Recent Advances in the Synthesis of Sulfides, Sulfoxides and Sulfones via C-S Bond Construction from Non-Halide Substrates

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Abstract: The construction of a C-S bond is a powerful strategy for the synthesis of sulfur containing compounds including sulfides, sulfoxides, and sulfones. Recent methodological developments have revealed lots of novel protocols for C-S bond formation, providing easy access to sulfur containing compounds. Unlike traditional Ullmann typed C-S coupling reaction, the recently developed reactions frequently use non-halide compounds, such as diazo compounds and simple arenes/alkanes instead of aryl halides as substrates. On the other hand, novel C-S coupling reaction pathways involving thiyl radicals have emerged as an important strategy to construct C-S bonds. In this review, we focus on the recent advances on the synthesis of sulfides, sulfoxides, and sulfones from non-halide substrates involving C-S bond construction.

Keywords: sulfide; sulfoxide; sulfone; C-S bond construction

1. Introduction

Sulfur containing compounds are a fundamental building block in organic synthesis [1,2]. They are also of great significance in bioactive natural products, pharmaceutical drugs, pesticides, and materials [3–8]. As shown in Figure 1, sulfur containing compounds, such as sulfides, sulfoxides, and sulfones, constitute a large portion of pharmaceutical drugs possessing a variety of biological activities. Given the great importance of sulfur containing compounds, intensive efforts have been made to develop effective synthetic methods. The construction of C-S bond is considered as an efficient strategy for the synthesis of sulfur containing compounds. The traditional Ullmann typed reaction, which has been well established in the past several decades, has provided lots of synthetic protocols for diaryl sulfides and aryl alkyl sulfides through transition-metal catalyzed C-S coupling reaction of aryl halides with aryl thiols or other sulfur precursors [9–11]. Methodological advancements in C-H functionalization have led to the development of a variety of useful transformations to prepare sulfur containing compounds directly from (hetero)arenes and alkanes [12,13]. In very recent years, many novel strategies involving thiyl radicals have emerged as powerful tools for constructing C-S bonds in very facile manner using non-halide substrates.

In this review, we highlight the very recent advances in novel strategies that take advantage of C-S bond construction for the synthesis of sulfides, sulfoxides, and sulfones from non-halide substrates.
2. Synthesis of Sulfides via C-S Bond Construction

Traditional preparation of diaryl sulfides or aryl alkyl sulfides relies on the coupling reaction of aryl halides with thiols, disulfides, elemental sulfur, or other sulfur containing reagents. These conversions often require the employment of transition-metal catalysts and ligand in the presence of base. Recent research demonstrates that non-halide substrates, such as diazo compounds and simple arenes/alkanes, successfully coupled with thiol or disulfides to afford the corresponding sulfides. On the other hand, the sulfur containing reagents such as thiol and elemental sulfur, which are often used in the Ullmann typed C-S coupling reactions, also found wide application in the synthesis of sulfides from non-halide substrates. In this part, the synthetic methods for sulfides will be classified and discussed by different sulfonylation sources.

2.1. Disulfide as Sulfonylation Source

The S-S bond of disulfide can be reversibly cleaved when exposed to heating, photoirradiation, or transition-metal catalyst such as rhodium catalyst [14]. The generated active organothio intermediates such as thiyl radicals and sulfoniums, can easily couple with partners to afford sulfides. In addition, disulfide is a favorable sulfonylation source as it is stable and odorless.

2.1.1. C(sp^3)-S Bond Formation with Disulfide

Diazoo compounds are widely used in organic synthesis for constructing new bonds. Generally, photoinitiated, thermal, or transition metal catalyzed decomposition of the diazo compound affords the corresponding carbene or metal carbenoid, which subsequently undergo various transformations [15,16].

In 2016, the Lee group reported a rhodium pivalate-catalyzed direct sulfonylation of diazooxindoles using disulfide as the sulfonylation reagent via intermolecular C-S bond formation to synthesize thioindoles (Scheme 1) [17]. In this conversion, the diazo compound was converted to rhodium carbenoid, which further reacted with disulfide to afford a C-S bond. The subsequent S-S cleavage led to the formation of sulfides and cyclohexenylsulfide, respectively. In the developed reaction system, diazooxindoles reacted with a variety of alkyl or aryl disulfides to afford sulfide or dithioacetals under very mild condition. Generally, the reaction of diazo compounds with alkyl disulfides predominantly gave monoalkylthioindoles. However, when the 5-position of diazooxindoles was substituted with chloro, the reactions gave dialkylthiooxindoles as products. The same results were observed with the reaction of diaryl disulfides, which afforded diarylthiooxindoles in the conversion. It is interesting that
the reaction of diazo compounds with dicycloalkyl disulfide provided both 3-cycloalkylthiooxindoles and 3-cycloalkenylthiooxindoles.

**Scheme 1.** C-S coupling reaction of diazo compounds and disulfides.

N-tosylhydrazone is a readily available alkenyl source through the generation of metal-carbene species. In 2018, the Zhang and Wang group developed a Cu(I) catalyzed protocol for the synthesis of dithioacetals via C-S coupling reaction of N-tosylhydrazones and disulfides (Scheme 2) [18]. N-tosylhydrazone functioned as a precursor of diazo compound in this reaction. The reaction occurred at 60 °C in the presence of CuI, 1,10-phenanthroline (1,10-phen) and t-BuONa in DMF. This protocol was featured by a broad substrate scope. A great variety of N-tosylhydrazones and disulfides were successfully converted into the corresponding dithioacetals in moderate to good yields.

**Scheme 2.** Copper catalyzed synthesis of dithioacetals from tosylhydrazones.

C–C bond fragmentation is a useful method to provide a fragment for the functionalization of molecules and has been extensively applied in organic synthesis [19]. In recent years, C-C bond cleavage/sulfenylation reactions via radical pathways have emerged as a practical tool for the synthesis of aryl alkyl sulfides [20].

In 2018, the Wu group reported a facile method for the synthesis of aryl methyl sulfides by treating di-tert-butyl peroxide (DTBP) with diaryl disulfide in acetonitrile (Scheme 3) [21]. In this conversion, di-tert-butyl peroxide gave methyl radicals under heating condition and the methyl radicals subsequently reacted with diaryl disulfide to afford aryl methyl sulfides.

