

Article

Copper-Catalyzed One-Pot Synthesis of *N*-Sulfonyl Amidines from Sulfonyl Hydrazine, Terminal Alkynes and Sulfonyl Azides

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Abstract: *N*-Sulfonyl amidines are developed from a Cu-catalyzed three-component reaction from sulfonyl hydrazines, terminal alkynes and sulfonyl azides in toluene at room temperature. Particularly, the intermediate *N*-sulfonylketenimines was generated via a CuAAC/ring-opening procedure and took a nucleophilic addition with the weak nucleophile sulfonyl hydrazines. In addition, the stability of the product was tested by a HNMR spectrometer.

Keywords: amidines; multicomponent reactions; CuAAC/ring-opening; *N*-sulfonylketenimines; nucleophilic addition



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1. Introduction

Amidine derivatives are important privileged scaffolds in medicinal chemistry [1–3], synthetic chemistry [4] and an important pharmacophore in drug discovery [5,6]. One subset of such compounds is *N*-sulfonyl amidine derivatives that show a prolific set of biological activities, including antifungal (I) [7], anticancer (II) [8], antiresorptive (III and IV) [9–11], antiproliferative (V) [12], dopamine transporter inhibitors (VI) [13] (Figure 1), etc. [14,15]. Therefore, the establishment of robust synthetic approaches for the preparation of *N*-sulfonyl amidines and their functionalizations is highly required.

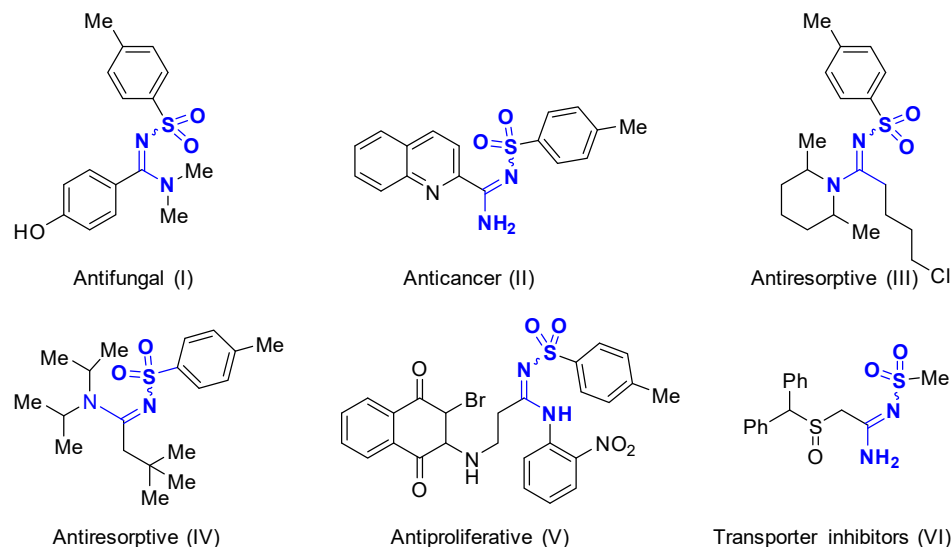
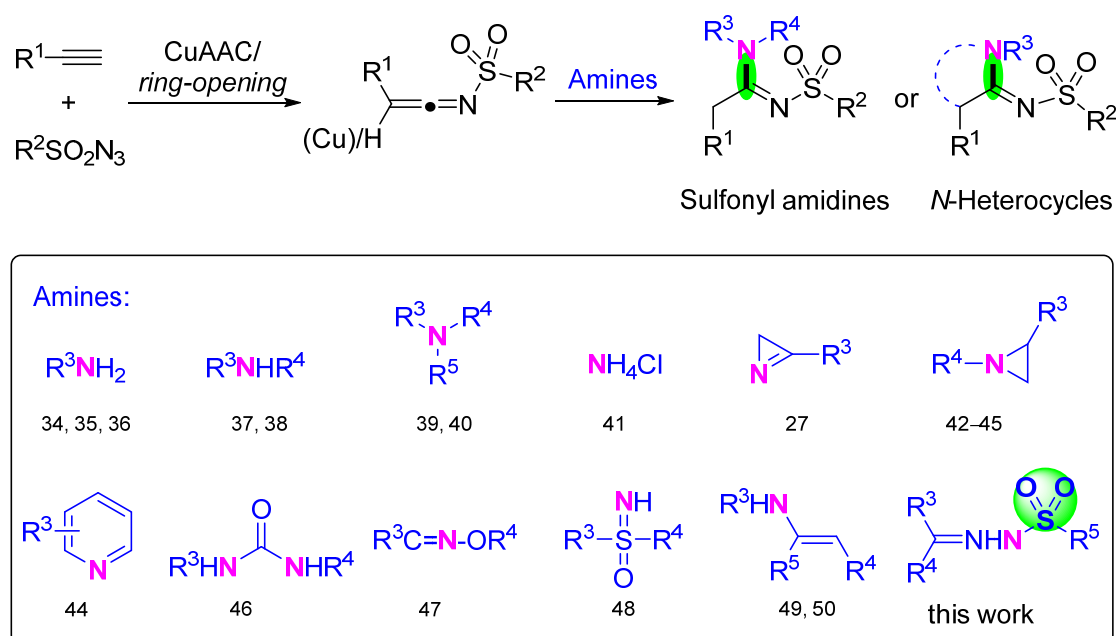


Figure 1. Part of the sulfonyl amidine drug candidates.

Classical types of reactions have focused on the preparation of *N*-sulfonyl amidines involved in the reaction of cyclic thioamides and thioacetamide derivatives with sulfonyl azides [14,16–18], the phosphite-mediated Beckmann-like coupling of oximes and *p*-toluenesulfonyl azide [19], sulfonamide derivatives condensation with DMF–DMA [20], the sulfonamide reaction with formamide [21] and the sulfonyl ynamide rearrangement [22]. The most efficient method is the Cu-catalyzed multicomponent reaction of terminal alkynes, sulfonyl azides and amines, which has been applied to synthesize numerous oxygen-containing and nitrogen-containing heterocyclic compounds [23–31]. The ketenimine intermediate generated by Cu-catalyzed alkynes and sulfonyl azides [31–33] could take a nucleophilic addition reaction with most amines, as show in Scheme 1, including aliphatic primary amines [34–36], aliphatic secondary amines [37,38], aliphatic tertiary amines [39,40], quaternary amine salts [41], imines [27], nitrogenous heterocyclic compounds [42–45], urea derivatives [46], oximes [47], sulfoximines [48] and enyl amine [49,50]. However, to our knowledge, there are few previous works that used the weak nucleophile sulfonyl hydrazines for this method. Herein, the Cu-catalyzed one-pot synthesis of *N*-sulfonyl amidines from sulfonyl hydrazine, terminal alkynes and sulfonyl azides was reported.




