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Cite this: *RSC Adv.*, 2017, 7, 12022

Received 5th January 2017  
Accepted 9th February 2017

DOI: 10.1039/c7ra00193b

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# Further insight into the photochemical behavior of 3-aryl-*N*-(arylsulfonyl)propiolamides: tunable synthetic route to phenanthrenes†

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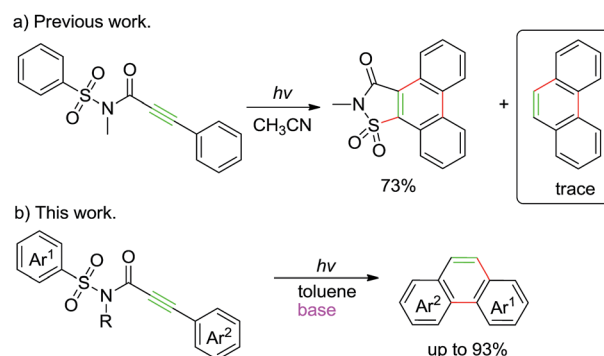
Reported herein is further insight into the photochemical behaviour of 3-aryl-*N*-(arylsulfonyl)propiolamides, which provides a straightforward way to access meaningful phenanthrenes. Mechanistic investigation indicated that aryl migration, C–C coupling, 1,3-hydrogen shift, desulfonylation and elimination were involved in the process. Moreover, this protocol allowed for scale-up using a flow reactor.

Phenanthrenes represent an important class of the simplest polycyclic aromatic hydrocarbons (PAHs) and they have received much interest in materials science and technology due to their utility in functional materials, especially in photoelectronic devices.<sup>1</sup> Moreover, compounds containing this structural core are often of great value in pharmaceutical chemistry because they generally display a broad spectrum of biological activities, such as anti-microbial, anti-tumor, anti-inflammatory, anti-malarial and others.<sup>2</sup> Additionally, phenanthrene motifs can be found in a large class of natural products.<sup>3</sup> Therefore, continuous effort has been devoted to the preparation of phenanthrene derivatives, and in the past decades, numerous relevant studies have been reported.<sup>4</sup> Among all the available strategies, the Lewis acid/base/metal mediated intramolecular annulation of alkynylated biaryls<sup>5</sup> and various metal induced intermolecular [4 + 2] cyclization of biaryls with alkynes<sup>6</sup>/bis(pinacolatoboryl) alkenes<sup>7</sup> represented the typical approaches to construct the phenanthrene skeleton. In addition, the preparation of stilbene followed by photocyclization, radical or metal catalyzed oxidative C–C coupling reactions has been extensively studied as well.<sup>8</sup> Besides, the construction of a phenanthrene ring could also be achieved *via* intramolecular ring-closing metathesis.<sup>9</sup> Despite the above, most of these methods suffered from certain limitations, such as the use of complex and expensive metal catalysts which may be unfriendly to the environment, not-easily accessible precursors, air- or moisture-sensitive reagents, and harsh reaction conditions. Thus, development of an efficient and convenient method for the construction of phenanthrene scaffold from readily prepared substrates is of important significance.

During our recent exploration of the ultraviolet light induced photochemical behavior of *N*-methyl-3-phenyl-*N*-(phenylsulfonyl)propiolamide in CH<sub>3</sub>CN, we disclosed that

along with the main product 2-methylphenanthro[9,10-*d*]isothiazol-3(2*H*)-one 1,1-dioxide through the Smiles rearrangement/Mallory reaction, a trace amount of phenanthrene was also observed after removal of the sulfonylamide group (Scheme 1a).<sup>10</sup> Such a result prompted us to explore suitable reaction conditions for the preferential preparation of phenanthrenes, and described in this paper are our continuous work on the photochemical behavior of 3-aryl-*N*-(arylsulfonyl)propiolamides to provide a straightforward way to a series of phenanthrenes tuned by base (Scheme 1b).

