



Advanced Synthesis & Catalysis

Accepted Article

Title: Iron-Mediated Oxidative C-H Alkylation of *S*,*S*-Functionalized Internal Olefins via C(sp²)-H/C(sp³)-H Cross-Coupling

Authors: Quannan Wang, Jiang Lou, Ping Wu, Kaikai Wu, and Zhengkun Yu

This manuscript has been accepted after peer review and appears as an Accepted Article online prior to editing, proofing, and formal publication of the final Version of Record (VoR). This work is currently citable by using the Digital Object Identifier (DOI) given below. The VoR will be published online in Early View as soon as possible and may be different to this Accepted Article as a result of editing. Readers should obtain the VoR from the journal website shown below when it is published to ensure accuracy of information. The authors are responsible for the content of this Accepted Article.

To be cited as: *Adv. Synth. Catal.* 10.1002/adsc.201700654

Link to VoR: <http://dx.doi.org/10.1002/adsc.201700654>

DOI: 10.1002/adsc.201700654((will be filled in by the editorial staff))

Iron-Mediated Oxidative C–H Alkylation of *S,S*-Functionalized Internal Olefins via C(sp²)–H/C(sp³)–H Cross-Coupling

Quannan Wang,^{a,b} Jiang Lou,^{a,b} Ping Wu,^{a,b} Kaikai Wu,^a and Zhengkun Yu^{*,a,c}^a Dalian Institute of Chemical Physics, Chinese Academy of Sciences, 457 Zhongshan Road, Dalian, Liaoning 116023, People's Republic of China^b University of Chinese Academy of Sciences, Beijing 100049, People's Republic of China^c State Key Laboratory of Organometallic Chemistry, Shanghai Institute of Organic Chemistry, Chinese Academy of Sciences, 354 Fenglin Road, Shanghai 200032, People's Republic of China
Fax: +86-411-8437-9227; e-mail: zkyu@dicp.ac.cn

Received: ((will be filled in by the editorial staff))

Supporting information for this article is available on the WWW under <http://dx.doi.org/10.1002/adsc.201700654>. ((Please delete if not appropriate))

Abstract. Catalytic amount of FeCl₃-mediated oxidative C–H alkylation of *S,S*-functionalized internal olefins, that is, α -oxo ketene dithioacetals and the analogs, was efficiently achieved by means of simple ethers and toluene derivatives as the coupling partners, di-*tert*-butyl peroxide (DTBP) as the oxidant, and DABCO·6H₂O as the additive. The alkylthio functionality is essential for the internal olefinic C–H bond to undergo such alkylation with the *O*-adjacent C(sp³)–H bonds of ethers and benzylic C–H bonds of toluene derivatives, respectively. Tetrasubstituted olefins were thus synthesized and further transformed to highly sub-

stituted pyrazoles and isoxazoles. The strategy to activate an internal olefinic C–H bond by polarizing its parent olefinic C=C bond with both the dialkylthio and an electron-withdrawing functionality was investigated. The mechanistic studies suggest a radical pathway for the C(sp²)–H/C(sp³)–H cross-coupling reactions. The present protocol provides a convenient route to tetrasubstituted olefins.

Keywords: iron; C–H functionalization; internal olefins; C–H/C–H cross-coupling; alkylation

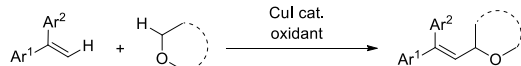
Introduction

Multisubstituted olefin motifs are abundant in numerous natural products, pharmaceuticals, and functional materials.^[1] Continuous efforts have been made to synthesize multisubstituted olefins, and transition-metal-catalyzed cross-coupling has been proven to be one of the most useful synthetic methods in this area.^[2] Although vinyl halides,^[3] vinyl pseudohalides such as vinyl boronic acids and boronates,^[4] vinyl-metals such as vinylzirconium,^[4a] vinylzinc,^[4c,5a] vinylmagnesium,^[5b] and vinylaluminum^[5c] compounds, and other reagents^[6] can be utilized for this purpose, more concise and alternative methods have been strongly desired. Recently, direct C–H functionalization has emerged as an important method for the construction of a C–C bond owing to atom-economy, operational simplicity, and environmental friendliness.^[7] In this regard, cross-dehydrogenative-coupling (CDC)^[8] has been paid particular attention, and transition-metal-catalyzed oxidative C(sp³)–H functionalization of ethers, toluene derivatives, cycloalkanes, and common alkanes have been documented.^[9] However, oxidative C(sp²)–H alkylation of olefins remains challenging and only limited examples have recently been reported. Yu, et al. reported palladium-catalyzed

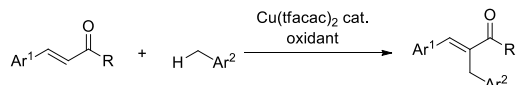
CDC reactions of terminal olefins via directed C(sp³)–H activation.^[10] Sanford, et al. realized pyridine-directed, palladium-catalyzed aerobic C(sp³)–H bond cleavage to establish an olefinic C(sp²)–C(sp³) bond.^[11] Wang and coworkers documented copper/organo-catalyzed asymmetric oxidative cross-coupling reactions between tertiary amines and olefins using molecular oxygen as the oxidant.^[12] Liu, et al. synthesized 1*H*-indenes through DDQ-mediated cross-coupling between vinylic C(sp²)–H bonds and benzylic C(sp³)–H bonds.^[13] Wei lab achieved copper-catalyzed oxidative direct alkylation of styrenes with unactivated alkanes.^[14] Lei, et al. developed Cu(I)-catalyzed oxidative cross-coupling between terminal olefinic C–H bonds and C(sp³)–H bonds of simple ethers for the synthesis of allylic ethers (Scheme 1a).^[15] Toluene derivatives were also employed for similar transformations of enone derivatives (Scheme 1b).^[16] Transition-metal-catalyzed oxidative cross-coupling of coumarins and flavones with unactivated C(sp³)–H bonds were recently reported.^[17] Unfortunately, transition-metal-catalyzed oxidative alkylation of olefins has been focused on terminal olefinic C(sp²)–H bonds, implicating that direct C–H alkylation of internal olefins still remains a challenge due to the steric hindrance and electronic impact from the internal olefin substrates.

During the ongoing investigation of the reactivities of internal olefins, we have recently developed a strategy to activate internal olefinic C–H bonds.^[18,19] Introduction of an electron-withdrawing group such as carbonyl, ester, or cyano, and one or two electron-donating alkylthio functionalities to the two ends of an olefinic C=C bond can intrinsically polarize the carbon-carbon double bond and thus activate its olefinic C–H bond. In this regard, we successfully furnished tetrasubstituted olefins through transition-metal-catalyzed oxidative C–H functionalization of ketene dithioacetals^[20] as the polarized internal olefin substructures,^[18b,18e,f] and an iron-catalyzed alkylation method was also established with styrenes as the coupling partners (Scheme 1c).^[21] Due to the advantages of iron catalysis in C–H functionalization,^[8e,9e,22] we reasonably envisioned that simple iron salts might be applied as the catalysts or mediators in the CDC reactions of such internal olefinic C–H bonds. Herein, we disclose FeCl₃-mediated oxidative C–H alkylation of ketene dithioacetals with simple ethers and toluene derivatives for the synthesis of tetrasubstituted olefins (Scheme 1d).^[23]

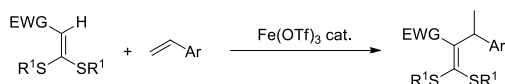
(a) Copper-catalyzed alkylation of terminal olefins with ethers^[15]



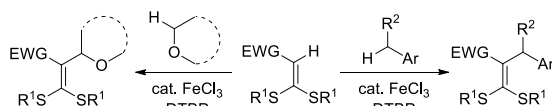
(b) Copper-catalyzed benzylation of enones with toluene derivatives^[16]



(c) Previous work^[21]



(d) This work



Scheme 1. Oxidative C–H alkylation of olefins.

Results and Discussion

Initially, the reaction of α -benzoyl ketene dithioacetal **1a** and 1,4-dioxane (**2a**) was conducted to screen the reaction conditions (Table 1). In the presence of 10 mol% FeCl₃ and 2 equivalents of di-*tert*-butyl peroxide (DTBP) as the oxidant, the reaction was continued in 1,4-dioxane (2 mL) at 120 °C for 24 h under a nitrogen atmosphere, forming the target C–H alkylation product **3a** in 9% yield (Table 1, entry 1). Addition of 1,5-diaza-bicyclo[5.4.0]-5-undecene (DBU) (10 mol%) obviously enhanced the yield to 40%, and use of 5 mol% FeCl₃ remarkably improved the yield to 67% (Table 1, entries 2 and 3). Moreover, replacement of DBU by 1,4-diaza[2.2.2]bicyclo-octane hydrate (DABCO•6H₂O) further increased the product yield (69%), presumably due to its better

stabilization of the iron(III) salt mediator during the reaction (Table 1, entry 4). Elevating the reaction temperature to 130 or 140 °C resulted in **3a** in 85–86% yields (Table 1, entries 5 and 6). The reaction could not efficiently undergo in benzene, and did not occur in both DMF and DMSO (Table 1, entries 7–9). An air atmosphere deteriorated the reaction efficiency to give **3a** in 71% yield, and the best yield (92%) was obtained in the case of using 3 equiv DTBP as the oxidant (Table 1, entries 10 and 11). It was noted that without either of FeCl₃ and the oxidant the reaction did not happen. Product **3a** was not detected in the reaction mixture of **1a** and **2a** by GC analysis in the absence of the FeCl₃ catalyst, revealing that DTBP itself could not promote the desired reaction without the catalyst (Table 1, entries 12 and 13).

Table 1. Screening of reaction conditions.^[a]

Entry	FeCl ₃ [mol%]	Additive [mol%]	Solvent	Temp [°C]	Yield ^[b] [%]
1	10		1,4-dioxane	120	9
2	10	DBU	1,4-dioxane	120	40
3	5	DBU	1,4-dioxane	120	67
4	5	DABCO•6H ₂ O	1,4-dioxane	120	69
5	5	DABCO•6H ₂ O	1,4-dioxane	130	86
6	5	DABCO•6H ₂ O	1,4-dioxane	140	85
7 ^c	5	DABCO•6H ₂ O	benzene	130	59
8 ^[c]	5	DABCO•6H ₂ O	DMF	130	0
9 ^[c]	5	DABCO•6H ₂ O	DMSO	130	0
10 ^[d]	5	DABCO•6H ₂ O	1,4-dioxane	130	71
11 ^[e]	5	DABCO•6H ₂ O	1,4-dioxane	130	92(80) ^[f]
12 ^[e]	0	DABCO•6H ₂ O	1,4-dioxane	130	0
13 ^[g]	5	DABCO•6H ₂ O	1,4-dioxane	130	0

^[a] Conditions: **1a** (0.3 mmol), additive (0.03 mmol), DTBP (0.6 mmol), solvent (2 mL), in a 25-mL sealed tube, 0.1 MPa N₂, 24 h.

^[b] Determined by GC analysis with mesitylene as the internal standard.

^[c] Using 1,4-dioxane (3 mmol).

^[d] Under an air atmosphere.

^[e] Using 3 equiv DTBP.

^[f] Isolated yield given in parentheses.

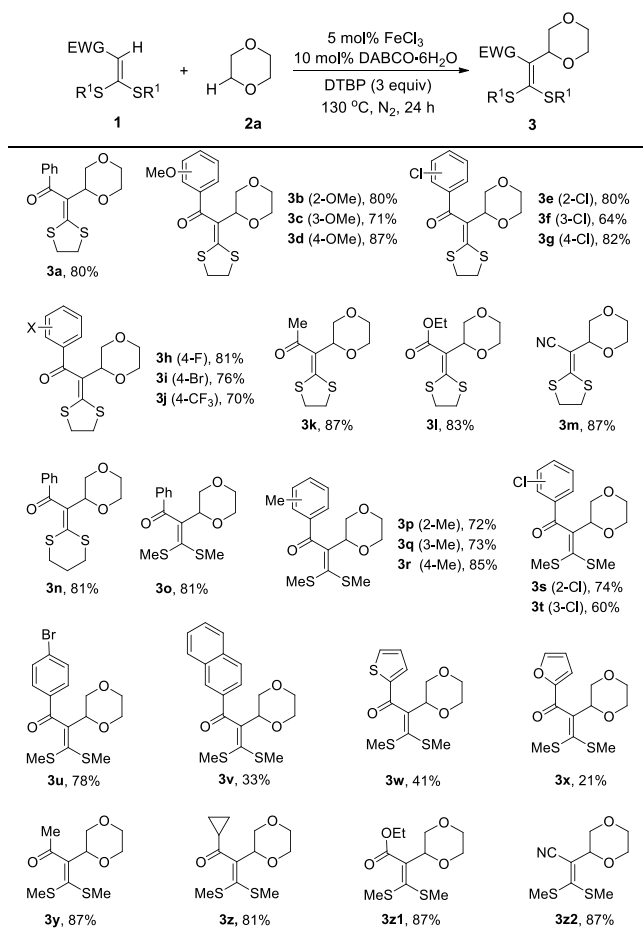
^[g] Without DTBP.

Under the optimal conditions the scope of ketene dithioacetals **1** was investigated by means of their reactions with 1,4-dioxane (**2a**) (Table 2). Cyclic α -benzoyl ketene dithioacetal (**1a**) reacted with **2a** to afford the target product **3a** in 80% isolated yield. In a similar fashion, the reactions of methoxy- and chloro-substituted aryl ketene dithioacetals with **2a** gave products **3b–3g** (64–87%). It is noted that 3-MeO and 3-Cl substituents exhibited a negative impact on the reaction efficiency. The *para*-substituents, i.e., 4-F, 4-Br, and 4-CF₃ on the aryl

moiety, demonstrated various electronic effects, leading to products **3h–3j** in 70–81% yields. Acetyl, ester, and cyano-functionalized ketene dithioacetals also efficiently underwent the reactions to generate

reactant and solvent for the reactions. The molecular structure of tetrasubstituted olefin product **3a** was further confirmed by the X-ray single crystallographic structural analysis (Figure 1).^[24]

Table 2. Scope of ketene dithioacetals **1**.^[a]

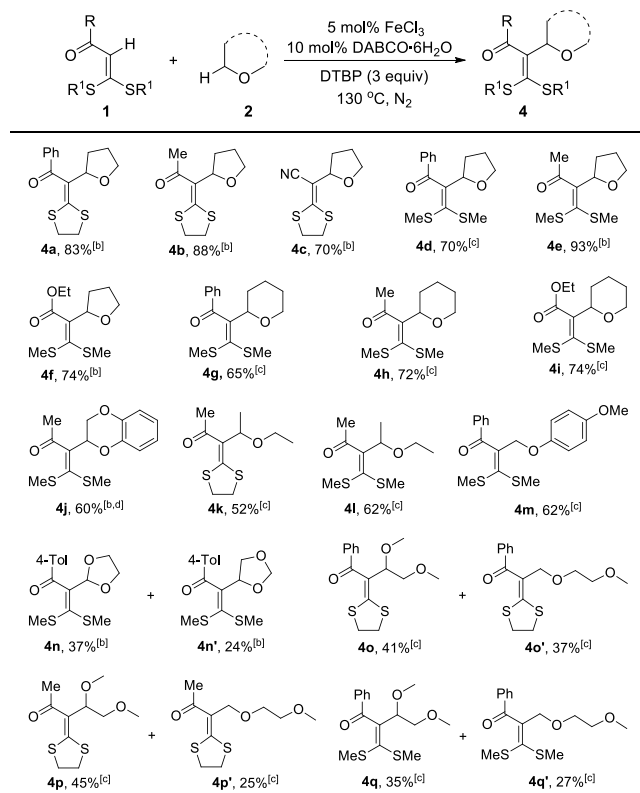


^[a] Conditions: **1** (0.5 mmol), **2a** (3 mL), FeCl₃ (0.025 mmol), DABCO·6H₂O (0.05 mmol), DTBP (1.5 mmol), 130 °C, in a 25-mL sealed tube, 0.1 MPa N₂, 24 h. Yields refer to the isolated products.

3k–3m (83–87%). Enlarging the dithioalkyl ring of **1** or using the open-chain ketene dithioacetal substrates did not affect the formation of the target products **3n** (81%) and **3o** (81%). In the cases of using the open-chain methyl-substituted α -benzoyl ketene di(methylthio)acetals products **3p–3r** (72–85%) were obtained, showing a steric effect from the 2-methyl substituent. The open-chain α -chlorobenzoyl ketene dithioacetals exhibited a reactivity lower than that of their cyclic analogs, forming **3s** (74%) and **3t** (60%), respectively. However, 4-Br-substituted di(methylthio)acetal exhibited a reactivity similar to its cyclic analog to produce **3u** in 78% yield. Bulky α -(2-naphthoyl) and α -heteroaryl (e.g., 2-furoyl and 2-thienoyl) ketene dithioacetals reacted with **2a** less efficiently, only yielding **3v–3x** in 21–41% yields. For the open-chain acetyl, cyclopropylcarbonyl, ester or cyano-functionalized ketene di(methylthio)acetals their oxidative C–H/C–H cross-coupling reactions with **2a** efficiently underwent to afford **3y–3z2** in 81–87% yields. It is noteworthy that 1,4-dioxane acted as both the

Figure 1. Molecular structure of compound **3a**.

Next, the generality of the protocol was explored by using various ethers **2** as the coupling partners (Table 3). With cyclic α -oxo and α -cyano ketene dithioacetals as the internal olefin substrates their reactions with tetrahydrofuran (THF) afforded **4a** (83%), **4b** (88%), and **4c** (70%), respectively. As compared to the corresponding reactions of the same olefin substrates with 1,4-dioxane (**2a**) as shown in Table 2, the formation of **4a** and **4b** was superior to that of **3a** (80%) and **3k** (87%), while the generation of **4c** was inferior to that of **3m** (87%). The reaction of α -acetyl ketene di(methylthio)acetal with THF efficiently formed **4e** in 93% yield, whereas in other cases THF reacted less efficiently than **2a** to give **4d** (70%) and **4f** (74%) over a longer period of 48 h. The reactivity of tetrahydropyran (THP) was lower than THF that its reactions with α -oxo ketene di(methylthio)acetals only produced **4g** (65%), **4h** (72%), and **4i** (74%) in moderate yields. 1,4-Benzodioxane also exhibited a lower reactivity than aliphatic 1,4-dioxane to react with α -acetyl ketene di(methylthio)acetal to form **4j** (60%). Acyclic ether, e.g., diethyl ether, could undergo the oxidative cross-coupling reactions via its methylene C–H bond, affording **4k** (52%) and **4l** (62%), respectively. 1,4-Dimethoxybenzene showed a moderate reactivity to form **4m** (62%), while anisole did not undergo the desired reaction under the stated conditions, which is presumably attributed to the negative electronic effect from the ether substrate. Two kinds of C(sp³)–H bonds are present in 1,3-dioxolane which participated in the reaction to generate two separable isomers **4n** (37%) and **4n'** (24%), demonstrating that the O,O-methylene C–H bond is more reactive than the O-methylene C–H bond. Glycol dimethyl ether reacted with the cyclic α -benzoyl ketene acetal, also affording two regioisomers **4o** (41%) and **4o'** (37%). Similar results were obtained from the reactions of α -acetyl and α -benzoyl ketene dithioacetals, yielding **4p** (45%) and **4p'** (25%), and **4q** (35%) and **4q'** (27%), respectively, revealing that the O-methylene C–H bond is more reactive than the methoxyl C–H in glycol dimethyl ether.

Table 3. Scope of ethers^{2[a]}

[a] Conditions: **1** (0.5 mmol), **2** (3 mL), FeCl₃ (0.025 mmol), DABCO·6H₂O (0.05 mmol), DTBP (1.5 mmol), 130 °C, in a 25-mL sealed tube, 0.1 MPa N₂, 24–48 h. Yields refer to the isolated products.

