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Recent advances in sulfur–nitrogen bond formation *via* cross-dehydrogenative coupling reactions

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This focus-review surveys literature methods for the construction of sulfur–nitrogen bonds through cross-dehydrogenative coupling reactions between thiols and N–H compounds with a particular emphasis on the mechanistic aspects of the reactions. The literature has been surveyed until the end of 2017.

1. Introduction

Needless to say, organosulfur compounds play a very important role in various fields such as organic synthesis, materials science, agriculture, and medicinal chemistry.^{1–4} Among them, compounds bearing sulfur–nitrogen bonds are important building blocks in a variety of biologically active substances. For instance, sultiame **1** (Fig. 1) with brand name Ospolot is a synthetic sulfa-based drug marketed worldwide for the treatment of epilepsy.^{5,6} The drug works by reducing the activity of carbonic anhydrase, an enzyme responsible for catalyzing the interconversion of carbon dioxide and water to carbonic acid by using a metal hydroxide nucleophilic mechanism.⁷ Topiramate

2 with trade name Topamax is a promising anticonvulsant medicine available in a number of countries worldwide. The drug is also used for migraine prevention in adults.⁸ Folpet **3** is a fungicide used for broad-spectrum control of fungal pathogens.⁹ Sulfoxaflor **4** is an insecticide with a sulfoximine motif, this compound is very active against a wide range of sap-feeding insect pests and widely used around the world.¹⁰ Thus, construction of biologically and synthetically important sulfur–nitrogen bonds is one of the essential transformations in organic synthesis. Conventional methods to achieve such linkages involve cross-coupling of S-halo compounds and N–H compounds in the presence of strong bases¹¹ or metal-catalyzed cross-coupling reactions between N-halogenated compounds and thiols/disulfides.¹² However, the limited availability of commercial starting materials, the formation of undesired by-products and/or harsh reaction conditions are the main drawbacks of these methods. Therefore, the search for novel straightforward and eco-friendly green approaches to the

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synthesis of sulfur–nitrogen bond-containing compounds remains a challenge.

Metal-catalyzed and metal-free cross-coupling reactions have been shown to be one of the most powerful methods for the construction of C–C, C–X, and X–X' bonds (X, X' = heteroatom).^{13–15} Cross-dehydrogenative coupling (often called oxidative coupling) reactions are green and sustainable alternatives to traditional coupling procedures which rely on the use of two unmodified coupling partners.^{15,16} Indeed, in these reactions the relatively inactive C(X)–H bonds can be treated as a functional group, similar to the traditionally used C(X)–(pseudo) halide bonds. Along this line, recently, oxidative coupling of thiols and N–H compounds has gained a lot of attention because of the atom-economic and waste-minimization characters. The most obvious benefit of this route for construction of sulfur–nitrogen bond is that there is no need for preparation and isolation of halogenated reagents, thus improving pot, atom, and step economy. An overview of this emerging area is

presented in this mini-review with a particular emphasis on the mechanistic aspects of the reactions. It is noted that the reactions were classified according to the type of products (Fig. 2). Literature has been surveyed until the end of 2017.

2. Synthesis of sulfenamides

In 2010, N. Taniguchi reported the first example of Cu(I)-catalyzed cross-dehydrogenative coupling of aryl thiols **5** with amines **6** in the presence of Cu(I)/bpy (2,2'-bipyridine) combination as a catalytic system in air as the terminal oxidant, which allows for preparation of a broad range of sulfenamides **7** in moderate to excellent yields (Scheme 1).^{17a} The reaction is noteworthy in that both primary and secondary amines are tolerated. The results demonstrated that aliphatic amines generally gave higher yields than arylamines. It should be mentioned that the electronic character of the substituents in aryl thiols had a strong effect on the facility of reaction. While electron-donating, mild electron-withdrawing and unsubstituted thiophenol gave the desired products in good yields, the strongly electron-withdrawing nitro-substituted thiophenol failed to afford the product. Interestingly, when the reaction was carried out under the oxygen atmosphere, the corresponding sulfenamides were obtained in moderate to high yields. Furthermore, the author observed that using PdCl₂/CuI binary catalyst, the reaction of aryl thiols with primary amines in air selectively afforded sulfenamides derivatives. The author proposed mechanism for the formation of sulfenamides is shown in Scheme 2. A possible process for the formation of sulfenamides **7** should involve the formation of intermediate **B** via oxidative addition of starting aryl thiol **5** with copper catalyst **A**, followed by oxidation of this intermediate with air oxygen to give the intermediate **C**, which after reaction with amine **6** delivers the intermediate **D**. Finally, reductive elimination of intermediate **D** affords the desired product **7** (cycle A, Scheme 2). In another possibility, oxidative homocoupling of thiol **5** can happen to give the disulfide **E**, which after reaction with amine



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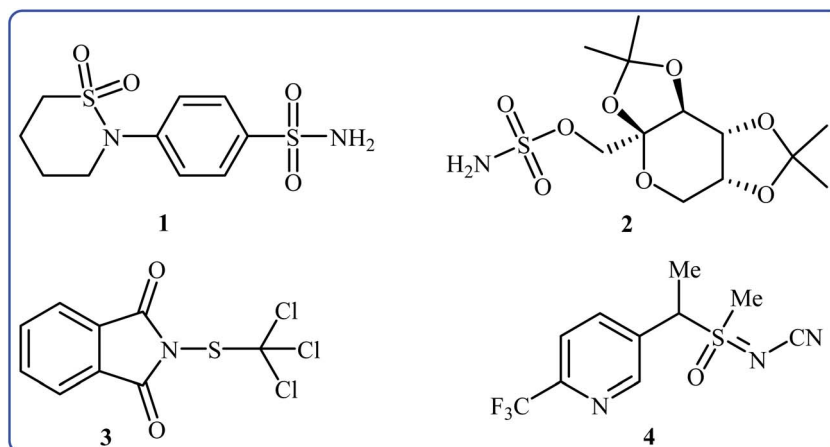


Fig. 1 Selected examples of biologically active sulfur–nitrogen bond-containing organosulfur compounds.

6 affords sulfenamide 7 with the generation of intermediate C (cycle B, Scheme 2).¹⁷

Four years later, the group of Wacharasindhu studied the possibility of synthesizing heterocyclic sulfenamides **10** through hypervalent iodine-catalyzed cross-dehydrogenative coupling of N-heterocyclic thiols **8** with amines **9**.¹⁸ Thus, the careful analysis of the reaction optimization results showed that the optimum condition for this cross-coupling reaction was the addition of DBU (2 equiv.) and amine (2 equiv.) at 0 °C, to a solution of DIB (1.1 equiv.) and thiol (1.0 equiv.) in ¹PrOH. The optimized protocol tolerated a variety of N-heterocyclic thiols, such as 2-mercaptopyridine, 4-mercaptopyridine, 2-mercaptopyrimidine, 2-mercaptobenzothiazol, and 2-mercaptoquinoline, and provided heterocyclic sulfenamides **10** in moderate to high yields with a mixture of corresponding disulfides **11** (Scheme 3). In this study, the authors

reported some limitations in their methodology, since considerable difficulties were found when they attempted to react the sterically hindered *tert*-butylamine and the less nucleophilic aniline. In these cases, they observed only disulfide products with no formation of sulfenamide product. The mechanism shown in Scheme 4 was proposed for the formation of sulfonamides **10**.