Scheme 1. Copper-catalyzed tandem reactions of the terminal alkynes, sulfonyl azides and amines.

2. Results

We began our investigation by examining the synthesis of 4-methyl-*N*-(2-phenyl-1-(2-(1-phenylethylidene)-1-tosylhydrazinyl) ethylidene)benzenesulfonamide **4a** via 4-methyl-*N'*-(1-phenylethylidene)benzenesulfonohydrazide **1a**, ethynylbenzene **2a** and *p*-tosyl azide **3a**. The reaction was carried out in the presence of CuI and Et₃N in CH₂Cl₂ at room temperature for 1 h, and **4a** was isolated in a 78% yield (Table 1, entry 1). Based on this finding, the reaction conditions were screened. First, several catalysts were screened, and most Cu-catalysts exhibited a high catalytic reactivity in this reaction, whether Cu^I-catalysts or Cu^{II}-catalysts (Table 1, entries 2–6). Other catalysts such as AgTFA failed to produce the desired product (Table 1, entries 7). Then, the effects of different bases were evaluated, and the screening results revealed that the use of Et₃N achieved a superior result compared to DMAP, DIPEA, pyridine and the other bases (Table 1, entries 8–12). Finally, the solvents were screened, and a lower or comparable yield was obtained when CHCl₃, DCE, MeCN, THF, DMSO and DMF were used as solvents, while toluene gave **4a** the highest yield of 84% (Table 1, entry 13–19). Encouraged by this promising result, we tracked the reaction

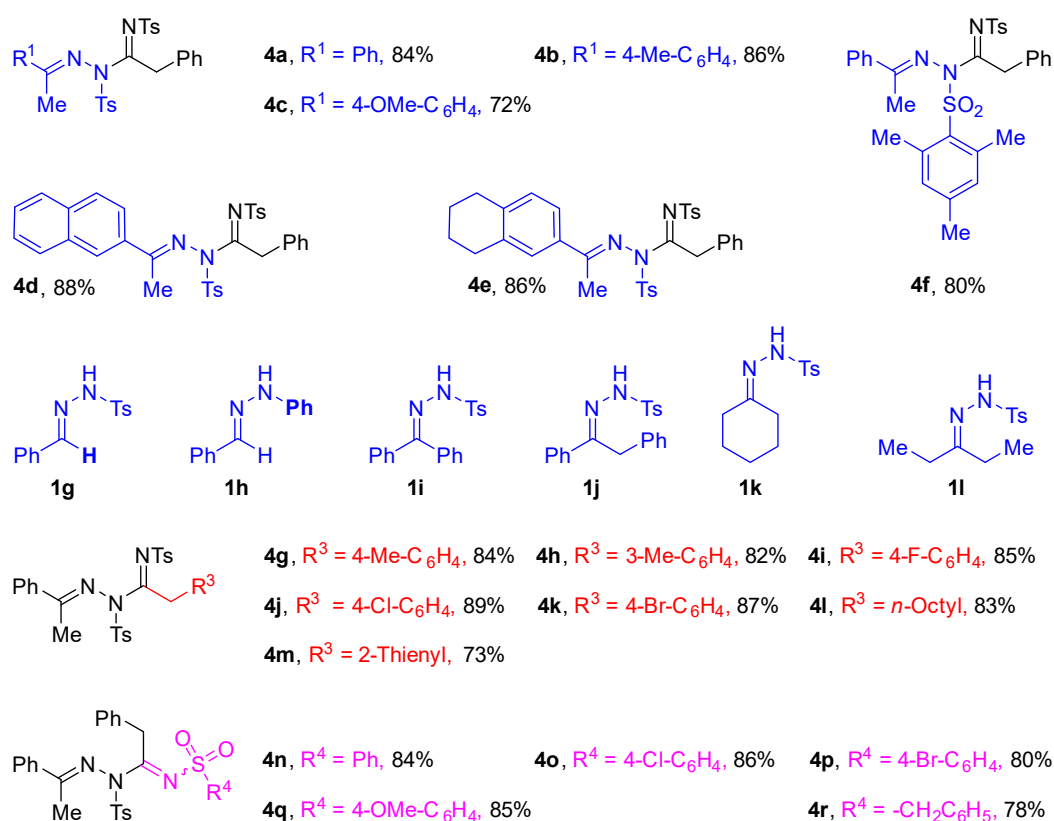
by TLC and found that it could be completed in less than an hour at room temperature (Table 1, entry 20–23).

Table 1. Optimization of the catalytic conditions ^a.