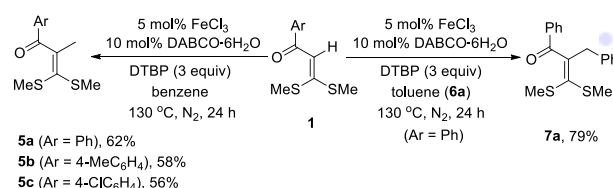
[b] 24 h.

[c] 48 h.

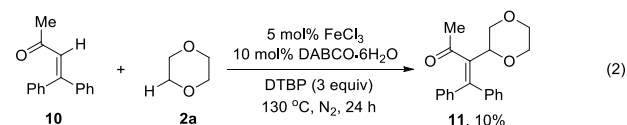
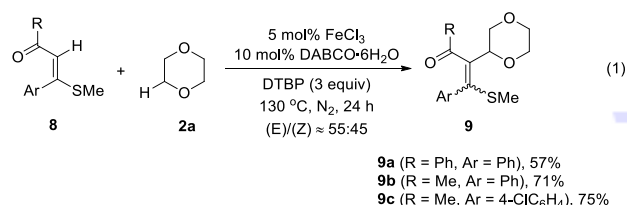
[d] 1.5 mmol of ether in benzene (3 mL).

During the screening of conditions for the reactions of **1** with **2**, control experiments were performed in the absence of **2** in benzene or toluene solvent under the standard conditions (Scheme 2). In benzene α -aroyl ketene dithioacetal substrates underwent olefinic C(sp²)–H methylation to form **5a–5c** (56–62%) with DTBP as the methylating agent,^[25] while in toluene benzylation occurred to yield olefinic C(sp²)–H benzylation product **7a** (79%). Thus, C–H benzylation of α -oxo ketene dithioacetals **1** was investigated by replacement of the ether substrates with toluene derivatives as the coupling partners (Table 4). The reactions underwent efficiently with toluene and its analogs to afford the C–H benzylated olefin products **7a–7h** (73–79%). Both electron-donating and withdrawing groups such as MeO, Me, F, Cl, and CO₂Me were tolerant on the aroyl moieties of the internal olefin substrates, and no obvious steric effect was observed. Multimethyl-substituted toluene derivatives such as *o*-, *m*-, and *p*-xylenes, and mesitylene efficiently reacted with α -benzoyl ketene di(methylthio)acetal to form the benzylation products **7i–7l** (80–87%). The reaction of 4-methoxy toluene gave two separable regioisomers **7m** (59%) and **7m'** (17%), revealing that the benzylic C–H bond is more reactive than the methoxy C–H bond in the substituted toluene substrate. 4-*tert*-Butyl

and 2-chlorotoluenes also reacted with the internal olefin, yielding **7n** (79%) and **7o** (79%), respectively. However, 3-Br, 4-F, and 4-acetyl-substituted toluene derivatives exhibited a moderate reactivity, and their reactions produced **7p–7r** in 51–69% yields. Both acyclic α -acetyl and cyclic α -benzoyl ketene dithioacetals reacted well with toluene and its derivatives to give **7s–7v** (68–88%). Unexpectedly, ethylbenzene could demonstrate a moderate reactivity to form the relatively sterically hindered benzylation products **7w** and **7x** (55–59%). These results have supplemented the previous work on C–H benzylation of electron-rich arenes^[26] and activated olefins^[22c] through iron-catalyzed oxidative C–H/C–H cross-coupling.

**Scheme 2.** Reactions of **1** in the absence of ethers.

In order to further probe into the C–H bond reactivity of the polarized internal olefins, the impact of alkylthio(s) was investigated. Reacting α -acetyl and α -benzoyl ketene (methylthio)acetals **8** with 1,4-dioxane (**2a**) under the standard conditions gave the oxidative C(sp²)–H/C(sp³)–H cross-coupling products **9a–9c** in 57–75% yields [eq (1)]. As compared to the corresponding di(methylthio)acetals, these monothioacetal substrates exhibited a lower reactivity



to **2a**. In the case of using 4,4-diphenylbut-3-en-2-one (**10**) as the internal olefin substrate, the desired reaction could not effectively occur under the stated conditions, only forming the target product **11** in 10% yield [eq (2)]. These results have unambiguously demonstrated that the alkylthio functionality is necessary for the C–H functionalization of internal olefins **1**.^[18b,18e,f]

Then, the potential applications of the target tetrasubstituted olefin products were explored (Scheme 3). Treatment of **3o**, **3y**, **3z**, and **7s** with excessive amount of hydrazine hydrate (10 equiv) in refluxing toluene afforded multisubstituted pyrazole derivatives **12a–12d** in excellent yields (87–96%).

Table 4. C–H benzylation of α -oxo ketene dithioacetals **1**.^[a]

1	6		7
----------	----------	--	----------

7a, 79%

7b, 75%

7c, 77%

7d, 77%

7e, 79%

7f, 74%

7g, 75%

7h, 73%

7i, 87%

7j, 85%

7k, 81%

7l, 80%

7m, 59%

7m', 17%

7n, 79%

7o, 79%

7p, 69%

7q, 69%

7r, 51%

7s, 76%

7t, 88%

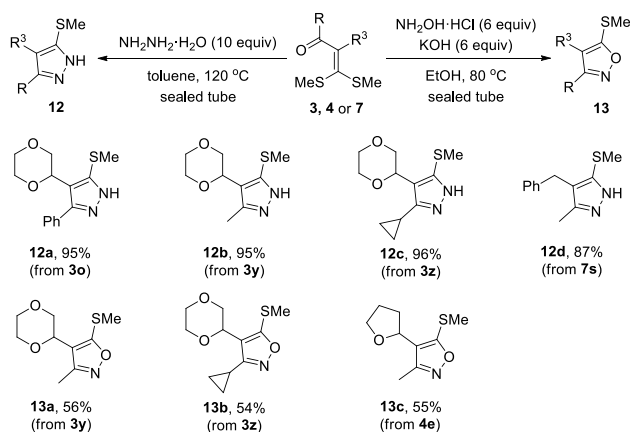
7u, 87%

7v, 68%

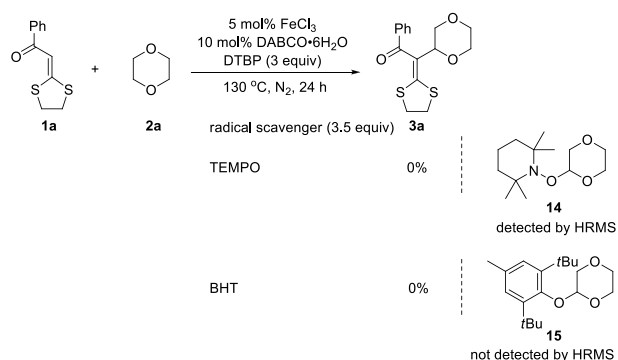
7w, 55%

7x, 59%

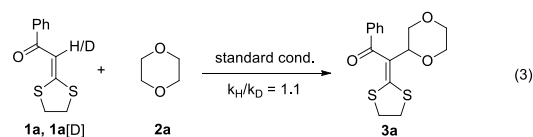
^[a] Conditions: **1** (0.5 mmol), **6** (3 mL), FeCl₃ (0.025 mmol), DABCO·6H₂O (0.05 mmol), DTBP (1.5 mmol), 130 °C, in a 25-mL sealed tube, 0.1 MPa N₂, 24 h. Yields refer to the isolated products.

**Scheme 3.** Synthetic applications of tetrasubstituted olefins.

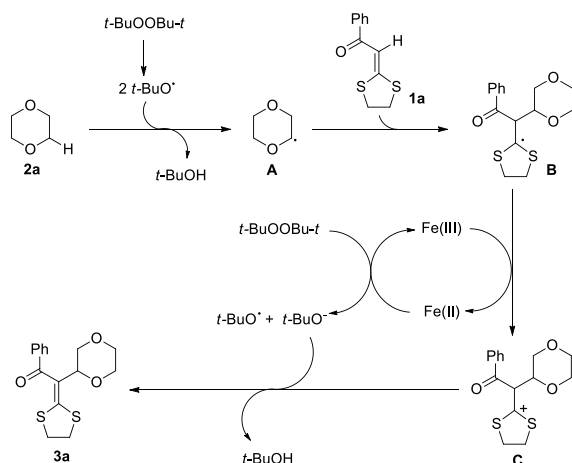
The reactions of **3y**, **3z**, and **4e** with hydroxylamine (6 equiv) in refluxing ethanol gave fully substituted isoxazoles **13a–13c** (54–56%). These results have depicted the potential applications of the resultant tetrasubstituted olefins in the synthesis of trisubstituted pyrazoles^[27] and fully functionalized isoxazoles.

**Scheme 4.** Radical trapping experiments.

Additional control experiments were carried out to probe into the reaction mechanism. Introduction of a radical scavenger such as 2,2,6,6-tetramethyl-1-piperidinyloxy (TEMPO) or 2,6-di-*tert*-butyl-4-methylphenol (BHT) (3.5 equiv) to the reaction of α -oxo ketene dithioacetal **1a** and 1,4-dioxane (**2a**) completely inhibited the oxidative C–H/C–H cross-coupling reaction (Scheme 4), suggesting that a radical pathway is involved in the catalytic cycle. The alkyl-scavenger adduct **14** was detected in the reaction mixture of **1a** with **2a** by HRMS determination, but it was not successfully isolated (see the Supporting Information). The kinetic isotope effect (KIE) experiments were also conducted under the standard conditions by means of the reactions of **1a** and its deuterated form **1a**[D] with **2a** [eq (3)]. A secondary isotope effect was observed with $k_{\text{H}}/k_{\text{D}} = 1.1$, which implicates that the internal olefinic C(sp²)–H bond cleavage is not involved in the rate-determining step in the overall catalytic cycle (see the Supporting Information for details).



A plausible mechanism is proposed in Scheme 5. Initially, the homolytic cleavage of DTBP generates two *tert*-butoxy radicals. Subsequent interaction of ether **2a** with *tert*-butoxy radical yields new radical species **A** with release of *tert*-butyl alcohol by hydrogen abstraction from the C–H bond adjacent to the ether oxygen atom. Addition of radical **A** to the internal olefinic C=C bond of α -oxo ketene dithioacetal **1a** results in radical **B** which then reacts with the high valent Fe(III) salt to form cationic species **C** and the low valent Fe(II) species through a single electron transfer (SET) process. Oxidation of the Fe(II) species with DTBP regenerates the Fe(III) mediator, and produces *tert*-butoxy anion as well as the *tert*-butoxy radical species. The alkylthio functionalities may help to stabilize the cationic species **C** due to their electron-donating property. Abstraction of a proton from species **C** by basic *tert*-butoxy anion generated from DTBP affords the target product **3a**.



Scheme 5. Proposed mechanism.

Conclusion

In summary, iron-mediated oxidative C–H alkylation of *S,S*-functionalized internal olefins has been efficiently realized by means of simple ethers and toluene derivatives as the cross-coupling partners. Tetrasubstituted olefins were thus synthesized and efficiently applied for the preparation of trisubstituted *NH*-pyrazoles and fully functionalized isoxazoles. The present synthetic protocol provides a convenient route to fully functionalized olefins.

Experimental Section

General Considerations

^1H and $^{13}\text{C}\{^1\text{H}\}$ NMR spectra were recorded on a 400 MHz spectrometer and all chemical shift values refer to CDCl_3 ($\delta(^1\text{H})$, 7.26 ppm and $\delta(^{13}\text{C})$, 77.16 ppm). X-Ray crystallographic analysis was achieved by the Analysis Center, Dalian Institute of Chemical Physics, Chinese Academy of Sciences. The HRMS analysis was obtained by ESI on a GC-TOF mass spectrometer. Column chromatographic purifications were performed on silica gel. All the chemical reagents were purchased from commercial sources and used as received unless otherwise indicated.

Typical Procedure for the C–H alkylation of ketene dithioacetals **1** with ethers **2**

Under a nitrogen atmosphere, a mixture of α -oxo ketene dithioacetal **1** (0.5 mmol), FeCl_3 (4 mg, 0.025 mmol), $\text{DABCO} \cdot 6\text{H}_2\text{O}$ (11 mg, 0.05 mmol), and DTBP (219 mg, 1.5 mmol) in 1,4-dioxane (**2a**) (3 mL) was stirred in a 25-mL sealed tube at 130 °C for 24 h. After cooled to ambient temperature, the mixture was evaporated all the volatiles under reduced pressure. The resultant residue was purified by silica gel column chromatography (eluent: petroleum ether (60–90 °C)/ethyl acetate = 4:1, v/v) to afford the target product **3** or **4**.

2-(1,4-Dioxan-2-yl)-2-(1,3-dithiolan-2-ylidene)-1-phenylethanone (3a): 123 mg, 80% yield; pale yellow solid, m.p. 129–131 °C. ^1H NMR (400 MHz, CDCl_3 , 25 °C, TMS) δ 7.69 (d, J = 7.1 Hz, 2 H, aromatic CH), 7.48 (t, 1 H, aromatic CH), 7.40 (t, 2 H, aromatic CH), 4.71 (dd, J = 10.5 and 3.0 Hz, 1 H, OCHCH_2O), 4.12 and 3.89 (m each, 1:1 H, OCHCH_2O), 3.72 (m, 2 H, $\text{CH}_2\text{OCH}_2\text{CH}_2\text{O}$), 3.62 (m) and 3.53 (dd, J = 11.7, 2.9 Hz) (1:1 H, $\text{CHOCH}_2\text{CH}_2\text{O}$), 3.42, 3.31 and 3.16 (m, 1:2:1 H, $\text{SCH}_2\text{CH}_2\text{S}$). ^{13}C NMR (100 MHz, CDCl_3 , 25 °C, TMS) δ 193.2 (Cq, C=O), 163.1 and 138.6 (Cq), 131.8, 128.5, and 128.4 (aromatic CH), 121.9 (Cq), 76.7 (OCHCH_2O), 67.2 (OCHCH_2O), 66.7 and 65.7 ($\text{OCH}_2\text{CH}_2\text{O}$), 37.4 and 37.2 ($\text{SCH}_2\text{CH}_2\text{S}$). HRMS (ESI) calcd for $\text{C}_{15}\text{H}_{16}\text{O}_3\text{S}_2$ [$\text{M}+\text{H}$] $^+$: 309.0619; found: 309.0617.

2-(1,4-Dioxan-2-yl)-2-(1,3-dithiolan-2-ylidene)-1-(2-methoxyphenyl)et-hanone (3b): 133 mg, 80% yield; pale yellow solid, m.p. 157–158 °C. ^1H NMR (400 MHz, CDCl_3 , 25 °C, TMS) δ 7.34 (m, 1 H, aromatic CH), 7.18 (d, J = 7.4 Hz, 1 H, aromatic CH), 6.95 (t, 1 H, aromatic CH), 6.89 (d, J = 8.3 Hz, 1 H, aromatic CH), 4.50 (dd, J = 10.6 and 3.0 Hz, 1 H, OCHCH_2O), 4.08 (t) and 3.85 (d, J = 10.7 Hz) (1:1 H, OCHCH_2O), 3.78 (s, 3 H, OCH_3), 3.67 (m, 2 H, $\text{CH}_2\text{OCH}_2\text{CH}_2\text{O}$), 3.57 (m) and 3.45 (dd, J = 11.6 and 3.0 Hz) (1:1 H, $\text{CH}_2\text{OCH}_2\text{CH}_2\text{O}$), 3.38 and 3.18 (m each, 2 H, $\text{SCH}_2\text{CH}_2\text{S}$). ^{13}C NMR (100 MHz, CDCl_3 , 25 °C, TMS) δ 190.8 (Cq, C=O), 168.4 (Cq), 156.1 (Cq), 131.0 (aromatic CH), 129.9 (Cq), 128.2 (aromatic CH), 122.3 (Cq), 120.8 and 111.0 (aromatic CH), 76.2 (OCHCH_2O), 67.1 (OCHCH_2O), 66.3 and 65.5 ($\text{OCH}_2\text{CH}_2\text{O}$), 55.5 (OCH_3), 37.3 and 36.9 ($\text{SCH}_2\text{CH}_2\text{S}$). HRMS (ESI) calcd for $\text{C}_{16}\text{H}_{18}\text{O}_4\text{S}_2$ [$\text{M}+\text{H}$] $^+$: 339.0725; found: 339.0729.

2-(1,4-Dioxan-2-yl)-2-(1,3-dithiolan-2-ylidene)-1-(3-methoxyphenyl)et-hanone (3c): 122 mg, 71% yield; pale yellow solid, m.p. 136–137 °C. ^1H NMR (400 MHz, CDCl_3 , 25 °C, TMS) δ 7.22 (m, 2 H, aromatic CH), 7.16 (d, J = 2.4 Hz, 1 H, aromatic CH), 6.95 (m, 1 H, aromatic CH), 4.65 (dd, J = 10.5 and 3.1 Hz, 1 H, OCHCH_2O), 4.06 (m) and 3.80 (dd, J = 13.5, 5.1 Hz) (1:1 H, OCHCH_2O), 3.73 (s, 3H, OCH_3), 3.65 (m, 2 H, $\text{CH}_2\text{OCH}_2\text{CH}_2\text{O}$), 3.55 (m) and 3.45 (dd, J = 11.7 and 3.0 Hz) (1:1 H, $\text{CH}_2\text{OCH}_2\text{CH}_2\text{O}$), 3.36, 3.25 and 3.09 (m each, 1:2:1 H, $\text{SCH}_2\text{CH}_2\text{S}$). ^{13}C NMR (100 MHz, CDCl_3 , 25 °C, TMS) δ 192.8 (Cq, C=O), 163.7, 159.5 and 139.9 (Cq), 129.4 (aromatic CH), 121.8 (Cq), 120.9 (s), 118.3 and 112.6 (aromatic CH), 76.5 (OCHCH_2O), 67.2 (OCHCH_2O), 66.6 and 65.7 ($\text{OCH}_2\text{CH}_2\text{O}$), 55.37 (OCH_3), 37.4 and 37.2 ($\text{SCH}_2\text{CH}_2\text{S}$). HRMS (ESI) calcd for $\text{C}_{16}\text{H}_{18}\text{O}_4\text{S}_2$ [$\text{M}+\text{H}$] $^+$: 339.0725; found: 339.0724.

2-(1,4-Dioxan-2-yl)-2-(1,3-dithiolan-2-ylidene)-1-(4-methoxyphenyl)et-hanone (3d): 147 mg, 87% yield; pale yellow solid, m.p. 103–105 °C. ^1H NMR (400 MHz, CDCl_3 , 25 °C, TMS) δ 7.75 (d, J = 8.7 Hz, 2 H, aromatic CH), 6.88 (d, J = 8.7 Hz, 2 H, aromatic CH), 4.65 (dd, J = 10.3, 2.6 Hz, 1 H, OCHCH_2O), 4.02 (t) and 3.79 (m) (1:4 H, OCH_3 and OCHCH_2O), 3.71 and 3.59 (m each, 2:2 H, $\text{OCH}_2\text{CH}_2\text{O}$), 3.38, 3.28 and 3.13 (m, 1:2:1 H, $\text{SCH}_2\text{CH}_2\text{S}$). ^{13}C (100 MHz, CDCl_3 , 25 °C, TMS) δ 192.6 (Cq, C=O), 163.0 (Cq of Ph), 156.9 (Cq), 131.4 (aromatic CH), 130.5 (Cq), 122.5 (Cq), 113.6 (aromatic CH), 77.2 (OCHCH_2O), 67.1 (OCHCH_2O), 67.0 and 65.7 ($\text{OCH}_2\text{CH}_2\text{O}$), 55.4

(OCH₃), 37.4 and 37.4 (SCH₂CH₂S). HRMS (ESI) calcd for C₁₆H₁₈O₄S₂ [M+H]⁺: 339.0725; found: 339.0719.