**Scheme 3.** Synthesis of aryl methyl sulfides using diaryl sulfides and DTBP.

In 2019, the Kwon group developed a dealkenylative thiolation method for the synthesis of aryl alkyl sulfides (Scheme 4) [22]. The isopropenyl group of substrate was easily removed via
C-C bond cleavage by the ozonolysis of alkenes at −78 °C and the followed Fe(II) salt mediated single electron transfer (SET)-based reductive cleavage resulted in the formation of alkyl radicals, which can be trapped with an aryl disulfide to form a C(sp³)–S bond. This method was featured by mild condition and high efficiency. The formation of C-S bond was accomplished within only 1 min. Moreover, this method is practical as it can be applied in the dealkenylation thiolation of (exo-methylene) cycloalkanes and methylocycloalkenes. Alkanes containing exo-methylene group were converted into the corresponding phenylthiol-containing methyl carboxylates and methylocycloalkenes were converted into phenylthio-aldehydes.

![Scheme 4.](image)

In 2019, the Abbasi group reported a decarbonylative thiolation of 2-phenylpropanal with disulfides in the presence of morpholine and K₂CO₃ in DMF at 130 °C (Scheme 5) [23]. This conversion provided the aryl benzyl sulfides in good to excellent yields. This reaction can be scaled up to 1 g with good yield. Moreover, a one-pot synthesis of aryl benzyl sulfides from simple aryl halides, CS₂ and KCN was also developed in their work.

![Scheme 5.](image)

C-H sulfenylation represents an atom-economic strategy for the synthesis of sulfides as it employs simple arenes or alkanes as substrates. In 2019, the Yang group reported an unprecedented site-selective C(sp³)-H sulfenylation of alkyl tosylamides at the δ-position (Scheme 6) [24]. The C-S coupling reaction occurred to N-fluoro-tosylamide and diaryl disulfide at 40 °C in the presence of Cu(acac)₂, 1,10-phen, Na₂HPO₄ and indium powder under the irradiation of blue LED light. This protocol was compatible with various diaryl disulfides, and even dibenzyl disulfides and diphenyl diselenides. It is noteworthy that C-H thiolation using this protocol exhibited high site-selectivity and substrate compatibility with N-fluoro-tosylamides, no matter the δ-C located at a chain or a cycle. It was suitable for the thiolation of 1°, 2°, and 3° C-H bonds, and the reactivity of hydrogen atoms decreased in the order tertiary>secondary>primary. On the other hand, the reactivity of aryl disulfide was higher than that of aliphatic disulfide, and electro-donating group was more favorable than electro-withdrawing group on the phenyl ring of diaryl disulfide. Examples of applying this protocol in the synthesis of complex biological active compounds were also reported in their work.
In the same year, the Meng group reported a Cu(I)-catalyzed sulfenylation of 1,3-dicarbonyl substrates with diaryl sulfides (Scheme 7) [25]. This protocol was featured by a mild condition, simple procedure, and satisfying yield. This coupling reaction occurred in the presence of CuI in acetonitrile at room temperature in the darkness, providing the corresponding sulfides in moderate to excellent yields. Moreover, this protocol was also compatible with the β-carbonyl amide and dialkyl sulfide.

Chalcogenium ions can be readily captured by nucleophile-tethered alkenes to afford functionalized chalcogenides. In 2018, the Zhao group developed an efficient and practical protocol for the synthesis of sulfides from nucleophile tethered alkenes and diaryl disulfides in the presence of TfOH and diethyl azodicarboxylate (DEAD) (Scheme 8) [26]. This conversion occurred at room temperature, tolerated various functional groups and provided sulfides in good to excellent yields. In this conversion, DEAD reacted with disulfide to generate N-sulfenylhydrazine and [PhS⁺][−OTf] in the presence of TfOH via the cleavage of S-S bonds. [PhS⁺][−OTf] was a very active intermediate, which can give a variety of sulfides when it was treated with carboxyl, hydroxyl, and sulfonamide-tethered alkenes, respectively. Moreover, aryl boronic acids and arenes were also suitable for this conversion. They were thiolated with disulfides to afford the corresponding sulfides when BF₃•OEt₂ was used instead of TfOH. This reaction system was also successfully applied in the synthesis of diaryl selenides from diphenyl diselenides.

In 2018, the Ouyang and Yang group found a very facile protocol to convert pentafluorobenzenes to dithiolated tetrafluorobenzenes (Scheme 9) [27]. The reaction occurred in the presence of KOH.
in DMF at room temperature, affording the corresponding dithiolated compounds. Mechanistic investigation indicated that the fluoro atom at 3-position of the pentafluorobenzene was initially thiolated by disulfide, and the C-H bond was subsequently thiolated to give the dithiolated product. Based on this result, they further used 1-aryl-2,3,5,6-tetrafluorobenzenes as substrate to synthesize the unsymmetrical dithiolated products.

Scheme 9. Synthesis of dithiolated tetrafluorobenzenes.

In 2019, the Yotphan group reported a copper catalyzed regioselective C-H thiolation and thiocyanation of uracils using disulfide and NH₄SCN, respectively [28]. The reaction readily proceeded in the presence of catalytic Cu(OTf)₂ and K₂S₂O₈ in acetonitrile (Scheme 10). A variety of functional groups were tolerated in the reaction system and good to excellent yields were obtained. Electron-rich diaryl disulfides were more favorable for this conversion than electron-deficient diaryl disulfides. For example, 1,2-bis(4-chlorophenyl)disulfane gave only 17% of the corresponding product. Control experiments indicated this conversion proceed through a radical process.

Scheme 10. C-H thiolation of uracils with disulfides.

2.2. Thiol as Sulfenylation Source

Thiol is an important sulfenylation reagent in the construction of sulfides. Traditional Ullmann typed C-S coupling reactions use aryl halides as substrates to couple with thiols in the presence of transition metal catalyst. Recent years have witnessed the fast development of direct C-S coupling of non-halide substrates with aryl/alkyl thiol.