Very recently, Yuan's research team developed an efficient synthesis of 2-benzothiazole-sulfenamides **14** *via* direct aerobic oxidative amination of 2-mercaptobenzothiazole **12** under metal-free conditions.¹⁹ In this transformation, 2,2,6,6-(tetramethylpiperidin-1-yl)oxyl (TEMPO) was utilized as the catalyst and O₂ as the oxidant without any co-oxidant. Solvent has a dramatic role in the reaction and among the various solvents such as MeOH, EtOH, DCM, DCE, DMF, MeCN, EtOAc, toluene, acetone, chlorobenzene, ethylbenzene; MeCN was the

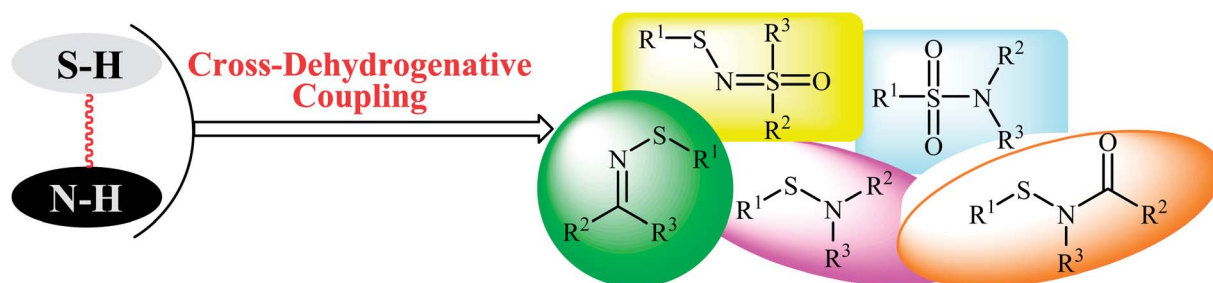
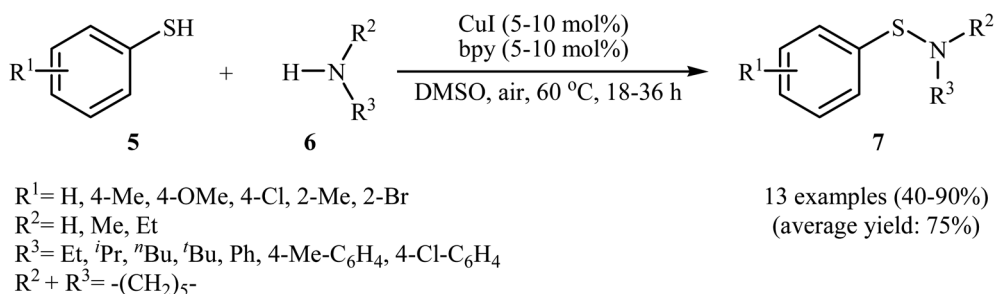
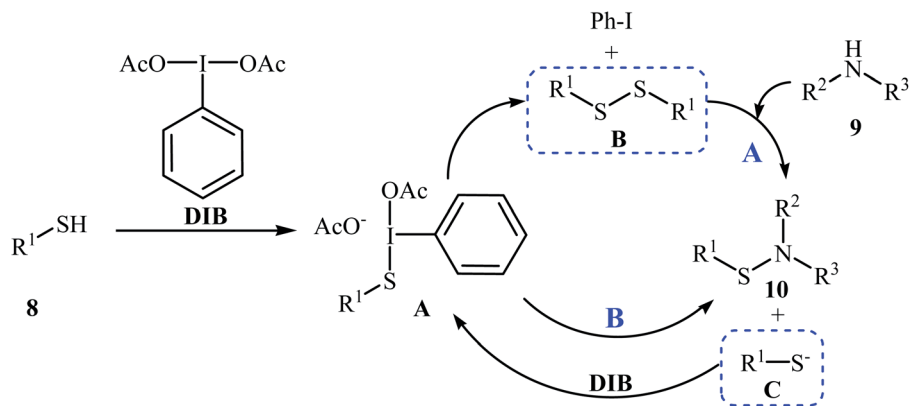


Fig. 2 Synthesis of a variety of sulfur–nitrogen bond-containing organosulfur compounds *via* cross-dehydrogenative coupling reactions.



Scheme 1 Cu(I)-catalyzed S–N coupling of aryl thiols **5** with amines **6**.





Scheme 4 Mechanistic proposal for the formation of 10.

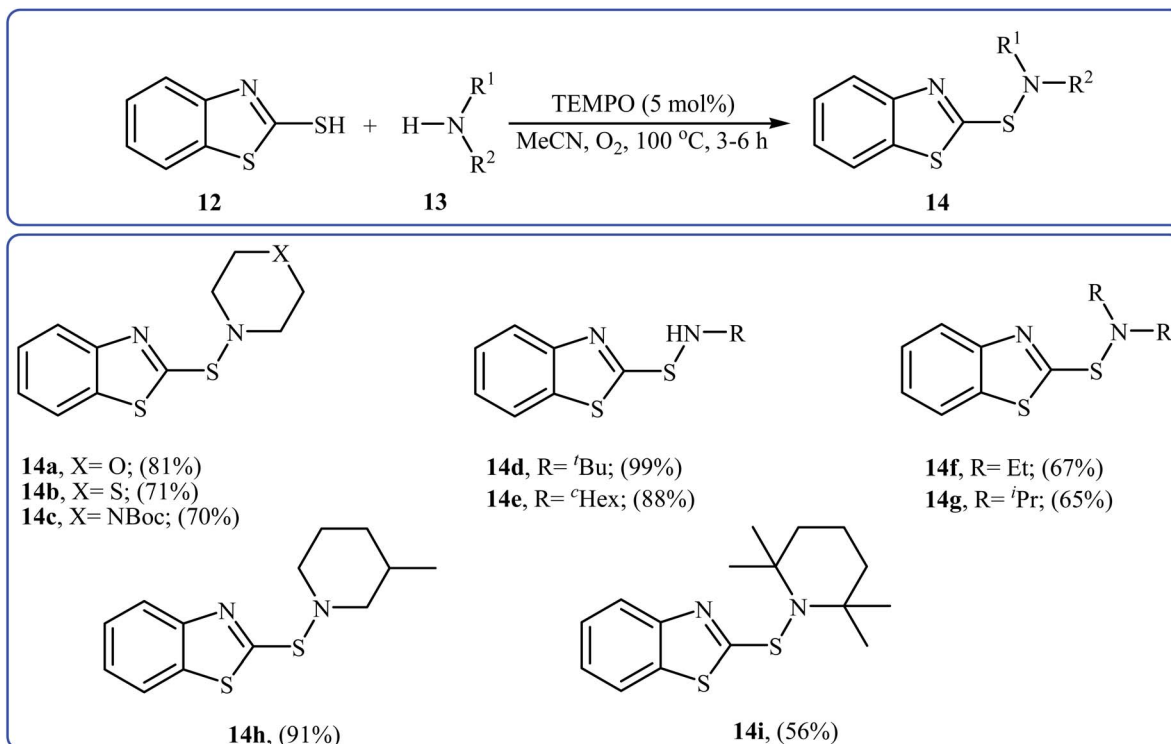
radicals **A** gives disulfide **B**; and (iv) reaction of disulfide **B** and amine **13** affords the expected sulfenamide **14** and mercaptide ion **C**, which can be oxidized *in situ* to regenerate the disulfide **B**.