At the outset of further investigation, compound **1a** was chosen as the model substrate and additives with different property such as acid/base/salt were employed to modify the reaction, respectively. The results turned out to be that, unlike the less effect of acid and salt, the addition of 1 equivalent of dimethylamine dramatically improved the yield of **A** in comparison to the reaction with no additives (Table 1, entries 1–4). Moreover, the increasing amount of dimethylamine to 3 equivalents totally yielded the product **A** with the retard of formation of **1b** (Table 1, entry 5). Based on this observation, screening on the bases, such as triethylamine, methyl amine,

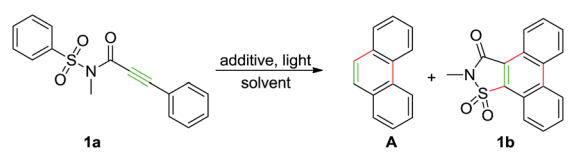


Scheme 1 Previous work and this work.

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† Electronic supplementary information (ESI) available: Experimental details, product characterization and NMR spectral data. See DOI: 10.1039/c7ra00193b



Table 1 Screening on the reaction conditions<sup>a</sup>


Entry	Solvent	Additive (equiv.)	Light	t (h)	Yield <sup>b</sup> (%)	
					A	1b
1	Toluene	No additive	300 nm	1.5	<5	71
2	Toluene	Acetic acid (1.0)	300 nm	1.5	<5	68
3	Toluene	Dimethylamine (1.0)	300 nm	1.0	40	45
4	Toluene	NaCl (1.0)	300 nm	1.3	<5	65
5	Toluene	Dimethylamine (3.0)	300 nm	1.0	70	0
6	Toluene	Et <sub>3</sub> N (3.0)	300 nm	1.0	68	0
7	Toluene	Methylamine (3.0)	300 nm	1.2	63	0
8	Toluene	Cyclohexamine (3.0)	300 nm	0.9	65	0
9	Toluene	Morpholine (3.0)	300 nm	0.7	72	0
10	Toluene	Morpholine (1.0)	300 nm	0.9	80	0
11	Toluene	Morpholine (0.5)	300 nm	1.2	55	20
12 <sup>c</sup>	Toluene	Morpholine (1.0)	300 nm	0.9	80	0
13	MeCN	Morpholine (1.0)	300 nm	1.2	15	60
14	MeOH	Morpholine (1.0)	300 nm	1.2	23	55
15	DMF	Morpholine (1.0)	300 nm	1.0	20	53
16	THF	Morpholine (1.0)	300 nm	1.3	72	0
17	Benzene	Morpholine (1.0)	300 nm	1.0	70	0
18	DCM	Morpholine (1.0)	300 nm	1.5	65	Trace
19	Toluene	Morpholine (1.0)	350 nm	12.0	58	Trace
20	Toluene	Morpholine (1.0)	HPML	4.0	60	Trace
21	Toluene	Morpholine (1.0)	MPML	6.0	62	Trace
22	Toluene	Morpholine (1.0)	300 nm	1.8	78	0

<sup>a</sup> Reaction conditions: **1a** (0.2 mmol), solvent (anhydrous 40 ml), under air atmosphere, irradiation at room temperature. <sup>b</sup> Isolated yield.

<sup>c</sup> Under N<sub>2</sub> atmosphere.

cyclohexamine and morpholine, was conducted which led to comparable yields (Table 1, entries 6–9). Remarkably, the decrease in the amount of morpholine to 1 equivalent gave the best yield of 80% (Table 1, entry 10), whereas further decrease to 0.5 equivalent led to lower yield and poor selectivity (Table 1, entry 11). Gladly, we found that reactions run equally efficiently under air/inert gas conditions (Table 1, entries 10 and 12). Then a set of solvents was screened and the reaction performed in toluene provided the best result than in other medium (Table 1, entries 13–18). It is noteworthy that the solvent plays a critical role in the result in which most of the tested substrates have been decomposed in strong polar solvents, leading to a low yield of the desired product. The screening of light source indicated that the suitable wavelength was proved to be 300 nm (Table 1, entries 10, 19–21). Finally, we found there was a little loss on yield (in comparison with entry 10) when the irradiation time was increased to 1.8 hours (Table 1, entries 22).