2.2.1. C(sp³)-S Bond Formation with Thiol

In 2018, the Lei group reported a radical-radical cross-coupling strategy for the construction of C(sp³)-S bond using aryl thiols [29]. A series of α-alkythionitriles were synthesized from aryl thiols and isobutyronitrile in the presence of copper(II) salt as a catalyst (Scheme 11). The thiol radical was generated from aryl thiol by oxidation by Cu(II) and stabilized by Cu(I). It further coupled with isobutyronitrile radical generated from azobisisobutyronitrile. The reaction was carried out in the presence of Cu(OTf)₂ as catalyst and 1,10-phen as ligand in 1,2-dichloroethane (DCE) at 80 °C. Both aryl thiols and heteroaryl thiol were smoothly converted into the corresponding C-S coupled products. Moreover, the cross-coupling product can be easily modified to valuable carboxylic acids, amides, and ketones.
Diazoo compound and diazo precursors such as hydrozene are also suitable for C-S coupling reaction with aryl thiols. In 2019, the Singh group reported an Eosin Y catalyzed, photoredox facilitated C-S bond formation for the synthesis of aryl benzyl sulfides from acetophenone hydrazines (Scheme 12) [30]. Acetophenone hydrazine coupled with aryl thiol to afford the corresponding sulfides in the presence of Eosin Y and NaOH under green LED (530 nm) irradiation. The addition of radical trap reagent 2,2,6,6-tetramethylpiiperidine-1-oxyl (TEMPO) suppressed the transformation, indicating that the conversion was involved in a radical pathway.

In 2020, the Huang and Baell group prepared TFMSA@SBA-15 by immobilizing trifluoromethanesulfonic acid on mesoporous silica SBA-15, and applied it as an efficient catalyst for the construction of C-C, C-O, C-N, and C-S bonds (Scheme 13) [31]. TFMSA@SBA-15 was effective in the coupling reaction of α-diazoester and aryl thiol to afford the corresponding phenylsulfenyl phenylacetate esters. Moreover, the reaction system was attractive that the reaction occurred under solvent-free and room temperature condition. In addition, the reaction system was very efficient as the conversion can complete within 5–50 min.
Scheme 14. Deaminative C-S coupling of benzyl amines and aryl thiols.

Dithioacetal is not only a traditional protecting group, but also an important precursor of ketones, alkyls, allkenes, and so on [33]. In 2018, the Wang group developed a facile protocol for the synthesis of dithioacetals through cascade C-S bond formation (Scheme 15) [34]. 3-Oxo-butanamides reacted with aryl thiols in the presence of NaOH and molecular oxygen in n-PrCN at 80 °C, providing dithioamides in good to excellent yields. Moreover, this protocol can be also applied in the synthesis of diselenoacetal amides.

Scheme 15. Synthesis of dithioacetals from aryl thiols and 3-oxo-butanamide.

In 2018, the Lee group reported a novel dehydrative C-S bond formation for the synthesis of sulfides from 2-(hydromethyl)phenol and thiols in the presence of I$_2$ (Scheme 16) [35]. The reaction system was mild and efficient. The reaction occurred at room temperature and accomplished within 4 h. It is noteworthy that both aryl thiol and aliphatic thiol were compatible in this protocol. In fact, in the competition experiment studies, slightly higher selectivity was observed with aliphatic thiols. Moreover, long chain aliphatic thiol can afford products in good yields in gram scale.

Scheme 16. Dehydrative C-S coupling of 2-(hydromethyl)phenols and thiols.

The Panda group reported another dehydrative C-S bond formation for the synthesis of triarylthioarenes from triaryl methanol and aryl thiols (Scheme 17) [36]. The reaction proceeded in the presence of indium (III) triflate as catalyst in THF at room temperature. This method allowed the reaction to complete within 15 min and give yields up to 76%. This protocol was also successfully applied in the synthesis of tetraaryl methane from triaryl methanol and heteroarenes such as indole and furan.
Scheme 17. Dehydrative C-S coupling of triaryl methanol and aryl thiols.

In 2018, the Yan and Yu group reported an oxidative disulfenylation/cyclization reaction of 2'-hydroxyacetophenones with thiophenols mediated by TBAI/K$_2$S$_2$O$_8$ for the synthesis of functionalized 2,2-dithio-benzofuran-3(2H)-ones (Scheme 18) [37]. This protocol was featured by metal-free condition, satisfying yields and broad substrate scope. A variety of functional groups, including methoxyl, halogen, and amide, were well endured. The proposed mechanism indicated that the conversion underwent through two steps including disulfenylation and intramolecular cyclization.

Scheme 18. Oxidative disulfenylation/cyclization reaction of 2'-hydroxyacetophenones.

In 2018, the Jang group found a novel synthetic method for allylic sulfides from α-methylstyrenes and aryl thiols via direct C-S bond formation (Scheme 19) [38]. The reaction was carried out in the presence of 2,3-dichloro-5,6-dicyano-1,4-benzoquinone (DDQ) and 1,4-naphthoquinone (NQ) in DMA at 100 °C. A variety of allylic hydrocarbons and aryl thiols were converted into the corresponding allylic sulfides in good yields. The mechanism investigation indicated that the radical process may be involved in this reaction and DDQ was employed to generate radicals.

Scheme 19. Dehydrogenative C-S coupling of α-methylstyrene and aryl thiol.

In 2019, the Xu group reported an iron-catalyzed C-H sulfenylation of oxindoles with aryl thiol for the direct synthesis of aryl sulfides (Scheme 20) [39]. The reaction was carried out in the presence of FeCl$_3$ and K$_2$CO$_3$ in DMSO at 50 °C. This method exhibited excellent substrate endurance and generally gave good to excellent yields. Besides oxindoles, phenylacetamides, pyrazolones, phenylacetonitriles, and ethyl cyanoacetates were smoothly thiolated with aryl thiols to afford the corresponding sulfides in good yields.
In 2019, the Gagnon group disclosed a direct S-cyclopropylation of thiophenols using tricyclopropylbismuth to synthesize aryl cyclopropyl sulfides (Scheme 21) [40]. The reaction was carried out in the presence of copper(II) acetate as the catalyst and pyridine as the base under mild conditions, providing the corresponding aryl cyclopropyl sulfides in moderate to good yields. A variety of functional groups including halogen, methoxyl, nitro, trifluoromethyl, and ester, were well tolerated.