<div style="text-align: center;">  </div>						
Entry	Cat. (10 mol%)	Base (10 mol%)	Solvent (10 mol%)	Temp. (°C)	Time (h)	Yield (%) ^b
entry 1	CuI	Et ₃ N	CH ₂ Cl ₂	rt	1.0	78
entry 2	CuBr	Et ₃ N	CH ₂ Cl ₂	rt	1.0	76
entry 3	CuCl	Et ₃ N	CH ₂ Cl ₂	rt	1.0	72
entry 4	CuBr ₂	Et ₃ N	CH ₂ Cl ₂	rt	1.0	64
entry 5	Cu(OAc) ₂	Et ₃ N	CH ₂ Cl ₂	rt	1.0	52
entry 6	Cu(OTf) ₂	Et ₃ N	CH ₂ Cl ₂	rt	1.0	21
entry 7	AgTFA	Et ₃ N	CH ₂ Cl ₂	rt	1.0	0
entry 8	CuI	DMAP	CH ₂ Cl ₂	rt	1.0	26
entry 9	CuI	DIPEA	CH ₂ Cl ₂	rt	1.0	75
entry 10	CuI	Pyridine	CH ₂ Cl ₂	rt	1.0	32
entry 11	CuI	<i>t</i> -BuONa	CH ₂ Cl ₂	rt	1.0	10
entry 12	CuI	K ₂ CO ₃	CH ₂ Cl ₂	rt	1.0	8
entry 13	CuI	Et ₃ N	CHCl ₃	rt	1.0	76
entry 14	CuI	Et ₃ N	DCE	rt	1.0	75
entry 15	CuI	Et ₃ N	Toluene	rt	1.0	84
entry 16	CuI	Et ₃ N	MeCN	rt	1.0	52
entry 17	CuI	Et ₃ N	THF	rt	1.0	80
entry 18	CuI	Et ₃ N	DMSO	rt	1.0	10
entry 19	CuI	Et ₃ N	DMF	rt	1.0	6
entry 20	CuI	Et ₃ N	Toluene	40	1.0	75
entry 21	CuI	Et ₃ N	Toluene	rt	0.5	80
entry 22	CuI	Et ₃ N	Toluene	rt	2.0	84
entry 23	CuI	Et ₃ N	Toluene	rt	3.0	84

^a Reaction conditions: To **1a** (0.5 mmol), Cat. 10 mol%, base 1.2 eq. in the solvent (3 mL) was added **2a** (1.2 eq.) and **3a** (1.2 eq.), stirred at specified temperatures and times. ^b Isolated yields.

With the optimized reaction conditions obtained, the substrate diversity with the sulfonyl hydrazines **1** was tested first. As shown in Scheme 2, the R¹ electron effects of the substituents **1** had slight influences. For example, substrates bearing 4-OMe-C₆H₄, 4-Me-C₆H₄, 2-naphthyl and 2-tetra-hydronaphthalyl were examined, and the 72–88% yields of **4a–4e** were isolated. The R² of substrates **1** bearing the 2,4,6-trimethylphenyl group also can obtain **4f** in a good yield of 80%. However, when changing the substrates **1** to other sulfonyl hydrazines, such as **1g–1k**, it could not obtain the desired products and give decomposed or complex compounds. Next, the scopes and limitations of terminal alkynes **2** and sulfonyl azides **3** were examined. An aryl-substituted, aliphatic or 2-thienyl terminal alkynes and aryl-substituted or aliphatic sulfonyl azides can smoothly obtain the corresponding products **4g–4m** with yields of 73–89% and **4n–4q** with yields of 78–86%, in which both the substituents led to high yields and were influenced slightly.



Scheme 2. The synthesis of products **4a–4r**.

The structure of **4a** was confirmed by X-ray crystallography (Figure 2, CCDC deposition number 2075031).

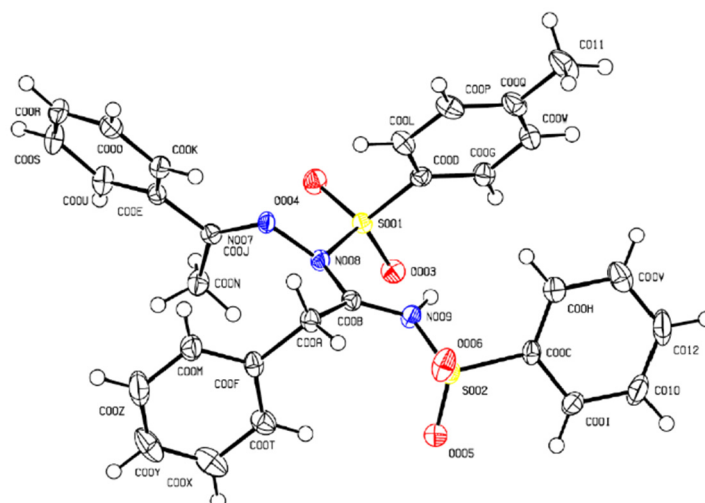


Figure 2. X-ray crystal structure of compound **4a**.

Curiously, we found that the separated products in the solvent were unstable and would decompose. Thus, the stability of product **4a** was tested by a HNMR spectrometer. As shown in Figure 3, the products dissolved in DMSO were relatively stable in the first four days, and the decomposition complex could be observed starting from the fifth day; then, the concentration of byproducts became thicker day by day. After a month, the system was relatively stable, and the decomposition was slow. Therefore, it is recommended that products **4a–4q** should be dried and stored at a low temperature.

