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# Transition-metal-free radical cleavage of a hydrazonyl N-S bond: tosyl radical-initiated cascade C(sp<sup>3</sup>)-OAr cleavage, sulfonyl rearrangement and atropisomeric cyclopropanation†

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For the first time, the combination of catalytic 1,10-phenanthroline and potassium carbonate enables a radical cleavage of a hydrazonyl N–S bond to deliver a tosyl radical and a diazo compound, along with releasing molecular hydrogen spontaneously. The tosyl radical and hydrogen evolution are witnessed by radical trapping, EPR experiment and gas chromatography. An innovative application of this strategy into metal-free coupling of *N*-tosylhydrazone and phosphinyl allene provides a one-step synthesis of novel atropisomeric 3-tosyl-1-enyl-cyclopropyl-diphenylphosphine oxide derivatives with excellent diastereoselectivity and *E*-selectivity. Moreover, these atropisomeric products, verified by <sup>1</sup>H-NMR, DFT calculations and X-ray crystallography, are controllably formed. Mechanistic studies indicate that the multistep cascade reaction occurs through radical hydrazonyl N–S bond cleavage, radical C(sp<sup>3</sup>)–OAr bond cleavage, sulfonyl rearrangement and atropisomeric cyclopropanation.

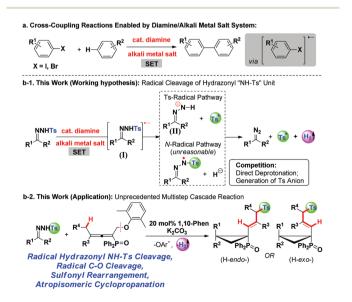
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# Introduction

In the past decade, the green and sustainable strategy for freeradical generation from a diamine/alkali metal salt system has contributed greatly to cross-coupling reactions, which received significant attention from scientific and industrial communities. Contributions by Hayashi, Shi, Lei, Murphy and others have witnessed the blooming of this chemistry, 2 lately classified as base-promoted homolytic aromatic substitution (BHAS),<sup>3</sup> in various inter/intramolecular C-C couplings between aryl halides and arenes or olefins (Scheme 1a). Although potassium t-butoxide alone was reported to enable the couplings, 4 a wide range of organic additives including 1,10-phenanthroline derivatives, aliphatic imines, amino acids, hydrazine derivatives, N-heterocyclic carbenes, and diphosphines have been used in initiating the radical chain process more efficiently. 2a-g,3b,5 From both experimental facts and theoretical calculation considerations, <sup>2i</sup>

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Scheme 1 Previous work with diamine/alkali metal salt combination (a) and this work (b-1, b-2).

a single electron transfer process has been verified, for example, 1,10-phenanthroline/t-BuOK initiated radical cleavage of the aryl–I bond will deliver an aryl radical and an iodide anion. We believe that this strategy should be ideally suitable for developing novel radical chain reactions other

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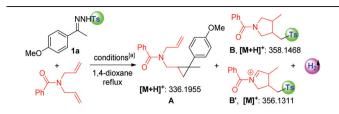
than C–C couplings; however, to the best of our knowledge, this field remains untouched so far.

The pursuit of N-tosylhydrazone chemistry dates back to the mid-20th century, as well-known as the Bamford-Stevens reaction and Shapiro reaction.<sup>6</sup> Since the first report of palladium-catalyzed coupling of N-tosylhydrazones with arylbromides in 2007, this chemistry has been attracting extensive interest in providing diazo precursors in building versatile functionalities for synthetic compounds, led by the groups of Barluenga and Valdés, Wang and others. Despite these great achievements, only a handful of methods have been regarding free-radical established generation N-tosylhydrazone, albeit confined to a transition metal-catalysis system. 10 For example, Xiao's group has established a photocatalytic hydroamination of N-tosylhydrazones with intramolecular alkenes, in which N-radical formation from single-electron oxidation of deprotonated N-tosylhydrazones was involved. 10a,b Very recently, cobalt III – carbene radicals, generated through metalloradical activation of N-tosylhydrazones by cobalt<sup>II</sup> complexes, were developed in the reactions of N-tosylhydrazones and olefins, alkynes or alkanes. 10d-f,10h-i Inspired by the aforementioned studies and our previous studies,11 we hypothesized that, a diamine/alkali metal salt excited hydrazonyl radical anion might undergo a probable Tsradical pathway (N-S bond cleavage)12 to furnish a diazo compound, tosyl radical and molecular hydrogen (Scheme 1b-1), and this unprecedented strategy should offer a powerful and important platform for discovering new radical chemistry of N-tosylhydrazones with allenes toward complex molecules.