**Scheme 21.** Synthesis of aryl cyclopropyl sulfide from aryl thiol.

In 2019, the Xiao and Deng group reported a multicomponent reaction of o-phenylenediamines, thiophenols, and paraformaldehyde for the synthesis of N-thiomethyl benzimidazoles via sequential C-N/C-S bond formation (Scheme 22) [41]. The reaction occurred in the presence of 4-aminodiphenylamine (4-ADPA) as base in a mixed solvent of 1,1,1,2-tetrachloroethane (TCE) and water at 130 °C. This reaction provided the 1-phenylthiomethyl benzimidazoles in moderate to good yields. Moreover, the addition of different aldehydes led to the successful introduction of various functional groups at the 2-position of the benzimidazole.

**Scheme 22.** Three-component reaction for the synthesis of 1-phenylthiomethyl benzimidazoles.

In 2019, the Wang group developed a unique C-S coupling reaction for the synthesis of α-sulfenylated ketones from thiols and TEMPO. The reaction proceeded in the presence of K₂CO₃ in N,N-dimethylacetamide (DMA) at 150 °C (Scheme 23) [42]. A variety of aryl thiols and TEMPO analogues coupled smoothly to give the corresponding α-sulfenylated ketones. In this transformation, TEMPO functioned as a C3 synthon.
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2.2.2. C(sp²)-S Bond Formation with Thiol

Decarboxylative C-S coupling reaction is an important strategy for the synthesis of sulfides. C-S bonds are readily constructed by the coupling reaction of carboxylic acid with aryl thiol, disulfides, sulfonyl hydrazides, sodium sulfinates, and sulfinic acids, respectively [43]. In 2016, the Hoover group reported a decarboxylative C-S bond formation for the synthesis of diaryl sulfides from 2-nitrobenzoic acid and aryl thiols (Scheme 24) [44]. The reaction was carried out at 140 °C in the presence of CuI as catalyst, 1,10-phen as ligand, K₂CO₃ as base and molecular oxygen as oxidant. Generally, good to excellent yields were obtained unless the 5-position of 2-nitrobenzoic acid was substituted. Heteroaryl thiols were converted into the corresponding sulfides in the same reaction system at 160 °C. Moreover, treating 2-nitrobenzoic acid with aryl disulfide afforded the thiolated compounds as well. It is interesting that pentafluorobenzoic acid was dithiolated under the optimal condition.

In 2019, the Jafarpour group reported a simple procedure for the synthesis of unsymmetrical diaryl sulfides from arylhydrazines and aryl thiols (Scheme 25) [45]. The reaction proceeded at room temperature in the presence of TBHP, KI, and Cs₂CO₃ in water, providing the corresponding sulfides in good yields. In this reaction, TBHP was converted into several types of radicals in the presence of iodide. The radicals converted arylhydrazine and aryl thiol into phenyl radical and disulfide, respectively. The final C-S coupling reaction of phenyl radical and disulfide afforded the products.

In 2019, the Moghaddam group developed a coupling reaction of arylboronic acid with aryl or alkyl thiols to prepare unsymmetrical sulfides using CoCuFe₂O₄ nanoparticles as catalyst (Scheme 26) [46]. The reaction proceeded in the presence of n-Bu₄NOH in DMF at 100 °C. This protocol was also applied in the C-N coupling reaction of arylboronic acid and anilines.
Scheme 26. CoCuFe$_2$O$_4$ nanoparticles catalyzed C-S coupling of aryl boronic acids and thiols.

In 2019, the Shankar group reported a BCl$_3$ mediated one-pot debenzylation/sulfenylation reaction for the synthesis of sulfides. 3-((Benzyloxy) methyl) imidazo[1,2-a]pyridine was converted into sulfides in moderate to good yields (Scheme 27) [47]. This method was compatible with both aryl thiols and aliphatic thiols. The proposed mechanism indicated that the reaction proceeded through two steps including BCl$_3$ mediated debenzylation and nucleophilic addition by thiols. In addition, thiols can be replaced by amine and alcohol to afford the corresponding amines and ethers, respectively.

Scheme 27. Synthesis of sulfides from 3-((benzyloxy) methyl) imidazo[1,2-a]pyridines and thiols.

In 2018, the Das group developed a C-H sulfenylation of 4-quinolones with thiols or disulfides in the presence of NaI and TBHP in DMSO (Scheme 28) [48]. The reactions of aryl thiols without a substituent at the para-position provided excellent yields while the reactions with para-substituted aryl thiols gave lower yields (about 60%).

Scheme 28. C-H sulfenylation of 4-quinolones with aryl thiols.

In 2018, the Mal group developed a C-S coupling reaction of aryl thiols and benzenes bearing multiple methyl and/or methoxy groups in the presence of PhI(OAc)$_2$ in HFIP (Scheme 29) [49]. It is interesting that the product was controlled by the amount of PhI(OAc)$_2$. When 1.0 equivalent of PhI(OAc)$_2$ was used, diaryl sulfides were predominantly formed. However, when 3.0 equivalent of PhI(OAc)$_2$ was used, the main products were diaryl sulfoxides. This reaction system was featured by a broad substrate scope. Functional groups, such as nitro, trifluoromethyl, halogen, ether, and formyl, were well endured. Moreover, aliphatic thiols were also tolerated in this protocol.
In 2019, the Yuan group reported a C-H thiolation strategy for the synthesis of aryl sulfide through the C-S coupling of thiols with functionalized arenes (Scheme 30) [50]. The protocol employed cobalt(II) phthalocyanine-tetrasodium sulfonate (CoPcS) as catalyst and the reaction occurred in water in the atmosphere of oxygen. Both naphthol and naphthylamine coupled smoothly with aryl thiols to afford the corresponding diaryl sulfides in good to excellent yields. However, the coupling of naphthol with aliphatic thiols gave relatively low yields. A variety of functional groups such as methoxy, amine, and halogen were well tolerated in this reaction system. In addition, C-H bonds of indole and 2-phenylimidazo[1,2-a]pyridine were successfully thiolated with aryl thiols using the developed protocol.