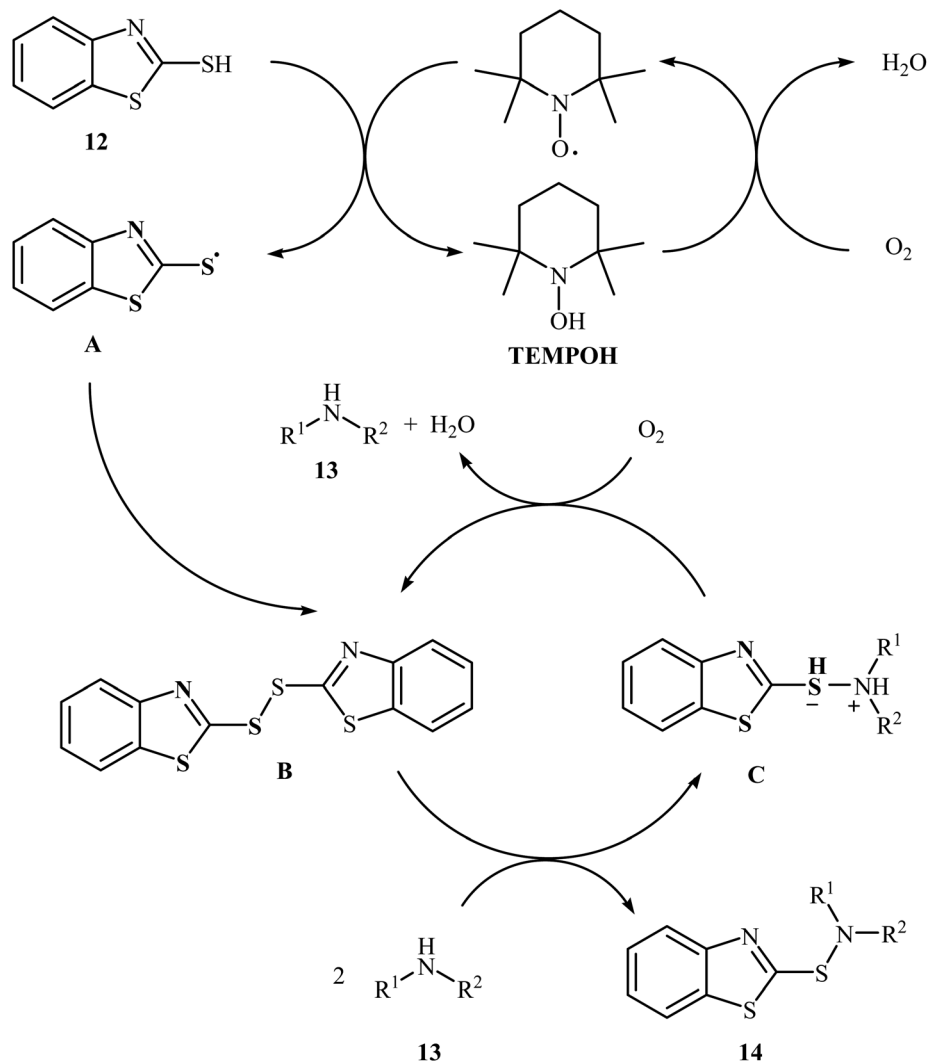
With the aim of designing a greener procedure to sulfenamides *via* aerobic oxidative coupling of thiols with amines, the same research group was able to demonstrate that a range of 2-benzothiazole-sulfenamides **16** could be obtained from the reaction of 2-mercaptobenzothiazole **12** with various aromatic and aliphatic amines **15** in the water media, employing Co(phcy)(SO₃Na)₄ as a catalyst under O₂ atmosphere (Scheme 7).²⁰ Beside good yields, broad substrate scope, scale-up ability, and easy work-up were the advantages, mentioned for this green synthetic approach. According to the author proposed

mechanism, the reaction proceeds along the similar mechanistic pathway that described in Scheme 6. The authors also nicely showed the application of this environment-friendly catalytic system for the oxidative homocoupling of a variety of aromatic and aliphatic thiols to the construction of corresponding disulfides in excellent to almost quantitative yields.

3. Synthesis of *N*-acylsulfenamides

In 2006, Tellitu and Domínguez along with co-workers synthesized a library of biologically important *N*-substituted benzo[*d*]isothiazol-3(2*H*)-one derivatives **18** *via* the hypervalent iodine reagent [phenyliodine(III)-bis(trifluoroacetate)] (PIFA) mediated

Scheme 5 Yuan's synthesis of 2-benzothiazole-sulfenamides **14**.



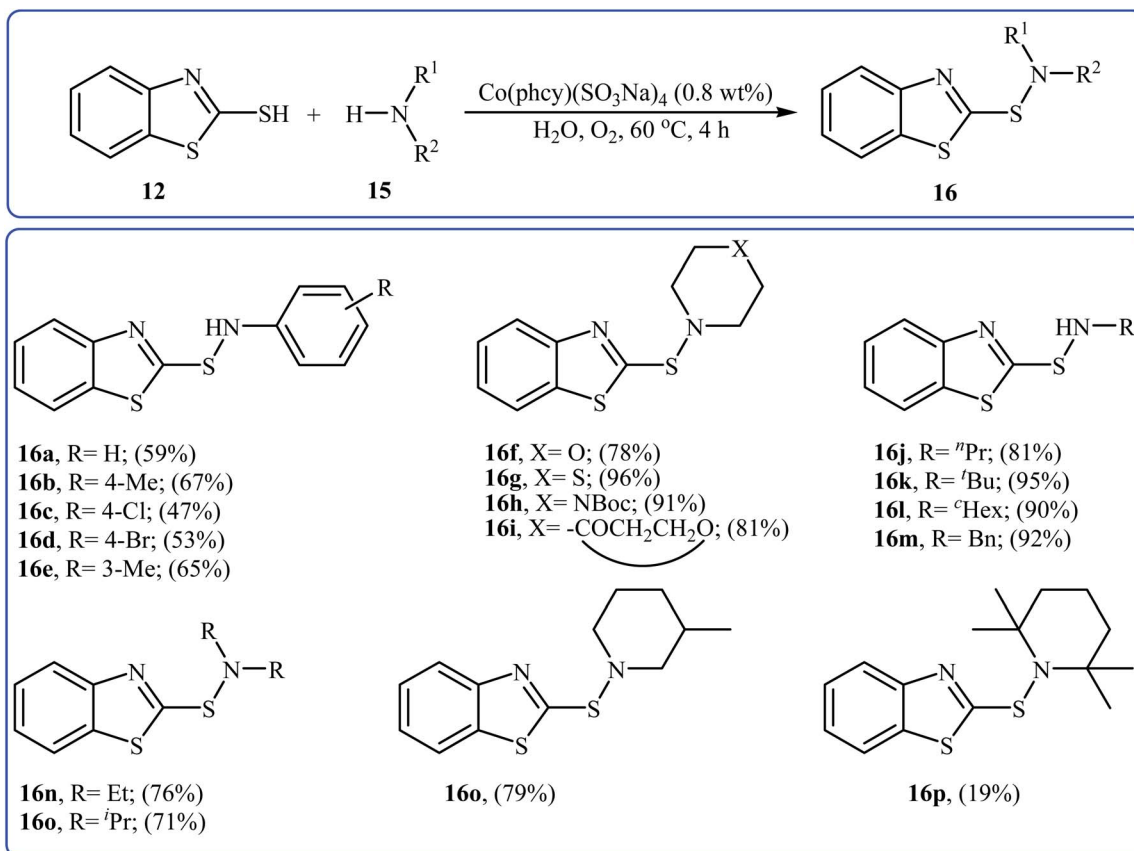
Scheme 6 Mechanism that accounts for the formation of 2-benzothiazole-sulfenamides 14.