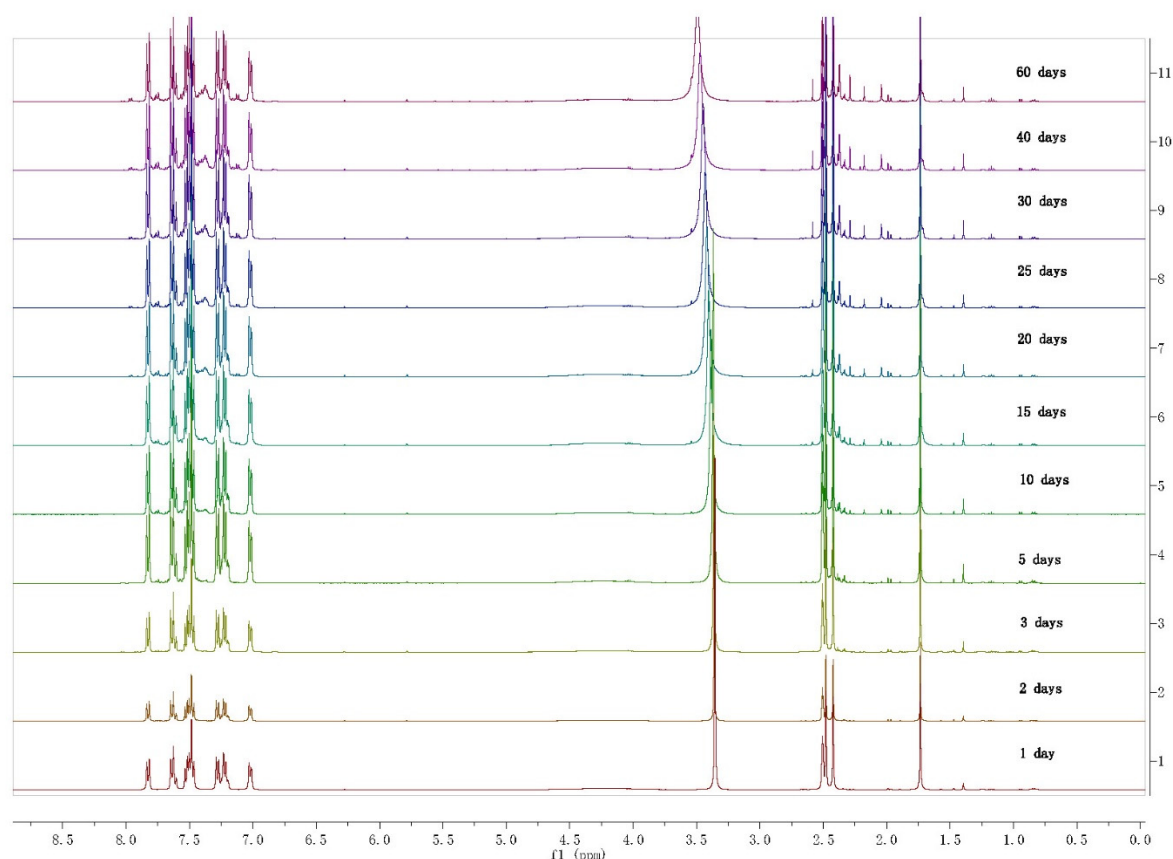


Figure 3. The stability of product **4a** tested by a ^1H NMR spectrometer.

3. Experimental

3.1. General Information

All melting points were determined on a Yanaco melting point apparatus and were uncorrected. IR spectra were recorded as KBr pellets on a Nicolet FT-IR 5DX spectrometer. All spectra of ^1H NMR (400 MHz) and ^{13}C NMR (100 MHz) were measured on a 400 MHz Bruker spectrometer using $\text{DMSO}-d_6$ or CDCl_3 as the solvent, with tetramethylsilane (TMS) as the internal standard, at room temperature. Chemical shifts are given in δ relative to TMS, and the coupling constants J are given in Hz. HRMS were obtained on a Bruker micrOTOF-Q II spectrometer. All commercially available reagents were purchased from Sigma-Aldrich, Acros, Aladdin, TCI, Alfa, Innochem in China and were used without further purification. All reactions were carried out in dried reaction tube (25 mL). The original ^1H and ^{13}C NMR spectra are available in supplementary material.

3.2. Compound Characterizations and Preparations

4-methyl-*N*-((*E*)-2-phenyl-1-(2-((*E*)-1-phenylethylidene)-1-oxylhydrazineyl) ethylidene) benzenesulfonamide (**4a**). 4-methyl-*N'*-(1-phenylethylidene) benzenesulfonylhydrazide (**1a**) (0.114 mg, 0.50 mmol) was mixed with CuI (9.5 mg, 0.05 mmol) in 1-mL toluene. Then, ethynylbenzene (**2a**) (76.5 mg, 0.75 mmol), TsN_3 (147.8 mg, 0.75 mmol) and TEA (101 mg, 1.0 mmol) were mixed in toluene (2 mL). After stirring at room temperature for 1 h and concentrated under reduced pressure, the mix was purified a flash chromatography (petroleum ether/ethyl acetate: 7:1) to give product **4a** as a white solid, mp 143–144 °C. IR (KBr) ν 3063, 1564, 1492, 1442, 1309, 1145, 1082 cm^{-1} ; ^1H NMR (400 MHz, $\text{DMSO}-d_6$) δ 7.82 (d, J = 8.0 Hz, 2H), 7.62 (t, J = 8.0 Hz, 3H), 7.53–7.46 (m, 6H), 7.28–7.21 (m, 5H), 7.01 (d, J = 6.8 Hz, 2H), 4.14 (s, 2H), 2.48 (s, 3H), 2.42 (s, 3H), 1.73 (s, 3H); ^{13}C NMR (100 MHz, $\text{DMSO}-d_6$) δ 182.7, 165.2, 145.6, 143.6, 138.6, 135.0, 134.0, 133.1, 132.4, 129.7 (2C), 129.6 (2C), 128.9 (2C),

128.8 (2C), 128.6, 128.5 (2C), 127.8 (2C), 127.2, 126.5 (3C), 21.3 (3C), 17.7; HRMS (ESI-TOF) (m/z). Calcd for $C_{30}H_{29}N_3O_4S_2$, $[M + H]^+$ 560.1672; found 560.1675.

The products **4b–4q** were prepared by a similar procedure.

4-methyl-*N*-((*E*)-2-phenyl-1-(2-((*E*)-1-(*p*-tolyl)ethylidene)-1-tosylhydrazineyl) ethylidene) benzenesulfonamide (**4b**). White solid, mp 153–155 °C. IR (KBr) ν 3062, 1594, 1568, 1307, 1172, 1147, 1084 cm^{-1} ; 1H NMR (400 MHz, DMSO- d_6) δ 7.72 (d, J = 8.0 Hz, 2H), 7.62 (d, J = 8.0 Hz, 2H), 7.47 (t, J = 7.8 Hz, 4H), 7.31 (d, J = 8.0 Hz, 2H), 7.27 (d, J = 8.0 Hz, 2H), 7.24–7.19 (m, 3H), 7.00 (d, J = 6.8 Hz, 2H), 4.19 (s, 2H), 2.47 (s, 3H), 2.42 (s, 3H), 2.39 (s, 3H), 1.69 (s, 3H); ^{13}C NMR (100 MHz, DMSO- d_6) δ 182.3, 165.3, 145.5, 143.5, 142.6, 138.6, 134.0, 133.1, 132.3, 129.7 (2C), 129.6 (2C), 129.3 (2C), 128.9 (2C), 128.6 (2C), 128.5 (2C), 127.8 (2C), 127.1, 126.5 (3C), 21.2 (3C), 17.7; HRMS (ESI-TOF) (m/z). Calcd for $C_{31}H_{31}N_3O_4S_2$, $[M + H]^+$ 574.1829; found 574.1831.