In 2018, the Lei and Tang group reported an electrochemical oxidative C-H sulfenylation of imidazopyridines with thiols (Scheme 31) [51]. The conversion was achieved in an undivided electrolytic cell using methanol and acetonitrile as cosolvent. Both aryl thiols and aliphatic thiols reacted smoothly with imidazopyridines, providing di(hetero)aryl sulfides in moderate to good yields. They further found that this C-H sulfenylation reaction can occur between imidazopyridines with diaryl disulfides.

Elemental sulfur is an attractive sulfur source for the construction of C-S bond due to the low toxicity, stability, ready availability, odorless, and economical price [52]. In 2018, the Adimurthy group reported a copper catalyzed C-H disulfenylation of imidazo[1,2-a]pyridine for the synthesis of diaryl sulfides. Multiple components reaction of heteroarene, elemental sulfur, and aryl iodide afforded the disulfenylated products in good to excellent yields (Scheme 32) [53]. The reaction was carried out at 130 °C in the presence of Cul as catalyst and KOr-Bu as a base in a mixed solvent of DMF and acetic acid. A variety of functional groups including halogen, methoxyl, boronic acid, and so on, were well...
tolerated in the developed reaction system. The mechanistic investigation showed this conversion underwent through two steps including the formation of aryl thiols from aryl iodides and elemental sulfur, and the C-H sulfonylation followed.

Scheme 32. Synthesis of disulfenylated imidazopyridines.

In 2018, the Zolfigol group reported a C-O bond activation/C-S coupling reaction for the synthesis of unsymmetrical diaryl sulfides from phenolic esters, aryl boronic acids and elemental sulfur using NiFe\(_2\)O\(_4\) nanoparticles as a catalyst (Scheme 33) [54]. The reaction was compatible with several phenolic esters including acetates, triflates, and tosylates, and occurred in PEG2000 at 90 °C to afford the products in good to excellent yields. Moreover, the three-component reaction of phenolic esters, Ph\(_3\)SnCl and sulfur also successfully afforded the unsymmetrical diaryl sulfides. They further found that the reaction of phenolic esters with sulfur can produce symmetrical diaryl sulfides in good to excellent yields at 120 °C using anhydrous DMF as a solvent or under the solvent-free condition. In their paper, they provided a detailed comparison of their developed methods with the previously reported methods in view of required agents and starting materials/coupling partners.

Scheme 33. C-S coupling reaction of phenolic esters and aryl boronic acids.

Later in 2019, the Rostami and Khakyzadeh group used phenols, sulfur, and aryl boronic acids as substrates and reported a Pd/Cu dual-metal catalyzed one-pot C-O bond activation/C-S coupling reaction for the synthesis of unsymmetrical diaryl sulfides (Scheme 34) [55]. Phenol was modified with sulfonyl or acetyl to afford the corresponding ester at the first step. Next, a synthesized novel SiO\(_2\) nanoparticles supported the Pd catalyst was used for C-O bond activation and the subsequent C-S bond formation. Compared to Zolfigol’s work [54], this protocol allowed the reaction to proceed under milder conditions. The second step reaction was completed with 6 h at room temperature, and afforded the unsymmetrical diaryl sulfides in good to excellent yields. It is noteworthy that some sensitive functional groups such as formyl and cyano group were well tolerated in the reaction system.

Scheme 34. Pd/Cu dual-metal catalyzed one-pot synthesis of diaryl sulfides.
2.4. Sulfonyl Derivative as the Sulfenylation Source

Sulfonyl derivatives such as sulfonyl chloride and sodium sulfonates are employed as sulfenylation source. They are often used in a combination of reductant to afford disulfide as ultimate sulfenylation intermediate in the conversion. In 2018, the Radosevich group developed an electrophilic sulfenylation reaction for the synthesis of sulfides from indoles and sulfonyl chlorides (Scheme 35) [56]. This conversion was catalyzed by a readily available phosphetane in conjunction with a hydrosilane terminal reductant to afford sulfenyl electrophiles, providing easy access to valuable trifluoromethyl-, perfluoroalkyl-, and heteroaryl derivatives. The reaction occurred at 40 °C in dioxane and accomplished within 12 h. This protocol was further featured by broad substrate scope. Various functional groups, such as methoxyl, halogen, aldehyde, ester, and nitro group, were well tolerated.

Scheme 35. Synthesis of sulfenylated indoles using sulfonyl chloride as a sulfenylative source.

In 2019, the Liu group reported a CuI-catalyzed C-S coupling reaction for the synthesis of diaryl sulfide from aryl boronic acid and arylsulfonyl chlorides (Scheme 36) [57]. This protocol used 1,10-phen as ligand and the conversion proceeded in the presence of K$_2$CO$_3$ and PPh$_3$ in THF/DMSO (2:1) at 50 °C. The reaction was completed within 4 h. This method was featured by broad substrate scope for both arylboronic acids and arylsulfonyl chlorides. It is worth mentioning that the conversion required oxygen as the yield dramatically decreased when the reaction was carried out in the atmosphere of nitrogen. Using arylsulfonyl chlorides as sulfenylation source, this reaction system was successfully applied in the thiolation of acetylacetone and β-naphthol. In this reaction, arylsulfonyl chloride functioned as a disulfide precursor as it was converted into diaryl disulfide by treating it with PPh$_3$. Disulfide subsequently formed a Cu(I)-S complex and coupled with arylboronic acid to afford the corresponding diaryl thioether.

Scheme 36. Synthesis of diaryl sulfides from sulfonyl chlorides and aryl boronic acids.