1-(2-Chlorophenyl)-2-(1,4-dioxan-2-yl)-2-(1,3-dithiolan-2-ylidene)eth-anone (3e): 137 mg, 80% yield; pale yellow solid, m.p. 130-131 °C. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.38 (dd, *J* = 7.9 and 1.2 Hz, 1 H, aromatic CH), 7.30 (m, 2 H, aromatic CH), 7.21 (dd, *J* = 7.4 and 1.6 Hz, 1 H, aromatic CH), 4.45 (dd, *J* = 10.6 and 3.1 Hz, 1 H, OCHCH₂O), 4.04 (t) and 3.84 (d, *J* = 8.8 Hz) (1:1 H, OCHCH₂O), 3.64 (m, 2 H, CH₂OCH₂CH₂O), 3.59 (m) and 3.50 (dd, *J* = 11.7 and 2.9 Hz) (1:1 H, CH₂OCH₂CH₂O), 3.42 and 3.23 (m each, 2:2 H, SCH₂CH₂S). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 189.4 (Cq, C=O), 171.4, 139.7 and 130.6 (Cq), 130.4, 129.8, 127.8 and 126.9 (aromatic CH), 121.1 (Cq), 76.0 (OCHCH₂O), 67.2 (OCHCH₂O), 66.2 and 65.6 (OCH₂CH₂O), 37.6 and 37.1 (SCH₂CH₂S). HRMS (ESI) calcd for C₁₅H₁₅O₃S₂Cl [M+H]⁺: 343.0229; found: 343.0232.

1-(3-Chlorophenyl)-2-(1,4-dioxan-2-yl)-2-(1,3-dithiolan-2-ylidene)eth-anone (3f): 109 mg, 64% yield; pale yellow solid, m.p. 125-127 °C. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.69 (s, 1 H, aromatic CH), 7.57 (d, *J* = 7.6 Hz, 1 H, aromatic CH), 7.46 (d, *J* = 8.0 Hz, 1 H, aromatic CH), 7.36 (t, 1 H, aromatic CH), 4.68 (dd, *J* = 10.5 and 2.9 Hz, 1 H, OCHCH₂O), 4.14 (t) and 3.91 (d, *J* = 9.4 Hz) (1:1 H, OCHCH₂O), 3.74 (m, 2 H, CH₂OCH₂CH₂O), 3.63 (m) and 3.53 (dd, *J* = 11.7 and 2.7 Hz) (1:1 H, CH₂OCH₂CH₂O), 3.46, 3.36 and 3.20 (m each, 1:2:1 H, SCH₂CH₂S). ¹³C {¹H} (100 MHz, CDCl₃, 25 °C, TMS) δ 191.6 (Cq, C=O), 165.2, 140.5 and 134.6 (Cq), 131.7, 129.8, 128.6 and 126.6 (aromatic CH), 121.5 (Cq), 76.6 (OCHCH₂O), 67.4 (OCHCH₂O), 66.7 and 65.8 (OCH₂CH₂O), 37.5 and 37.4 (SCH₂CH₂S). HRMS (ESI) calcd for C₁₅H₁₅O₃S₂Cl [M+H]⁺: 343.0229; found: 342.0230.

1-(4-Chlorophenyl)-2-(1,4-dioxan-2-yl)-2-(1,3-dithiolan-2-ylidene)eth-anone (3g): 140 mg, 82% yield; pale yellow solid, m.p. 149-151 °C. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.66 (d, *J* = 8.3 Hz, 2 H, aromatic CH), 7.39 (d, *J* = 8.3 Hz, 2 H, aromatic CH), 4.65 (dd, *J* = 10.4 and 2.8 Hz, 1 H, OCHCH₂O), 4.11 (t) and 3.88 (d, *J* = 9.4 Hz) (1:1 H, OCHCH₂O), 3.73 (m, 2 H, CH₂OCH₂CH₂O), 3.62 and 3.53 (m each, 1:1 H, CH₂OCH₂CH₂O), 3.43, 3.33 and 3.18 (m each, 1:2:1 H, SCH₂CH₂S). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 192.1 (Cq, C=O), 163.2, 138.2 and 136.9 (Cq), 130.2 and 128.8 (aromatic CH), 121.7 (Cq), 76.8 (OCHCH₂O), 67.3 (OCHCH₂O), 66.8 and 65.8 (OCH₂CH₂O), 37.5 and 37.3 (SCH₂CH₂S). HRMS (ESI) calcd for C₁₅H₁₅O₃S₂Cl [M+H]⁺: 343.0229; found: 343.0232.

2-(1,4-Dioxan-2-yl)-2-(1,3-dithiolan-2-ylidene)-1-(4-fluorophenyl)etha-none (3h): 133 mg, 81% yield; pale yellow solid, m.p. 135-137 °C. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.76 (dd, *J* = 8.6 and 5.5 Hz, 2 H, aromatic CH), 7.09 (t, 2 H, aromatic CH), 4.66 (dd, *J* = 10.4 and 3.0 Hz, 1 H, OCHCH₂O), 4.10 (t) and 3.88 (d, *J* = 10.0 Hz) (1:1 H, OCHCH₂O), 3.71 (m, 2 H, CH₂OCH₂CH₂O), 3.64 (m) and 3.55 (dd, *J* = 11.7 and 2.9 Hz) (1:1 H, CH₂OCH₂CH₂O), 3.43, 3.33 and 3.17 (m each, 1:2:1 H, SCH₂CH₂S). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 192.0 (Cq, C=O), 165.1 (d, *J* = 253.3 Hz, Cq of Ph), 161.7

(Cq), 134.6 (d, *J* = 3.1 Hz, Cq of Ph), 131.4 (d, *J* = 9.1 Hz, aromatic CH), 121.9 (Cq), 115.6 (d, *J* = 21.8 Hz, aromatic CH), 76.9 (OCHCH₂O), 67.3 (OCHCH₂O), 66.8 and 65.8 (OCH₂CH₂O), 37.5 and 37.3 (SCH₂CH₂S). HRMS (ESI) calcd for C₁₅H₁₅O₃S₂F [M+H]⁺: 327.0525; found: 327.0517.

1-(4-Bromophenyl)-2-(1,4-dioxan-2-yl)-2-(1,3-dithiolan-2-ylidene)eth-anone (3i): 146 mg, 76% yield; pale yellow solid, m.p. 141-143 °C. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.56 (q, *J* = 8.5 Hz, 4 H, aromatic CH), 4.64 (dd, *J* = 10.4 and 2.9 Hz, 1 H, OCHCH₂O), 4.10 (t) and 3.87 (d, *J* = 9.3 Hz) (1:1 H, OCHCH₂O), 3.71 (m, 2 H, CH₂OCH₂CH₂O), 3.61 (m) and 3.52 (dd, *J* = 11.7 and 2.8 Hz) (1:1 H, CH₂OCH₂CH₂O), 3.42, 3.32 and 3.17 (m each, 1:2:1 H, SCH₂CH₂S). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 192.2 (Cq, C=O), 163.5 and 137.4 (Cq), 131.8 and 130.3 (aromatic CH), 126.7 and 121.7 (Cq), 76.8 (OCHCH₂O), 67.4 (OCHCH₂O), 66.8 and 65.8 (OCH₂CH₂O), 37.6 and 37.4 (SCH₂CH₂S). HRMS (ESI) calcd for C₁₅H₁₅O₃S₂Br [M+H]⁺: 386.9724; found: 386.9722.

2-(1,4-Dioxan-2-yl)-2-(1,3-dithiolan-2-ylidene)-1-(4-(trifluoromethyl)phenyl)ethanone (3j): 132 mg, 70% yield; pale yellow solid, m.p. 133-135 °C. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.78 (d, *J* = 8.0 Hz, 2 H, aromatic CH), 7.69 (d, *J* = 8.1 Hz, 2 H, aromatic CH), 4.65 (dd, *J* = 10.5 and 2.9 Hz, 1 H, OCHCH₂O), 4.15 (t) and 3.91 (d, *J* = 9.6 Hz) (1:1 H, OCHCH₂O), 3.72 (m, 2 H, CH₂OCH₂CH₂O), 3.63 and 3.52 (m each, 1:1 H, CH₂OCH₂CH₂O), 3.47, 3.37 and 3.22 (m, 1:2:1 H, SCH₂CH₂S). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 191.6 (Cq, C=O), 166.7 and 142.2 (Cq), 133.0 (q, *J* = 32.7 Hz, Cq of Ph), 128.7 (aromatic CH), 125.5 (q, *J* = 3.7 Hz, aromatic CH), 123.8 (q, *J* = 272.6 Hz, CF₃), 121.3 (Cq), 76.5 (OCHCH₂O), 67.4 (OCHCH₂O), 66.6 and 65.8 (OCH₂CH₂O), 37.5 and 37.4 (SCH₂CH₂S). HRMS (ESI) calcd for C₁₆H₁₅O₃S₂F₃ [M+H]⁺: 377.0493; found: 377.0491.

1-(1,4-Dioxan-2-yl)-1-(1,3-dithiolan-2-ylidene)propan-2-one (3k): 107 mg, 87% yield; white solid, m.p. 119-121 °C. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 4.83 (dd, *J* = 10.7 and 3.1 Hz, 1 H, OCHCH₂O), 3.93 (m, 2 H, OCHCH₂O), 3.84, 3.73 and 3.63 (m, 1:2:1 H, OCH₂CH₂O), 3.34 and 3.20 (m each, 2:2 H, SCH₂CH₂S), 2.36 (s, 3 H, COCH₃). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 193.2 (Cq, C=O), 166.3 and 121.5 (Cq), 77.0 (OCHCH₂O), 67.5 (OCHCH₂O), 66.8 and 65.8 (OCH₂CH₂O), 38.0 and 36.2 (SCH₂CH₂S), 28.9 (COCH₃). HRMS (ESI) calcd for C₁₀H₁₄O₃S₂ [M+H]⁺: 247.0463; found: 247.0454.

Ethyl 2-(1,4-dioxan-2-yl)-2-(1,3-dithiolan-2-ylidene)-acetate (3l): 114 mg, 83% yield; white solid, m.p. 91-93 °C. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 4.81 (dd, *J* = 10.3 and 2.8 Hz, 1 H, OCHCH₂O), 4.16 (dd, *J* = 13.0 and 5.9 Hz, 2 H, OCH₂CH₃), 3.73 (m, 6 H, CH₂OCH₂CH₂O), 3.32 and 3.19 (m, 2:2 H, SCH₂CH₂S), 1.25 (t, 3 H, OCH₂CH₃). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 165.2 (Cq, C=O), 165.1 and 113.0 (Cq), 75.3 (OCHCH₂O), 67.2 (OCHCH₂O), 66.6 and 65.8 (OCH₂CH₂O), 60.6 (OCH₂CH₃), 37.5 and 37.1 (SCH₂CH₂S), 14.3 (OCH₂CH₃). HRMS (ESI) calcd for C₁₁H₁₆O₄S₂ [M+H]⁺: 277.0568; found: 277.0567.

2-(1,4-Dioxan-2-yl)-2-(1,3-dithiolan-2-ylidene)acetonitrile (3m): 100 mg, 87% yield; white solid, m.p. 122–124 °C. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 4.30 (dd, *J* = 10.0 and 2.8 Hz, 1 H, OCHCH₂O), 3.80, 3.69 and 3.63 (m each, 3:2:1 H, CH₂OCH₂CH₂O), 3.56 (m, 4 H, SCH₂CH₂S). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 166.8, 116.9 and 94.3 (Cq), 75.3 (OCHCH₂O), 68.5 (OCHCH₂O), 66.9 and 66.1 (OCH₂CH₂O), 39.6 and 38.3 (SCH₂CH₂S). HRMS (ESI) calcd for C₉H₁₁NO₂S₂ [M+Na]⁺: 252.0120; found: 252.0120.

2-(1,4-Dioxan-2-yl)-2-(1,3-dithian-2-ylidene)-1-phenylethanone (3n): 130 mg, 81% yield; pale yellow solid, m.p. 128–130 °C. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.94 (d, *J* = 7.5 Hz, 2 H, aromatic CH), 7.53 (t, 1 H, aromatic CH), 7.43 (t, 2 H, aromatic CH), 4.85 (dd, *J* = 9.1 and 4.1 Hz, 1 H, OCHCH₂O), 3.77 and 3.69 (m each, 2:2 H, CH₂OCH₂CH₂O), 3.59 and 3.51 (m each, 1:1 H, CH₂OCH₂CH₂O), 2.98 (m, 2 H, SCH₂), 2.74 (m, 2 H, SCH₂), 2.09 (m, 2 H, SCH₂CH₂CH₂S). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 195.0 (Cq, C=O), 138.6, 137.7 and 134.4 (C=CCH), 133.2, 129.5 and 128.6 (aromatic CH), 75.6 (OCHCH₂O), 68.2 (OCHCH₂O), 66.9 and 66.0 (OCH₂CH₂O), 29.1 and 28.8 (SCH₂), 23.92 (SCH₂CH₂CH₂S). HRMS (ESI) calcd for C₁₆H₁₈O₃S₂ [M+H]⁺: 323.0776; found: 323.0774.

2-(1,4-Dioxan-2-yl)-3,3-bis(methylthio)-1-phenylprop-2-en-1-one (3o): 125 mg, 81% yield; white solid, m.p. 111–112 °C. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.91 (d, *J* = 7.6 Hz, 2 H, aromatic CH), 7.53 (t, 1 H, aromatic CH), 7.44 (t, 2 H, aromatic CH), 5.05 (dd, *J* = 10.2 and 2.5 Hz, 1 H, OCHCH₂O), 3.83, 3.68 and 3.48 (m each, 1:4:1 H, CH₂OCH₂CH₂O), 2.35 (s, 3 H, SCH₃), 2.04 (s, 3 H, SCH₃). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 194.7 (Cq, C=O), 143.4, 137.6 and 137.2 (Cq), 133.1 (aromatic CH), 129.4 and 128.4 (aromatic CH), 76.6 (OCHCH₂O), 68.7 (OCHCH₂O), 66.8 and 66.1 (OCH₂CH₂O), 17.1 (SCH₃), 16.4 (SCH₃). HRMS (ESI) calcd for C₁₅H₁₈O₃S₂ [M+H]⁺: 311.0776; found: 311.0774.

2-(1,4-Dioxan-2-yl)-3,3-bis(methylthio)-1-o-tolylprop-2-en-1-one (3p): 117 mg, 72% yield; white solid, m.p. 93–94 °C. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.69 (d, *J* = 7.6 Hz, 1 H, aromatic CH), 7.33 (t, 1 H, aromatic CH), 7.20 (t, 2 H, aromatic CH), 5.03 (dd, *J* = 10.0 and 2.9 Hz, 1 H, OCHCH₂O), 3.79 (m, 4 H, CH₂OCH₂CH₂O), 3.63 and 3.53 (m each, 1:1 H, CH₂OCH₂CH₂O), 2.60 (s, 3 H, SCH₃), 2.32 (s, 3 H, SCH₃), 1.89 (s, 3 H, Ph-CH₃). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 196.6 (Cq, C=O), 145.5, 134.0, 137.9 and 137.8 (Cq), 131.6, 131.6, 131.2 and 125.1 (aromatic CH), 76.9 (OCHCH₂O), 68.8 (OCHCH₂O), 66.8 and 66.1 (OCH₂CH₂O), 21.5 (Ph-CH₃), 16.7 (SCH₃), 16.4 (SCH₃). HRMS (ESI) calcd for C₁₆H₂₀O₃S₂ [M+H]⁺: 325.0932; found: 325.0930.

2-(1,4-Dioxan-2-yl)-3,3-bis(methylthio)-1-m-tolylprop-2-en-1-one (3q): 118 mg, 73% yield; white solid, m.p. 93–95 °C. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.76 (s, 1 H, aromatic CH), 7.68 (d, *J* = 6.9 Hz, 1 H, aromatic CH), 7.33 (q, *J* = 7.5 Hz, 2 H, aromatic CH), 5.05 (dd, *J* = 10.2 and 2.6 Hz, 1 H, OCHCH₂O), 3.83 (dd, *J* = 11.5 and 2.4 Hz), 3.68 (m) and 3.49 (m) (1:4:1 H, CH₂OCH₂CH₂O), 2.40 (s, 3 H, SCH₃), 2.36 (s, 3 H, SCH₃), 2.06 (s, 3 H, Ph-CH₃). ¹³C NMR (100 MHz, CDCl₃, 25 °C,

TMS) δ 194.9 (Cq, C=O), 143.6, 138.3, 137.5 and 136.9 (Cq), 134.1 (aromatic CH), 129.5 (aromatic CH), 128.3 (aromatic CH), 127.1 (aromatic CH), 76.6 (OCHCH₂O), 68.7 (OCHCH₂O), 66.8 and 66.1 (OCH₂CH₂O), 21.5 (Ph-CH₃), 17.2 (SCH₃), 16.4 (SCH₃). HRMS (ESI) calcd for C₁₆H₂₀O₃S₂ [M+H]⁺: 325.0932; found: 325.0934.

2-(1,4-Dioxan-2-yl)-3,3-bis(methylthio)-1-(p-tolyl)prop-2-en-1-one (3r): 138 mg, 85% yield; white solid, m.p. 112–114 °C. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.81 (d, *J* = 8.1 Hz, 2 H, aromatic CH), 7.24 (t, 2 H, aromatic CH), 5.04 (dd, *J* = 10.2 and 2.8 Hz, 1 H, OCHCH₂O), 3.81 (dd, *J* = 11.5 and 2.6 Hz), 3.65 (m) and 3.47 (m) (1:4:1 H, CH₂OCH₂CH₂O), 2.39 (s, 3 H, Ph-CH₃), 2.34 (s, 3 H, SCH₃), 2.08 (s, 3 H, SCH₃). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 194.3 (Cq, C=O), 144.0, 143.5, 136.8 and 135.1 (Cq), 129.5 and 129.2 (aromatic CH), 76.5 (OCHCH₂O), 68.6 (OCHCH₂O), 66.7 and 66.0 (OCH₂CH₂O), 21.8 (Ph-CH₃), 17.1 (SCH₃), 16.3 (SCH₃). HRMS (ESI) calcd for C₁₆H₂₀O₃S₂ [M+H]⁺: 325.0932; found: 325.0935.

2-(1,4-Dioxan-2-yl)-1-(4-methoxyphenyl)-3,3-bis(methylthio)prop-2-en-1-one (3s): 128 mg, 74% yield; white solid, m.p. 108–110 °C. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.76 (d, *J* = 7.6 Hz, 1 H, aromatic CH), 7.42 (d, *J* = 7.8 Hz, 1 H, aromatic CH), 7.36 (dd, *J* = 10.7 and 4.3 Hz, 1 H, aromatic CH), 7.28 (t, 1 H, aromatic CH), 5.02 (dd, *J* = 9.1 and 3.9 Hz, 1 H, OCHCH₂O), 3.79 (m, 4 H, CH₂OCH₂CH₂O), 3.64 (dd, *J* = 11.6 and 2.1 Hz) and 3.55 (m) (1:1 H, CH₂OCH₂CH₂O), 2.33 (s, 3 H, SCH₃), 1.91 (s, 3 H, SCH₃). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 192.8 (Cq, C=O), 144.2, 140.4, 137.6 and 133.8 (Cq), 132.1, 131.4, 131.1 and 126.1 (aromatic CH), 76.9 (OCHCH₂O), 68.6 (OCHCH₂O), 66.8 and 66.1 (OCH₂CH₂O), 16.7 (SCH₃), 16.4 (SCH₃). HRMS (ESI) calcd for C₁₅H₁₇O₃S₂Cl [M+H]⁺: 345.0386; found: 345.0387.