intramolecular oxidative cyclization of corresponding *N*-substituted 2-mercapto-benzamides 17 employing TFA as an additive in DCM at 0 °C.²¹ As shown in Scheme 8 the reaction showed good functional group tolerance, including bromo, iodo, and methoxy functionalities and gave corresponding cyclic products in moderate to good yields. However, *N*-allyl substituted 2-mercapto-benzamide failed to participate in this cyclization. It is noted that starting 2-mercapto-benzamides can be easily synthesized from the reaction of commercially available methyl thiosalicylate with primary amines. According to the authors proposed mechanism, the key step of the reaction involves the formation of a reactive *N*-acylnitrenium ion intermediate A. Shortly afterwards, the group of Cosford employed this interesting synthetic approach to access a series of novel drug-like benzoisothiazolones with significant inhibitory activities against phosphomannose isomerase (PMI), an enzyme that converts mannose-6-phosphate (Man-6-P) into fructose-6-phosphate (Fru-6-P).²²

Inspired by these works, Wang and co-workers also showed that *N*-substituted benzoisothiazolones 20 were successfully

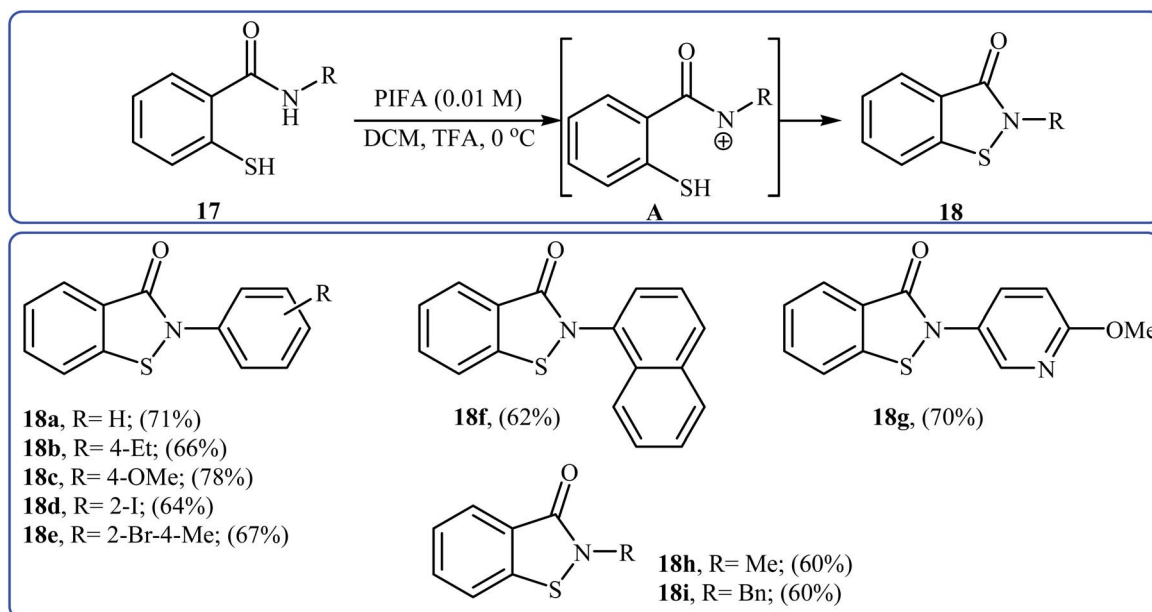
formed from corresponding 2-mercapto-benzamides 19 in a simple process employing commercially available and inexpensive CuI as a catalyst and oxygen as an environmentally benign oxidant, in DMF as solvent and at 70 °C.²³ This method afforded the desired benzoisothiazolones in high to quantitative yields with various alkyl, aryl, and allyl substituents on nitrogen atom (Scheme 9). The process can also be scaled up to provide multigram quantities of the target compound without obvious loss in the yield or outcome. The authors nicely applied this methodology to the high yielding synthesis of piroxicam 23, a nonsteroidal anti-inflammatory drug, precursor (Scheme 10). The following mechanism has been proposed to explain the formation of compounds 20 (Scheme 11). The complex A is first formed by the coordination of 2-mercapto-benzamide 19 to a copper atom. A subsequent oxidative formation of a Cu–S bond *via* the elimination of H₂O provides the intermediate B, which then undergoes oxidative formation of a Cu–N bond *via* the elimination of H₂O₂ to form intermediate C that, after reductive elimination affords the final product 20. In their subsequent study, they reinvestigated the same reaction by

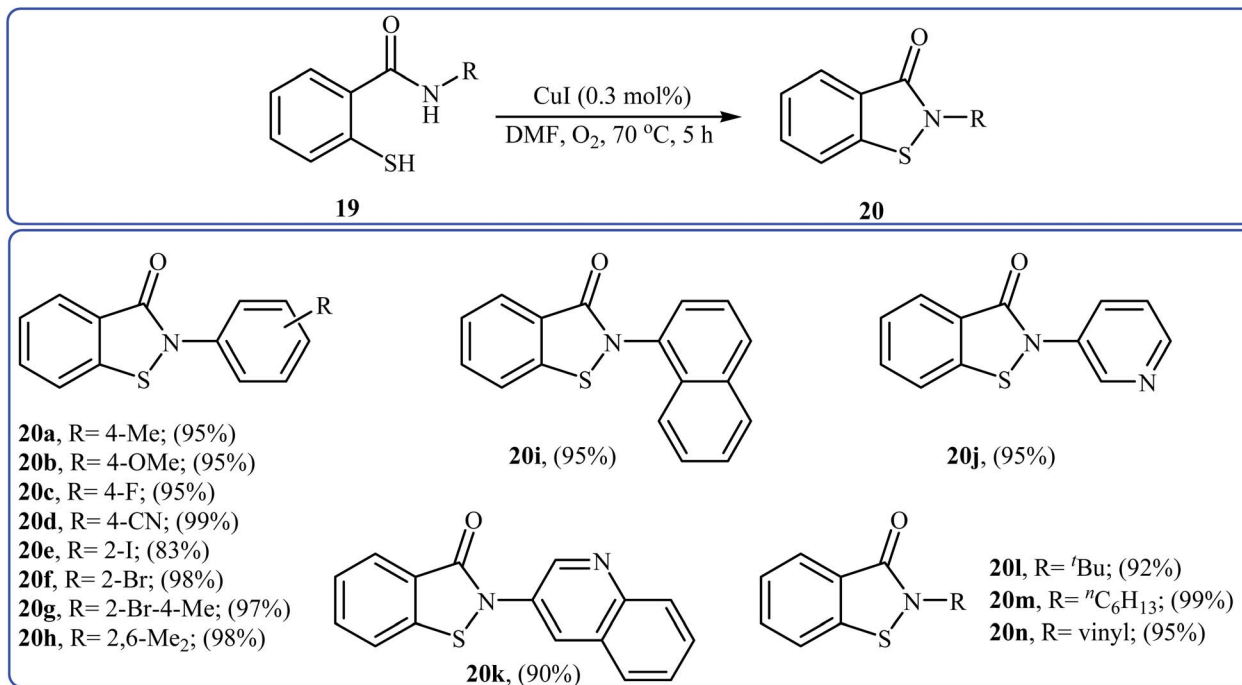


Scheme 7 Co(phcy)(SO₃Na)₄ catalyzed synthesis of 2-benzothiazole-sulfenamides **16** reported by Yuan.²⁰