The Zhao and Deng group reported a facile protocol for the sulfenylation of aliphatic ethers using sodium sulfonates in the presence of iodine, di-tert-butyl peroxide (TBP) and diethyl phosphite. The reaction occurred at 130 °C in the air (Scheme 37) [58]. Initially, 1,4-dioxane was used as model substrate to react with sodium sulfinates, providing a variety of aryl sulfides in moderate to excellent yields. They further broaden the substrate scope to common cyclic and chained ethers. Moreover, sulfonyl hydrazides can also function as a thiolating reagent in this reaction system. In this conversion, TBP was converted to tert-butoxy radical under heating, and the radical grasped hydrogen from 1,4-dioxane to provide a pivotal alkyl disulfide. On the other hand, sodium sulfonate was converted to disulfide by treating with iodine and diethyl phosphite. The radical intermediate coupled with disulfide to give the final product.
2.5. Other Sulfenylation Sources

In 2018, the Li and Wen group reported a very facile and efficient protocol for the synthesis of 1,3-benzothiazepines from thioamides via dehydrogenative C-S bond formation (Scheme 38) [59]. In the presence of 4-F-HTIB (hydroxyl(4-fluorophenyl) iodotosylate), thioamides were readily converted to the corresponding 1,3-benzothiazepines in good to excellent yields. The conversion proceeded in HFIP at room temperature and were accomplished within 1 min. It is very interesting that, when 4-methoxyphenethylthioamides were employed as starting materials, a new type of thiazaspiro[5.5]undecanones other than 1,3-benzothiazepines were afforded in good yields.

Scheme 38. Intramolecular thiolation of thioamides for the synthesis of 1,3-benzothiazepines.

In 2019, the Yamaguchi group developed a Pd-catalyzed synthesis of alkenyl thioether from thioester and N-tosyl hydrazones (Scheme 39) [60]. The reaction employed Pd(OAc)$_2$ as catalyst, P(n-Bu)$_3$ as ligand, LiO(t-Bu) as base and occurred at 60 °C in THF. In the developed reaction system, thioester was the best sulfur source because other sulfur-containing reagents such as thiophenol and diphenyl disulfide resulted in lower yield and poor reaction selectivity. This method exhibited broad substrate scope as it was compatible with various functional groups, such as methoxyl, halogen, ester, amide, trifluoromethyl, etc.

Scheme 39. Synthesis of aryl alkenyl sulfides from thioesters and N-tosyl hydrazones.

In 2020, the Singh group reported an iodine-catalyzed synthesis of 3-arylthioindoles from indoles employing a 1-aryltriazene/CS$_2$ combination as the sulfenylation source (Scheme 40) [61]. The reaction was performed in ethanol at 70 °C. The corresponding diaryl sulfides were obtained in good to excellent yields. Broad substrate scope was exhibited by this protocol as functional groups including methoxyl, halogen, cyan, trifluoromethyl, and nitro group were well tolerated. Moreover, this protocol was also applied in the sulfenylation of trimethoxybenzene and 2-hydroxynaphthalene, yielding the corresponding sulfides.
2.6. Trifluoromethylthio Reagents as Sulfenylation Sources

Molecules containing trifluoromethylthio group are a unique type of sulfides and have attracted extensive attention because they have found wide application in pharmaceuticals, agrochemicals, and materials. Due to the strong electron-withdrawing effect and hydrophobic property, trifluoromethyl sulfides are important building blocks in drug design. The traditional method for the synthesis of trifluoromethylthiolated compounds is the nucleophilic trifluoromethylthiolation of aryl halides with trifluoromethylthiolated metals, such as Hg(SCF₃), Ag(SCF₃), Cu(SCF₃), Cs(SCF₃), and (bpy)Cu(SCF₃) [62]. In recent years, several novel organic trifluoromethylthio surrogates have been developed and enabled the trifluoromethylthiolation to occur to non-halide substrates.

In 2016, the Hopkinson and Glorius group reported the first visible-light-promoted radical trifluoromethylthiolation of styrenes using 2-((trifluoromethyl)thio)isoindoline-1,3-dione as trifluoromethylthio source (Scheme 41) [63]. The reaction proceeded in the presence of fac-[Ir(ppy)₃] as photocatalyst and nBu₄NBr as an activator in acetonitrile at room temperature. A variety of styrenes were smoothly converted into the corresponding trifluoromethylthiolated products in satisfying yields.

In 2017, the Xu group found a novel SCF₃ source, S-trifluoromethyl 4-methylbenzenesulfonothioate, and developed a dual gold and photoredox catalyzed reaction for the phenylsulfonyl-trifluoromethylthio-1,2-difunctionalization of alkenes (Scheme 42) [64]. S-trifluoromethyl 4-methylbenzenesulfonothioate functioned as bifunctional reagent responsible for both trifluoromethylthiolative and phenylsulfonylative reagent. Alkenes including styrenes, internal alkenes, and dienes were converted into the corresponding products in the presence of IPrAuCl, AgSbF₆, and Ru(bpy)₃Cl₂ in DCE at room temperature under irradiation with 100 W blue LED at the N₂ atmosphere. A series of functional groups, such as ester, cyano, triuromethyl, and halogen, were well tolerated under the optimized reaction condition. Moreover, it is noteworthy that the reaction system was also compatible with PhSO₂SR (R= aryl or alkyl), affording various phenylsulfonyl sulfides.
In 2017, the Dagousset and Magnier group reported the visible light driven synthesis of trifluoromethylthiol-substituted oxindoles from \( N \)-methyl-\( N \)-phenylmethacrylamides using \( N \)- trifluoromethylosoaccharin as an SCF\(_3\) radical precursor (Scheme 43) [65]. This conversion employed Ir(ppy)\(_3\) as photocatalyst and was carried out at room temperature under the irradiation of blue LEDs. Using different alkenes as substrates, the same reaction system led to the synthesis of SCF\(_3\)-containing oxindole, trifluoromethylthiolated isoquinolinediones, and diarylethyl trifluoromethyl sulfides in good yields. The reaction system was compatible with lots of functional groups, such as halogen, cyan group, acetyl, ester, and BocNH.

Later in 2018, the Zhao group applied \( S \)-trifluoromethyl 4-methylbenzenesulfonothioate as SCF\(_3\) radical precursor for the same conversion (Scheme 44) [66]. The reaction occurred in the presence of AgF and \( K_2S_2O_8\) in DMSO at 20 °C. The mechanistic studies indicated that AgSCF\(_3\) was generated from \( S \)-trifluoromethyl 4-methylbenzenesulfonothioate and AgF, and SCF\(_3\) radicals were subsequently formed by the oxidation with \( K_2S_2O_8\).