1-(3-Chlorophenyl)-2-(1,4-dioxan-2-yl)-3,3-bis(methylthio)prop-2-en-1-one (3t): 104 mg, 60% yield; colorless liquid. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.89 (s, 1 H, aromatic CH), 7.76 (d, *J* = 7.7 Hz, 1 H, aromatic CH), 7.49 (m, 1 H, aromatic CH), 7.38 (t, 1 H, aromatic CH), 5.04 (dd, *J* = 10.2 and 2.8 Hz, 1 H, OCHCH₂O), 3.82 (dd, *J* = 11.5 and 2.7 Hz), 3.68 (m) and 3.49 (m) (1:4:1 H, CH₂OCH₂CH₂O), 2.35 (s, 3 H, SCH₃), 2.05 (s, 3 H, SCH₃). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 193.5 (Cq, C=O), 142.8, 139.4, 138.0 and 134.8 (Cq), 139.8, 129.8, 129.1 and 127.6 (aromatic CH), 76.6 (OCHCH₂O), 68.6 (OCHCH₂O), 66.8 and 66.1 (OCH₂CH₂O), 17.1 (SCH₃), 16.4 (SCH₃). HRMS (ESI) calcd for C₁₅H₁₇O₃S₂Cl [M+H]⁺: 345.0386; found: 345.0389.

1-(4-Bromophenyl)-2-(1,4-dioxan-2-yl)-3,3-bis(methylthio)prop-2-en-1-one (3u): 152 mg, 78% yield; white solid, m.p. 131–133 °C. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.76 (m, 2 H, aromatic CH), 7.57 (d, *J* = 7.0 Hz, 2 H, aromatic CH), 5.03 (d, *J* = 10.1 Hz, 1 H, OCHCH₂O), 3.81 (d, *J* = 11.5 Hz), 3.66 (m), and 3.47 (t, *J* = 11.5 Hz) (1:4:1 H, CH₂OCH₂CH₂O), 2.34 (s, 3 H, SCH₃), 2.05 (s, 3 H, SCH₃). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 193.7 (Cq, C=O), 142.8, 137.6 and 136.5 (Cq), 131.7 and 130.8 (aromatic CH), 128.3 (Cq), 76.5 (OCHCH₂O), 68.5 (OCHCH₂O), 66.7 and 66.0 (OCH₂CH₂O), 17.1 (SCH₃),

16.3 (SCH₃). HRMS (ESI) calcd for C₁₅H₁₇O₃S₂Br [M+H]⁺: 388.9881; found: 388.9883.

2-(1,4-Dioxan-2-yl)-3,3-bis(methylthio)-1-(naphthalen-2-yl)prop-2-en-1-one (3v): 60 mg, 33% yield; pale yellow solid, m.p. 128–130 °C. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 8.42 (s, 1 H, aromatic CH), 8.01 (m, 2 H, aromatic CH), 7.88 (m, 2 H, aromatic CH), 7.56 (m, 2 H, aromatic CH), 5.14 (dd, *J* = 10.2 and 2.8 Hz, 1 H, OCHCH₂O), 3.90 (dd, *J* = 11.5 and 2.6 Hz), 3.70 (m) and 3.51 (m) (1:4:1 H, CH₂OCH₂CH₂O), 2.39 (s, 3 H, SCH₃), 2.05 (s, 3 H, SCH₃). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 194.8 (Cq, C=O), 143.6, 137.4, 135.8, 135.1 and 132.7 (Cq), 131.6, 129.9, 128.5, 128.4, 127.9, 126.7 and 124.7 (aromatic CH), 76.8 (OCHCH₂O), 68.8 (OCHCH₂O), 66.8 and 66.1 (OCH₂CH₂O), 17.2 (SCH₃), 16.4 (SCH₃). HRMS (ESI) calcd for C₁₉H₂₀O₃S₂ [M+H]⁺: 361.0932; found: 361.0936.

2-(1,4-Dioxan-2-yl)-3,3-bis(methylthio)-1-(thiophen-2-yl)prop-2-en-1-one (3w): 64 mg, 41% yield; pale yellow solid, m.p. 105–107 °C. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.64 (d, *J* = 4.9 Hz, 1 H, aromatic CH), 7.58 (d, *J* = 3.7 Hz, 1 H, aromatic CH), 7.09 (m, 1 H, aromatic CH), 5.01 (dd, *J* = 10.2 and 2.8 Hz, 1 H, OCHCH₂O), 3.65 (m, 6 H, CH₂OCH₂CH₂O), 2.35 (s, 3 H, SCH₃), 2.12 (s, 3 H, SCH₃). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 186.9 (Cq, C=O), 145.1, 143.0 and 138.7 (Cq), 134.3, 134.1 and 128.1 (aromatic CH), 76.2 (OCHCH₂O), 68.6 (OCHCH₂O), 66.8 and 66.0 (OCH₂CH₂O), 17.5 (SCH₃), 16.4 (SCH₃). HRMS (ESI) calcd for C₁₃H₁₆O₃S₃ [M+H]⁺: 317.0340; found: 317.0339.

2-(1,4-Dioxan-2-yl)-1-(furan-2-yl)-3,3-bis(methylthio)prop-2-en-1-one (3x): 31 mg, 21% yield; pale yellow solid, m.p. 116–118 °C. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.60 (s, 1 H, aromatic CH), 7.10 (d, *J* = 3.5 Hz, 1 H, aromatic CH), 6.53 (dd, *J* = 3.1 and 1.1 Hz, 1 H, aromatic CH), 5.01 (dd, *J* = 10.2 and 2.8 Hz, 1 H, OCHCH₂O), 3.74 (m, 4 H, CH₂OCH₂CH₂O), 3.61 and 3.51 (m each, 1:1 H, CH₂OCH₂CH₂O), 2.35 (s, 3 H, SCH₃), 2.14 (s, 3 H, SCH₃). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 182.3 (Cq, C=O), 153.3 (Cq), 147.1 (aromatic CH), 142.1 (Cq), 139.4 (Cq), 119.2 (aromatic CH), 112.5 (aromatic CH), 76.4 (OCHCH₂O), 68.7 (OCHCH₂O), 66.9 and 66.1 (OCH₂CH₂O), 17.8 (SCH₃), 16.4 (SCH₃). HRMS (ESI) calcd for C₁₃H₁₆O₄S₂ [M+H]⁺: 301.0568; found: 301.0569.

3-(1,4-Dioxan-2-yl)-4,4-bis(methylthio)but-3-en-2-one (3y): 108 mg, 87% yield; white solid, m.p. 101–102 °C. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 4.87 (dd, *J* = 10.2, 3.0 Hz, 1 H, OCHCH₂O), 3.60 (m, 6 H, CH₂OCH₂CH₂O), 2.34 (s, 3 H, SCH₃), 2.27 (s, 3 H, SCH₃), 2.21 (s, 3 H, CH₃CO). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 202.8 (Cq, C=O), 146.7 and 134.0 (Cq), 76.3 (OCHCH₂O), 68.2 (OCHCH₂O), 66.6 and 66.0 (OCH₂CH₂O), 32.3 (CH₃CO), 17.3 (SCH₃), 16.3 (SCH₃). HRMS (ESI) calcd for C₁₀H₁₆O₃S₂ [M+H]⁺: 249.0619; found: 249.0617.

1-Cyclopropyl-2-(1,4-dioxan-2-yl)-3,3-bis(methylthio)prop-2-en-1-one (3z): 111 mg, 81% yield; pale yellow solid, m.p. 81–83 °C. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 4.89 (dd, *J* = 10.1 and 3.0 Hz, 1 H, OCHCH₂O), 3.73 (dd, *J* = 8.1 and 2.0 Hz), 3.68 (dd, *J* = 11.5 and 2.9 Hz) and 3.56 (m) (2:1:3 H, CH₂OCH₂CH₂O),

2.30 (s, 3 H, SCH₃), 2.24 (s, 3 H, SCH₃), 2.09 (m, 1 H, CH₂CHCH₂), 1.17 and 0.96 (m, 2 H, CHCH₂CH₂). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 204.6 (Cq, C=O), 146.6 and 136.2 (Cq), 76.2 (OCHCH₂O), 68.6 (OCHCH₂O), 66.8 and 66.1 (OCH₂CH₂O), 23.2 (CH₂CHCH₂), 17.4 (SCH₃), 16.4 (SCH₃), 12.8 and 12.3 (CHCH₂CH₂). HRMS (ESI) calcd for C₁₂H₁₈O₃S₂ [M+H]⁺: 275.0776; found: 275.0775.

Ethyl 2-(1,4-dioxan-2-yl)-3,3-bis(methylthio)acrylate (3z1): 121 mg, 87% yield; colorless liquid. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 4.79 (t, 1 H, OCHCH₂O), 4.17 (m, 2 H, OCH₂CH₃), 3.64, 3.56 and 3.46 (m each, 2:3:1 H, CH₂OCH₂CH₂O), 2.23 (s, 3 H, SCH₃), 2.17 (s, 3 H, SCH₃), 1.23 (t, 3 H, OCH₂CH₃). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 165.8 (Cq, C=O), 138.4 and 137.6 (Cq), 75.2 (OCHCH₂O), 68.1 (OCHCH₂O), 66.5 and 65.7 (OCH₂CH₂O), 61.0 (OCH₂CH₃), 17.2 (SCH₃), 16.0 (SCH₃), 14.0 (OCH₂CH₃). HRMS (ESI) calcd for C₁₃H₁₄O₂S₂ [M+H]⁺: 279.0725; found: 279.0725.

2-(1,4-Dioxan-2-yl)-3,3-bis(methylthio)acrylonitrile (3z2): 101 mg, 87% yield; pale yellow liquid. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 4.74 (dd, *J* = 10.2 and 2.8 Hz, 1 H, OCHCH₂O), 3.70 (m, 6 H, CH₂OCH₂CH₂O), 2.47 and 2.44 (s each, 3:3 H, SCH₃). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 161.0, 115.9 and 111.6 (Cq), 73.5 (OCHCH₂O), 68.4 (OCHCH₂O), 66.8 and 65.9 (OCH₂CH₂O), 18.3 (SCH₃), 17.7 (SCH₃). HRMS (ESI) calcd for C₉H₁₃NO₂S₂ [M+H]⁺: 232.0460; found: 232.0462.

2-(1,3-Dithiolan-2-ylidene)-1-phenyl-2-(tetrahydrofuran-2-yl)ethanon-e (4a): 121 mg, 83% yield; pale yellow solid, m.p. 106–108 °C. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.68 (m, 2 H, aromatic CH), 7.41 (m, 3 H, aromatic CH), 4.83 (m, 1 H, OCHCH₂), 4.02 and 3.71 (m each, 1:1 H, OCH₂CH₂), 3.41, 3.28 and 3.13 (m each, 1:2:1 H, SCH₂CH₂S), 2.17 and 1.91 (m each, 1:3 H, OCHCH₂CH₂). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 193.6 (Cq, C=O), 160.6 and 139.0 (Cq), 131.3, 128.5 and 128.2 (aromatic CH), 124.9 (Cq), 79.6 (OCHCH₂), 67.9 (OCH₂CH₂), 37.4 and 36.9 (SCH₂CH₂S), 29.8 (CHCH₂CH₂), 26.6 (CHCH₂CH₂). HRMS (ESI) calcd for C₁₅H₁₆O₂S₂ [M+H]⁺: 293.0670; found: 293.0671.

1-(1,3-Dithiolan-2-ylidene)-1-(tetrahydrofuran-2-yl)propan-2-one (4b): 101 mg, 88% yield; pale yellow liquid. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 4.94 (dd, *J* = 9.4 and 6.3 Hz, 1 H, OCHCH₂), 4.05 and 3.77 (m each, 1:1 H, OCH₂CH₂), 3.28 and 3.10 (m each, 2:2 H, SCH₂CH₂S), 2.21 (s, 3 H, COCH₃), 1.99 (m, 4 H, CHCH₂CH₂). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 193.3 (Cq, C=O), 162.9 and 124.3 (Cq), 79.6 (OCHCH₂), 67.6 (OCH₂CH₂), 37.5 and 36.2 (SCH₂CH₂S), 23.0 (CHCH₂CH₂), 28.5 (CHCH₂CH₂), 26.4 (COCH₃). HRMS (ESI) calcd for C₁₀H₁₄O₂S₂ [M+H]⁺: 231.0513; found: 231.0516.

2-(1,3-Dithiolan-2-ylidene)-2-(tetrahydrofuran-2-yl)acetonitrile (4c): 75 mg, 70% yield; colorless liquid. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 4.58 (t, 1 H, OCHCH₂), 3.96 and 3.79 (dd each, 1:1 H, OCH₂CH₂), 3.51 (m, 4 H, SCH₂CH₂S), 2.04 (m, 4 H, CHCH₂CH₂). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 163.4 (Cq, CN), 117.5 and 99.2 (Cq), 78.8 (OCHCH₂), 68.7 (OCH₂CH₂), 39.4 and 37.9 (SCH₂CH₂S), 31.2 and 26.0 (CHCH₂CH₂).

HRMS (ESI) calcd for $C_9H_{11}NOS_2$ $[M+H]^+$: 214.0360; found: 214.0359.

3,3-Bis(methylthio)-1-phenyl-2-(tetrahydrofuran-2-yl)prop-2-en-1-one (4d): 103 mg, 70% yield; colorless liquid. 1H NMR (400 MHz, $CDCl_3$, 25 °C, TMS) δ 7.93 (d, J = 8.1 Hz, 2 H, aromatic CH), 7.48 (m, 3 H, aromatic CH), 5.23 (t, 1 H, $OCHCH_2$), 3.63 (dq, J = 22.1 and 7.6 Hz, 2 H, OCH_2CH_2), 2.33 (s, 3 H, SCH_3), 2.24 and 2.07 (m each, 1:4 H, SCH_3 and $OCHCH_2CH_2$), 1.83 (m, 2 H, $CH_2CH_2CH_2$). ^{13}C NMR (100 MHz, $CDCl_3$, 25 °C, TMS) δ 195.8 (Cq, C=O), 148.9, 137.6 and 133.4 (Cq), 133.0 (aromatic CH), 129.2 (aromatic CH), 128.5 (aromatic CH), 79.2 ($OCHCH_2$), 68.9 (OCH_2CH_2), 31.9 ($CHCH_2CH_2$), 26.2 ($CHCH_2CH_2$), 17.1 (SCH_3), 16.2 (SCH_3). HRMS (ESI) calcd for $C_{14}H_{16}O_2S_2$ $[M+H]^+$: 295.0826; found: 295.0828.

4,4-Bis(methylthio)-3-(tetrahydrofuran-2-yl)but-3-en-2-one (4e): 108 mg, 93% yield; white solid, m.p. 50–51 °C. 1H NMR (400 MHz, $CDCl_3$, 25 °C, TMS) δ 5.00 (t, 1 H, $OCHCH_2$), 3.71 (t, 2 H, OCH_2CH_2), 2.34 (s, 3 H, $COCH_3$), 2.25 (s, 3 H, SCH_3), 2.20 (s, 3 H, SCH_3), 2.15 and 1.85 (m each, 1:3 H, $CHCH_2CH_2$). ^{13}C NMR (100 MHz, $CDCl_3$, 25 °C, TMS) δ 203.9 (Cq, C=O), 152.5 and 130.2 (Cq), 79.0 ($OCHCH_2$), 68.8 (OCH_2CH_2), 32.4 ($CHCH_2CH_2$), 31.8 ($CHCH_2CH_2$), 26.1 ($COCH_3$), 17.3 (SCH_3), 16.2 (SCH_3). HRMS (ESI) calcd for $C_{10}H_{16}O_2S_2$ $[M+H]^+$: 233.0670; found: 233.0668.

Ethyl 3,3-bis(methylthio)-2-(tetrahydrofuran-2-yl)acrylate (4f): 97 mg, 74% yield; colorless liquid. 1H NMR (400 MHz, $CDCl_3$, 25 °C, TMS) δ 5.05 (t, J = 7.4 Hz, 1 H, $OCHCH_2$), 4.24 (m, 2 H, OCH_2CH_3), 3.77 (m, 2 H, OCH_2CH_2), 2.28 (s, 3 H, SCH_3), 2.25 (s, 3 H, SCH_3), 2.14 and 1.89 (m each, 1:3 H, $OCHCH_2CH_2$), 1.30 (t, J = 7.1 Hz, 3 H, OCH_2CH_3). ^{13}C NMR (100 MHz, $CDCl_3$, 25 °C, TMS) δ 166.9 (Cq, C=O), 143.5 and 135.2 (Cq), 78.2 ($OCHCH_2$), 69.1 (OCH_2CH_2), 61.0 (OCH_2CH_3), 32.0 ($CHCH_2CH_2$) and 26.3 ($CHCH_2CH_2$), 17.5 (SCH_3), 16.2 (SCH_3), 14.2 (OCH_2CH_3). HRMS (ESI) calcd for $C_{11}H_{18}O_3S_2$ $[M+H]^+$: 263.0776; found: 263.0774.

3,3-Bis(methylthio)-1-phenyl-2-(tetrahydro-2H-pyran-2-yl)prop-2-en-1-one (4g): 100 mg, 65% yield; colorless liquid. 1H NMR (400 MHz, $CDCl_3$, 25 °C, TMS) δ 7.95 (d, J = 7.4 Hz, 2 H, aromatic CH), 7.53 (t, 1 H, aromatic CH), 7.44 (t, 2 H, aromatic CH), 4.71 (dd, J = 10.9 and 2.5 Hz, 1 H, $OCHCH_2$), 3.90 and 3.46 (m each, 1:1 H, OCH_2CH_2), 2.34 (s, 3 H, SCH_3), 2.04 (s, 3 H, SCH_3), 1.72 and 1.49 (m each, 3:3 H, $CHCH_2CH_2CH_2CH_2$). ^{13}C NMR (100 MHz, $CDCl_3$, 25 °C, TMS) δ 195.3 (Cq, C=O), 148.4, 137.7 and 134.3 (Cq), 133.0, 129.6 and 128.4 (aromatic CH), 78.6 ($OCHCH_2$), 68.7 (OCH_2CH_2), 30.4 ($CHCH_2CH_2CH_2$), 25.5 ($CHCH_2CH_2CH_2$), 23.7 ($CHCH_2CH_2CH_2$), 17.2 (SCH_3), 16.2 (SCH_3). HRMS (ESI) calcd for $C_{16}H_{20}O_2S_2$ $[M+H]^+$: 309.0983; found: 309.0980.

4,4-Bis(methylthio)-3-(tetrahydro-2H-pyran-2-yl)but-3-en-2-one (4h): 89 mg, 72% yield; white solid, m.p. 65–67 °C. 1H NMR (400 MHz, $CDCl_3$, 25 °C, TMS) δ 4.57 (dd, J = 8.8 and 4.7 Hz, 1 H, $OCHCH_2$), 3.92 (dd, J = 11.1 and 3.6 Hz) and 3.40 (m) (1:1 H, OCH_2CH_2), 2.34 (s, 3 H, $COCH_3$), 2.26 (s, 3 H, SCH_3), 2.21 (s, 3 H, SCH_3), 1.90 and 1.52 (m, 1:5 H, $CHCH_2CH_2CH_2CH_2$). ^{13}C NMR (100 MHz, $CDCl_3$, 25 °C, TMS) δ 203.3 (Cq, C=O), 152.0 and 130.6 (Cq), 78.4 ($OCHCH_2$), 68.6 (OCH_2CH_2), 32.4

($COCH_3$), 30.2 ($CHCH_2CH_2CH_2$), 25.5 ($CHCH_2CH_2CH_2$), 23.4 ($CHCH_2CH_2CH_2$), 17.4 (SCH_3), 16.2 (SCH_3). HRMS (ESI) calcd for $C_{11}H_{18}O_2S_2$ $[M+H]^+$: 247.0826; found: 247.0824.