N-((*E*)-1-(2-((*E*)-1-(4-methoxyphenyl)ethylidene)-1-tosylhydrazineyl)-2-phenylethylidene)-4-methylbenzenesulfonamide (**4c**). White solid, mp 141–143 °C. IR (KBr) ν 3063, 1590, 1494, 1289, 1173, 1141, 1085 cm^{-1} ; 1H NMR (400 MHz, DMSO- d_6) δ 7.81 (d, J = 8.8 Hz, 2H), 7.62 (d, J = 8.0 Hz, 2H), 7.47 (t, J = 7.8 Hz, 4H), 7.27 (d, J = 8.0 Hz, 2H), 7.23–7.18 (m, 3H), 7.05–6.99 (m, 4H), 4.49 (s, 2H), 3.85 (s, 3H), 2.47 (s, 3H), 2.42 (s, 3H), 1.67 (s, 3H); ^{13}C NMR (100 MHz, DMSO- d_6) δ 181.5, 165.3, 162.6, 145.5, 143.5, 138.7, 134.0, 133.1, 132.3, 129.7 (2C), 129.6 (2C), 129.5 (2C), 128.9 (2C), 128.6, 128.5 (2C), 127.3, 127.1, 126.5 (3C), 114.1, 55.6, 21.2 (3C), 17.2; HRMS (ESI-TOF) (m/z). Calcd for $C_{31}H_{31}N_3O_5S_2$, $[M + H]^+$ 590.1778; found 590.1782.

4-methyl-*N*-((*E*)-1-(2-((*E*)-1-(naphthalen-2-yl)ethylidene)-1-tosylhydrazineyl)-2-phenylethylidene)benzenesulfonamide (**4d**). White solid, mp 172–173 °C. IR (KBr) ν 3056, 1590, 1574, 1494, 1359, 1305, 1144, 1084 cm^{-1} ; 1H NMR (400 MHz, DMSO- d_6) δ 8.39 (s, 1H), 8.07 (d, J = 8.0 Hz, 1H), 8.02 (t, J = 7.2 Hz, 3H), 7.68–7.61 (m, 4H), 7.53–7.46 (m, 4H), 7.29 (d, J = 8.0 Hz, 2H), 7.25–7.17 (m, 3H), 7.02 (d, J = 7.2, 2H), 4.34 (s, 2H), 2.48 (s, 3H), 2.43 (s, 3H), 1.87 (s, 3H); ^{13}C NMR (100 MHz, DMSO- d_6) δ 182.3, 165.3, 145.6, 143.6, 138.6, 134.7, 134.0, 133.1, 132.4 (2C), 129.6 (2C), 129.4 (2C), 129.3 (2C), 128.9 (2C), 128.6, 128.5 (2C), 128.3, 128.2, 127.7, 127.2, 127.0, 126.5 (3C), 123.7, 21.2 (3C), 17.6; HRMS (ESI-TOF) (m/z). Calcd for $C_{34}H_{31}N_3O_4S_2$, $[M + H]^+$ 610.1829; found 610.1832.

4-methyl-*N*-((*E*)-2-phenyl-1-(2-((*E*)-1-(5,6,7,8-tetrahydronaphthalen-2-yl)ethylidene)-1-tosylhydrazineyl)ethylidene)benzenesulfonamide (**4e**). White solid, mp 173–174 °C. IR (KBr) ν 3062, 3030, 1590, 1494, 1370, 1176, 1145, 1083 cm^{-1} ; 1H NMR (400 MHz, DMSO- d_6) δ 7.61 (d, J = 8.0 Hz, 2H), 7.50 (d, J = 8.0 Hz, 1H), 7.46 (d, J = 6.4 Hz, 5H), 7.28–7.16 (m, 6H), 6.99 (d, J = 8.0, 2H), 4.02 (s, 2H), 2.78 (s, 4H), 2.47 (s, 3H), 2.42 (s, 3H), 1.76 (s, 4H), 1.69 (s, 3H); ^{13}C NMR (100 MHz, DMSO- d_6) δ 182.6, 165.3, 145.6, 143.6, 141.8, 138.6, 137.2, 134.0, 133.1, 132.4, 129.7 (2C), 129.6 (2C), 129.3, 128.9 (2C), 128.7 (2C), 128.6 (2C), 128.4, 127.2, 126.5 (2C), 124.9, 28.9 (2C), 22.6, 22.5, 21.3, 21.2, 17.6 (2C); HRMS (ESI-TOF) (m/z). Calcd for $C_{34}H_{35}N_3O_4S_2$, $[M + H]^+$ 614.2142; found 614.2145.

N-(1-(1-(mesitylsulfonyl)-2-((*E*)-1-phenylethylidene)hydrazineyl)-2-phenylethylidene)-4-methylbenzenesulfonamide (**4f**). White solid, mp 181–183 °C. IR (KBr) ν 3062, 1600, 1551, 1354, 1304, 1141, 1088 cm^{-1} ; 1H NMR (400 MHz, DMSO- d_6) δ 7.77 (d, J = 7.6 Hz, 2H), 7.60 (d, J = 7.2 Hz, 1H), 7.50 (t, J = 7.8 Hz, 2H), 7.34–7.17 (m, 7H), 7.03 (d, J = 7.2 Hz, 2H), 6.93 (s, 2H), 4.58 (s, 2H), 2.43 (s, 6H), 2.34 (s, 3H), 2.32 (s, 3H), 1.82 (s, 3H); ^{13}C NMR (100 MHz, DMSO- d_6) δ 181.3, 164.7, 143.9, 143.2, 140.3, 138.5, 135.0, 133.1, 132.4, 132.3, 132.0, 131.9, 129.4 (2C), 128.8, 128.7 (2C), 128.5 (2C), 127.9, 127.7 (2C), 127.1 (2C), 126.3 (2C), 21.8 (2C), 21.0 (2C), 20.7, 18.5; HRMS (ESI-TOF) (m/z). Calcd for $C_{32}H_{33}N_3O_4S_2$, $[M + H]^+$ 590.1985; found 590.1988.