**Scheme 43.** Visible light driven synthesis of trifluoromethylthiol-substituted oxindoles.

**Scheme 44.** Synthesis of trifluoromethylthiol-substituted oxindoles from \( N \)-phenylmethacrylamides.

### 3. Synthesis of Sulfoxides via C-S Bond Construction

In 2018, the Bolm group reported a transition-metal-free arylation of \( \beta \)-sulfinyl esters with diaryliodonium salts to provide a wide range of \( S \), \( S \)-diaryl and \( S \)-alkyl \( S \)-aryl sulfoxides (Scheme 45) [67]. The reaction occurred at room temperature in the presence of KOH in the cosolvent of toluene and water. In this process, sulfenate anion was generated from \( \beta \)-sulfinyl esters and form a C-S bond with aryl group of diaryliodonium. In general, this protocol provided the corresponding diaryl sulfoxides in good to excellent yields. The yields decreased when alkyl sulfinyl esters or heteroaryliodonium were used as substrates.

**Scheme 45.** Synthesis of sulfoxides from \( \beta \)-sulfinyl esters.
In 2018, the Wang group found a facile protocol for the synthesis of diaryl sulfoxides and \( m \)-arylsulfinyl sulfoxides from arylsulfinic acids and dimethoxybenzenes via BF\(_3\) mediated C-S bond formation (Scheme 46) [68]. The reaction easily proceeded in CH\(_2\)Cl\(_2\) without any other additives. The reaction temperature was very important for the reaction selectivity. The diaryl sulfoxides were predominantly afforded when the reaction was carried out at 30 °C. Diaryl sulfoxides were further sulfonylated by arylsulfinic acid to provide \( m \)-arylsulfinyl sulfoxides at 80 °C.

![Scheme 46. Synthesis of diaryl sulfoxides from arylsulfinic acids and arenes.](image)

4. Synthesis of Sulfoxides via C-S Bond Construction

Sulfonylhydrazides and sulfonylhydrazones are frequently used as sulfonyl source for the synthesis of sulfoxides via C-S bond formation. In 2016, the Zhao group reported a denitrogenative C-S bond formation for the synthesis of disubstituted methylsulfoxides from sulfonylhydrazones (Scheme 47) [69]. Sulfonylhydrazones were readily prepared from ketones and sulfonyl hydrazides. Sulfonylhydrazones decomposed to release nitrogen and afford the methylsulfoxides in the presence of FePcCl in dioxane at 80 °C. Using this protocol, methylsulfoxides were obtained in good to excellent yields.

![Scheme 47. Denitrogenative synthesis of disubstituted methylsulfoxides from sulfonylhydrazones.](image)

In 2018, the Kleij and Yang group used \( N \)-sulfonylhydrazone as sulfonyl source and reported a synthetic method for allylic sulfoxides via decarboxylative C-S bond formation (Scheme 48) [70]. Vinylene carbonate was used as substrate. C-S coupling reaction occurred to afford allylic sulfoxides at 80 °C in acetonitrile in the presence of Pd\(_2\)(dba)\(_3\) as catalyst and K\(_2\)CO\(_3\) as base. It is worth noting that the products were obtained with high regio-selectivity. A wide range of functional groups were tolerated under the optimal condition. Moreover, changing the catalyst from Pd\(_2\)(dba)\(_3\) to Pd(PPh\(_3\))\(_4\) predominantly afforded the allylic hydrazones as products. The developed protocol was successfully applied in the functionalization of some biologically important scaffolds.

![Scheme 48. Decarboxylative C-S bond formation between vinylene carbonate and \( N \)-sulfonylhydrazone.](image)
In 2018, the Wang and Ji group reported a Cu(II)/Ag(I)-catalyzed cascade reaction for the synthesis of 2-aryl-3-sulfonyl substituted quinoline from sulfonyl hydrazine with anthraniis (Scheme 49) [71]. It is interesting that new C-C, C-N and C-S bonds were constructed in one step. The conversion proceeded at 110 °C in the presence of Cu(OAc)$_2$ and AgOTf in DCE without any other additives. No product was observed in the absence of either Cu(OAc)$_2$ or AgOTf. Phenyl sulfonyl hydrazines gave moderate yields while heteroaryl sulfonyl hydrazines were not favorable for this conversion.

Scheme 49. Synthesis of 2-aryl-3-sulfonyl substituted quinoline using sulfonyl hydrazine as sulfonyl source.

In 2018, the Liang group developed a novel protocol for the synthesis of 2-sulfonated pyrrolo[1,2-α]indoles using toluenesulfonyl hydrazide as a sulfonyl reagent (Scheme 50) [72]. N-propargyl-substituted indoles were used as substrates and the reaction occurred in the presence of Cu(NO$_3$)$_2$·3H$_2$O as catalyst, TBHP and AgTFA in DCE at 90 °C. Toluensulfonyl hydrazide proved as a good tosyl source while sodium tosylate and tosyl chloride were not able to furnish this conversion under the standard condition. The proposed mechanism indicated that the reaction of sulfonyl hydrazide with tert-butoxyl or tert-butyperoxyl radical generated from TBHP gave the sulfonyl radical. Then, an addition reaction of the sulfonyl radical occurred to the alkynyl moiety give the vinyl radical. The followed 1,2-aryl migration cascade process produced the final product.

Scheme 50. Synthesis of 2-sulfonated pyrrolo[1,2-α]indoles using sulfonyl hydrazine as a sulfonyl source.

In the same year, the Liu and Zhu group used the same sulfonylation reagent and found a new method for the synthesis of diaryl methylsulfones via sulfonylation of para-quinone methides (Scheme 51) [73]. The reaction occurred smoothly in the cosolvent of ethanol and water at 50 °C without the use of any catalyst. This protocol was further featured by broad substrate scope and good to excellent yields.

Scheme 51. C-H sulfonylation of para-quinone methides with sulfonyl hydrazides.
Later in 2019, the Du group further applied sulfonyl hydrazides as sulfonyl reagent in the one-pot synthesis of 3-sulfonylbenzofurans from o-hydroxybenzyl alcohols (Scheme 52) [74]. This reaction system was compatible with a variety of functional groups and provided the diaryl sulfones in good yields. It is noteworthy that the 3-sulfonylbenzofurans can be readily elaborated to diaryl thioethers, diarylmethanones, and Suzuki coupling products.

Scheme 52. Synthesis of 3-sulfonylbenzofurans from o-hydroxybenzyl alcohols and sulfonyl hydrazines.