Subsequently, a series of substrates were subjected to the optimal conditions to investigate the reaction scope, and the results were summarized in Tables 2 and 3. As expected, substrate with the same aryl groups tethered to the sulfonyl or the alkynyl resulted in the same adduct regardless of divergent

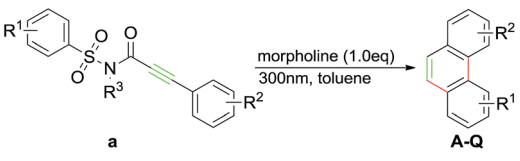
R<sup>3</sup> substituent on nitrogen atom (Table 2, **A**, **B**, **C** and Table 3, **R**), it should be noted that the non-substituted phenanthrene **A**, 3-methylphenanthrene **B**, 3-methoxyphenanthrene **C** and the heterocyclic aromatic compound **R** could be readily prepared in moderate to excellent yields. Substrates with electrophilic groups such as the halogen atom, trifluoromethyl and cyano on the *para*-position of aryl ring were perfectly tolerant with the reaction conditions, generating the corresponding products in 30–72% yields (Table 2, **D–G**, **J**). Sterically bulkier group such as butyl- or phenyl-substituted substrate **15a** and **16a** could furnish the corresponding products as well in the yield of 68% and 56%, respectively (Table 2, **H** and **I**). Substrate **18a** with *p*-methyl on both benzene rings afforded the single product **K** in 67% yield (Table 2, **K**). The reaction of compound **19a** bearing *meta*-methyl on aryl ring delivered a mixture of regioisomers in a ratio of 4 : 3 (Table 2, **L** and **L'**), while the *meta*-fluoro substituted substrate **20a** gave two regioselective products **M** and **M'** in a ratio of 8 : 1.<sup>4c</sup> As for the photoreaction of *ortho*-methyl substituted substrates **21a**, this protocol allowed access to both the desired 1-methylphenanthrene **N** and the unexpected 1-methyl-2-(phenylethynyl)benzene **N'** in a ratio of 3 : 5. In view of our previous work, the arising of **N'** was supposed possibly from the Smiles rearrangement of **21a** followed by the elimination of sulfonylamide group. Replacing the methyl group with F, Cl atom and trifluoromethyl group equally afforded corresponding structural isomers in different ratios (Table 2, **N–Q**). More importantly, other polycyclic compounds, such as benzo[*c*]phenanthrene and chrysene, could also be prepared *via* this synthetic protocol (Table 3, **T** and **U**). In addition, the reaction was also applicable for the synthesis of heteroaromatic compounds and poly-substituted phenanthrene (Table 3, **R**, **S** and **V**). Particularly, the yield of compound **V** reached to 50% even the amount of **30a** increased to 0.31 gram.


Notably, among all the prepared products, compounds **A**, **B**, **L'**, **N**, **K**, **U** and **V** were identified as natural products and have been reported in previous literatures,<sup>11–17</sup> which indicated the inherent synthetic utility of this novel protocol.

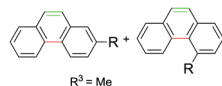
Before the reaction mechanism was proposed, some control experiments were conducted using **7a** as representative compound. First, **7a** was irradiated under the standard conditions for 0.2 h, fortunately, except for the product **B**, both the rearrangement product **7c** and a new compound **7d** which was quite labile and could be rapidly converted into **B** was isolated (Scheme 2, eqn (1)). Further exploration disclosed that **7d** was derived from compound **7c**, thus suggesting that product **B** was probably resulted from a tandem three-step photo transformation (Scheme 2, eqn (2)), and all these transformations only took place under the irradiation of UV light.