Ethyl 3,3-bis(methylthio)-2-(tetrahydro-2H-pyran-2-yl)acrylate (4i): 102 mg, 74% yield; colorless liquid. 1H NMR (400 MHz, $CDCl_3$, 25 °C, TMS) δ 4.58 (dd, J = 11.2 and 1.8 Hz, 1 H, $OCHCH_2$), 4.27 (m, 2 H, OCH_2CH_3), 3.96 (dd, J = 10.9 and 2.8 Hz) and 3.44 (m) (1:1 H, OCH_2CH_2), 2.29 (s, 3 H, SCH_3), 2.24 (s, 3 H, SCH_3), 1.78 and 1.51 (m each, 2:4 H, $CHCH_2CH_2CH_2CH_2$), 1.31 (t, 3 H, OCH_2CH_3). ^{13}C NMR (100 MHz, $CDCl_3$, 25 °C, TMS) δ 166.7 (Cq, C=O), 143.2 and 135.2 (Cq), 77.6 ($OCHCH_2$), 68.7 (OCH_2CH_2), 61.1 (OCH_2CH_3), 30.2 ($CHCH_2CH_2CH_2$), 25.5 ($CHCH_2CH_2CH_2$), 23.5 ($CHCH_2CH_2CH_2$), 17.5 (SCH_3), 16.1 (SCH_3), 14.3 (OCH_2CH_3). HRMS (ESI) calcd for $C_{12}H_{20}O_3S_2$ $[M+H]^+$: 277.0932; found: 277.0935.

3-(2,3-Dihydrobenzo[b][1,4]dioxin-2-yl)-4,4-bis(methylthio)but-3-en-2-one (4j): 89 mg, 60% yield; yellow liquid. 1H NMR (400 MHz, $CDCl_3$, 25 °C, TMS) δ 6.85 (m, 4 H, aromatic CH), 5.46 (dd, J = 9.0 and 2.6 Hz, 1 H, $OCHCH_2O$), 4.25 (dd, J = 11.4 and 2.6 Hz) and 4.13 (dd, J = 11.4 and 9.0 Hz) (1:1 H, $OCHCH_2O$), 2.49 (s, 3 H, SCH_3), 2.36 (s, 3 H, SCH_3), 2.32 (s, 3 H, $COCH_3$). ^{13}C NMR (100 MHz, $CDCl_3$, 25 °C, TMS) δ 202.2 (Cq, C=O), 144.1, 143.0, 142.9 and 137.5 (Cq), 121.9, 121.6, 117.4 and 117.3 (aromatic CH), 74.1 ($OCHCH_2O$), 65.8 ($OCHCH_2O$), 32.4 ($COCH_3$), 17.3 (SCH_3), 16.5 (SCH_3). HRMS (ESI) calcd for $C_{14}H_{16}O_3S_2$ $[M+H]^+$: 297.0619; found: 297.0622.

3-(1,3-Dithiolan-2-ylidene)-4-ethoxypentan-2-one (4k): 60 mg, 52% yield; colorless liquid. 1H NMR (400 MHz, $CDCl_3$, 25 °C, TMS) δ 4.69 (q, J = 6.7 Hz, 1 H, CH_3CHO), 3.41 and 3.2 (m each, 1:3 H, CH_3CH_2O and SCH_2), 3.22 (m, 2 H, SCH_2), 2.32 (s, 3 H, $COCH_3$), 1.46 (d, J = 6.7 Hz, 3 H, CH_3CHO), 1.18 (t, 3 H, CH_3CH_2O). ^{13}C NMR (100 MHz, $CDCl_3$, 25 °C, TMS) δ 194.2 (Cq, C=O), 162.6 and 126.3 (Cq), 76.5 (CH_3CHO), 64.1 (CH_3CH_2O), 37.9 and 36.1 (SCH_2CH_2S), 28.5 ($COCH_3$), 19.8 (CH_3CHO), 15.4 (CH_3CH_2O). HRMS (ESI) calcd for $C_{10}H_{17}O_2S_2$ $[M+Na]^+$: 255.0489; found: 255.0487.

3-(Bis(methylthio)methylene)-4-ethoxypentan-2-one (4l): 73 mg, 62% yield; colorless liquid. 1H NMR (400 MHz, $CDCl_3$, 25 °C, TMS) δ 4.75 (q, J = 6.6 Hz, 1 H, CH_3CHO), 3.47 (m, 2 H, CH_3CH_2O), 2.33 (s, 3 H, $COCH_3$), 2.28 (s, 3 H, SCH_3), 2.24 (s, 3 H, SCH_3), 1.31 (d, J = 6.6 Hz, 3 H, CH_3CHO), 1.14 (t, 3 H, CH_3CH_2O). ^{13}C NMR (100 MHz, $CDCl_3$, 25 °C, TMS) δ 203.1 (Cq, C=O), 152.4 and 132.6 (Cq), 75.2 (CH_3CHO), 64.6 (CH_3CH_2O), 32.0 ($COCH_3$), 20.4 (CH_3CHO), 17.4 (SCH_3), 16.3 (SCH_3), 15.3 (CH_3CH_2O). HRMS (ESI) calcd for $C_{10}H_{18}O_2S_2$ $[M+Na]^+$: 257.0646; found: 257.0640.

2-((4-Methoxyphenoxy)methyl)-3,3-bis(methylthio)-1-phenylprop-2-en-1-one (4m): 112 mg, 62% yield; colorless liquid. 1H NMR (400 MHz, $CDCl_3$, 25 °C, TMS) δ 7.91 (d, J = 7.2 Hz, 2 H, aromatic CH), 7.54 (t, 1 H, aromatic CH), 7.44 (t, 2 H, aromatic CH), 6.78 (m, 4 H, aromatic CH), 4.99 (s, 2 H, OCH_2), 3.74 (s, 3 H, OCH_3), 2.39 and 2.11 (s each, 3:3 H, SCH_3). ^{13}C NMR (100 MHz, $CDCl_3$, 25 °C, TMS) δ 196.3 (Cq, C=O), 154.2, 152.5, 142.9 and 141.3 (Cq), 137.7 (C=CCH), 133.1, 129.3, 128.5, 116.2 and 114.6 (aromatic CH), 68.9 (OCH_2), 55.8 (OCH_3),

17.2 and 16.7 (SCH₃). HRMS (ESI) calcd for C₁₉H₂₀O₃S₂ [M+Na]⁺: 383.0752; found: 383.0747.

2-(1,3-Dioxolan-2-yl)-3,3-bis(methylthio)-1-(p-tolyl)-prop-2-en-1-one (4n): 57 mg, 37% yield; colorless liquid. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.86 (d, *J* = 8.1 Hz, 2 H, aromatic CH), 7.25 (d, *J* = 7.8 Hz, 2 H, aromatic CH), 6.13 (s, 1 H, OCHO), 3.81 (m, 4 H, OCH₂CH₂O), 2.40 (s, 3 H, SCH₃), 2.37 (s, 3 H, SCH₃), 2.14 (s, 3 H, Ph-CH₃). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 193.8 (Cq, C=O), 144.2, 143.3, 140.7 and 134.7 (Cq), 129.7 (aromatic CH), 129.3 (aromatic CH), 101.4 (OCHO), 65.5 (OCH₂CH₂O), 21.9 (Ph-CH₃), 17.1 (SCH₃), 16.5 (SCH₃). HRMS (ESI) calcd for C₁₅H₁₈O₃S₂ [M+H]⁺: 311.0776; found: 311.0775.

2-(1,3-Dioxolan-4-yl)-3,3-bis(methylthio)-1-(p-tolyl)-prop-2-en-1-one (4n'): 38 mg, 24% yield; colourless liquid. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.84 (d, *J* = 8.2 Hz, 2 H, aromatic CH), 7.27 (d, *J* = 7.9 Hz, 2 H, aromatic CH), 5.37 (t, 1 H, OCHCH₂O), 4.84 and 4.73 (s each, 1:1 H, OCH₂O), 4.21 (dd, *J* = 8.3 and 6.9 Hz) and 3.97 (dd, *J* = 8.3 and 7.1 Hz) (1:1 H, OCHCH₂O), 2.41 (s, 3 H, Ph-CH₃), 2.37 (s, 3 H, SCH₃), 2.12 (s, 3 H, SCH₃). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 194.7 (Cq, C=O), 144.9, 144.4, 136.1 and 134.9 (Cq), 129.5 and 129.4 (aromatic CH), 95.9 (OCH₂O), 75.6 (OCHCH₂O), 69.2 (OCHCH₂O), 21.9 (Ph-CH₃), 17.2 (SCH₃), 16.4 (SCH₃). HRMS (ESI) calcd for C₁₅H₁₈O₃S₂ [M+H]⁺: 311.0776; found: 311.0779.

2-(1,3-Dithiolan-2-ylidene)-3,4-dimethoxy-1-phenylbutan-1-one (4o): 64 mg, 41% yield; pale yellow liquid. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.59 (m, 2 H, aromatic CH), 7.44 (m, 3 H, aromatic CH), 4.59 (dd, *J* = 8.2 and 4.3 Hz, 1 H, CHOCH₃), 3.79 (dd, *J* = 10.4 and 8.2 Hz) and 3.45 (m) (1:1 H, CHCH₂OCH₃), 3.40 and 3.34 (m each, 4:1 H, OCH₃ and SCH₂), 3.27 and 3.16 (m each, 4:1 H, OCH₃ and SCH₂). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 194.0 (Cq, C=O), 161.7 and 139.5 (Cq), 131.2, 128.5 and 128.1 (aromatic CH), 122.3 (Cq), 81.0 (CHOCH₃), 72.9 (CHCH₂OCH₃), 59.0 (OCH₃), 57.0 (OCH₃), 37.5 and 37.4 (SCH₂CH₂S). HRMS (ESI) calcd for C₁₅H₁₈O₃S₂ [M+Na]⁺: 333.0595; found: 333.0594.

2-(1,3-Dithiolan-2-ylidene)-3-(2-methoxyethoxy)-1-phenylpropan-1-one (4o'): 58 mg, 37% yield; pale yellow liquid. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.76 (m, 2 H, aromatic CH), 7.42 (m, 3 H, aromatic CH), 4.34 (s, 2 H, CH₂OCH₂CH₂), 3.59 (s, 4 H, OCH₂CH₂O), 3.38 (m, 7 H, OCH₃ and SCH₂CH₂S). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 191.9 (Cq, C=O), 172.3 and 139.2 (Cq), 131.0, 128.3 and 128.1 (aromatic CH), 121.1 (Cq), 71.8 (CH₂OCH₂CH₂), 71.4 (CH₂OCH₂CH₂), 69.3 (CH₂OCH₂CH₂), 59.1 (OCH₃), 38.9 and 36.0 (SCH₂CH₂S). HRMS (ESI) calcd for C₁₅H₁₈O₃S₂ [M+H]⁺: 333.0595; found: 333.0596.

3-(1,3-Dithiolan-2-ylidene)-4,5-dimethoxypentan-2-one (4p): 56 mg, 45% yield; pale yellow liquid. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 4.65 (dd, *J* = 8.5, 3.9 Hz, 1 H, CHOCH₃), 3.78 (m, 1 H), and 3.39 (dd, *J* = 10.4 and 3.9 Hz) (1:1 H, CHCH₂OCH₃), 3.34 (s, 3 H, OCH₃), 3.30 (m, 2 H, SCH₂), 3.25 (s, 3 H, OCH₃), 3.17 (m, 2 H, SCH₂), 2.27 (s, 3 H, COCH₃). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 193.9 (Cq, C=O), 165.5 and 121.6 (Cq), 81.7 (CHOCH₃), 73.4 (CHCH₂OCH₃), 59.2 (OCH₃), 56.8

(OCH₃), 37.9 and 36.2 (SCH₂CH₂S), 28.5 (COCH₃). HRMS (ESI) calcd for C₁₀H₁₆O₃S₂[M+Na]⁺: 271.0439; found: 271.0438.

3-(1,3-Dithiolan-2-ylidene)-4-(2-methoxyethoxy)-butan-2-one (4p'): 25 mg, 52% yield; colorless liquid. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 4.52 (s, 2 H, CH₂OCH₂CH₂), 3.63 and 3.56 (m each, 2:2 H, SCH₂CH₂S), 3.38 (m, 5 H, OCH₃ and CH₂OCH₂CH₂), 3.32 (m, 2 H, CH₂OCH₂CH₂), 2.34 (s, 3 H, COCH₃). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 194.5 (Cq, C=O), 167.4 and 121.4 (Cq), 72.0 (CH₂OCH₂CH₂), 71.4 (CH₂OCH₂CH₂), 69.0 (CH₂OCH₂CH₂), 59.1 (OCH₃), 39.3 and 35.8 (SCH₂CH₂S), 27.2 (COCH₃). HRMS (ESI) calcd for C₁₀H₁₆O₃S₂ [M+Na]⁺: 271.0439; found: 271.0434.

2-(Bis(methylthio)methylene)-3,4-dimethoxy-1-phenylbutan-1-one (4q): 54 mg, 35% yield; colorless liquid. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.85 (d, *J* = 7.3 Hz, 2 H, aromatic CH), 7.47 (t, 1 H, aromatic CH), 7.38 (t, 2 H, aromatic CH), 4.83 (dd, *J* = 7.0 and 4.7 Hz, 1 H, CHOCH₃), 3.48 (dd, *J* = 10.2 and 7.1 Hz) and 3.40 (m) (1:4 H, CH₃OCHCH₂ and OCH₃), 3.22 (s, 3 H, OCH₃), 2.30 (s, 3 H, SCH₃), 2.04 (s, 3 H, SCH₃). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 194.7 (Cq, C=O), 144.4, 138.5 and 137.3 (Cq), 133.0, 129.1 and 128.4 (aromatic CH), 80.8 (CHOCH₃), 74.0 (CHCH₂OCH₃), 59.0 (OCH₃), 58.1 (OCH₃), 17.1 (SCH₃), 16.5 (SCH₃). HRMS (ESI) calcd for C₁₅H₂₀O₃S₂ [M+Na]⁺: 335.0752; found: 335.0755.

2-((2-Methoxyethoxy)methyl)-3,3-bis(methylthio)-1-phenylprop-2-en-1-one (4q'): 42 mg, 27% yield; pale yellow liquid. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.91 (d, *J* = 7.6 Hz, 2 H, aromatic CH), 7.53 (t, 1 H, aromatic CH), 7.44 (t, 2 H, aromatic CH), 4.56 (s, 2 H, CH₂OCH₂CH₂), 3.59 and 3.43 (m each, 2:2 H, OCH₂CH₂O), 3.26 (s, 3 H, OCH₃), 2.36 (s, 3 H, SCH₃), 2.08 (s, 3 H, SCH₃). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 196.5 (Cq, C=O), 143.4, 140.7 and 137.5 (Cq), 133.0, 129.3 and 128.5 (aromatic CH), 71.8 (CH₂OCH₂CH₂), 71.0 (CH₂OCH₂CH₂), 70.2 (CH₂OCH₂CH₂), 59.1 (OCH₃), 17.2 (SCH₃), 16.6 (SCH₃). HRMS (ESI) calcd for C₁₅H₂₀O₃S₂ [M+Na]⁺: 335.0752; found: 335.0752.

Typical procedure for the methylation of ketene dithioacetals 1

Under a nitrogen atmosphere, a mixture of α-oxo ketene dithioacetal **1** (0.5 mmol), FeCl₃ (4 mg, 0.025 mmol), DABCO·6H₂O (11 mg, 0.05 mmol), and DTBP (219 mg, 1.5 mmol) in benzene (3 mL) was stirred in a 25-mL sealed tube at 130 °C for 24 h. After cooled to ambient temperature, the mixture was evaporated all the volatiles under reduced pressure. The resultant residue was purified by silica gel column chromatography (eluent: petroleum ether (60-90 °C)/ethyl acetate = 40:1, v/v) to afford the target product **5**.

2-Methyl-3,3-bis(methylthio)-1-phenylprop-2-en-1-one (5a): 74 mg, 62% yield; pale yellow liquid. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.86 (m, 2 H, aromatic CH), 7.56 (t, 1 H, aromatic CH), 7.46 (t, 2 H, aromatic CH), 2.36 (s, 3 H, SCH₃), 2.20 (s, 3 H, C=CCH₃), 2.05 (s, 3 H, SCH₃). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 197.5 (Cq, C=O), 143.6 (CSMe), 136.3 (Cq of Ph), 133.5

(C=CCH₃), 133.3, 129.1 and 128.7 (aromatic CH), 20.1 (C=CCH₃), 17.2 (SCH₃), 16.2 (SCH₃). HRMS (ESI) calcd for C₁₂H₁₄OS₂ [M+H]⁺: 239.0564; found: 239.0561.

2-Methyl-3,3-bis(methylthio)-1-(p-tolyl)prop-2-en-1-one (5b): 73 mg, 58% yield; pale yellow liquid. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.77 (d, *J* = 8.2 Hz, 2 H, aromatic CH), 7.26 (m, 2 H, aromatic CH), 2.41 (s, 3 H, Ph-CH₃), 2.35 (s, 3 H, SCH₃), 2.18 (s, 3 H, C=CCH₃), 2.07 (s, 3 H, SCH₃). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 197.3 (Cq, C=O), 144.2, 143.9 and 133.6 (Cq), 132.3 (Cq), 129.5 and 129.3 (aromatic CH), 21.8 (Ph-CH₃), 20.1 (C=CCH₃), 17.3 (SCH₃), 16.2 (SCH₃). HRMS (ESI) calcd for C₁₃H₁₆OS₂ [M+H]⁺: 253.0721; found: 253.0724.

1-(4-Chlorophenyl)-2-methyl-3,3-bis(methylthio)-prop-2-en-1-one (5c): 76 mg, 56% yield; pale yellow liquid. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.79 (d, *J* = 8.5 Hz, 2 H, aromatic CH), 7.43 (d, *J* = 8.5 Hz, 2 H, aromatic CH), 2.35 (s, 3 H, SCH₃), 2.18 (s, 3 H, C=CCH₃), 2.05 (s, 3 H, SCH₃). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 196.3 (Cq, C=O), 142.8, 139.6, 134.8 and 134.3 (Cq), 130.4 and 129.1 (aromatic CH), 20.1 (C=CCH₃), 17.3 (SCH₃), 16.2 (SCH₃). HRMS (ESI) calcd for C₁₂H₁₃ClOS₂ [M+H]⁺: 273.0175; found: 273.0175.

Typical procedure for the C–H alkylation of ketene dithioacetals **1** with toluene derivatives **6**

Under a nitrogen atmosphere, a mixture of α-oxo ketene dithioacetal **1** (0.5 mmol), FeCl₃ (4 mg, 0.025 mmol), DABCO·6H₂O (11 mg, 0.05 mmol), and DTBP (219 mg, 1.5 mmol) in toluene derivative **6** (3 mL) was stirred in a 25-mL sealed tube at 130 °C for 24 h. After cooled to ambient temperature, the mixture was evaporated all the volatiles under reduced pressure. The resultant residue was purified by silica gel column chromatography (eluent: petroleum ether (60–90 °C)/ethyl acetate = 100:1, v/v) to afford the target product **7**.

2-Benzyl-3,3-bis(methylthio)-1-phenylprop-2-en-1-one (7a): 124 mg, 79% yield; white solid, m.p. 60–62 °C. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.59 (d, *J* = 7.4 Hz, 2 H, aromatic CH), 7.43 (t, 1 H, aromatic CH), 7.30 (t, 2 H, aromatic CH), 7.17 (m, 5 H, aromatic CH), 4.09 (s, 2 H, PhCH₂), 2.39 and 2.00 (s each, 3:3 H, SCH₃). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 197.3 (Cq, C=O), 146.0, 137.7, 137.6 and 136.4 (Cq), 132.7, 129.2, 128.9, 128.6, 128.3 and 126.5 (aromatic CH), 40.86 (PhCH₂), 17.1 and 16.5 (SCH₃). HRMS (ESI) calcd for C₁₈H₁₈OS₂ [M+H]⁺: 315.0877; found: 315.0878.