4-methyl-*N*-((*E*)-1-(2-((*E*)-1-phenylethylidene)-1-tosylhydrazineyl)-2-(*p*-tolyl)ethylidene) benzenesulfonamide (**4g**). White solid, mp 159–160 °C. IR (KBr) ν 3062, 2920, 1596, 1566, 1367, 1174, 1142, 1085 cm^{-1} ; 1H NMR (400 MHz, DMSO- d_6) δ 7.83 (d, J = 7.6 Hz, 2H), 7.61

(d, $J = 8.0$ Hz, 3H), 7.54–7.45 (m, 6H), 7.26 (d, $J = 8.0$ Hz, 2H), 7.02 (d, $J = 7.6$ Hz, 2H), 6.90 (d, $J = 8.0$ Hz, 2H), 4.19 (s, 2H), 2.47 (s, 3H), 2.41 (s, 3H), 2.26 (s, 3H), 1.74 (s, 3H); ^{13}C NMR (100 MHz, DMSO- d_6) δ 182.7, 165.4, 145.5, 143.5, 143.2, 138.6, 136.4, 135.1, 134.0, 132.3, 130.0, 129.6 (2C), 129.5 (2C), 129.1 (2C), 128.8 (2C), 128.5 (2C), 127.8 (2C), 126.5 (3C), 21.2 (2C), 20.7 (2C), 17.8; HRMS (ESI-TOF) (m/z). Calcd for $\text{C}_{31}\text{H}_{31}\text{N}_3\text{O}_4\text{S}_2$, $[\text{M} + \text{H}]^+$ 574.1829; found 574.1832.

4-methyl-*N*-((*E*)-1-(2-((*E*)-1-phenylethylidene)-1-tosylhydrazineyl)-2-(*m*-tolyl)ethylidene) benzenesulfonamide (**4h**). White solid, mp 146–148 °C. IR (KBr) ν 3062, 2920, 1598, 1569, 1489, 1359, 1367, 1294, 1142, 1087 cm^{-1} ; ^1H NMR (400 MHz, DMSO- d_6) δ 7.84 (d, $J = 7.6$ Hz, 2H), 7.62 (d, $J = 8.0$ Hz, 3H), 7.53–7.45 (m, 6H), 7.28 (d, $J = 8.0$ Hz, 2H), 7.11 (t, $J = 7.6$ Hz, 1H), 7.02 (d, $J = 7.6$ Hz, 1H), 6.87 (d, $J = 7.6$ Hz, 1H), 6.66 (s, 1H), 4.21 (s, 2H), 2.47 (s, 3H), 2.42 (s, 3H), 1.99 (s, 3H), 1.74 (s, 3H); ^{13}C NMR (100 MHz, DMSO- d_6) δ 182.6, 165.2, 145.6, 138.6, 137.7, 134.9, 134.0, 133.0, 132.4, 130.5 (2C), 129.6 (2C), 129.5 (2C), 128.7 (2C), 128.5 (2C), 127.8 (2C), 127.6, 126.5 (3C), 125.7, 21.2 (3C), 20.7, 17.6; HRMS (ESI-TOF) (m/z). Calcd for $\text{C}_{31}\text{H}_{31}\text{N}_3\text{O}_4\text{S}_2$, $[\text{M} + \text{H}]^+$ 574.1829; found 574.1830.

N-((*E*)-2-(4-fluorophenyl)-1-(2-((*E*)-1-phenylethylidene)-1-tosylhydrazineyl) ethylidene)-4-methylbenzenesulfonamide (**4i**). White solid, mp 157–159 °C. IR (KBr) ν 3062, 1595, 1564, 1375, 1308, 1190, 1083 cm^{-1} ; ^1H NMR (400 MHz, DMSO- d_6) δ 7.85 (d, $J = 8.0$ Hz, 2H), 7.62 (d, $J = 8.0$ Hz, 3H), 7.54–7.45 (m, 6H), 7.27 (d, $J = 8.0$ Hz, 2H), 7.09–7.05 (m, 4H), 4.20 (s, 2H), 2.47 (s, 3H), 2.42 (s, 3H), 1.86 (s, 3H); ^{13}C NMR (100 MHz, DMSO- d_6) δ 182.5, 165.0, 161.2 (d, $J = 256.7$ Hz), 145.7, 143.6, 138.5, 135.0, 133.9, 132.4, 130.7 (2C), 129.7 (2C), 129.6 (2C), 129.2 (d, $J = 3.1$ Hz), 128.8 (2C), 128.5 (2C), 127.8 (2C), 126.5 (3C), 115.5 (d, $J = 21.8$ Hz), 21.1 (2C), 21.1 (d, $J = 7.7$ Hz), 17.9; HRMS (ESI-TOF) (m/z). Calcd for $\text{C}_{30}\text{H}_{28}\text{FN}_3\text{O}_4\text{S}_2$, $[\text{M} + \text{H}]^+$ 578.1578; found 578.1581.

N-((*E*)-2-(4-chlorophenyl)-1-(2-((*E*)-1-phenylethylidene)-1-tosylhydrazineyl) ethylidene)-4-methylbenzenesulfonamide (**4j**). White solid, mp 153–155 °C. IR (KBr) ν 3064, 1593, 1562, 1444, 1345, 1272, 1122, 1081 cm^{-1} ; ^1H NMR (400 MHz, DMSO- d_6) δ 7.84 (d, $J = 8.4$ Hz, 2H), 7.62 (d, $J = 8.0$ Hz, 3H), 7.53–7.45 (m, 6H), 7.30–7.27 (m, 4H), 7.02 (d, $J = 8.8$ Hz, 2H), 4.24 (s, 2H), 2.47 (s, 3H), 2.42 (s, 3H), 1.90 (s, 3H); ^{13}C NMR (100 MHz, DMSO- d_6) δ 182.5, 164.8, 145.7, 143.7, 138.4, 135.0, 133.8, 132.4, 132.1, 132.0, 130.6 (2C), 129.7 (2C), 129.6 (2C), 128.8 (2C), 128.5 (2C), 127.8 (2C), 126.5 (3C), 38.0, 21.1 (2C), 21.1, 18.0; HRMS (ESI-TOF) (m/z). Calcd for $\text{C}_{30}\text{H}_{28}\text{ClN}_3\text{O}_4\text{S}_2$, $[\text{M} + \text{H}]^+$ 594.1283; found 594.1285.