Based upon the above results, a tentative mechanism for this photoreaction was proposed as shown in Scheme 3. The alkynyl group of substrate **a** was first excited to 1,2-biradicals which initiated radical Smiles rearrangement/C–S bonding cascade reaction to form the isolatable intermediate **c**,<sup>10</sup> then subsequently oxidative cyclization delivered product **b** (Scheme 3, path a).<sup>10</sup> And cyclization reaction occurred to form the intermediate **II** which underwent 1,3-H shift process with the morpholine served as an assistant agent<sup>18</sup> to form compound **d** that

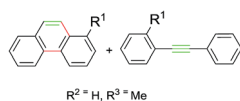


Table 2 The photoinduced synthesis of phenanthrenes<sup>a</sup>


Subs	R <sup>1</sup>	R <sup>2</sup>	R <sup>3</sup>	t (h)	Product	Yield <sup>b</sup> (%)
1a	H	H	Me	0.9	A	80
2a	H	H	Bn	1.2	A	75
3a	H	H	Propargyl	1.0	A	70
4a	H	H	Allyl	1.1	A	75
5a	H	H		1.0	A	87
6a	Me	H	Me	1.0	B	85
7a	H	Me	Me	1.2	B	90
8a	Me	H	Allyl	1.2	B	80
9a	OMe	H	Me	2.5	C	55
10a	H	OMe	Me	2.5	C	65
11a	F	H	Me	1.2	D	55
12a	Cl	H	Me	1.2	E	72
13a	Br	H	Me	2.5	F	30
14a	CF <sub>3</sub>	H	Me	1.5	G	70
15a	Ph	H	Me	2.5	H	56
16a	<i>t</i> -Butyl	H	Me	1.2	I	84
17a	CN	H	Me	2.0	J	55
18a	Me	Me	Me	1.2	K	67



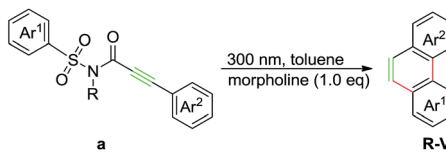
Subs	R <sup>1</sup>	R <sup>2</sup>	R	t (h)	Product	Yield <sup>b</sup> (%)
19a	H	Me	Me	2.0	L/L' (4/3) <sup>c</sup>	77 <sup>d</sup>
20a	F	H	F	1.0	M/M' (8/1) <sup>c</sup>	74 <sup>d</sup>



Subs	R <sup>1</sup>	t (h)	Product <sup>c</sup>	Yield <sup>b</sup> (%)
21a	Me	2.3	N/N' (3/5) <sup>c</sup>	45 <sup>d</sup>
22a	F	1.0	O/O' (3/1) <sup>c</sup>	93 <sup>d</sup>
23a	Cl	1.8	P/P' (6/5) <sup>c</sup>	70 <sup>d</sup>
24a	CF <sub>3</sub>	2.0	Q/Q' (1/7) <sup>c</sup>	50 <sup>d</sup>

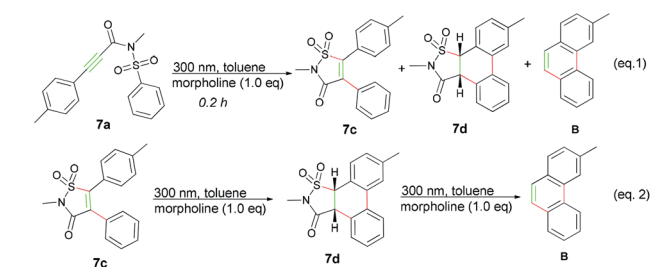
<sup>a</sup> All reactions are carried out under the standard conditions. <sup>b</sup> Isolated yield. <sup>c</sup> The ratio was determined by GC. <sup>d</sup> The total yield of both isomers.

was identified by the isolation of **7d** as shown in Scheme 2. Then a ring opening reaction of **d** occurred and afforded the 1,4-biradical intermediate **III** after exclusion of sulfur dioxide. The final phenanthrene product was formed from intermediate **III** after elimination of isocyanic intermediate **e**, which could be