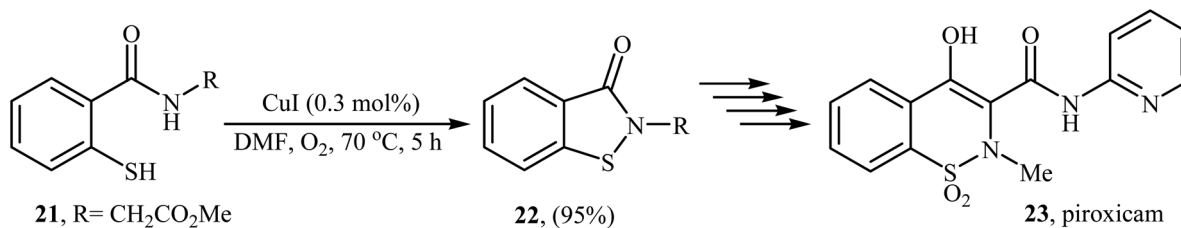
using KBr as a green catalyst under air condition. A diverse range of benzo[*d*]isothiazol-3(2*H*)-ones (14 examples) in very impressive yields (up to 95%) with good functional group tolerance were obtained.²⁴

The first copper-catalyzed intermolecular cross-dehydrogenative coupling reaction between thiols and amides was reported by Jang's group in 2015.²⁵ In this study, various copper salts (*e.g.*, CuI, CuBr, CuOAc, and CuBr₂) and base (*e.g.*,

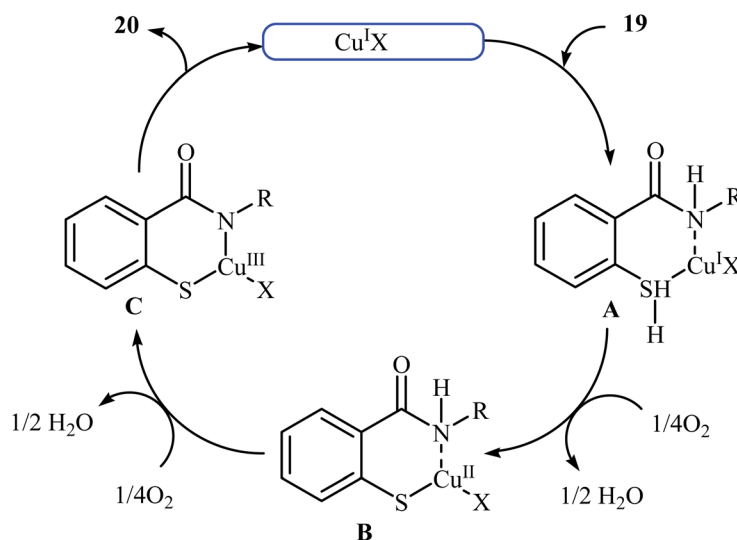
Scheme 8 Tellitu's synthesis of benzo[*d*]isothiazol-3(2*H*)-one derivatives **18**.



Scheme 9 Cu(I)-catalyzed synthesis of benzoisothiazolones **20** through intramolecular S–N cross-dehydrogenative coupling of 2-mercapto-benzamides **19**.

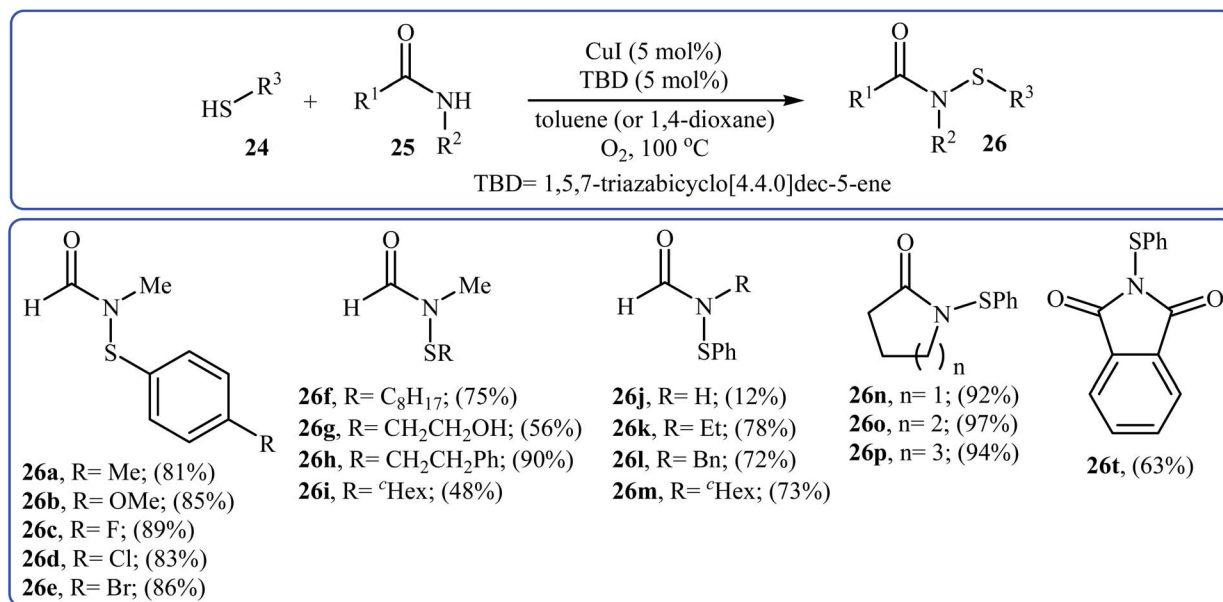


Scheme 10 Wang's synthesis of piroxicam **23**.



Scheme 11 Mechanistic proposal for the reactions in Scheme 9.

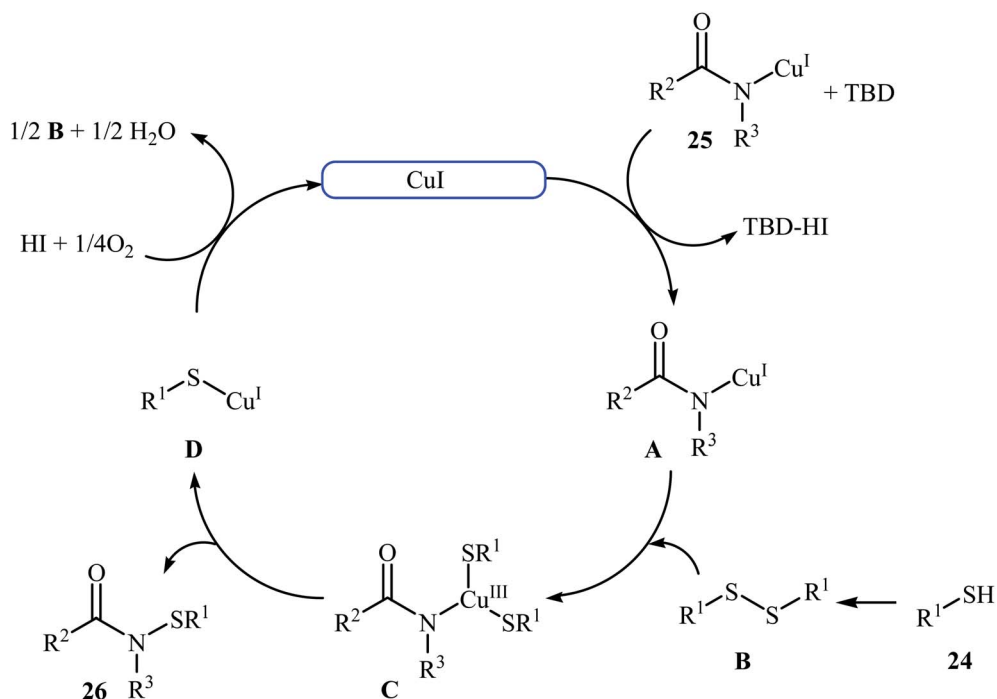




Scheme 12 Cu(I)-catalyzed synthesis of *N*-formyl/acylsulfenamides **26** reported by Jang.²⁵

