scheme 47). Fig. 10 Analogs of thieno[2,3-b]indole.

3-Aminothiophene-2-carboxylate **87** was synthesized in two steps. Initially, methyl ketone **86** (R = aryl, styryl or thien-2-yl) was treated with Vilsmeier reagent (POCl₃–DMF complex) and NH₂OH·HCl at 30–60 °C to give 3-substituted-3-chloroacrylonitrile,^{61–64} which then reacted with methyl thioglycolate in NaOMe in CH₃OH solution⁶⁵ to form the product following Fiesselmann method for making the thiophene ring⁶⁶ (Scheme 48).

In the earlier method for the preparation of thieno[3,2-b] indole, DMSO was used for the saponification of 3-



Scheme 51 Plausible mechanism for the formation of the product.



hydroxythiophene-2-carboxylate. However, the problem was the high temperature required for the reaction due to the initial occurrence of sodium-2-(methoxycarbonyl)thiophen-3-olate. Therefore, the direct conversion of 3-aminothiophene-2carboxylate was carried out using NaOH in aqueous i-PrOH, without the isolation of the intermediate formed, 3-aminothiophene.



Scheme 53 Mechanism for the synthesis of thieno[2,3-b]indole via nanoparticles.

HOOC

n-C₄H

107



Fig. 12 Examples of thienoindole analogs having biological activity.



Fig. 13 Thienodolin 103, a natural derivative of thieno[2,3-*b*]indole.

When 3-aminothiophene-2-carboxylates were substituted at the C-5 position by either aromatic or thien-2-yl groups, 55–77% of 2-substituted thieno[3,2-*b*]indole **73** was obtained (Scheme 49).

2.11 Metal-free synthesis of thieno[2,3-*b*]indoles using elemental sulfur

Thieno[2,3-*b*]indoles have been synthesized *via* a base-assisted metal-free approach using cheap and readily available elemental sulfur.⁶⁷ The development of attractive and valuable routes for forming carbonyl group-containing thieno[2,3-*b*] indoles represents several challenges. Here, β -indolyl ketone derivatives were used as effective and easily accessible substrates for the synthesis of carbonyl group-containing thieno [2,3-*b*]indoles through a C=S bond formation reaction.

3-(1*H*-indol-3-yl)-3-phenyl-1-(*o*-tolyl)-propan-1-one **94** treated with elemental sulfur and base NaO*t*Bu using anhydrous DMSO as the solvent at 120 °C under N₂ for 24 h gave product **99** in 97% yield. Moreover, other organic bases, *e.g.*, DBU, gave the product in a minute amount, whereas DABCO yielded the final

product in 90%. The quantity of elemental sulfur did not affect the yield of the product⁶⁸ (Scheme 50).

Fig. 15 Thienoindole-based compounds having electrical activities in

HOOC

106

push-pull dyes.

Substrate 3-(1*H*-indol-3-yl)-3-phenyl-1-(*o*-tolyl)-propan-1-one **94** was converted to intermediate **95** and remained in proper tautomeric equilibrium with **96**. Further, the base abstracted the proton and carbon at the α -position of the CO group was electrophilically attacked by the elemental sulfur, giving intermediate **97**. Later, intramolecular nucleophilic cyclization of **97** gave **98** by eliminating elemental sulfur (S_{*n*-1}) and oxidative aromatization of elemental sulfur formed 2-substituted-3phenyl-8*H*-thieno[2,3-*b*]indole **99** (Scheme 51). Hence, a novel synthetic route to poly-substituted thienoindoles was achieved, where S₈ promoted C=S bond formation to form the desired product.

Moreover, the yield of the product decreased when an electron-withdrawing group was substituted at the *p*-position of an extra benzene ring (**99a**, Fig. 10) and increased in the case of an electron-donating group (**99b**, Fig. 10).

Furthermore, the presence of methyl, chloride, phenyl and nitrile groups at the *p*-position of the aromatic ring substituted at the C-3 position of the thienoindole derivative resulted in



104 Donor- acceptor π conjugated polymer





105 Organic Dye MKZ-40



Scheme 54 Synthetic approach of push-pull molecules having thieno[2,3-b]indole.



Fig. 16 Diagram showing electron-donating and electron-with-drawing parts of push-pull $D-\pi-A$ molecules.

a high yield of the corresponding derivative products **99c–99f** (Fig. 11).