In 2017, the Reiser group reported a temperature controlled selective C-S or C-C formation from heteroarenes and aryl sulfonyl chloride for the synthesis of diaryl sulfones or aryl heteroarenes (Scheme 53) [75]. The reaction employed $\text{fac}[\text{Ir}(ppy)_3]$ as photocatalyst, and was carried out in anhydrous acetonitrile under the irradiation of LED455. Heteroarenes, such as thiophene, pyrrole, and indole were well tolerated in this conversion. It is very interesting that the reaction at room temperature gave diaryl sulfones as a product, but the reaction at 60 $^\circ$C provided aryl heteroarenes as the product.

Scheme 53. C-H sulfonylation of heteroarenes with phenylsulfonyl chloride.

In 2019, the Deng group developed a three-component reaction for the preparation of diarylmethyl sulfones from aldehydes, arenes, and sodium sulfinates via tandem C-C/C-S bond formation (Scheme 54) [76]. The reaction proceeded at 60 $^\circ$C in water in the presence of ion exchange resin amberlyst 15. Compared with previously reported methods, this protocol was featured by simple procedure, no requirement of metal catalysts and use of recyclable heterogeneous catalyst, and organic solvent-free condition, providing a green method to synthesize diarylmethyl sulfones.

Scheme 54. Three-component reaction for the preparation of diarylmethyl sulfones.

In 2017, the Willis group developed a direct photoinduced sulfonylation of $N$-alkylated anilines with sodium sulfinates using an iridium complex as catalyst (Scheme 55) [77]. The reactions were carried out under the irradiation of blue LEDs in the presence of $\text{Bu}_4\text{NHSO}_4$ and $\text{K}_2\text{S}_2\text{O}_8$ in the cosolvent of acetonitrile and water. This protocol exhibited regio-selectivity and provided good overall
yields. Very broad substrate scope and good functional group endurance were observed with this protocol. Both alkylsulfonates and arylsulfonates were well tolerated. Moreover, they showed the produced sulfones were readily converted into diverse derivatives. The mechanism investigation indicated that oxidation of the aniline and the sulfinate salt by the iridium catalytic cycle afforded aniline radical cation and sulfonyl radical, respectively. The two radicals coupled to provide the sulfone product via C-S bond formation.

\[ \text{R}_1\text{R}_2\text{R}_3 + \text{S}^–\text{Na} \rightarrow \text{S}\text{R}_4^– \rightarrow \text{R}_1\text{R}_2\text{R}_3\text{S}\text{R}_4 \]

Scheme 55. Photoinduced sulfonylation of N-alkylated anilines with sodium sulfonates.

In 2019, the Waldvogel group reported an electrochemical sulfonylation of arenes containing electron donating group using sodium sulfonates as a sulfonylation reagent (Scheme 56) [78]. The reaction employed BDD as electrode and HFIP (1,1,1,3,3,3-hexafluoropropan-2-ol)/water as cosolvent. The sulfonates were used as both coupling component and supporting electrolyte. The arenes containing electron donating groups, such as methoxy, methyl, dimethylamino and acetylamino, were readily sulfonated with either sodium arylsulfonates or sodium alkylsulfonates.

\[ \text{R}_1\text{R}_2\text{R}_3 + \text{S}^–\text{Na} \rightarrow \text{S}\text{R}_4^– \rightarrow \text{R}_1\text{R}_2\text{R}_3\text{S}\text{R}_4 \]

Scheme 56. Electrochemical sulfonylation of arenes with sodium sulfonates.

In 2018, the Song group developed the first example of Rh(II)/phosphine-cocatalyzed reaction system for the synthesis of dithioketals from diazo compounds through simultaneous construction of two different C-S Bonds (Scheme 57) [79]. Diazoacetate reacted with $S$-methyl benzenesulfonylthioate at room temperature in the presence of $\text{Rh}_2(\text{OAc})_4$ and dpdp in DCE, providing dithioketal derivatives in moderate to good yields.

\[ \text{N}_2\text{CO}_2\text{R}_1 + \text{S}\text{O}_\text{S}\text{R}_4 \rightarrow \text{R}_1\text{R}_2\text{R}_3\text{S}\text{O}_\text{S}\text{R}_4 \]

Scheme 57. Synthesis of dithioketals from diazo compounds.

In 2020, the Reddy group developed an interesting strategy for the synthesis of benzyl disulfanylsulfones from $S$-benzyl thiosulfonates via phenylboronic acid catalyzed tandem construction of S-S and C-S Bonds (Scheme 58) [80]. The reaction occurred in the presence of $\text{K}_2\text{CO}_3$ in PhCl at 90 °C, providing the corresponding benzyl disulfanylsulfones in good to excellent yields.
5. Conclusions and Prospects

The research on the synthetic methods for organosulfur compounds is of great importance in both academic research and industry. In the past several years, significant progresses have been made in the field of direct C-S bond construction for the synthesis of organosulfur compounds. Various synthetic methods have been developed to provide easy accesses to sulfides, sulfoxides, and sulfones from the non-halide substrates. Unlike C-S coupling reaction of halide substrates, a radical pathway was often involved in the C-S coupling reaction of non-halide substrates. It is satisfying that many reactions are featured by mild conditions, high efficiency and excellent selectivity by the employment of radical-radical coupling strategies. The Diazo compound is frequently used as carbon radical precursor to couple with the thiyl radical; however, this transformation may be limited by pre-functionalization of the substrate and poor atom-economy. C-H functionalization represents a promising strategy to produce carbon radicals from simple arenes and alkanes. A variety of C(sp^3)-H and C(sp^2)-H bonds are successfully transformed to the corresponding sulfides, sulfoxides, and sulfones. On the other hand, in addition to reactions catalyzed by traditional transition metals, novel protocols involving photocatalysts have been extensively developed in very recent years. Electrolysis mediated reactions also found application in the C-S coupling reaction for the synthesis of sulfides and sulfones from (hetero)arenes. These novel protocols enable the C-S coupling reaction to occur in a simple, economical, and efficient reaction system. In the future, C-S bond formation should get more benefits from photo-catalyzed and electrochemical reaction, and more novel protocols in this field can be expected.

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