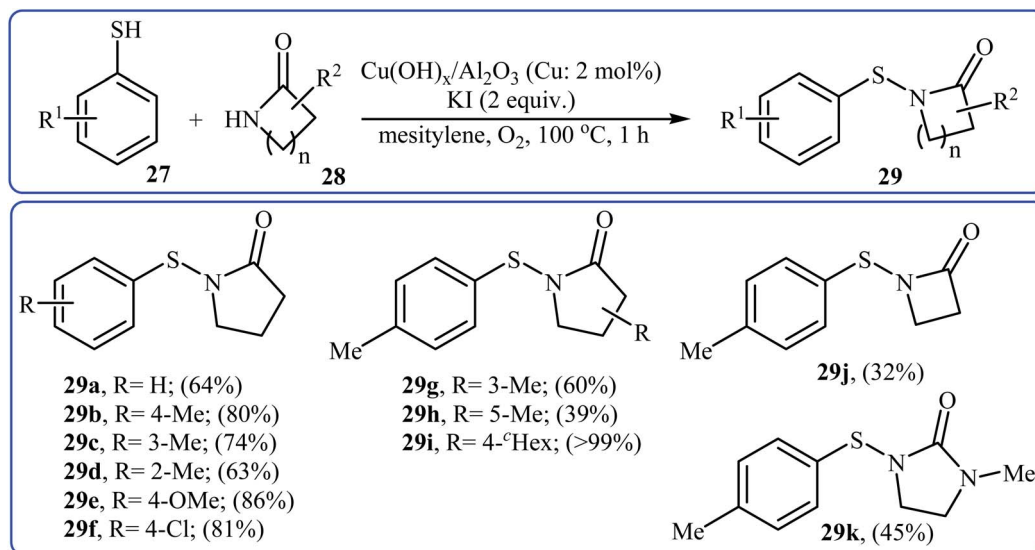
K₂CO₃, ^tBuOK, TBD, and DBU) were examined and the system CuI/TBD was found to be optimal for this transformation. Various thiols **24** and secondary amides **25** were reacted well under the optimized reaction conditions to produce the corresponding *N*-carbonyl sulfenamides **26** in moderate to excellent yields (Scheme 12). However, this protocol for *N*-sulfenylation of primary amides was considerably less efficient. It is noted that this CuI-catalyzed reaction is equally efficient for both the aromatic and aliphatic thiols. A wide range of sensitive functional groups including ester, fluoro, chloro, bromo, alkoxy, and

hydroxy are tolerated by the reaction conditions employed. This procedure offers scope for further manipulation of products. Mechanistic investigations revealed that the reaction starts with the generation of amide–CuI complex **A** via coordination of amide **25** to a copper ion, and then the reaction of intermediate **A** with *in situ* generated disulfide **B** from homocoupling of corresponding thiol **24** furnishes copper complex **C**. Subsequently, intermediate **C** undergoes reductive elimination to provide the expected sulfenamide **26** and generates PhS–CuI complex **D**. Finally, under aerobic oxidation conditions, PhS–



Scheme 13 Mechanistic proposal for the formation of **26**.



Scheme 14 Sakagami's synthesis of *N*-acylsulfenamides **29**.

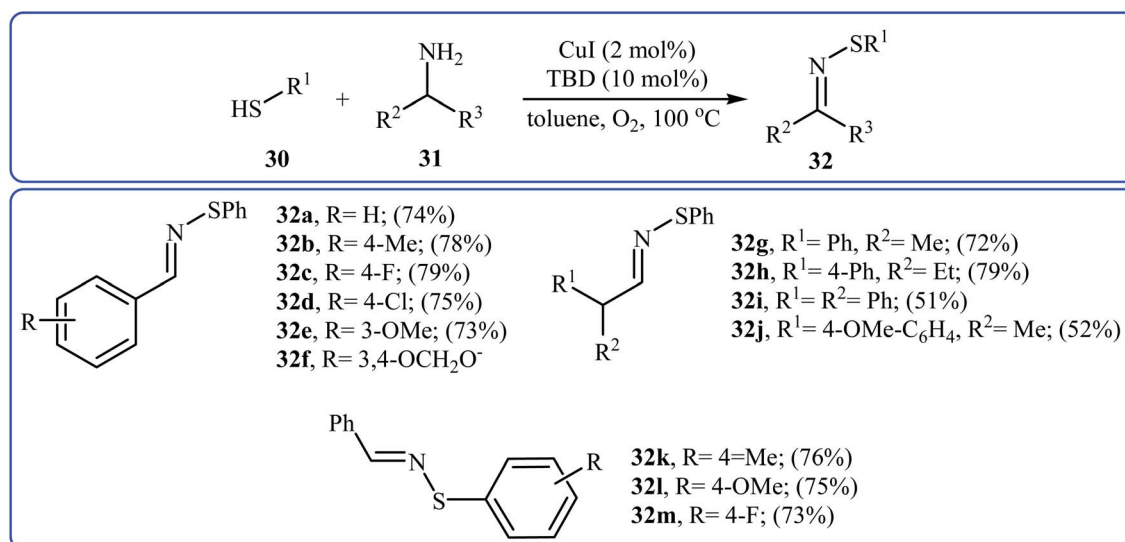
CuI complex **D** converts into a catalytically more reactive CuI complex and disulfide **B** (Scheme 13).