N-((*E*)-2-(4-bromophenyl)-1-(2-((*E*)-1-phenylethylidene)-1-tosylhydrazineyl) ethylidene)-4-methylbenzenesulfonamide (**4k**). White solid, mp 158–160 °C. IR (KBr) ν 3062, 1592, 1560, 1486 1369, 1282, 1142, 1082 cm^{-1} ; ^1H NMR (400 MHz, DMSO- d_6) δ 7.83 (d, $J = 7.2$ Hz, 2H), 7.62 (d, $J = 8.0$ Hz, 3H), 7.53–7.41 (m, 8H), 7.28 (d, $J = 8.0$ Hz, 2H), 6.95 (d, $J = 8.0$ Hz, 2H), 4.21 (s, 2H), 2.47 (s, 3H), 2.42 (s, 3H), 1.91 (s, 3H); ^{13}C NMR (100 MHz, DMSO- d_6) δ 182.4, 164.7, 145.7, 143.6, 138.4, 135.0, 133.8, 132.5, 132.4, 131.5, 132.0, 130.8 (2C), 129.7 (2C), 129.6 (2C), 128.8 (2C), 128.5 (2C), 127.8 (2C), 126.5 (3C), 120.3, 21.1 (2C), 18.0; HRMS (ESI-TOF) (m/z). Calcd for $\text{C}_{30}\text{H}_{28}\text{BrN}_3\text{O}_4\text{S}_2$, $[\text{M} + \text{H}]^+$ 638.0778; found 638.0779.

4-methyl-*N*-((*E*)-1-(2-((*E*)-1-phenylethylidene)-1-tosylhydrazineyl)octylidene) benzenesulfonamide (**4l**). White solid, mp 103–105 °C. IR (KBr) ν 3063, 2864, 1595, 1338, 1264, 1155, 1076 cm^{-1} ; ^1H NMR (400 MHz, DMSO- d_6) δ 8.00 (d, $J = 7.2$ Hz, 2H), 7.63 (t, $J = 7.6$ Hz, 1H), 7.57–7.53 (m, 6H), 7.40 (d, $J = 8.4$ Hz, 2H), 7.29 (d, $J = 8.0$ Hz, 2H), 2.75 (d, $J = 7.6$ Hz, 2H), 2.56 (s, 3H), 2.44 (s, 3H), 2.40 (s, 3H), 1.39 (s, 2H), 1.17–1.08 (m, 8H), 0.75 (t, $J = 6.8$ Hz, 3H); ^{13}C NMR (100 MHz, DMSO- d_6) δ 181.5, 167.9, 145.6, 143.3, 138.9, 135.4, 134.1, 132.4, 129.7 (2C), 129.6 (2C), 128.9 (2C), 128.4 (2C), 127.8 (2C), 126.3 (2C), 32.5, 30.9, 28.8, 27.8, 24.9, 21.9, 21.3, 21.1, 18.7, 13.9; HRMS (ESI-TOF) (m/z). Calcd for $\text{C}_{30}\text{H}_{37}\text{N}_3\text{O}_4\text{S}_2$, $[\text{M} + \text{H}]^+$ 568.2298; found 568.2231.

4-methyl-*N*-((*E*)-1-(2-((*E*)-1-phenylethylidene)-1-tosylhydrazineyl)-2-(thiophen-2-yl)ethylidene)benzenesulfonamide (**4m**). Yellow solid, mp 67–69 °C. IR (KBr) ν 3062, 2927, 2866, 1590, 1369, 1307, 1153, 1087 cm^{-1} ; ^1H NMR (400 MHz, DMSO- d_6) δ 7.85 (t, $J = 6.8$ Hz,

4H), 7.65 (d, $J = 9.2$ Hz, 3H), 7.48 (d, $J = 7.8$ Hz, 2H), 7.36 (d, $J = 7.8$ Hz, 2H), 7.11 (d, $J = 7.8$ Hz, 3H), 6.86–6.82 (m, 2H), 4.58 (s, 2H), 2.50 (s, 3H), 2.41 (s, 3H), 2.00 (s, 3H); ^{13}C NMR (100 MHz, DMSO- d_6) δ 183.3, 163.9, 145.4, 143.4, 139.2, 135.8, 134.4, 134.2, 132.2, 129.4 (2C), 129.3 (2C), 129.2 (2C), 128.8 (2C), 128.1, 127.9 (2C), 127.1 (2C), 127.0, 125.4, 33.8, 21.9, 21.8, 18.2; HRMS (ESI-TOF) (m/z). Calcd for $\text{C}_{28}\text{H}_{27}\text{N}_3\text{O}_4\text{S}_3$, $[\text{M} + \text{H}]^+$ 565.1237; found 565.1239.

N-(2-phenyl-1-(2-((*E*)-1-phenylethylidene)-1-tosylhydrazineyl)ethylidene) benzenesulfonamide (**4n**). White solid, mp 149–151 °C. IR (KBr) ν 3062, 1589, 1561, 1494, 1365, 1282, 1140, 1085 cm^{-1} ; ^1H NMR (400 MHz, DMSO- d_6) δ 7.82 (d, $J = 8.0$ Hz, 2H), 7.76 (d, $J = 6.8$ Hz, 3H), 7.70–7.60 (m, 3H), 7.53–7.46 (m, 4H), 7.27–7.20 (m, 5H), 7.02 (d, $J = 6.8$ Hz, 2H), 4.23 (s, 2H), 2.41 (s, 3H), 1.74 (s, 3H); ^{13}C NMR (100 MHz, DMSO- d_6) δ 182.7, 165.7, 145.6, 141.3, 135.0, 133.9, 133.1, 133.0, 132.4, 129.6, 129.3 (2C), 128.9 (2C), 128.8 (2C), 128.6 (2C), 128.5 (2C), 127.8 (2C), 127.2, 126.4 (3C), 21.2 (2C), 17.7; HRMS (ESI-TOF) (m/z). Calcd for $\text{C}_{29}\text{H}_{27}\text{N}_3\text{O}_4\text{S}_2$, $[\text{M} + \text{H}]^+$ 546.1516; found 546.1519.