2-Benzyl-1-(2-methoxyphenyl)-3,3-bis(methylthio)-prop-2-en-1-one (7b): 123 mg, 75% yield; pale yellow liquid. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.34 (m, 2 H, aromatic CH), 7.21 (m, 4 H, aromatic CH), 7.15 (m, 1 H, aromatic CH), 6.85 (dd, *J* = 7.8, 5.3 Hz, 2 H, aromatic CH), 4.08 (s, 2 H, PhCH₂), 3.69 (s, 3 H, OCH₃), 2.35 (s, 3 H, SCH₃), 1.95 (s, 3 H, SCH₃). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 195.8 (Cq, C=O), 158.6, 148.5, 138.6 and 137.2 (Cq), 133.2, 130.8 and 129.2 (aromatic CH), 128.7 (Cq), 128.4, 126.2, 120.1 and 111.5 (aromatic CH), 55.5 (OCH₃), 40.3 (PhCH₂), 16.93 (SCH₃), 16.6 (SCH₃). HRMS (ESI) calcd for C₁₉H₂₀O₂S₂ [M+H]⁺: 345.0983; found: 345.0982.

2-Benzyl-1-(3-methoxyphenyl)-3,3-bis(methylthio)-prop-2-en-1-one (7c): 132 mg, 77% yield; pale yellow liquid. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.15 (m, 8 H, aromatic CH), 6.98 (m, 1 H, aromatic CH), 4.08 (s, 2 H, PhCH₂), 3.73 (s, 3 H, OCH₃), 2.39 (s, 3 H, SCH₃), 2.01 (s, 3 H, SCH₃). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 197.1 (Cq, C=O), 159.6, 146.1, 139.0, 137.8, 136.3 (Cq), 129.2, 128.6, 126.6, 121.8, 119.7 and 112.4 (aromatic CH), 55.4 (OCH₃), 40.9 (PhCH₂), 17.1 (SCH₃), 16.5 (SCH₃). HRMS (ESI) calcd for C₁₉H₂₀O₂S₂ [M+H]⁺: 345.0983; found: 345.0982.

2-Benzyl-1-(4-methoxyphenyl)-3,3-bis(methylthio)-prop-2-en-1-one (7d): 132 mg, 77% yield; pale yellow liquid. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.59 (d, *J* = 8.8 Hz, 2 H, aromatic CH), 7.15 (m, 5 H, aromatic CH), 6.79 (d, *J* = 8.9 Hz, 2 H, aromatic CH), 4.07 (s, 2 H, PhCH₂), 3.80 (s, 3 H, OCH₃), 2.38 (s, 3 H, SCH₃), 2.06 (s, 3 H, SCH₃). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 196.1 (Cq, C=O), 163.3, 146.2, 137.8 and 134.9 (Cq), 131.4 (aromatic CH), 130.2 (Cq), 129.1, 128.5, 126.5 and 113.5 (aromatic CH), 55.5 (OCH₃), 40.8 (PhCH₂), 17.1 (SCH₃), 16.4 (SCH₃). HRMS (ESI) calcd for C₁₉H₂₀O₂S₂ [M+H]⁺: 345.0983; found: 345.0979.

2-Benzyl-3,3-bis(methylthio)-1-(p-tolyl)prop-2-en-1-one (7e): 130 mg, 79% yield; colorless liquid. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.51 (d, *J* = 8.1 Hz, 2 H, aromatic CH), 7.20 (m, 4 H, aromatic CH), 7.13 (m, 3 H, aromatic CH), 4.08 (s, 2 H, PhCH₂), 2.39 (s, 3 H, PhCH₃), 2.34 (s, 3 H, SCH₃), 2.04 (s, 3 H, SCH₃). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 196.7 (Cq, C=O), 146.1, 143.5, 137.8, 135.6 and 134.9 (Cq), 129.1, 129.0, 128.5 and 126.5 (aromatic CH), 40.8 (PhCH₂), 21.8 (PhCH₃), 17.1 (SCH₃), 16.4 (SCH₃). HRMS (ESI) calcd for C₁₉H₂₀OS₂ [M+H]⁺: 329.1034; found: 329.1034.

2-Benzyl-1-(2-fluorophenyl)-3,3-bis(methylthio)-prop-2-en-1-one (7f): 123 mg, 74% yield; yellow liquid. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.38 (m, 1 H, aromatic CH), 7.31 (m, 1 H, aromatic CH), 7.23 (m, 4 H, aromatic CH), 7.16 (m, 1 H, aromatic CH), 7.03 (m, 2 H, aromatic CH), 4.12 (s, 2 H, PhCH₂), 2.38 (s, 3 H, SCH₃), 1.97 (s, 3 H, SCH₃). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 193.5 (Cq, C=O), 161.2 (d, *J* = 256.8 Hz, Cq of Ph), 146.9 (Cq), 139.4 (d, *J* = 2.4 Hz, Cq), 138.0 (Cq), 133.6 (d, *J* = 8.8 Hz, aromatic CH), 130.5, 129.2 and 128.6 (aromatic CH), 127.60 (d, *J* = 10.3 Hz, Cq), 126.5 (aromatic CH), 123.8 (d, *J* = 3.7 Hz, aromatic CH), 116.4 (d, *J* = 22.2 Hz, aromatic CH), 40.4 (PhCH₂), 17.0 (SCH₃), 16.6 (SCH₃). HRMS (ESI) calcd for C₁₈H₁₇FOS₂ [M+H]⁺: 333.0783; found: 333.0783.

2-Benzyl-1-(3-chlorophenyl)-3,3-bis(methylthio)-prop-2-en-1-one (7g): 130 mg, 75% yield; white solid, m.p. 84–86 °C. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 7.55 (s, 1 H, aromatic CH), 7.38 (m, 2 H, aromatic CH), 7.18 (m, 6 H, aromatic CH), 4.08 (s, 2 H, PhCH₂), 2.40 (s, 3 H, SCH₃), 2.00 (s, 3 H, SCH₃). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 195.9 (Cq, C=O), 145.2, 139.5, 137.5, 137.4 and 134.5 (Cq), 132.4, 129.4, 129.0, 128.7, 126.8 and 126.7 (aromatic CH), 40.9 (PhCH₂), 17.0 (SCH₃), 16.5 (SCH₃). HRMS (ESI) calcd for C₁₈H₁₇ClOS₂ [M+H]⁺: 349.0488; found: 349.0484.

Methyl 4-(2-benzyl-3,3-bis(methylthio)acryloyl)benzoate (7h): 136 mg, 73% yield; pale yellow liquid. ¹H

NMR (400 MHz, CDCl_3 , 25 °C, TMS) δ 7.94 (d, J = 8.5 Hz, 2 H, aromatic CH), 7.56 (d, J = 8.4 Hz, 2 H, aromatic CH), 7.16 (m, 5 H, aromatic CH), 4.10 (s, 2 H, PhCH_2), 3.89 (s, 3 H, OCH_3), 2.39 (s, 3 H, SCH_3), 1.94 (s, 3 H, SCH_3). ^{13}C NMR (100 MHz, CDCl_3 , 25 °C, TMS) δ 196.7 (Cq, C=O), 166.4 (Cq, C=O), 145.4, 141.6, 137.9 and 137.5 (Cq of Ph), 133.1 (Cq), 129.4, 129.1, 128.7, 128.5 and 126.7 (aromatic CH), 52.5 (OCH_3), 41.0 (PhCH_2), 16.9 (SCH_3), 16.5 (SCH_3). HRMS (ESI) calcd for $\text{C}_{20}\text{H}_{20}\text{O}_3\text{S}_2$ $[\text{M}+\text{H}]^+$: 373.0932; found: 373.0931.

2-(2-Methylbenzyl)-3,3-bis(methylthio)-1-phenylprop-2-en-1-one (7i): 142 mg, 87% yield; colorless solid, m.p. 71–73 °C. ^1H NMR (400 MHz, CDCl_3 , 25 °C, TMS) δ 7.64 (d, J = 7.4 Hz, 2 H, aromatic CH), 7.45 (t, 1 H, aromatic CH), 7.32 (t, 2 H, aromatic CH), 7.16 (m, 1 H, aromatic CH), 7.03 (m, 3 H, aromatic CH), 4.08 (s, 2 H, PhCH_2), 2.40 (s, 3 H, SCH_3), 2.26 (s, 3 H, Ph-CH_3), 2.08 (s, 3 H, SCH_3). ^{13}C NMR (100 MHz, CDCl_3 , 25 °C, TMS) δ 196.9 (Cq, C=O), 146.0, 137.1, 136.5, 135.7 and 135.1 (Cq), 132.8, 130.3, 129.7, 128.8, 128.3, 126.7 and 126.1 (aromatic CH), 37.8 (PhCH_2), 20.0 (Ph-CH_3), 17.3 (SCH_3), 16.5 (SCH_3). HRMS (ESI) calcd for $\text{C}_{19}\text{H}_{20}\text{OS}_2$ $[\text{M}+\text{H}]^+$: 329.1034; found: 329.1029.

2-(3-Methylbenzyl)-3,3-bis(methylthio)-1-phenylprop-2-en-1-one (7j): 140 mg, 85% yield; pale yellow liquid. ^1H NMR (400 MHz, CDCl_3 , 25 °C, TMS) δ 7.62 (m, 2 H, aromatic CH), 7.44 (t, 1 H, aromatic CH), 7.32 (t, 2 H, aromatic CH), 7.09 (t, 1 H, aromatic CH), 6.97 (m, 3 H, aromatic CH), 4.08 (s, 2 H, PhCH_2), 2.40 (s, 3 H, SCH_3), 2.23 (s, 3 H, Ph-CH_3), 2.02 (s, 3 H, SCH_3). ^{13}C NMR (100 MHz, CDCl_3 , 25 °C, TMS) δ 197.3 (Cq, C=O), 145.9, 138.0, 137.5, 137.4 and 136.1 (Cq), 132.6, 129.9, 128.9, 128.3, 128.1, 127.2 and 126.0 (aromatic CH), 40.7 (PhCH_2), 21.3 (Ph-CH_3), 17.0 (SCH_3), 16.4 (SCH_3). HRMS (ESI) calcd for $\text{C}_{19}\text{H}_{20}\text{OS}_2$ $[\text{M}+\text{H}]^+$: 329.1034; found: 329.1035.

2-(4-Methylbenzyl)-3,3-bis(methylthio)-1-phenylprop-2-en-1-one (7k): 133 mg, 81% yield; white solid, m.p. 87–89 °C. ^1H NMR (400 MHz, CDCl_3 , 25 °C, TMS) δ 7.62 (d, J = 7.3 Hz, 2 H, aromatic CH), 7.45 (t, 1 H, aromatic CH), 7.31 (dd, J = 20.4 and 12.8 Hz, 2 H, aromatic CH), 7.04 (dd, J = 29.4 and 7.9 Hz, 4 H, aromatic CH), 4.06 (s, 2 H, PhCH_2), 2.39 (s, 3 H, SCH_3), 2.26 (s, 3 H, Ph-CH_3), 2.01 (s, 3 H, SCH_3). ^{13}C NMR (100 MHz, CDCl_3 , 25 °C, TMS) δ 197.3 (Cq, C=O), 146.2, 137.6, 136.1, 135.9 and 134.6 (Cq), 132.6 (s), 129.2, 129.0, 128.9 and 128.2 (aromatic CH), 40.4 (PhCH_2), 21.1 (Ph-CH_3), 17.0 (SCH_3), 16.4 (SCH_3). HRMS (ESI) calcd for $\text{C}_{19}\text{H}_{20}\text{OS}_2$ $[\text{M}+\text{H}]^+$: 329.1034; found: 329.10354.

2-(3,5-Dimethylbenzyl)-3,3-bis(methylthio)-1-phenylprop-2-en-1-one (7l): 137 mg, 80% yield; white solid, m.p. 88–89 °C. ^1H NMR (400 MHz, CDCl_3 , 25 °C, TMS) δ 7.63 (d, J = 7.5 Hz, 2 H, aromatic CH), 7.45 (t, 1 H, aromatic CH), 7.31 (dd, J = 20.2, 12.6 Hz, 2 H, aromatic CH), 6.77 (d, J = 8.7 Hz, 3 H, aromatic CH), 4.02 (s, 2 H, PhCH_2), 2.40 (s, 3 H, SCH_3), 2.19 (s, 6 H, Ph-CH_3), 2.02 (s, 3 H, SCH_3). ^{13}C NMR (100 MHz, CDCl_3 , 25 °C, TMS) δ 197.4 (Cq, C=O), 146.2, 137.9, 137.6, 137.4 and 135.9 (Cq), 132.6, 129.0, 128.2, 128.1 and 127.0 (aromatic CH), 40.7 (PhCH_2), 21.3 (Ph-CH_3), 17.0 (SCH_3), 16.4 (SCH_3). HRMS (ESI) calcd for $\text{C}_{20}\text{H}_{22}\text{OS}_2$ $[\text{M}+\text{H}]^+$: 343.1194; found: 343.1194.

2-(4-Methoxybenzyl)-3,3-bis(methylthio)-1-phenylprop-2-en-1-one (7m): 101 mg, 59% yield; white solid, m.p. 87–89 °C. ^1H NMR (400 MHz, CDCl_3 , 25 °C, TMS) δ 7.62 (m, 2 H, aromatic CH), 7.46 (t, 1 H, aromatic CH), 7.32 (dd, J = 18.6 and 11.0 Hz, 2 H, aromatic CH), 7.11 (d, J = 8.6 Hz, 2 H, aromatic CH), 6.75 (d, J = 8.6 Hz, 2 H, aromatic CH), 4.05 (s, 2 H, PhCH_2), 3.74 (s, 3 H, OCH_3), 2.41 (s, 3 H, SCH_3), 2.02 (s, 3 H, SCH_3). ^{13}C NMR (100 MHz, CDCl_3 , 25 °C, TMS) δ 197.4 (Cq, C=O), 158.2, 146.4, 137.6 and 135.8 (Cq), 132.7 and 130.1 (aromatic CH), 129.8 (Cq), 128.9, 128.3 and 114.0 (aromatic CH), 55.3 (OCH_3), 39.9 (PhCH_2), 17.0 (SCH_3), 16.5 (SCH_3). HRMS (ESI) calcd for $\text{C}_{19}\text{H}_{20}\text{O}_2\text{S}_2$ $[\text{M}+\text{H}]^+$: 345.0983; found: 345.0984.

3,3-Bis(methylthio)-1-phenyl-2-((p-tolylloxy)methyl)prop-2-en-1-one (7m'): 30 mg, 17% yield; pale yellow liquid. ^1H NMR (400 MHz, CDCl_3 , 25 °C, TMS) δ 7.92 (m, 2 H, aromatic CH), 7.54 (t, 1 H, aromatic CH), 7.44 (t, 2 H, aromatic CH), 7.02 (d, J = 8.4 Hz, 2 H, aromatic CH), 6.75 (d, J = 8.5 Hz, 2 H, aromatic CH), 5.01 (s, 2 H, PhOCH_2), 2.39 (s, 3 H, SCH_3), 2.25 (s, 3 H, Ph-CH_3), 2.11 (s, 3 H, SCH_3). ^{13}C NMR (100 MHz, CDCl_3 , 25 °C, TMS) δ 196.3 (Cq, C=O), 156.3, 142.9, 141.4 and 137.7 (Cq), 133.0 (aromatic CH), 130.5 (Cq), 129.9, 129.3, 128.5 and 114.9 (aromatic CH), 68.3 (PhOCH_2), 20.6 (PhCH_3), 17.2 (SCH_3), 16.7 (SCH_3). HRMS (ESI) calcd for $\text{C}_{19}\text{H}_{20}\text{O}_2\text{S}_2$ $[\text{M}+\text{Na}]^+$: 367.0802; found: 367.0803.

2-(4-(tert-Butyl)benzyl)-3,3-bis(methylthio)-1-phenylprop-2-en-1-one (7n): 146 mg, 79% yield; pale yellow liquid. ^1H NMR (400 MHz, CDCl_3 , 25 °C, TMS) δ 7.29 (m, 9 H, aromatic CH), 4.03 (s, 2 H, PhCH_2), 2.37 (s, 3 H, SCH_3), 1.98 (s, 3 H, SCH_3), 1.22 (s, 9 H, $\text{C}(\text{CH}_3)_3$). ^{13}C NMR (100 MHz, CDCl_3 , 25 °C, TMS) δ 197.5 (Cq, C=O), 149.3, 146.4, 137.6 and 135.7 and 134.6 (Cq), 132.6, 129.0, 128.8, 128.1 and 125.5 (aromatic CH), 40.3 (PhCH_2), 34.4 ($\text{C}(\text{CH}_3)_3$), 31.4 ($\text{C}(\text{CH}_3)_3$), 17.0 (SCH_3), 16.5 (SCH_3). HRMS (ESI) calcd for $\text{C}_{22}\text{H}_{26}\text{OS}_2$ $[\text{M}+\text{H}]^+$: 371.1503; found: 371.1497.

2-(2-Chlorobenzyl)-3,3-bis(methylthio)-1-phenylprop-2-en-1-one (7o): 138 mg, 79% yield; white solid, m.p. 92–94 °C. ^1H NMR (400 MHz, CDCl_3 , 25 °C, TMS) δ 7.69 (d, J = 7.5 Hz, 2 H, aromatic CH), 7.43 (t, 1 H, aromatic CH), 7.27 (m, 4 H, aromatic CH), 7.05 (m, 2 H, aromatic CH), 4.19 (s, 2 H, PhCH_2), 2.37 (s, 3 H, SCH_3), 2.07 (s, 3 H, SCH_3). ^{13}C NMR (100 MHz, CDCl_3 , 25 °C, TMS) δ 196.5 (Cq, C=O), 144.3, 137.0, 136.5, 135.2 and 134.2 (Cq), 132.9, 131.3, 129.4, 128.8, 128.3, 128.1 and 126.90 (aromatic CH), 38.0 (PhCH_2), 17.2 (SCH_3), 16.5 (SCH_3). HRMS (ESI) calcd for $\text{C}_{18}\text{H}_{17}\text{ClOS}_2$ $[\text{M}+\text{H}]^+$: 349.0488; found: 349.0490.

2-(3-Bromobenzyl)-3,3-bis(methylthio)-1-phenylprop-2-en-1-one (7p): 135 mg, 69% yield; pale yellow liquid. ^1H NMR (400 MHz, CDCl_3 , 25 °C, TMS) δ 7.63 (m, 2 H), 7.46 (m, 1 H, aromatic CH), 7.36 (dd, J = 12.3 and 4.5 Hz, 3 H, aromatic CH), 7.27 (m, 1 H, aromatic CH), 7.07 (m, 2 H, aromatic CH), 4.05 (s, 2 H, PhCH_2), 2.39 (s, 3 H, SCH_3), 2.04 (s, 3 H, SCH_3). ^{13}C NMR (100 MHz, CDCl_3 , 25 °C, TMS) δ 196.9 (Cq, C=O), 144.6, 140.0, 137.6 and 137.3 (Cq), 132.8, 132.1, 130.1, 129.6, 128.9, 128.4 and 127.8 (aromatic CH), 122.5 (Cq), 40.2 (PhCH_2), 17.1 (SCH_3), 16.5 (SCH_3). HRMS (ESI) calcd for $\text{C}_{18}\text{H}_{17}\text{BrOS}_2$ $[\text{M}+\text{H}]^+$: 392.9982; found: 392.9988.