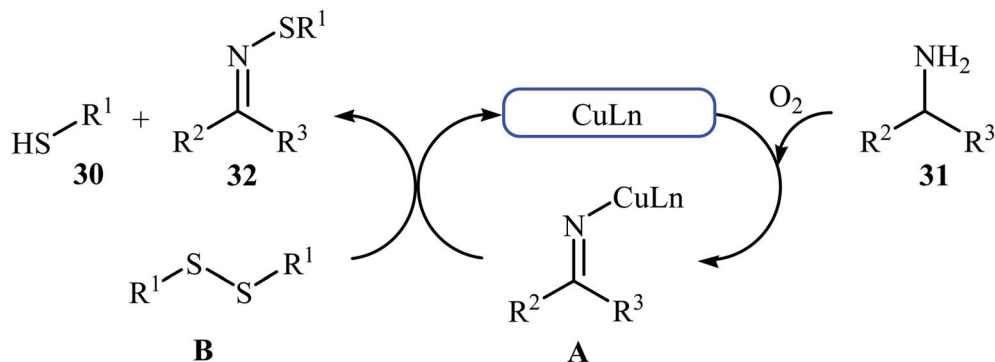
Recently, an efficient method for the synthesis of *N*-acylsulfenamides **29** via Cu -catalyzed cross-coupling of aryl thiols **27** and lactams **28** has been developed by Sakagami and co-workers (Scheme 14). This intermolecular thiolation was achieved in mesitylene using a supported copper hydroxide catalyst $\text{Cu}(\text{OH})_x/\text{Al}_2\text{O}_3$ (Cu: 2 mol%) at 100 °C for 1 h under an oxygen atmosphere to afford *N*-acylsulfenamides **3** in moderate to excellent yields.²⁶

4. Synthesis of sulfenylimines

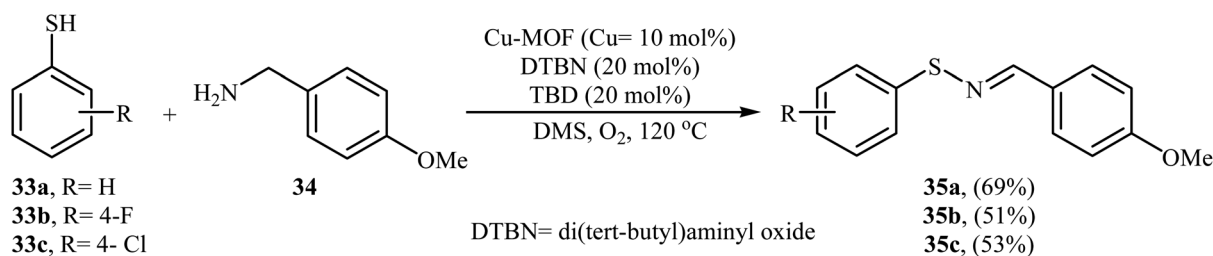
In 2015, Lee, Wang, and Jang developed an elegant $\text{Cu}(\text{i})$ -catalyzed oxidative cross-coupling reaction between thiols **30** and benzyl amines **31** for the construction of synthetically important

N-sulfenylimines **32** (Scheme 15).²⁷ The best conversion efficiency was obtained for the reactions containing CuI (2 mol%) and TBD (10 mol%) in toluene at 100 °C under O_2 atmosphere. This approach provides the first highly practical procedure for synthesizing *N*-sulfenylimines from commercially available and inexpensive thiols and amines. Functional groups including fluoro, chloro, and ether are tolerated by the reaction conditions employed. It is noteworthy that this $\text{Cu}(\text{i})$ -catalyzed reaction is equally efficient for both the primary and secondary amines. According to the authors proposed mechanism, the reaction starts with the generation of an imine- Cu complex **A** via the aerobic oxidation of benzyl amine **31** by Cu catalyst. Meanwhile, homocoupling of starting thiol **30** affords disulfide intermediate **B**. Finally, the reaction of complex **A** with disulfide **B** yields the expected *N*-sulfenylimines **32** (Scheme 16).

Scheme 15 Synthesis of *N*-sulfenylimines **32** from thiols **30** and benzyl amines **31**.



Scheme 16 Plausible mechanism for the formation of 32.

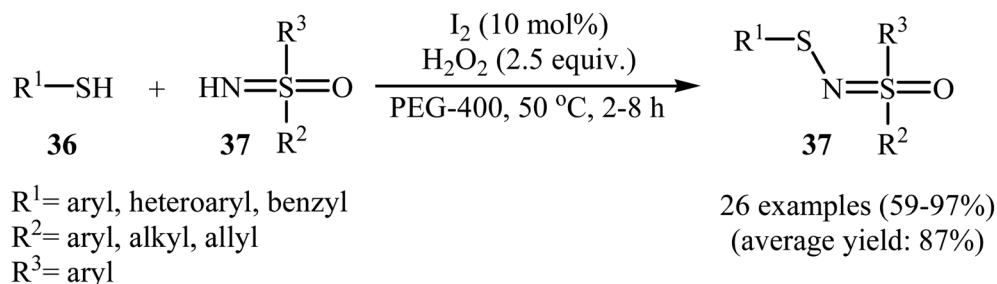
Scheme 17 Qiu-He's synthesis of *N*-sulfenylimines 35.

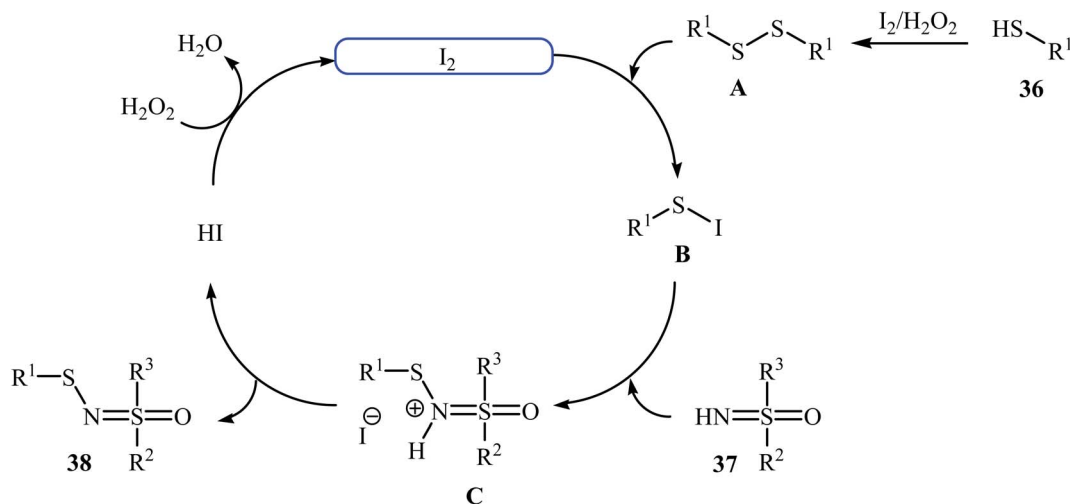
Shortly after this report, Qiu and He along with their co-workers showed that copper-based metal-organic frameworks with *pcu*-topology (*pcu*-MOF) are efficient heterogeneous catalyst for the oxidative coupling of thiols and amines. Thus, by using *pcu*-MOF/DTBN/TBD as a catalytic system in DMSO under an O₂ atmosphere, the coupling of thiophenols 33 with 3-methoxybenzylamine 34 afforded corresponding *N*-sulfenylimines 35 in moderate yields (Scheme 17).²⁸

5. Synthesis of sulfenylsulfoximines

Very recently, Yang and co-workers showed that *N*-sulfenylsulfoximine derivatives 38 could be synthesized *via* dehydrocoupling reaction of thiols 36 with sulfoximines 37 in the presence of 1.0 mol% of molecular iodine as the catalyst and 2.5 equiv. of H₂O₂ as an oxidant in eco-friendly polyethylene glycol-400 (PEG-400) at 50 °C. Generally, target *N*-sulfenylsulfoximines were obtained in good to excellent yields (Scheme 18).²⁹ The

reaction showed excellent functional group tolerance, including fluoro, chloro, bromo, methoxy, and nitro groups that would allow further elaboration of the products. This I₂-catalyzed reaction is applicable for both the (hetero)aromatic as well as benzylic thiols and for various substituted sulfoximines. It should be mentioned that the optimized reaction condition was also applied for the cross-dehydrogenative coupling reactions of thiophenol with a diverse set of functionalized anilines, and the reaction gave good yields of the expected sulfonamides. A possible mechanistic pathway was also proposed (Scheme 19), whereby the reaction is initiated with homocoupling of thiol 36 to give disulfide intermediate A, which undergoes reaction with I₂ affording sulfenyl iodide intermediate B. Subsequently, nucleophilic attack of sulfoximine 37 onto the activated S-I bond in intermediate B affords the cationic *N*-sulfenylsulfoximine C, which by deprotonation convert to the desired products 38. The generated hydriodic acid is then oxidized by H₂O₂ to restore the catalyst I₂, and water is the sole by-product. The

Scheme 18 Synthesis of *N*-sulfenylsulfoximines 37 through metal-free N-H/S-H cross-dehydrogenative coupling reaction.