### Results and discussion

With an aim to rationalize our hypothesis, an EPR experiment for the reaction of 1-(1-(4-methoxyphenyl)ethylidene)-2-tosylhydrazine (1a), 20 mol% of 1,10-phen and 1.5 equivalents of t-BuOK was conducted to monitor the formation of radical species, which indicated the formation of the tosyl radical (see more details in the ESI†).13 The tosyl radical could also be captured by TEMPO as monitored by HR-MS (in ESI†). As shown in Table 1, N,N-diallylbenzamide was subsequently applied to trap the fragments cleaved from N-tosylhydrazone (1a). In preliminary results, 20 mol% of 1,10-phen and 1.5 equivalents of t-BuOK or K2CO3 successfully enabled radical cleavage of the "NHTs" unit (entries 1 and 2), which was strongly supported by the detection of adducts A, B and B', deriving from the diazo precursor and the tosyl radical, respectively. Interestingly, a catalytic amount of potassium carbonate afforded the same result of fragment abundance in the HRMS spectrum, even for the case with high purity reagents (entries 3 and 4). A parallel experiment without 1,10-phen gave only the cyclopropanation product (A, entry 5). The observations in entries 3 and 4 deserve to be highlighted since the tosyl anion could be generated in the presence of excess base, and a single-electron oxidation of the tosyl anion by metal catalysts has been reported to afford the tosyl radical;<sup>14</sup> however, the

**Table 1** Studies on cleavage of *N*-tosylhydrazones: evidence of tosyl radical and molecular hydrogen evolution<sup>a</sup>



| Entry          | Radical initiator   | HR-MS<br>result            | $H_2$ evolution |
|----------------|---|----------------------------|-----------------|
| 1              | 20 mol% 1,10-phen/1.5 equiv. t-BuOK   | A + B + B'                 | Yes             |
| 2              | 20 mol% 1,10-phen/1.5 equiv. K <sub>2</sub> CO <sub>3</sub>                                       | A + B + B'                 | Yes             |
| 3              | 20 mol% 1,10-phen/15 mol% K <sub>2</sub> CO <sub>3</sub>  | A + B + B'                 | Yes             |
| $4^b$          | 20 mol% 1,10-phen/15 mol% K <sub>2</sub> CO <sub>3</sub>  | A + B + B'                 | Yes             |
| 5              | 1.5 equiv. K <sub>2</sub> CO <sub>3</sub>   | Only A                     | No              |
| 6 <sup>c</sup> | 20 mol% 1,10-phen/15 mol% K <sub>2</sub> CO <sub>3</sub>  | _ `                        | No              |
| 7              | 0.2 equiv. TBAI/2 equiv. TBHP   | $\mathbf{B} + \mathbf{B'}$ | No              |
| 8              | 5 mol% Pd(OAc) <sub>2</sub> /20 mol%  | A + B + B'                 | Yes             |
|                | 1,10-phen/15 mol% K <sub>2</sub> CO <sub>3</sub>  |                            |                 |
| 9              | 5 mol% Pd(PPh <sub>3</sub> ) <sub>2</sub> Cl <sub>2</sub> /15 mol% K <sub>2</sub> CO <sub>3</sub> | Only A                     | No              |
|                |   |                            |                 |

<sup>a</sup> N-Tosylhydrazone (1a, 0.1 mmol), N,N-diallylbenzamide (0.12 mmol), catalyst or oxidant in 3 mL refluxing 1,4-dioxane for 2 hours. <sup>b</sup> High purity reagents: 1,10-phen (anhydrous, purity > 99 + %), K<sub>2</sub>CO<sub>3</sub> (purity: 99.997%). <sup>c</sup> TsNa was used instead of 1a.

negative result of sodium 4-methylbenzene-sulfinate and *N,N*-diallyl-benzamide entirely ruled out the possibility of forming the tosyl radical from the tosyl anion under the current conditions (entry 6). Although TBAI/TBHP evidenced a positive result in producing the trapping products (**B** and **B**′) from the tosyl radical as monitored by HR-MS, it was found that the diazo precursor decomposed rapidly (entry 7). The results in entries 8 and 9 showed that a palladium catalyst would not inhibit the radical process but did not deliver the tosyl radical without the participation of 1,10-phen. Intriguingly, molecular hydrogen evolution was detected by gas chromatography in entries 1–4 and 8, indicating that the hydrogen radical or hydrogen anion might be produced in the system (in ESI†).