2-(4-Fluorobenzyl)-3,3-bis(methylthio)-1-phenylprop-2-en-1-one (7q): 113 mg, 69% yield; pale yellow liquid. ^1H NMR (400 MHz, CDCl_3 , 25 °C, TMS) δ 7.58 (m, 2 H, aromatic CH), 7.45 (t, 1 H, aromatic CH), 7.32 (t, 2 H, aromatic CH), 7.14 (dd, J = 8.5 and 5.5 Hz, 2 H, aromatic CH), 6.88 (dd, J = 12.1 and 5.3 Hz, 2 H, aromatic CH), 4.04 (s, 2 H, PhCH_2), 2.38 (s, 3 H, SCH_3), 2.01 (s, 3 H, SCH_3). ^{13}C NMR (100 MHz, CDCl_3 , 25 °C, TMS) δ 197.2 (Cq, C=O), 161.0 (d, J = 244.6 Hz, Cq of Ph), 145.5, 137.5 and 136.7 (Cq), 133.5 (d, J = 3.2 Hz, Cq of Ph), 132.8 (aromatic CH), 130.6 (d, J = 7.9 Hz, aromatic CH), 128.88 and 128.35 (aromatic CH), 115.4 (d, J = 21.3 Hz, aromatic CH), 39.9 (PhCH_2), 17.0 (SCH_3), 16.4 (SCH_3). HRMS (ESI) calcd for $\text{C}_{18}\text{H}_{17}\text{FOS}_2$ $[\text{M}+\text{H}]^+$: 333.0783; found: 333.0783.

2-(4-Acetylbenzyl)-3,3-bis(methylthio)-1-phenylprop-2-en-1-one (7r): 92 mg, 51% yield; pale yellow liquid. ^1H NMR (400 MHz, CDCl_3 , 25 °C, TMS) δ 7.75 (d, J = 8.3 Hz, 2 H, aromatic CH), 7.55 (m, 2 H, aromatic CH), 7.40 (dd, J = 10.5 and 4.3 Hz, 1 H, aromatic CH), 7.25 (m, 4 H, aromatic CH), 4.08 (s, 2 H, PhCH_2), 2.48 (s, 3 H, SCH_3), 2.35 (s, 3 H, COCH_3), 1.97 (s, 3 H, SCH_3). ^{13}C NMR (100 MHz, CDCl_3 , 25 °C, TMS) δ 197.9 (Cq, C=O), 196.9 (Cq, C=O), 144.3 (CSMe), 143.5, 138.0 and 137.4 (Cq of Ph), 135.5 (C=CCH₂), 132.9, 129.4, 128.9, 128.8 and 128.4 (aromatic CH), 40.7 (PhCH_2), 26.7 (COCH_3), 17.2 (SCH_3), 16.5 (SCH_3). HRMS (ESI) calcd for $\text{C}_{20}\text{H}_{20}\text{O}_2\text{S}_2$ $[\text{M}+\text{H}]^+$: 357.0983; found: 357.0987.

3-Benzyl-4,4-bis(methylthio)but-3-en-2-one (7s): 96 mg, 76% yield; colourless liquid. ^1H NMR (400 MHz, CDCl_3 , 25 °C, TMS) δ 7.34 (t, 2 H, aromatic CH), 7.25 (m, 3 H, aromatic CH), 4.05 (s, 2 H, PhCH_2), 2.42 and 2.39 (s each, 3:3 H, SCH_3 and COCH_3), 2.21 (s, 3 H, SCH_3). ^{13}C NMR (100 MHz, CDCl_3 , 25 °C, TMS) δ 204.1 (Cq, C=O), 148.1, 138.0 and 136.5 (Cq), 128.7, 128.6 and 126.5 (aromatic CH), 40.0 (PhCH_2), 31.1 (COCH_3), 17.5 (SCH_3), 16.7 (SCH_3). HRMS (ESI) calcd for $\text{C}_{13}\text{H}_{16}\text{OS}_2$ $[\text{M}+\text{H}]^+$: 253.0721; found: 253.0723.

3-(4-Methylbenzyl)-4,4-bis(methylthio)but-3-en-2-one (7t): 117 mg, 88% yield; pale yellow liquid. ^1H NMR (400 MHz, CDCl_3 , 25 °C, TMS) δ 7.06 (q, J = 8.1 Hz, 4 H, aromatic CH), 3.93 (s, 2 H, PhCH_2), 2.35 (s, 3 H, SCH_3), 2.32 and 2.30 (s each, 3:3 H, Ph-CH_3 and COCH_3), 2.15 (s, 3 H, SCH_3). ^{13}C NMR (100 MHz, CDCl_3 , 25 °C, TMS) δ 204.3 (Cq, C=O), 148.5, 136.1, 136.0 and 134.8 (Cq), 129.4 and 128.5 (aromatic CH), 39.6 (PhCH_2), 31.2 (COCH_3), 21.1 (Ph-CH_3), 17.5 (SCH_3), 16.7 (SCH_3). HRMS (ESI) calcd for $\text{C}_{14}\text{H}_{18}\text{OS}_2$ $[\text{M}+\text{H}]^+$: 267.0877; found: 267.0877.

3-(3,5-Dimethylbenzyl)-4,4-bis(methylthio)but-3-en-2-one (7u): 122 mg, 87% yield; pale yellow solid, m.p. 53–54 °C. ^1H NMR (400 MHz, CDCl_3 , 25 °C, TMS) δ 6.83 (s, 1 H, aromatic CH), 6.76 (s, 2 H, aromatic CH), 3.91 (s, 2 H, PhCH_2), 2.36 and 2.33 (s each, 3:3 H, SCH_3 and Ph-CH_3), 2.27 (s, 6 H, COCH_3 and Ph-CH_3), 2.15 (s, 3 H, SCH_3). ^{13}C NMR (100 MHz, CDCl_3 , 25 °C, TMS) δ 204.4 (Cq, C=O), 148.5, 138.2, 137.8 and 136.1 (Cq), 128.3 and 126.5 (aromatic CH), 40.0 (PhCH_2), 31.3 (COCH_3), 21.4 (Ph-CH_3), 17.5 (SCH_3), 16.8 (SCH_3). HRMS (ESI) calcd for $\text{C}_{15}\text{H}_{20}\text{OS}_2$ $[\text{M}+\text{H}]^+$: 281.1034; found: 281.1037.

2-(1,3-Dithiolan-2-ylidene)-1,3-diphenylpropan-1-one (7v): 106 mg, 68% yield; pale yellow solid, m.p. 112–

114 °C. ^1H NMR (400 MHz, CDCl_3 , 25 °C, TMS) δ 7.52 (m, 3 H, aromatic CH), 7.35 (m, 5 H, aromatic CH), 7.20 (d, J = 7.3 Hz, 2 H, aromatic CH), 4.14 (s, 2 H, PhCH_2), 3.45 (m, 4 H, $\text{SCH}_2\text{CH}_2\text{S}$). ^{13}C NMR (100 MHz, CDCl_3 , 25 °C, TMS) δ 193.6 (Cq, C=O), 164.4, 139.8, 138.8 (Cq of Ph), 130.6, 128.5, 128.2, 128.1, 127.6 and 126.2 (aromatic CH), 122.9 (Cq), 40.4 (PhCH_2), 39.2 and 36.4 ($\text{SCH}_2\text{CH}_2\text{S}$). HRMS (ESI) calcd for $\text{C}_{18}\text{H}_{16}\text{OS}_2$ $[\text{M}+\text{H}]^+$: 313.0721; found: 313.0728.

2-(Bis(methylthio)methylene)-1,3-diphenylbutan-1-one (7w): 90 mg, 55% yield; yellow solid, m.p. 62–64 °C. ^1H NMR (400 MHz, CDCl_3 , 25 °C, TMS) δ 7.61 (d, J = 7.4 Hz, 2 H, aromatic CH), 7.41 (t, 1 H, aromatic CH), 7.27 (dd, J = 15.2 and 7.6 Hz, 4 H, aromatic CH), 7.15 (t, 2 H, aromatic CH), 7.07 (t, 1 H, aromatic CH), 4.74 (q, J = 7.3 Hz, 1 H, CHCH_3), 2.32 (s, 3 H, SCH_3), 2.00 (s, 3 H, SCH_3), 1.49 (d, J = 7.3 Hz, 3 H, CHCH_3). ^{13}C NMR (100 MHz, CDCl_3 , 25 °C, TMS) δ 196.0 (Cq, C=O), 150.9, 142.8, 137.6 and 133.3 (Cq), 132.7, 129.1, 128.3, 128.2, 128.1, and 126.5 (aromatic CH), 43.8 (PhCH), 19.4 (CHCH_3), 17.1 (SCH_3), 16.4 (s). HRMS (ESI) calcd for $\text{C}_{19}\text{H}_{20}\text{OS}_2$ $[\text{M}+\text{H}]^+$: 329.1034; found: 329.1035.

3-(Bis(methylthio)methylene)-4-phenylpentan-2-one (7x): 87 mg, 59% yield; white solid, m.p. 65–67 °C. ^1H NMR (400 MHz, CDCl_3 , 25 °C, TMS) δ 7.25 (m, 5 H, aromatic CH), 4.65 (q, J = 7.2 Hz, 1 H, C=CCH), 2.33 and 2.26 (s each, 3:3 H, SCH_3 and COCH_3), 1.86 (s, 3 H, SCH_3), 1.49 (d, J = 7.2 Hz, 3 H, CHCH_3). ^{13}C NMR (100 MHz, CDCl_3 , 25 °C, TMS) δ 204.4 (Cq, C=O), 155.3, 142.1 and 130.3 (Cq), 128.6, 127.8 and 126.8 (aromatic CH), 42.7 (PhCH_2), 32.2 (COCH_3), 18.2 (CHCH_3), 17.4 (SCH_3), 16.5 (SCH_3). HRMS (ESI) calcd for $\text{C}_{14}\text{H}_{18}\text{OS}_2$ $[\text{M}+\text{H}]^+$: 267.0877; found: 267.0874.

2-(1,4-Dioxan-2-yl)-3-(methylthio)-1,3-diphenylprop-2-en-1-one (9a): 97 mg, 57% yield; white solid, m.p. 137–139 °C. ^1H NMR (400 MHz, CDCl_3 , 25 °C, TMS) δ for (*E*)-isomer: 7.62 (m, 2 H, aromatic CH), 7.49 (m, 2 H, aromatic CH), 7.31 (m, 1 H, aromatic CH), 7.18 (t, 2 H, aromatic CH), 7.04 (m, 3 H, aromatic CH), 5.10 (dd, J = 10.2 and 2.7 Hz, 1 H, OCHCH_2O), 4.06 (dd, J = 11.5 and 2.7 Hz) and 3.65 (m) (1:5 H, $\text{CH}_2\text{OCH}_2\text{CH}_2\text{O}$), 1.85 (s, 3 H, SCH_3); for (*Z*)-isomer: 8.11 (m, 2 H, aromatic CH), 7.57 (t, 1 H, aromatic CH), 7.49 (m, 3 H, aromatic CH), 7.40 (m, 1 H, aromatic CH), 7.04 (m, 3 H, aromatic CH), 4.35 (dd, J = 10.0 and 2.9 Hz, 1 H, OCHCH_2O), 3.65 (m, 6 H, $\text{CH}_2\text{OCH}_2\text{CH}_2\text{O}$), 1.62 (s, 3 H, SCH_3). ^{13}C NMR (100 MHz, CDCl_3 , 25 °C, TMS) δ for (*E*)-isomer: 197.1 (Cq, C=O) [190.0 for (*Z*)-isomer], 143.2 (Cq) [141.62 for (*Z*)-isomer], 137.9 (Cq) [137.8 for (*Z*)-isomer], 136.0 (Cq) [136.2 and 135.6 for (*Z*)-isomer], 132.6, 129.8, 129.3, 128.4, 128.1 and 127.8 (aromatic CH) [133.1, 129.4, 129.0, 128.8, 128.9 and 128.5 for (*Z*)-isomer], 76.6 (OCHCH_2O) [76.3 for (*Z*)-isomer], 69.2 (s), 68.6 ($\text{CH}_2\text{OCH}_2\text{CH}_2\text{O}$) [69.2 for (*Z*)-isomer], 67.0 and 66.2 ($\text{CH}_2\text{OCH}_2\text{CH}_2\text{O}$) [66.5 and 65.9 for (*Z*)-isomer], 15.6 (SCH_3) [15.4 for (*Z*)-isomer]. HRMS (ESI) calcd for $\text{C}_{20}\text{H}_{20}\text{O}_3\text{S}$ $[\text{M}+\text{H}]^+$: 341.1211; found: 341.1211.

3-(1,4-Dioxan-2-yl)-4-(methylthio)-4-phenylbut-3-en-2-one (9b): 99 mg, 71% yield; white solid, m.p. 73–75 °C. ^1H NMR (400 MHz, CDCl_3 , 25 °C, TMS) δ for (*E*)-isomer: 7.39 (m, 3 H, aromatic CH), 7.27 (m, 2 H, aromatic CH), 4.18 (dd, J = 10.3 and 2.7 Hz, 1 H, OCHCH_2O), 3.87, 3.67

and 3.51 (m each, 2:2:2 H, $\text{CH}_2\text{OCH}_2\text{CH}_2\text{O}$), 2.54 (s, 3 H, SCH_3), 1.79 (s, 3 H, COCH_3); for (Z)-isomer: 7.39 (m, 3 H, aromatic CH), 7.27 (m, 2 H, aromatic CH), 4.95 (dd, $J = 9.9$ and 3.0 Hz, 1 H, OCHCH_2O), 3.87, 3.67 and 3.51 (m each, 2:2:2 H, $\text{CH}_2\text{OCH}_2\text{CH}_2\text{O}$), 1.86 (s, 3 H, SCH_3), 1.76 (s, 3 H, COCH_3). ^{13}C NMR (100 MHz, CDCl_3 , 25 °C, TMS) δ for (E)-isomer: 203.3 (Cq, C=O) [203.8 for (Z)-isomer], 141.8 (Cq) [142.1 for (Z)-isomer], 137.4 (Cq) [138.6 for (Z)-isomer], 135.4 (Cq) [136.3 for (Z)-isomer], 128.7, 128.6 and 128.5 (aromatic CH) [129.3, 128.8 and 128.7 for (Z)-isomer], 75.7 (OCHCH_2O) [76.6 for (Z)-isomer], 68.5 (OCHCH_2O) [67.9 for (Z)-isomer], 66.5 and 65.8 ($\text{OCH}_2\text{CH}_2\text{O}$) [66.8 and 66.0 for (Z)-isomer], 31.9 (COCH_3) [32.0 for (Z)-isomer], 15.4 (SCH_3). HRMS (ESI) calcd for $\text{C}_{15}\text{H}_{18}\text{O}_3\text{S}$ $[\text{M}+\text{H}]^+$: 279.1055; found: 279.1048.

4-(4-Chlorophenyl)-3-(1,4-dioxan-2-yl)-4-(methylthio)but-3-en-2-one (9c): 117 mg, 75% yield; white solid, m.p. 86–88 °C. ^1H NMR (400 MHz, CDCl_3 , 25 °C, TMS) for (E)-isomer: 7.37 (d, $J = 8.3$ Hz, 2 H, aromatic CH), 7.21 (d, $J = 8.3$ Hz, 2 H, aromatic CH), 4.08 (dd, $J = 10.3$ and 2.8 Hz, 1 H, OCHCH_2O), 3.86 (dd, $J = 11.4$ and 2.8 Hz) and 3.75 (m) (1:1 H, $\text{CH}_2\text{OCH}_2\text{CH}_2\text{O}$), 3.54 (m, 4 H, $\text{OCH}_2\text{CH}_2\text{O}$), 2.48 (s, 3 H, SCH_3), 1.76 (s, 3 H, COCH_3); for (Z)-isomer: 7.30 (d, $J = 8.4$ Hz, 2 H, aromatic CH), 7.16 (d, $J = 8.4$ Hz, 2 H, aromatic CH), 4.89 (dd, $J = 10.1$ and 2.9 Hz, 1 H, OCHCH_2O), 3.75 (m, 2 H, $\text{CH}_2\text{OCH}_2\text{CH}_2\text{O}$), 3.54 (m, 4 H, $\text{OCH}_2\text{CH}_2\text{O}$), 1.82 (s, 3 H, SCH_3), 1.78 (s, 3 H, COCH_3). ^{13}C NMR (100 MHz, CDCl_3 , 25 °C, TMS) δ for (E)-isomer: 203.4 (Cq, C=O) [203.8 for (Z)-isomer], 139.8 (Cq) [140.5], 138.4 and 134.8 (Cq) [139.9 and 135.0 for (Z)-isomer], 134.0 (Cq) [134.8 for (Z)-isomer], 130.1 and 129.1 (aromatic CH) [130.8 and 129.0 for (Z)-isomer], 75.6 (OCHCH_2O) [76.5 for (Z)-isomer], 68.5 (OCHCH_2O) [68.0 for (Z)-isomer], 66.6 and 65.9 ($\text{OCH}_2\text{CH}_2\text{O}$) [66.9 and 66.1 for (Z)-isomer], 31.9 (COCH_3) [32.3 for (Z)-isomer], 15.60 (SCH_3). HRMS (ESI) calcd for $\text{C}_{15}\text{H}_{18}\text{ClO}_3\text{S}$ $[\text{M}+\text{H}]^+$: 313.0665; found: 313.0670.

3-(1,4-Dioxan-2-yl)-4,4-diphenylbut-3-en-2-one (11): 16 mg, 10% yield; white solid, m.p. 94–96 °C. ^1H NMR (400 MHz, CDCl_3 , 25 °C, TMS) δ 7.31 (m, 3 H, aromatic CH), 7.20 (m, 5 H, aromatic CH), 7.17 (m, 2 H, aromatic CH), 4.39 (dd, $J = 10.2$ and 2.7 Hz, 1 H, OCHCH_2O), 3.90 (dd, $J = 11.5$ and 10.3 Hz) and 3.69 (d, $J = 10.4$ Hz) (1:1 H, OCHCH_2O), 3.55 (m, 4 H, $\text{OCH}_2\text{CH}_2\text{O}$), 1.86 (s, 3 H, COCH_3). ^{13}C NMR (100 MHz, CDCl_3 , 25 °C, TMS) δ 206.9 (Cq, C=O), 144.9 (CSMe), 140.3 and 139.8 (Cq of Ph), 138.5 (C=CCH), 129.4, 129.0, 128.8, 128.7, 128.6 and 128.2 (aromatic CH), 77.0 (OCHCH_2O), 68.9 ($\text{CH}_2\text{OCH}_2\text{CH}_2\text{O}$), 66.6 and 66.1 ($\text{OCH}_2\text{CH}_2\text{O}$), 32.7 (COCH_3). HRMS (ESI) calcd for $\text{C}_{20}\text{H}_{20}\text{O}_3$ $[\text{M}+\text{H}]^+$: 309.1491; found: 309.1491.

Typical procedure for the synthesis of trisubstituted pyrazoles 12

A mixture of **3** or **7** (0.2 mmol) and $\text{NH}_2\text{NH}_2\cdot\text{H}_2\text{O}$ (114 mg, 2.0 mmol, aq 85%) in toluene (2 mL) was stirred in a 10-mL sealed tube at 120 °C for 24 h. After cooled to ambient temperature, the mixture was evaporated all the volatiles under reduced pressure. The resultant residue was

purified by silica gel column chromatography (eluent: dichloromethane) to afford the target product.