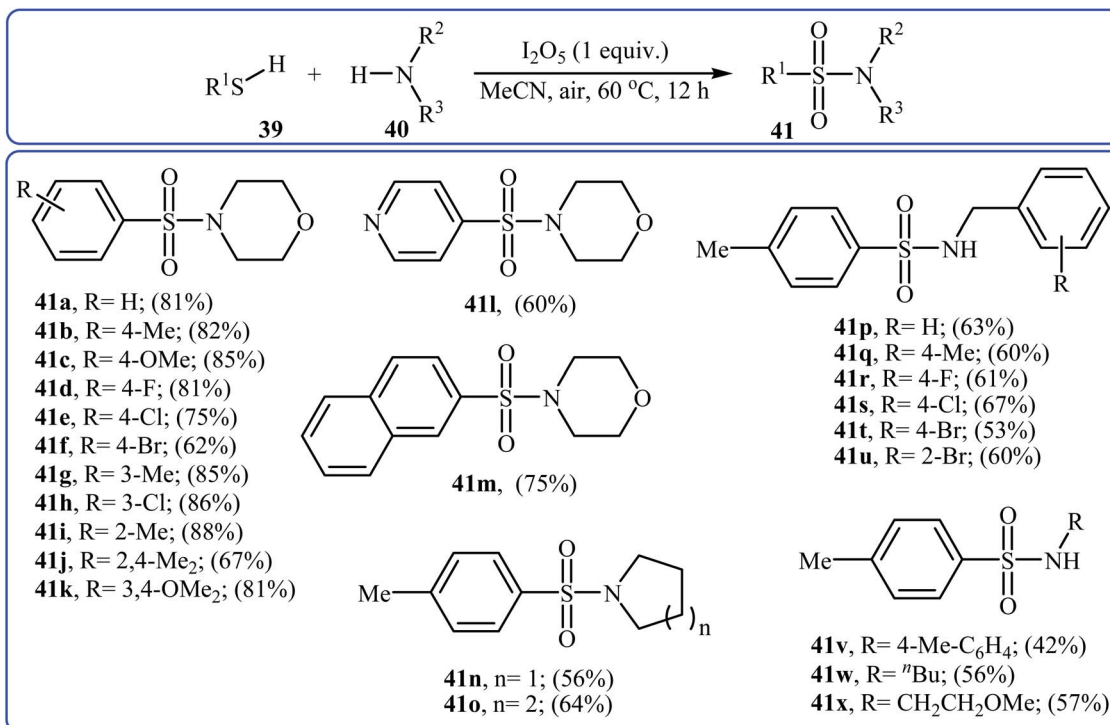
Scheme 19 Mechanistic proposal for the reaction in Scheme 18.

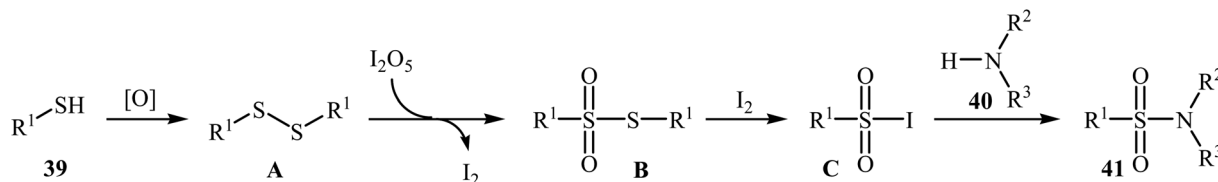
authors claimed that their report is the first example of the direct cross-dehydrogenative coupling reaction to construct *N*-sulfenylsulfoximines.

6. Synthesis of sulfonamides

Very recently, the Wei-Wang group have developed an efficient and straightforward approach for the synthesis of biologically important secondary and tertiary sulfonamides **41** *via* I_2O_5 -mediated oxidative coupling of aryl thiols **39** and amines **40** (Scheme 20).³⁰ The reaction was performed in acetonitrile at 60 °C, tolerated various functional groups, and generally

provided corresponding sulfonamides in moderate to high yields. It is noted that the reaction is applicable for various amines like aromatic, benzylic, and aliphatic amines. This protocol was also successfully applied to the preparation of gram quantities of the desired sulfonamide without sacrificing the yield or outcome of the methodology. The proposed reaction mechanism is outlined in Scheme 21 and comprises the following basic steps: (i) initial formation of disulfide **A** *via* dimerization of thiol **39**; (ii) oxidation of disulfide **A** by I_2O_5 to give arylsulfonothioate **B** with the concomitant formation of molecular iodine; (iii) interaction of intermediate **B** with iodine to produce sulfonyl iodide **C**; and (iv) nucleophilic substitution

Scheme 20 I_2O_5 -mediated direct construction of sulfonamides **41** from thiols **39** and amines **40**.



Scheme 21 Mechanism that accounts for the formation of sulfonamides 41.

of sulfonyl iodide C by amine 40 and formation of the sulfonamide 41. To the best of our knowledge, this is the only example of direct construction of sulfonamides from readily-available thiols and amines reported so far.

7. Outlook and conclusion

Sulfur–nitrogen bond-containing compounds have extensive application in organic and medicinal chemistry. Synthetically, such organosulfur compounds are commonly prepared *via* cross-coupling of S-halo compounds and N–H compounds or cross-coupling reactions between N-halogenated compounds and thiols. However, the limited availability of commercial starting materials, the formation of undesired by-products and/or harsh reaction conditions are the main drawbacks of these methods. Recently, cross-dehydrogenative coupling reaction has become an attractive alternative to traditional S–N cross-coupling reactions due to the waste-minimization and the costs associated with the preparation of starting materials. Shorter synthetic routes and high atom economy are the key features of these eco-friendly green reactions. This new page of sulfur–nitrogen bond formation has been successfully applied in the synthesis of a variety of S–N bond-containing compounds such as sulfenamides, sulfonamides, sulfonylimines, and sulfonyl-sulfoximines. Despite the significant achievements during the past few years in this interesting research arena, there are still many problems remained: (i) almost all of the catalysts and oxidants are limited to the use of copper and hypervalent iodine, respectively. Thus the exploration of different catalytic systems is highly desirable; (ii) the number of reported examples in some reactions such as synthesis of sulfonamides and sulfonylsulfoximines from S–H and N–H compounds are narrow and there is an urgent need to study the scope and limitations of these reactions; and (iii) other reactions such as coupling of thiols with urethanes and ureas should be explored. It is anticipated that the insight provided in this review will be beneficial for eliciting further research in this domain.

Conflicts of interest

There are no conflicts to declare.

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