Subsequently, we applied this strategy in the coupling of N-tosylhydrazones and phosphinyl allenes (Scheme 1b-2). This choice was based on our previous report of a transition-metalcatalyzed coupling of N-tosylhydrazones with phosphinyl allenes to form phosphinyl [3]dendralenes, 11a which we believe would be a model reaction suitable for providing an expeditious platform for investigating the reactivity of the tosyl radical and diazo compound in situ generated from N-tosylhydrazones and differentiating the 1,10-phen/K2CO3 system from metal contamination. 14,15 The study was initiated by examining the coupling of 1a with phosphinyl allene (2a). As shown in Table 2, for a preliminary experiment with a catalytic amount of 1,10-phen and t-BuOK, 6% yield of (E)-(2-(4methoxy phenyl)-2-methyl-1-(2-methyl-3-tosylprop-1-en-1-yl) cyclopropyl)diphenyl-phosphine oxide (3a) was obtained with high diastereoselectivity, along with 50% yield of tosylated 1,3diene (4a) (entry 2). The ratios of 3a/4a changed in proportion to the amount of base used (entries 3-5), in which 2.5 equiva-

Table 2 Metal-free coupling of N-tosylhydrazone (1a) with phosphinyl allenes (2a): screening of conditions<sup>a</sup>

| Entry           | Catalyst (20 mol%)                                 | Base  | Yield (%) $(3a/4a)^b$ |
|-----------------|--|---|-----------------------|
| 1               | 1,10-Phen  | t-BuOK (15 mol%)                            | 0/14                  |
| 2               | 1,10-Phen  | t-BuOK (40 mol%)                            | 6/50                  |
| 3               | 1,10-Phen  | K <sub>2</sub> CO <sub>3</sub> (15 mol%)    | <5/85                 |
| 4               | 1,10-Phen  | $K_2CO_3$ (1.5 equiv.)                      | 35/30                 |
| 5               | 1,10-Phen  | $K_2CO_3$ (2.5 equiv.)                      | 74/0                  |
| 6               | 1,10-Phen  | NaHCO <sub>3</sub> (2.5 equiv.)             | 56/0                  |
| 7               | 1,10-Phen  | CsF (2.5 equiv.)                            | 53/0                  |
| 8               | 1,10-Phen  | $Cs_2CO_3$ (2.5 equiv.)                     | 17/0                  |
| 9               | 1,10-Phen  | DBU (2.5 equiv.)                            | 0/0                   |
| 10              | 2,2'-Bipyridine                                    | $K_2CO_3$ (2.5 equiv.)                      | 38/13                 |
| 11              | 2,9-Dimethyl-1,<br>10-phenanthroline               | $K_2CO_3$ (2.5 equiv.)                      | 36/11                 |
| 12              | 2,2':6',2"-Terpyridine                             | K <sub>2</sub> CO <sub>3</sub> (2.5 equiv.) | 42/9                  |
| 13              | 4,7-Dimethoxy-1,<br>10-phenanthroline              | $K_2CO_3$ (2.5 equiv.)                      | 50/0                  |
| $14^{c,d}$      | Pd(PPh <sub>3</sub> ) <sub>2</sub> Cl <sub>2</sub> | K <sub>2</sub> CO <sub>3</sub> (2.5 equiv.) | 0/0                   |
| $15^c$          | Pd(OAc) <sub>2</sub> /1,10-phen                    | $K_2CO_3$ (2.5 equiv.)                      | 40/0                  |
| 16 <sup>c</sup> | Cu(OAc) <sub>2</sub> /1,10-phen                    | $K_2CO_3$ (2.5 equiv.)                      | 0/0                   |
| 17 <sup>c</sup> | CoCl <sub>2</sub> /1,10-phen                       | $K_2CO_3$ (2.5 equiv.)                      | 21/0                  |
| $18^e$          | 1,10-Phen  | $K_2CO_3$ (2.5 equiv.)                      | 75/0                  |

<sup>a</sup> Reaction conditions: N-Tosylhydrazone (1a, 0.9 mmol), phosphinyl allene (2a, 0.3 mmol), catalyst (20 mol%) and base in 3 mL refluxing 1,4-dioxane for 8 hours. b Isolated yield by column chromatography (dr > 20:1, only *E*-isomer for all cases). °5 mol% palladium catalyst. The major product was phosphinyl[3]dendralenes, see our previous work in ref. 11a. <sup>e</sup> High purity reagents: 1,10-phen (anhydrous, purity > 99 + %), K<sub>2</sub>CO<sub>3</sub> (purity: 99.997%).