2.12 Synthesis of thieno[2,3-*b*]indoles *via* green approach using magnetic nanoparticle-supported [urea]₄[ZnCl₂] deep eutectic solvent (DES@MNP)

Nguyen *et al.*⁶⁹ reported a green approach for the one-pot threecomponent synthesis of functionalized thieno[2,3-*b*]indoles *via* the use of a magnetic nanoparticle-supported $[urea]_4 [ZnCl_2]$ deep eutectic solvent.

The reaction was conducted in the presence of economical and widely accessible reagents, *i.e.*, sulfur, acetophenone **38** and indole **39a**, using the magnetic recyclable nanoparticle DES@MNP catalyst in *N*,*N*-dimethylformamide at 140 °C for 12 h. The deep eutectic solvent was prepared with 1.2 g urea in 0.68 g zinc chloride at 100 °C and further mixed with silicacoated Fe₃O₄ nanoparticles at 100 °C for 18 h and dried under low pressure at 60 °C for 6 h to form the DES@MNP catalyst. The deep eutectic solvent-covered nanoparticles were beneficial for easy handling, separation and recycling. Moreover, the magnetic nanoparticles have the advantages of easy preparation, high stability, low cost, availability, high surface area and easy separation by a magnet for reuse (Scheme 52).

The proposed mechanism for the synthesis of thieno[2,3-*b*] indole from acetophenone **38**, *N*,*N*-dimethylformamide and sulfur aided by the DES@MNP catalyst suggests that the reaction takes place through the formation of 1-(dimethylamino)-2-phenyl-1*H*-thiiren-1-ium **47**, which further forms **47a** (65%). The resulting intermediate **47** further reacts with indole **39a** *via*



Scheme 37 Synthetic route for methyl 5-aryl-3-hydroxythiophene-2-carboxylates.



Scheme 43 Synthesis of thieno[2,3-b]indole via RC adduct.

a ring-opening addition mechanism, ring closure and elimination of dimethyl-amine to form 47 in 87% yield. Furthermore, indole reacts with the reaction mixture, forming 87% product. Moreover, the proposed research led to an efficient protocol having merits, which include simple and efficient recyclable heterogeneous catalyst, vast substrate scope and regioselective product in high yield (Scheme 53).

3 Applications of thienoindoles

3.1 Biological activities of thienoindoles

Thienoindole derivatives show a broad spectrum of biological activities such as anti-bacterial, anti-inflammatory, antiallergic, anti-viral, anti-tuberculosis activities **100**, 5-HT5A receptor binding inhibition **101**, antitumor and anti-infective activities **102** (Fig. 12).

The alkaloid thienodolin⁷⁰ **103** is a natural derivative of thieno[2,3-*b*]indole obtained by the fermentation mixture of *Streptomyces albogriseolus* (Fig. 13). Kanbe *et al.* characterized its activity for plant growth regulation. Furthermore, some thienoindoles are used to treat diseases of the central nervous system and some are potential inhibitors of acetylcholine esterase and butyrylcholine esterase.

3.2 Chemical activities of thienoindoles

Besides their therapeutic properties, thienoindoles are also used for designing molecules of photosensitive and photovoltaic devices because of their π -extended conjugation from electron-rich systems. They are reported to be effective photosensitizers for photothermal and photodynamic therapies and polymers such as PTTICN, PTTIF, and PTTI. Moreover, TI-DTBT3 **104** is a donor–acceptor π -conjugated polymer with high charge carrier mobility.⁷¹ This moiety is interestingly available in several organic dyes such as MKZ-40 **105** and DPP-r-TI. They are also used as precursors of polymers used in solar cell applications (Fig. 14).

3.3 Electrical activities of thienoindoles

Derivatives of thieno[2,3-*b*]indole are used to design photo- and electroactive compounds, which have been recently assessed in dye-sensitized solar cells (DSSCs). Push–pull dyes IK-1,2 *viz.*, **106** and **107** have been recently reported to be synthesized as a donor part of DSSCs⁷² (Fig. 15).

N-Methyl **109a**- or *N*-phenyl **109b**-substituted thieno[2,3-*b*] indoles are used as the electron donor part of push-pull $D-\pi$ -A

molecules, whereas 2,2-dicyanovinylmethyl (DCV) **115** and (1-(dicyanomethylene)-3-oxo-1-inden-2-ylidene)-methyl (DCI) **114** are used as the electron acceptor⁷³ (Scheme 54 and Fig. 16).