Table 3 The synthesis of PAHs and polysubstituted phenanthrenes<sup>a</sup>


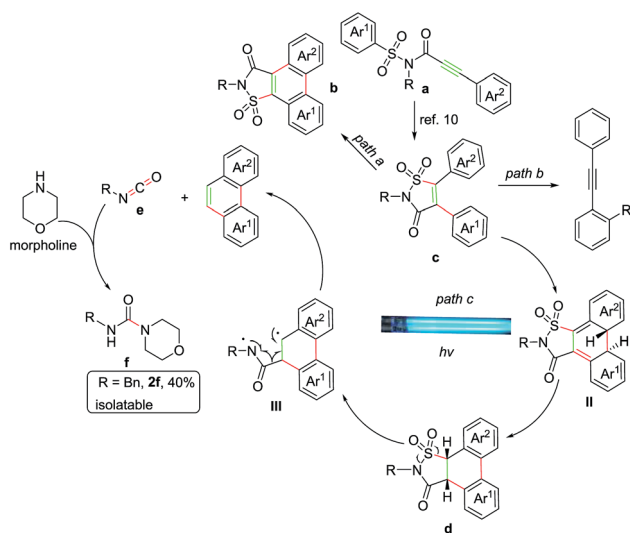
Subs	Ar <sup>1</sup>	Ar <sup>2</sup>	R <sup>3</sup>	t (h)	Yield <sup>b</sup> (%)
25a	2-thienyl	Ph	Me	2.5	35%
26a	Ph	2-thienyl	Me	3.0	50%
27a	2-thienyl	2-thienyl	Me	2.5	57%
28a	2-naphthyl	Ph	Me	3.0	37%
29a	1-naphthyl	Ph	Me	2.5	25%
30a	Ph	Ph	Me	3.5	50%

<sup>a</sup> All reactions are carried out under the standard conditions. <sup>b</sup> Isolated yield. <sup>c</sup> Reaction is carried out on the amount of 0.31 gram.



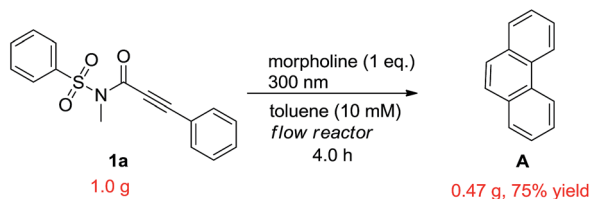
Scheme 2 Control experiments.

trapped by morpholine to produce morpholine-4-carboxamide **f** (Scheme 3, path c). Furthermore, to add more credence to the existence of compound **f**, *N*-benzylmorpholine-4-carboxamide



Scheme 3 The proposed reaction mechanism.





Scheme 4 Gram scale study with a flow reactor.

(R<sup>3</sup> = Bn) was isolated in 40% yield during the reaction of substrate **2a**.<sup>19</sup> It was reasonable to assume that some of the intermediate **c**, which was arisen from the substrate containing *ortho*-substituent on aryl ring, could not effectively undergo the C–C coupling due to the steric hindrance, but be converted into the diphenylethyne product instead after removing the sulfonylamide group with the aid of morpholine in the manner as described above (Scheme 3, path b).

In order to demonstrate the synthetic value and utility of this protocol, we turned our attention toward the preparation of phenanthrenes on a large scale.<sup>20</sup> With this aim in mind, we carried out the photoreaction in a continuous flow reactor with compound **1a**, of which the loading was increased to 1.0 gram and concentration to 10 mM, as the subject substrate. Then the reaction was run for 4 hours to afford the target product **A** in 75% with little loss on yield (Scheme 4).

In conclusion, the photoreaction of easily accessible 3-aryl-*N*-(arylsulfonyl)propiolamides to prepare phenanthrene derivatives *via* radical Smiles rearrangement/elimination has been developed. The addition of morpholine enables the 1,3-H shift which is crucial to the regioselective formation of phenanthrenes. Moreover, reasonable mechanism was elaborated in detail based on the isolated intermediates. It's worthy to point out that, using our established protocol, the desired substituents on phenanthrene rings could be simply introduced in advance on the aromatic groups of the starting material. Besides, the availability and efficiency of this approach makes it appealing for the syntheses of certain natural products.

## Acknowledgements

We are grateful for the financial support from China NSFC (No. 21372055, 21472030 and 21672047), SKLUWRE (No. 2015DX01), the Fundamental Research Funds for the Central Universities (Grant No. HIT.BRETIV.201310) and HLJNSF (B201406).

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