lents of K<sub>2</sub>CO<sub>3</sub> improved the isolated yield of 3a up to 74% (Note: this observation does not conclude that stoichiometric base is necessary to generate the tosyl radical, instead, excess base plays a vital role in converting 4a into 3a in a sulfonyl rearrangement step). A series of bases, including inorganic and organic bases, were then screened. Weaker inorganic bases, such as Na<sub>2</sub>CO<sub>3</sub>, NaHCO<sub>3</sub> and CsF, enabled the cascade reactions with yields ranging from 26 to 56%, whereas organic bases entirely inhibited the reaction (see the ESI†). Other catalysts of pyridine or phenanthroline derivatives were tested, which exhibited considerable activities as well (entries 10-13). The cascade reaction was also susceptible to solvents and catalyst loadings (in ESI†). Control experiments under transitionmetal catalysis were conducted as a comparison. Transition metals, including palladium, cobalt, nickel, rhodium, copper, zinc, manganese, iron and iridium, were investigated; however, all of the results were not comparable (entries 14–17, also in the ESI†). Most importantly, a high-impurity of 1,10phenanthroline and potassium carbonate delivered the final product 3a in a slightly better yield of 75% (entry 18), excluding the possibility of a metal-catalytic pathway and keeping in line with the HR-MS evaluations given in Table 1. The above observations indicated that the tosyl radical produced by 1,10phenanthroline and an alkali metal cation plays a key role in cleavage of the  $C(sp^3)$ -OAr bond, probably via a radical pathway, achieving a multistep cascade reaction with high efficiency.

With the optimal conditions in hand, we then evaluated the substrate scopes of various N-tosylhydrazones (1) with phosphinyl allenes (2). As summarized in Table 3, the cascade reactions gave adducts (3) in yields ranging from acceptable to medium with excellent diastereoselectivities E-selectivities, relying on the structure and electronic properties of substituents. For instance, electron-neutral N-tosylhydrazones and para-electron-rich ones afforded the products 3a-3c with medium yields. Otherwise, electrondeficient substituents, including p-F, p-Cl, and p-Br, impaired the efficiency to some extent, with yields dropping to below 50%. N-Tosylhydrazones bearing strong electron-deficient substituents of p-CF3 and p-NO2 performed very sluggishly without formation of the corresponding products. 56% yield of **3h** was obtained for β-naphthyl substituted substrates. Notably, the X-ray crystal structure of 3h unambiguously displayed the location of the tosyl group and the dimensional orientations of anchored moieties, 16 with methyl and diphenylphosphine oxide groups at syn-positions. Ethyl, n-butyl, and benzyl derived N-tosylhydrazones were also applicable to the system with acceptable to medium yields (3i-3m). Nevertheless, diphenyl derived N-tosyl-hydrazone performed

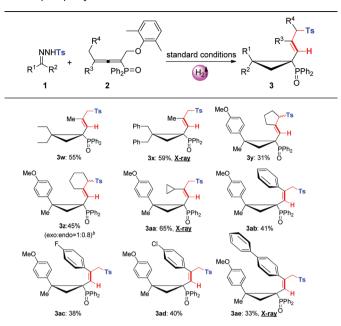
Table 3 1,10-Phen/K<sub>2</sub>CO<sub>3</sub> catalyzed coupling of N-tosyl-hydrazones (1) with phosphinyl allenes (2): formation of H-endo-isomers<sup>a</sup>

<sup>&</sup>lt;sup>a</sup> Isolated yield by column chromatography, dr > 20:1, only *E*-isomer.