4-chloro-*N*-(2-phenyl-1-(2-((*E*)-1-phenylethylidene)-1-tosylhydrazineyl)ethylidene) benzenesulfonamide (**4o**). White solid, mp 141–143 °C. IR (KBr) ν 3067, 1592, 1554, 1493, 1341, 1308, 1146, 1081 cm^{-1} ; ^1H NMR (400 MHz, DMSO- d_6) δ 7.83 (d, $J = 8.0$ Hz, 2H), 7.75 (t, $J = 9.6$ Hz, 4H), 7.62 (t, $J = 7.6$ Hz, 1H), 7.51 (t, $J = 8.0$ Hz, 4H), 7.29–7.20 (m, 5H), 7.02 (t, $J = 6.8$ Hz, 2H), 4.15 (s, 2H), 2.42 (s, 3H), 1.77 (s, 3H); ^{13}C NMR (100 MHz, DMSO- d_6) δ 182.8, 165.5, 145.7, 140.2, 138.0, 135.0, 134.0, 133.0, 132.4, 129.6 (2C), 129.4 (3C), 128.8, 128.7 (2C), 128.6 (2C), 128.4 (2C), 128.3 (2C), 127.8 (2C), 127.2, 21.2 (2C), 17.8; HRMS (ESI-TOF) (m/z). Calcd for $\text{C}_{29}\text{H}_{26}\text{ClN}_3\text{O}_4\text{S}_2$, $[\text{M} + \text{H}]^+$ 580.1126; found 580.1128.

4-bromo-*N*-(2-phenyl-1-(2-((*E*)-1-phenylethylidene)-1-tosylhydrazineyl)ethylidene) benzenesulfonamide (**4p**). White solid, mp 139–140 °C. IR (KBr) ν 3066, 1594, 1554, 1493, 1374, 1309, 1145, 1083 cm^{-1} ; ^1H NMR (400 MHz, DMSO- d_6) δ 7.89 (d, $J = 8.4$ Hz, 2H), 7.83 (t, $J = 7.6$ Hz, 2H), 7.68 (t, $J = 7.6$ Hz, 2H), 7.62 (t, $J = 7.2$ Hz, 1H), 7.53–7.49 (m, 4H), 7.29–7.20 (m, 5H), 7.01 (t, $J = 7.2$ Hz, 2H), 4.23 (s, 2H), 2.42 (s, 3H), 1.76 (s, 3H); ^{13}C NMR (100 MHz, DMSO- d_6) δ 182.8, 165.5, 145.7, 140.6, 135.0, 134.0, 133.0, 132.4 (3C), 129.6 (2C), 128.8 (4C), 128.7 (2C), 128.4 (3C), 128.3 (2C), 127.2 (2C), 127.0, 21.2 (2C), 17.8; HRMS (ESI-TOF) (m/z). Calcd for $\text{C}_{29}\text{H}_{26}\text{BrN}_3\text{O}_4\text{S}_2$, $[\text{M} + \text{H}]^+$ 624.0621; found 624.0622.

4-methoxy-*N*-(2-phenyl-1-(2-((*E*)-1-phenylethylidene)-1-tosylhydrazineyl)ethylidene) benzenesulfonamide (**4q**). White solid, mp 143–145 °C. IR (KBr) ν 3010, 1592, 1561, 1492, 1367, 1296, 1144, 1082 cm^{-1} ; ^1H NMR (400 MHz, DMSO- d_6) δ 7.82 (d, $J = 7.6$ Hz, 2H), 7.69 (t, $J = 8.4$ Hz, 2H), 7.62 (t, $J = 7.2$ Hz, 1H), 7.51 (t, $J = 8.0$ Hz, 4H), 7.29 (d, $J = 8.0$ Hz, 2H), 7.24–7.17 (m, 5H), 7.01 (d, $J = 6.8$ Hz, 2H), 4.24 (s, 2H), 3.92 (s, 3H), 2.42 (s, 3H), 1.73 (s, 3H); ^{13}C NMR (100 MHz, DMSO- d_6) δ 183.0, 165.4, 163.1, 146.0, 135.5, 134.4, 133.6, 132.7, 130.0, 129.3 (2C), 129.2 (3C), 129.1 (4C), 129.0 (2C), 128.9 (2C), 128.2 (2C), 127.5, 114.8, 56.3, 21.7 (2C), 18.1; HRMS (ESI-TOF) (m/z). Calcd for $\text{C}_{30}\text{H}_{29}\text{N}_3\text{O}_5\text{S}_2$, $[\text{M} + \text{H}]^+$ 576.1622; found 576.1621.

1-phenyl-*N*-(2-phenyl-1-(2-((*E*)-1-phenylethylidene)-1-tosylhydrazineyl)ethylidene) methanesulfonamide (**4r**). White solid, mp 125–127 °C. IR (KBr) ν 3063, 2972, 1590, 1576, 1493, 1365, 1293, 1173, 1086 cm^{-1} ; ^1H NMR (400 MHz, DMSO- d_6) δ 7.85–7.79 (m, 4H), 7.62 (t, $J = 7.2$ Hz, 1H), 7.56–7.50 (m, 4H), 7.21 (t, $J = 6.8$ Hz, 3H), 7.01 (d, $J = 7.2$ Hz, 2H), 4.18 (s, 2H), 3.04 (t, $J = 7.6$ Hz, 2H), 2.46 (s, 3H), 1.75 (s, 3H), 1.69 (s, 2H), 1.02 (t, $J = 7.6$ Hz, 3H); ^{13}C NMR (100 MHz, DMSO- d_6) δ 182.5, 165.5, 145.7, 135.1, 134.6, 133.1, 132.3, 129.9 (3C), 128.9 (2C), 128.8 (3C), 128.6 (2C), 128.3 (3C), 127.8 (3C), 127.1, 56.0 (2C), 21.2, 17.6, 16.8, 12.6; HRMS (ESI-TOF) (m/z). Calcd for $\text{C}_{30}\text{H}_{29}\text{N}_3\text{O}_4\text{S}_2$, $[\text{M} + \text{H}]^+$ 560.1672; found 560.1676.

4. Conclusions

We developed an effective copper-catalyzed three-component one-pot synthesis of *N*-sulfonyl amidines from terminal alkynes, sulfonyl azides and weak nucleophilic sulfonyl hydrazine. The synthetic pathway extended the applications of the CuAAC/ring-opening

reaction, and we expect that this methodology and *N*-sulfonyl amidines products could be applied to organic synthesis.

Supplementary Materials: The following are available online, The original ^1H and ^{13}C NMR spectra are available in supplementary material.

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