MTI-DCV **113** is the smallest push-pull molecule of the series. It has a high absorption coefficient, high thermal stability, good hole transport properties and good absorption in the visible region, and because of all these properties, a bilayer solar cell mostly composed of MTI-DCV showed a power conversion efficiency of more than 1%.

4 Conclusions

Thieno[2,3-b]indole and thieno[3,2-b]indole molecules have been extensively studied because of their wide range of acceptable biological and pharmaceutical applications and other characteristic uses such as in photothermal and organic photovoltaic cells (OPV), making them valuable heterocycles in synthetic organic chemistry, materials chemistry and chemical engineering. Over time, researchers have overcome multiple problems associated with their synthesis such as tedious product separation, problematic catalyst recovery and need for large stoichiometric amounts of solvent. Moreover, the use of highly volatile, toxic and explosive substrates, solvents, and additives for the preparation of thiophene-fused indoles limit their widespread application. Some synthetic protocols also use toxic and corrosive bases such as DABCO, DBU, and LDA and hazardous solvents such as 1,4-dioxane and DMF for the synthesis of thienoindoles. Moreover, a few of the reported methods for their synthesis require the use of functionalized furans, indoles, thiophenes, etc. as precursors, which are synthesized via multiple steps, making them inefficient. Thus, to overcome the aforementioned issues, newer and convenient synthetic strategies need to be developed, which involve environment-friendly solvents such as water, ethanol, and PEG and reduction in the use of workup solvents. Further, the electronic push-pull mechanism and extended conjugation indicate that thieno[2,3-b]indole-based polymers and dyes that display a large variation in properties are still to be discovered. Finally, the future prospects in the arena of thieno[2,3-b]indoles synthesis depends on the progress of competent synthetic procedures that can solve the above-mentioned concerns, while keeping environmental-friendliness a priority.

Conflicts of interest

There are no conflicts to declare.

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References

- 1 H. Jiang, J. Sun and J. Zhang, A Review on Synthesis of Carbazole-based Chromophores as Organic Light-emitting Materials, 2012, vol. 16.
- 2 A. N. Grinev, Y. I. Trofimkin, E. V Lomanova, G. N. Pershin,
 L. M. Polukhina, I. S. Nikolaeva, T. V Pushkina, L. N. Filitis,
 E. A. Golovanova and O. V Okinshevich, *Pharm. Chem. J.*, 1982, 16, 827–830.
- 3 S. A. Al-Trawneh, J. A. Zahra, M. R. Kamal, M. M. El-Abadelah, F. Zani, M. Incerti, A. Cavazzoni, R. R. Alfieri, P. G. Petronini and P. Vicini, *Bioorg. Med. Chem.*, 2010, 18, 5873–5884.
- 4 M. Shiozaki, K. Maeda, T. Miura, M. Kotoku, T. Yamasaki, I. Matsuda, K. Aoki, K. Yasue, H. Imai and M. Ubukata, *J. Med. Chem.*, 2011, **54**, 2839–2863.
- 5 S. A. Al-Trawneh, M. M. El-Abadelah, J. A. Zahra, S. A. Al-Taweel, F. Zani, M. Incerti, A. Cavazzoni and P. Vicini, *Bioorg. Med. Chem.*, 2011, **19**, 2541–2548.
- 6 M. S. C. Pedras and M. Suchy, *Bioorg. Med. Chem.*, 2006, 14, 714–723.
- 7 X.-H. Zhang, Y. Cui, R. Katoh, N. Koumura and K. Hara, J. Phys. Chem. C, 2010, **114**, 18283–18290.
- 8 Z.-S. Huang, T. Hua, J. Tian, L. Wang, H. Meier and D. Cao, *Dyes Pigm.*, 2016, **125**, 229–240.
- 9 Y. K. Eom, S. H. Kang, I. T. Choi, Y. Yoo, J. Kim and H. K. Kim, *J. Mater. Chem. A.*, 2017, 5, 2297–2308.
- 10 J. Kim, J. Lee, S. Chae, J. Y. Shim, D. Y. Lee, I. Kim, H. J. Kim, S. H. Park and H. Suh, *Polymer*, 2016, **83**, 50–58.
- 11 I. Jeong, S. Chae, A. Yi, J. Kim, H. H. Chun, J. H. Cho, H. J. Kim and H. Suh, *Polymer*, 2017, **109**, 115–125.
- 12 P. P. Singh, A. K. Yadav, H. Ila and H. Junjappa, *Chem.–Eur. J.*, 2011, **2011**, 4001–4007.
- 13 E. M. Beccalli, A. Marchesini and T. Pilati, J. Chem. Soc., Perkin Trans. 1, 1994, 579–587.
- 14 K. Albertshofer, B. Tan and C. F. Barbas III, *Org. Lett.*, 2012, 14, 1834–1837.
- 15 S.-H. Cao, X.-C. Zhang, Y. Wei and M. Shi, *Eur. J. Org. Chem.*, 2011, 2668–2672.
- 16 A. Shafiee and S. Sattari, *J. Heterocycl. Chem.*, 1982, **19**, 227–231.
- 17 P. A. S. Smith and J. H. Boyer, J. Am. Chem. Soc., 1951, 73, 2626–2629.
- 18 K. Takamatsu, K. Hirano, T. Satoh and M. Miura, *Org. Lett.*, 2014, **16**, 2892–2895.
- 19 P. Appukkuttan, E. Van der Eycken and W. Dehaen, *Synlett*, 2005, **01**, 127–133.
- 20 T. Q. Hung, T. T. Dang, A. Villinger, T. Van Sung and P. Langer, *Org. Biomol. Chem.*, 2012, **10**, 9041–9044.