sluggishly under current conditions. The phosphinyl allenes could be expanded to other aliphatic substitutions. For instance, the allenes with distal n-propyl, n-hexyl substituents furnished the target products (30, 3p) in yields of 40% and 31%, respectively. Other N-sulfonylhydrazone derivatives, including p-methoxyphenyl, m-methylphenyl, p-bromophenyl, polyaromatic and aliphatic ones, maintained certain reactivities in the radical cleavage and sulfonyl rearrangement step, giving exclusively E-isomers (3r-3v) with slightly lower yields. Moreover, it is worth mentioning that most of the proton chemical shifts of products (3a-3p, 3r-3v) locate in a range of 5.20-5.46 ppm, with alkenyl-H pointing to the top side of the three-membered ring (denoted as the H-endo-isomer). An exceptional case was found in the reaction of allenes bearing cycloheptyl substitution with 1a. Interestingly, 42% yield of H-endo-isomer 3q and 9% yield of H-exo-isomer 3'q were both isolated, and <sup>1</sup>H-NMR spectra of 3'q showed a significant downfield shift to 6.04 ppm. No reciprocal transformation was detected for these two isomers under thermal conditions, indicating that the cyclopropanation step should be decisive to obtain either atropisomer. Besides, this spatial configuration could be attributed to the congestion of moieties on a threemembered ring, in which aryl and diphenylphosphine oxide substituents strongly restricted the single bond rotation of the alkenyl moiety. DFT calculations showed that 3'q possesses higher energy than 3q with an energy gap of 20 kJ mol<sup>-1</sup>, and the torsion energy associated with the dihedral angle between 3q and 3'q is as high as  $59 \text{ kJ mol}^{-1}$  (see details in the ESI†).

To our delight, dominant H-exo-isomers appeared while altering the substitutions on N-tosylhydrazones and phosphinyl allenes (Table 4). In the case of diethyl and dibenzyl derived N-tosyl-hydrazones, products 3w and 3x were characterized as H-exo-configurations, evidenced by distinct alkene-H chemical shifts (5.62, 5.86 ppm) and an X-ray crystal structure of 3x.16 The ORTEP drawing of 3x unambiguously manifested the spatial proximity of the alkene-H atom to the P=O moiety, thus the deshielding effect resulted in an apparent downfield chemical shift. The scope of phosphinyl allenes was subsequently investigated. In general, the allenes bearing distal cyclopropyl, cyclopentyl and various aryl substituents furnished the corresponding H-exo-isomers exclusively in yields of 31-65% (3y-3ae), except for the cyclohexyl one (3z, exo: endo = 1:0.8). Meanwhile, the DFT calculation results were also well consistent with the ratio of product configurations (in the ESI†). X-ray crystal structures of 3aa and 3ae supported the formation of single atropisomers as well. Distinct from the substituents on hydrazones, electron effects were not apparent for allenes with terminal aromatics. Electron-deficient groups of para-fluoro and para-chloro gave indistinctive results compared to that of the electron-neutral substrate. Note that the development of atropisomers is currently a central task and hot research topic in organic chemistry. 17 However, as far as we know, atropisomeric cyclopropanation has never been reported and this cascade reaction provides highly efficient construction of multi-substituted atropisomers in one pot.

Table 4 1,10-Phen/K<sub>2</sub>CO<sub>3</sub> catalyzed coupling of N-tosyl-hydrazones (1) with phosphinyl allenes (2): formation of H-exo-isomers<sup>a,b</sup>



<sup>a</sup> Isolated yield by column chromatography, dr > 20:1, H-exo: H-endo > 20:1 with an exception of 3z. b The ratio of H-exo- and H-endo-isomers was determined by <sup>1</sup>H NMR, which cannot be separated even by preparative HPLC.

Next, in order to understand the reaction mechanism, control experiments were performed. Delightfully, while shortening the reaction time to 1 hour, we isolated the key intermediate, tosylated 1,3-dienes (4a), in a good yield of 85% (eqn (a), Scheme 2). Kinetic studies on this step verified that it proceeds through a radical pathway, and the aromatic ether moiety left in the radical form is trapped by BHT (in ESI†). The parallel reaction of tosyl sodium and allene 2a left all substrates intact (eqn (b)), implying that the tosyl anion cannot cleave the C-O bond to initiate the cascade reaction. Eqn (c) presented that the intermediate 4a could be converted into a sulfonyl rearrangement product 5a,18 but stoichiometric K<sub>2</sub>CO<sub>3</sub> turned out to be necessary, otherwise, this step severely retarded the cascade process. A crossover experiment with sodium 4-methoxybenzenesulfinate in eqn (d) indicated that the sulfonyl rearrangement step occurs inter-molecularly. After that, 5a underwent atropisomeric cyclopropanation rapidly to afford final product 3a in 60% yield. Besides, deuterium labelling experiments were conducted to investigate the kinetic isotope effects (KIE) (eqn (e)). The reaction of 1a and 2a-D<sup>6</sup> afforded deuterated product 4a-D<sup>6</sup> in 84% yield, which transformed into 5a-D in 60% yield with deuterium atoms distributed over all of the alkenyl positions. The KIE value from two parallel reactions between 4a and 4a-D<sup>6</sup> was found to be  $k_{\rm H}/k_{\rm D}$ = 1.5, suggesting that the rate-determining step probably involves in the sulfonyl rearrangement step, but does not contain a C(sp<sup>3</sup>)-H bond cleavage. Therefore, the mechanism studies and results in Table 1 collectively pointed out that the

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Scheme 2 Mechanism studies

cascade reaction consists of four steps: 1,10-phen/ $K_2CO_3$  catalysed cleavage of the NH-Ts bond to generate the tosyl radical, radical cleavage of the  $C(sp^3)$ -OAr bond, base-promoted sulfonyl rearrangement, and atropisomeric cyclopropanation.