4-(1,4-Dioxan-2-yl)-5-(methylthio)-3-phenyl-1H-pyrazole (12a): 52 mg, 95% yield; white solid, m.p. 149–151 °C. ^1H NMR (400 MHz, CDCl_3 , 25 °C, TMS) δ 7.45 (m, 5 H, aromatic CH), 4.72 (dd, $J = 10.7$ and 2.8 Hz, 1 H, OCHCH_2O), 3.99 (t) and 3.89 (d, $J = 11.3$ Hz) (1:1 H, OCHCH_2O), 3.80 (m), 3.70 (d, $J = 5.8$ Hz) and 3.60 (dd, $J = 11.7$ and 2.7 Hz) (1:2:1 H, $\text{OCH}_2\text{CH}_2\text{O}$), 2.44 (s, 3 H, SCH_3). ^{13}C NMR (100 MHz, CDCl_3 , 25 °C, TMS) δ 145.8, 144.8 and 130.1 (Cq), 129.0, 128.8 and 128.7 (aromatic CH), 114.5 (Cq), 70.9 (OCHCH_2O), 69.0 (OCHCH_2O), 67.4 and 66.2 ($\text{OCH}_2\text{CH}_2\text{O}$), 16.7 (SCH_3). HRMS (ESI) calcd for $\text{C}_{14}\text{H}_{16}\text{N}_2\text{O}_2\text{S}$ $[\text{M}+\text{H}]^+$: 277.1011; found: 277.1008.

4-(1,4-Dioxan-2-yl)-3-methyl-5-(methylthio)-1H-pyrazole (12b): 41 mg, 95% yield; white solid, m.p. 88–90 °C. ^1H NMR (400 MHz, CDCl_3 , 25 °C, TMS) δ 7.72 (s, 1 H, NH), 4.67 (dd, $J = 10.4$ and 3.0 Hz, 1 H, OCHCH_2O), 3.75 (m, 6 H, $\text{OCH}_2\text{CH}_2\text{OCH}_2$), 2.40 (s, 3 H, SCH_3), 2.33 (s, 3 H, CH_3). ^{13}C NMR (100 MHz, CDCl_3 , 25 °C, TMS) δ 143.5, 141.2 and 115.2 (Cq), 71.4 (OCHCH_2O), 70.0 (OCHCH_2O), 67.4 and 66.4 ($\text{OCH}_2\text{CH}_2\text{O}$), 17.5 (SCH_3), 11.1 (CH_3). HRMS (ESI) calcd for $\text{C}_9\text{H}_{14}\text{N}_2\text{O}_2\text{S}$ $[\text{M}+\text{H}]^+$: 215.0854; found: 215.0851.

3-Cyclopropyl-4-(1,4-dioxan-2-yl)-5-(methylthio)-1H-pyrazole (12c): 46 mg, 96% yield; white solid, m.p. 128–130 °C. ^1H NMR (400 MHz, CDCl_3 , 25 °C, TMS) δ 8.26 (s, 1 H, NH), 4.71 (dd, $J = 10.6$ and 2.7 Hz, 1 H, OCHCH_2O), 3.75 (m, 6 H, $\text{OCH}_2\text{CH}_2\text{OCH}_2$), 2.35 (s, 3 H, SCH_3), 1.94 (m, 1 H, CH_2CHCH_2), 0.82 and 0.63 (m each, 3:1 H, CH_2CHCH_2). ^{13}C NMR (100 MHz, CDCl_3 , 25 °C, TMS) δ 147.2, 142.9 and 115.9 (Cq of pyrazole), 71.3 (OCHCH_2O), 69.9 (OCHCH_2O), 67.4 and 66.4 ($\text{OCH}_2\text{CH}_2\text{O}$), 17.3 (SCH_3), 7.5 (CH_2CHCH_2), 6.7 and 6.6 (CH_2CHCH_2). HRMS (ESI) calcd for $\text{C}_{11}\text{H}_{16}\text{N}_2\text{O}_2\text{S}$ $[\text{M}+\text{H}]^+$: 241.1011; found: 241.1012.

4-Benzyl-3-methyl-5-(methylthio)-1H-pyrazole (12d): 38 mg, 87% yield; colourless liquid. ^1H NMR (400 MHz, CDCl_3 , 25 °C, TMS) δ 8.65 (s, 1 H, NH), 7.21 (m, 5 H, aromatic CH), 3.82 (s, 2 H, PhCH_2), 2.35 (s, 3 H, SCH_3), 2.19 (s, 3 H, CH_3). ^{13}C NMR (100 MHz, CDCl_3 , 25 °C, TMS) δ 142.7, 141.7 and 140.5 (Cq), 128.5, 128.3 and 126.1 (aromatic CH), 118.3 (Cq), 29.3 (PhCH_2), 17.6 (SCH_3), 10.9 (CH_3). HRMS (ESI) calcd for $\text{C}_{12}\text{H}_{14}\text{N}_2\text{S}$ $[\text{M}+\text{H}]^+$: 219.0956; found: 219.0956.

Typical procedure for the synthesis of isoxazoles 13

A mixture of **3** (50 mg, 0.2 mmol), $\text{NH}_2\text{OH}\cdot\text{HCl}$ (83 mg, 1.2 mmol), and KOH (79 mg, 1.2 mmol, 85%) in EtOH (2 mL) was stirred in a 10-mL sealed tube at 120 °C for 24 h. After cooled to ambient temperature, the mixture was evaporated all the volatiles under reduced pressure. The resultant residue was purified by silica gel column chromatography (eluent: petroleum ether (60–90 °C)/ethyl acetate = 10:1, v/v) to afford the target product.

4-(1,4-Dioxan-2-yl)-3-methyl-5-(methylthio)isoxazole (13a): 24 mg, 56% yield; colorless liquid. ^1H NMR (400 MHz, CDCl_3 , 25 °C, TMS) δ 4.53 (dd, $J = 8.0$ and 5.5 Hz, 1 H, OCHCH_2O), 3.76 (m, 6 H, $\text{OCH}_2\text{CH}_2\text{OCH}_2$), 2.55 (s, 3 H, SCH_3), 2.30 (s, 3 H, $\text{CH}_3\text{C}=\text{N}$). ^{13}C NMR (100 MHz,

CDCl₃, 25 °C, TMS) δ 163.7, 159.8 and 112.6 (Cq of isoxazole), 70.1 (OCHCH₂O), 69.3 (OCHCH₂O), 67.2 and 66.4 (OCH₂CH₂O), 14.9 (SCH₃), 11.1 (CH₃C=N). HRMS (ESI) calcd for C₉H₁₃NO₃S [M+H]⁺: 216.0694; found: 216.0690.

3-Cyclopropyl-4-(1,4-dioxan-2-yl)-5-(methylthio)isoxazole (13b): 26 mg, 54% yield; white solid, m.p. 71-73 °C. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 4.65 (dd, *J* = 10.4 and 2.9 Hz, 1 H, OCHCH₂O), 3.78 (m, 6 H, OCH₂CH₂OCH₂), 2.55 (s, 3 H, SCH₃), 1.94 (m, 1 H, CH₂CHCH₂), 1.08 and 0.94 (m each, 1:3 H, CH₂CHCH₂). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 165.0, 163.7 and 113.1 (Cq of isoxazole), 70.1 (OCHCH₂O), 69.5 (OCHCH₂O), 67.3 and 66.4 (OCH₂CH₂O), 14.9 (SCH₃), 8.5 (CH₂CHCH₂), 7.3 and 6.5 (CH₂CHCH₂). HRMS (ESI) calcd for C₁₁H₁₅NO₃S [M+H]⁺: 242.0851; found: 242.0854.

3-Methyl-5-(methylthio)-4-(tetrahydrofuran-2-yl)isoxazole (13c): 22 mg, 55% yield; colorless liquid. ¹H NMR (400 MHz, CDCl₃, 25 °C, TMS) δ 4.74 (dd, *J* = 8.6 and 6.7 Hz, 1 H, OCHCH₂), 4.02 and 3.84 (m each, 1:1 H, OCH₂CH₂), 2.55 (s, 3 H, SCH₃), 2.25 (s, 3 H, CH₃C=N), 2.08 and 1.85 (m each, 3:1 H, OCHCH₂CH₂). ¹³C NMR (100 MHz, CDCl₃, 25 °C, TMS) δ 162.2, 159.7 and 116.6 (Cq of isoxazole), 72.5 (OCHCH₂), 68.6 (OCH₂CH₂), 32.1 and 26.6 (OCHCH₂CH₂), 15.1 (SCH₃), 11.0 (CH₃C=N). HRMS (ESI) calcd for C₉H₁₃NO₂S [M+H]⁺: 200.0745; found: 200.0741.

Acknowledgements

We are grateful to the National Natural Science Foundation of China (21472185) and the National Basic Research Program of China (2015CB856600) for support of this research.

References

- [1] E.-I. Negishi, Z. H. Huang, G. W. Wang, S. Mohan, C. Wang, H. S. Hattori, *Acc. Chem. Res.* **2008**, *41*, 1474.
- [2] a) I. P. Beletskaya, A. V. Cheprakov, *Chem. Rev.* **2000**, *100*, 3009; b) X. J. Shang, Z. Q. Liu, *Chem. Soc. Rev.* **2013**, *42*, 3253.
- [3] a) Y. H. Takeda, M. Shimizu, T. Hiyama, *Angew. Chem. Int. Ed.* **2007**, *46*, 8659; b) S. G. Newman, M. Lautens, *J. Am. Chem. Soc.* **2010**, *132*, 11416.
- [4] a) Y. Nishihara, M. Miyasaka, M. Okamoto, H. Inoue, E. Takahashi, K. Tanemura, K. Takagi, *J. Am. Chem. Soc.* **2007**, *129*, 12634; b) M. Shimizu, I. Nagao, Y. Tomioka, T. Hiyama, *Angew. Chem. Int. Ed.* **2008**, *47*, 8096; c) C. Wang, T. Tobrman, Z. Q. Xu, E.-I. Negishi, *Org. Lett.* **2009**, *11*, 4092.
- [5] a) X. Bresser, P. Knochel, *Angew. Chem. Int. Ed.* **2011**, *50*, 1914; b) R. Egris, M. Villacampa, J. C. Menéndez, *Chem.-Eur. J.* **2009**, *15*, 10930; c) K. Akiyama, F. Gao, A. H. Hoveyda, *Angew. Chem., Int. Ed.* **2010**, *49*, 419–423.
- [6] a) K. Itami, Y. Ohashi, J.-i. Yoshida, *J. Org. Chem.* **2005**, *70*, 2778; b) M. Movassaghi, A. E. Ondrus, *J. Org. Chem.* **2005**, *70*, 8638; c) C. L. Sun, Y. Wang, X. Zhou, Z. H. Wu, B. J. Li, B. T. Guan, Z. J. Shi, *Chem.-Eur. J.* **2010**, *16*, 5844; d) Y. Ashida, Y. Sato, T. Ueno, K. Suzuki, K.-I. Kai, H. Nakatsuji, Y. Tanabe, *Chem.-Eur. J.* **2015**, *21*, 5934.
- [7] For selected recent reviews on C–H activation (functionalization), see: a) L. Yang, H. M. Huang, *Chem. Rev.* **2015**, *115*, 3468; b) M. Moselage, J. Li, L. Ackermann, *ACS Catal.* **2016**, *6*, 498; c) T. Gensch, M. N. Hopkinson, F. Glorius, J. W. Delord, *Chem. Soc. Rev.* **2016**, *45*, 2900.
- [8] a) C.-J. Li, *Acc. Chem. Res.* **2009**, *42*, 335; b) C. Liu, H. Zhang, W. Shi, A. W. Lei, *Chem. Rev.* **2011**, *111*, 1780; c) F. Jia, Z. P. Li, *Org. Chem. Front.* **2014**, *1*, 194; d) S. A. Girard, T. Knauber, C.-J. Li, *Angew. Chem. Int. Ed.* **2014**, *53*, 74; e) R. Vanjari, K. N. Singh, *Chem. Soc. Rev.* **2015**, *44*, 8062; f) L. Y. Lv, Z. P. Li, *Top. Curr. Chem.* **2016**, *374*, 38.
- [9] Selected recent reports on transition-metal-catalyzed oxidative C(sp³)–H functionalization, see: a) Z. P. Li, L. Cao, C. J. Li, *Angew. Chem. Int. Ed.* **2007**, *46*, 6505; b) Y. H. Zhang, C. J. Li, *Eur. J. Org. Chem.* **2007**, 4654; c) Z. P. Li, R. Yu, H. J. Li, *Angew. Chem. Int. Ed.* **2008**, *47*, 7497; d) K. Z. Li, G. Y. Tan, J. S. Huang, F. J. Song, J. S. You, *Angew. Chem. Int. Ed.* **2013**, *52*, 12942; e) D. Liu, C. Liu, H. Li, A. W. Lei, *Angew. Chem. Int. Ed.* **2013**, *52*, 4453; f) W. T. Wei, M.-B. Zhou, J. H. Fan, W. Liu, R.-J. Song, Y. Liu, M. Hu, P. Xie, J. H. Li, *Angew. Chem. Int. Ed.* **2013**, *52*, 3638; g) A. Modak, U. Dutta, R. Kancherla, S. Maity, M. Bhadra, S. M. Mobin, D. Maiti, *Org. Lett.* **2014**, *16*, 2602; h) K. Z. Li, Q. Wu, J. B. Lan, J. S. You, *Nat. Commun.* **2015**, *6*, 8404; i) M. Hu, J. H. Fan, Y. Liu, X. H. Ouyang, R. J. Song, J. H. Li, *Angew. Chem. Int. Ed.* **2015**, *54*, 9577; j) M. Brzozowski, J. A. Forni, G. P. Savage, A. Polyzos, *Chem. Commun.* **2015**, *51*, 334; k) S. Tang, P. Wang, H. R. Li, A. W. Lei, *Nat. Commun.* **2016**, *7*, 11676; l) H. Aruri, U. Singh, M. Kumar, S. Sharma, S. K. Aithagani, V. K. Gupta, S. Mignan, R. A. Vishwakarma, P. P. Singh, *J. Org. Chem.* **2017**, *82*, 1000.
- [10] a) S. H. Li, G. Cheng, C. G. Feng, W. Gong, J. Q. Yu, *J. Am. Chem. Soc.* **2014**, *136*, 5267; b) H. Jiang, J. He, T. Liu, J. Q. Yu, *J. Am. Chem. Soc.* **2016**, *138*, 2055.
- [11] K. J. Stowers, K. C. Fortner, M. S. Sanford, *J. Am. Chem. Soc.* **2011**, *133*, 6541.
- [12] G. Zhang, Y. X. Ma, S. L. Wang, Y. H. Zhang, R. Wang, *J. Am. Chem. Soc.* **2012**, *134*, 12334.
- [13] H. Wang, Y. L. Zhao, L. Li, S. S. Li, Q. Liu, *Adv. Synth. Catal.* **2014**, *356*, 3157.
- [14] Y. F. Zhu, Y. Y. Wei, *Chem. Sci.* **2014**, *5*, 2379.
- [15] D. Liu, C. Liu, H. Li, A. W. Lei, *Chem. Commun.* **2014**, *50*, 3623.
- [16] G. P. Qin, X. N. Chen, L. Yang, H. M. Huang, *ACS Catal.* **2015**, *5*, 2882.
- [17] a) B. Niu, W. N. Zhao, Y. C. Ding, Z. G. Bian, C. U. Pittman, A. H. Zhou, H. B. Ge, *J. Org. Chem.* **2015**, *80*, 7251; b) B. A. Mir, A. Banerjee, S. K. Santra, S. Rajamanickam, B. K. Patel, *Adv. Synth. Catal.* **2016**, *358*, 3471; c) L. Y. Dian, H. Zhao, D. Z. Negrier, Y. F. Du, *Adv. Synth. Catal.* **2016**, *358*, 2422.
- [18] a) H. F. Yu, Z. K. Yu, *Angew. Chem. Int. Ed.* **2009**, *48*, 2929; b) H. F. Yu, W. W. Jin, C. L. Sun, J. P. Chen, W. M. Du, S. B. He, Z. K. Yu, *Angew. Chem.*

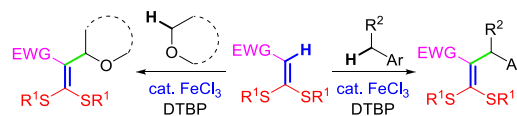
- Int. Ed.* **2010**, 49, 5792; c) W. W. Jin, W. M. Du, Q. Yang, H. F. Yu, J. P. Chen, Z. K. Yu, *Org. Lett.* **2011**, 13, 4272; d) Z. F. Mao, F. Huang, H. F. Yu, J. P. Chen, Z. K. Yu, Z. Q. Xu, *Chem.-Eur. J.* **2014**, 20, 3439; e) W. W. Jin, Q. Yang, P. Wu, J. P. Chen, Z. K. Yu, *Adv. Synth. Catal.* **2014**, 356, 2097; f) X. G. Yang, Z. Q. Liu, C. L. Sun, J. P. Chen, Z. K. Yu, *Chem.-Eur. J.* **2015**, 21, 14085; g) P. Wu, L. D. Wang, K. K. Wu, Z. K. Yu, *Org. Lett.* **2015**, 17, 868; h) X. G. Yang, K. K. Wu, P. Wu, J. P. Chen, C. L. Sun, Z. K. Yu, *Chem.-Eur. J.* **2015**, 21, 9323; i) T. L. Guo, Q. B. Jiang, Z. K. Yu, *Adv. Synth. Catal.* **2016**, 358, 3450.
- [19] L. D. Wang, W. He, Z. K. Yu, *Chem. Soc. Rev.* **2013**, 42, 599.
- [20] L. Pan, X. H. Bi, Q. Liu, *Chem. Soc. Rev.* **2013**, 42, 1251.
- [21] Q. Yang, P. Wu, J. P. Chen, Z. K. Yu, *Chem. Commun.* **2014**, 50, 6337.
- [22] Selected recent reports on iron-catalyzed C–H functionalization, see: a) C. L. Sun, B. J. Li, Z. J. Shi, *Chem. Rev.* **2011**, 111, 1293; b) J. H. Fan, M. B. Zhou, Y. Liu, W. T. Wei, X. H. Ouyang, R. J. Song, J. H. Li, *Synlett* **2014**, 25, 657; c) L. Iies, T. Matsubara, S. Ichikawa, S. Asako, E. Nakamura, *J. Am. Chem. Soc.* **2014**, 136, 13126; d) I. Bauer, H.-J. Knölker, *Chem. Rev.* **2015**, 115, 3170.
- [23] During the review period a communication on DTBP-promoted C–H alkylation of α -oxo ketene dithioacetals was published online. See: J. W. Wen, F. Zhang, W. Y. Shi, A. W. Lei, *Chem. - Eur. J.* **2017**, DOI:10.1002/chem.201701664.
- [24] CCDC 1486033 for **3a** contains the supplementary crystallographic data for this paper. These data can be obtained free of charge from The Cambridge Crystallographic Data Centre via www.ccdc.cam.ac.uk/data_request/cif.
- [25] a) Q. Dai, J. T. Yu, Y. Jiang, S. J. Guo, H. T. Yang, J. Cheng, *Chem. Commun.* **2014**, 50, 3865; b) N. B. Zhu, J. G. Zhao, H. L. Bao, *Chem. Sci.* **2017**, 8, 2081; c) P. Z. Zhang, J. A. Li, L. Zhang, A. Shoberu, J. P. Zou, W. Zhang, *Green Chem.* **2017**, 19, 919.
- [26] Y. Z. Li, B. J. Li, X. Y. Lu, S. Lin, Z. J. Shi, *Angew. Chem. Int. Ed.* **2009**, 48, 3817.
- [27] W. W. Jin, H. F. Yu, Z. K. Yu, *Tetrahedron Lett.* **2011**, 52, 5884.

FULL PAPER

Iron-Mediated Oxidative C–H Alkylation of *S,S*-
Functionalized Internal Olefins via
C(sp²)–H/C(sp³)–H Cross-Coupling

Adv. Synth. Catal. **Year**, *Volume*, Page – Page

Quannan Wang, Jiang Lou, Ping Wu, Kaikai Wu,
and Zhengkun Yu*



Accepted Manuscript