- K. Nozaki, K. Takahashi, K. Nakano, T. Hiyama, H. Tang, M. Fujiki, S. Yamaguchi and K. Tamao, *Angew. Chem.*, 2003, 115, 2097–2099.
- 22 S.-M. T. Toguem, A. Villinger and P. Langer, *Synlett*, 2009, **20**, 3311–3314.
- 23 J. I. G. Cadogan, M. Cameron-Wood, R. K. Mackie and R. J. G. Searle, *J. Chem. Soc.*, 1965, 4831–4837.
- 24 P. Ni, B. Li, H. Huang, F. Xiao and G.-J. Deng, *Green Chem.*, 2017, **19**, 5553–5558.
- 25 V. Estevez, M. Villacampa and J. C. Menendez, *Chem. Soc. Rev.*, 2010, **39**, 4402–4421.
- 26 H. Y. Cho and J. P. Morken, *Chem. Soc. Rev.*, 2014, **43**, 4368–4380.
- 27 Z.-Y. Gu, J.-J. Cao, S.-Y. Wang and S.-J. Ji, *Chem. Sci.*, 2016, 7, 4067–4072.
- 28 Y. Kasano, A. Okada, D. Hiratsuka, Y. Oderaotoshi, S. Minakata and M. Komatsu, *Tetrahedron*, 2006, 62, 537– 542.
- 29 C. Willgerodt, Ber. Dtsch. Chem. Ges., 1887, 20, 2467-2470.
- 30 S. R. Pathipati, V. Singh, L. Eriksson and N. Selander, Org. Lett., 2015, 17, 4506–4509.
- 31 B. Li, P. Ni, H. Huang, F. Xiao and G. J. Deng, *Adv. Synth. Catal.*, 2017, **359**, 4300–4304.
- 32 S. Chen, Y. Li, P. Ni, B. Yang, H. Huang and G.-J. Deng, *J. Org. Chem.*, 2017, **82**, 2935–2942.
- 33 B. P. Mundy, M. G. Ellerd and F. G. Favaloro Jr, *Name reactions and reagents in organic synthesis*, John Wiley & Sons. 2005.
- 34 Y. Hayashi, K. Okano and A. Mori, Org. Lett., 2018, 20, 958– 961.
- 35 K. Shono, Y. Sumino, S. Tanaka, S. Tamba and A. Mori, *Org. Chem. Front.*, 2014, **1**, 678–682.
- 36 E. Negishi, Angew. Chem., Int. Ed., 2011, 50, 6738-6764.
- 37 R. Levine and W. C. Fernelius, Chem. Rev., 1954, 54, 449-573.
- 38 K. Snégaroff, S. Komagawa, F. Chevallier, P. C. Gros, S. Golhen, T. Roisnel, M. Uchiyama and F. Mongin, *Chem.– Eur J.*, 2010, **16**, 8191–8201.
- 39 N. Miyaura, T. Yanagi and A. Suzuki, *Synth. Commun.*, 1981, 11, 513–519.
- 40 N. Miyaura and A. Suzuki, Chem. Rev., 1995, 95, 2457-2483.
- 41 F. Paul, J. Patt and J. F. Hartwig, J. Am. Chem. Soc., 1994, **116**, 5969–5970.
- 42 A. S. Guram and S. L. Buchwald, *J. Am. Chem. Soc.*, 1994, **116**, 7901–7902.
- 43 B. C. Hamann and J. F. Hartwig, *J. Am. Chem. Soc.*, 1998, **120**, 12706.
- 44 K. Okano, K. Sunahara, Y. Yamane, Y. Hayashi and A. Mori, *Chem.–Eur. J.*, 2016, **22**, 16450–16454.
- 45 R. A. Irgashev, A. S. Steparuk and G. L. Rusinov, *Org. Biomol. Chem.*, 2018, **16**, 4821–4832.
- 46 B. Robinson, Chem. Rev., 1963, 63, 373-401.
- 47 G. W. Gribble, *Indole ring synthesis: From natural products to drug discovery*, John Wiley & Sons. 2016.
- 48 J. J. Li, Tebbe reagent, in *Name Reactions*, Springer. 2014, pp. 599–600.
- 49 H. Fiesselmann and F. Thoma, *Chem. Ber.*, 1956, **89**, 1907–1912.