Based on our experimental facts and previous reports, 2a, 10g, 19 a proposed mechanism is outlined in Scheme 3. Initially, single electron transfer (SET) from the 1,10-phen/ K<sub>2</sub>CO<sub>3</sub> complex to N-tosylhydrazone (1) gives a radical anion (I), which undergoes N-S bond cleavage to result in a tosyl radical and an N-anion intermediate (II) ((II) could be trapped by the CF<sub>3</sub> cation source and detected by HR-MS, see details in the ESI†). The N-anion intermediate (II) will leave the hydrogen anion rapidly to give a diazo compound [C]. Molecular hydrogen evolution comes from the deprotonation of 1 by the in situ generated hydrogen anion. Meanwhile, the accompanying tosyl anion helps to quench the associated radical cation (IV) to end the catalytic cycle and releases another portion of the tosyl radical. This proposed catalytic cycle keeps in line with the result of catalytic amounts of 1,10phen and K<sub>2</sub>CO<sub>3</sub> in eqn (a), Scheme 2. Afterwards, the tosyl radical attacks allene 2 to yield tosylated 1,3-butadienes (4). Base-promoted sulfonyl rearrangement of 4 leads to the formation of thermally reversible (E)-S-trans or (E)-S-cis isomers (5/5'), depending on the nature of substituents on the olefin double bond. Eventually, these two isomers undertake 1,3dipolar cycloaddition with diazo intermediate [C] to give atropisomers 3 in the H-exo or H-endo configuration, along with the extrusion of nitrogen. Above all, the equilibrium ratios of 5 and 5' are associated directly with the dominant stereoconfiguration of atropisomers. For instance, when the R<sup>3</sup>

Scheme 3 Proposed mechanism and rationalization of atropisomer formation.

group is a bulky phenyl group, an (E)-S-cis isomer is favourably formed and rapidly transforms into an H-exo-isomer (3ab).

# Experimental

unfavourable

#### General procedure

To a 15 mL sealed reaction tube were added *N*-tosylhydrazones (1a, 286 mg, 0.9 mmol), phosphinyl allenes (2a, 120 mg, 0.3 mmol), 1,10-phen (12 mg, 20 mol%),  $K_2CO_3$  (103.5 mg, 0.75 mmol), and 3 mL degassed 1,4-dioxane. The reaction mixture was then heated to reflux for 8 hours until the complete consumption of 2a monitored by TLC. After all of the volatiles were removed under vacuum, the crude product was purified by flash chromatography (eluent: 1:2:1 (v/v) of ethyl acetate/petroleum ether) to afford product 3a as a white solid.

# Conclusions

In conclusion, we disclosed, for the first time, a transitionmetal-free radical cleavage of a hydrazonyl "NH-Ts" unit with a catalytic amount of 1,10-phenanthroline and potassium carbonate. The cleavage of N-tosylhydrazone through a radical anion pathway toward a tosyl radical, diazo compound and molecular hydrogen was rationalized by DFT calculations and verified by radical trapping, EPR experiment and gas chromatography. Further application of this novel strategy into metal-free coupling of N-tosylhydrazones and phosphinyl allenes provides expeditious synthesis of unprecedented atropisomeric 3-tosyl-1-enyl-cyclopropyldiphenyl-phosphine oxide derivatives with excellent diastereoselectivity and E-selectivity. A variety of atropisomeric products, confirmed by <sup>1</sup>H NMR, DFT calculations, and X-ray crystallography, were controllably formed depending on the substitutions of N-tosyl-hydrazone and phosphinyl allenes. Mechanistic studies indicate that the cascade reaction proceeds through a tosyl radical initiated successive C(sp<sup>3</sup>)-OAr cleavage, sulfonyl rearrangement and atropisomeric cyclopropanation. We expect that this strategy will enrich the blooming radical chemistry and offer new scaffolds for the development of potential chiral molecules.

# Conflicts of interest

Research Article

There are no conflicts to declare.

# Acknowledgements

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