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- 50 Y. Sato, Y. Onozaki, T. Sugimoto, H. Kurihara, K. Kamijo, C. Kadowaki, T. Tsujino, A. Watanabe, S. Otsuki and M. Mitsuya, *Bioorg. Med. Chem. Lett.*, 2009, **19**, 4673–4678.
- 51 H. KESKIN, R. E. MILLER and F. F. Nord, J. Org. Chem., 1951, 16, 199–206.
- 52 V. Mane, T. V Baiju and I. N. N. Namboothiri, *ACS Omega*, 2018, 3, 17617–17628.
- 53 V. Declerck, J. Martinez and F. Lamaty, *Chem. Rev.*, 2009, **109**, 1-48.
- 54 G. Masson, C. Housseman and J. Zhu, *Angew. Chem., Int. Ed.*, 2007, **46**, 4614–4628.
- 55 P. Xie and Y. Huang, Eur. J. Org. Chem., 2013, 28, 6213-6226.
- 56 M. Dadwal, S. M. Mobin and I. N. N. Namboothiri, Org. Biomol. Chem., 2006, 4, 2525–2528.
- 57 P. Shanbhag, P. R. Nareddy, M. Dadwal, S. M. Mobin and I. N. N. Namboothiri, *Org. Biomol. Chem.*, 2010, 8, 4867–4873.
- 58 R. A. Irgashev, A. S. Steparuk and G. L. Rusinov, *Tetrahedron* Lett., 2019, **60**, 1–3.
- 59 H. Zhang, M. S. Bednarz, N.-K. Lim, G. Hernandez and W. Wu, Org. Lett., 2014, 16, 2522–2525.
- 60 E. Campaigne and P. A. Monroe, *J. Am. Chem. Soc.*, 1954, **76**, 2447–2450.
- 61 J. Liebscher, B. Neumann and H. Hartmann, *J. Prakt. Chem.*, 1983, **325**, 915–918.

- 62 M. Pellet and F. Huet, Tetrahedron, 1988, 44, 4463-4468.
- 63 R. S. Al-awar, S. P. Joseph and D. L. Comins, *Tetrahedron Lett.*, 1992, 33, 7635–7638.
- 64 Z. Mirjafary, M. Abdoli, H. Saeidian, S. Boroon and A. Kakanejadifard, *RSC Adv.*, 2015, 5, 79361–79384.
- 65 H. Hartmann and J. Liebscher, Synth, 1984, 275–276.
- 66 H. Fiesselmann and G. Pfeiffer, *Chem. Ber.*, 1954, **87**, 848–856.
- 67 J. Liu, Y. Zhang, Y. Yue, Z. Wang, H. Shao, K. Zhuo, Q. Lv and Z. Zhang, J. Org. Chem., 2019, 84, 12946–12959.
- 68 L. Meng, T. Fujikawa, M. Kuwayama, Y. Segawa and K. Itami, J. Am. Chem. Soc., 2016, 138, 10351–10355.
- 69 T. T. Nguyen and P. H. Tran, RSC Adv., 2020, 10, 9663-9671.
- 70 R. Engqvist, A. Javaid and J. Bergman, *Eur. J. Org. Chem.*, 2004, **12**, 2589–2592.
- 71 H. Huang, Q. Li, M. Qiu, Z. Wang, X. Zhang, S. Liu, N. Fu,
 B. Zhao, R. Yang and W. Huang, *RSC Adv.*, 2016, 6, 45873–45883.
- 72 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.
- 73 L. Ackermann and A. Althammer, Synfacts, 2007, 586.