

Research Article

Selective Dealkenylative Functionalization of Styrenes via C-C Bond Cleavage

Jianzhong Liu,¹ Jun Pan,¹ Xiao Luo,¹ Xu Qiu,¹ Cheng Zhang,¹ and Ning Jiao^{1,2} 

¹State Key Laboratory of Natural and Biomimetic Drugs, Peking University, 100191 Beijing, China

²State Key Laboratory of Organometallic Chemistry, Chinese Academy of Sciences, Shanghai 200032, China

Correspondence should be addressed to Ning Jiao; jiaoning@bjmu.edu.cn

Received 15 July 2020; Accepted 8 October 2020; Published 10 November 2020

Copyright © 2020 Jianzhong Liu et al. Exclusive Licensee Science and Technology Review Publishing House. Distributed under a Creative Commons Attribution License (CC BY 4.0).

As a readily available feedstock, styrene with about 25 million tons of global annual production serves as an important building block and organic synthon for the synthesis of fine chemicals, polystyrene plastics, and elastomers. Thus, in the past decades, many direct transformations of this costless styrene feedstock were disclosed for the preparation of high-value chemicals, which to date, generally performed on the functionalization of styrenes through the allylic C-H bond, C(*sp*²)-H bond, or the C=C double bond cleavage. However, the dealkenylative functionalization of styrenes via the direct C-C single bond cleavage is so far challenging and still unknown. Herein, we report the novel and efficient C-C amination and hydroxylation reactions of styrenes for the synthesis of valuable aryl amines and phenols via the site-selective C(Ar)-C(alkenyl) single bond cleavage. This chemistry unlocks the new transformation and application of the styrene feedstock and provides an efficient protocol for the late-stage modification of substituted styrenes with the site-directed dealkenylative amination and hydroxylation.

1. Introduction

Styrenes are readily available bulk chemicals [1, 2] (produced globally ~25 million tons per year) and widely used in synthesis as a very common building blocks [3, 4]. In the past decades, the development of new direct transformations of styrenes has always been an attractive topic, because it represents the potential industrial application due to the readily available and costless properties of the styrene feedstock. Thus, some classical reactions including the traditional wacker oxidation [5, 6], alkene difunctionalization [7–14], oligomerization or polymerization [15], intramolecular cyclization [16, 17], oxidative cleavage of alkene [18–20], and Heck-type reactions [21, 22], as well as olefin metathesis [23–25], have been well developed and widely applied in chemical synthesis. Generally, these disclosed protocols rely on the functionalization of the C=C double bond [26], the C(*sp*²)-H bond [27–30], and the allylic C(*sp*³)-H bond [31–35] (Figure 1(a)). Although dealkenylative hydrogenation and thiylation of C(*sp*³)-C(*sp*²) bonds were significantly developed by Kwon and coworkers [36, 37], the dealkenylative C-C single bond functionalization of styrene is still unknown and remains an unmet challenging issue

due to its high thermodynamic stability (the BDE of the C(Ar)-C(alkenyl) single bond is 116.9 kcal/mol [38]) (Figure 1(b)). Thereby, the exploration of a new type of C-C bond activation [39–46] mode and strategy of styrene is undoubtedly very attractive, which may provide an alternative advance in the chemical synthesis and open new avenues for future research of alkene chemistry.

To address the above unsolved dealkenylative C-C single bond functionalization, we proposed a cascade activation strategy via the initial C=C double bond preactivation to break the conjugate structure of styrene and generate the active intermediate for the subsequent C(Ar)-C₁ bond cleavage. However, the intrinsic C₁-C₂ bond cleavage reactivity in styrene chemistry would be a challenging competitive pathway [18–20] (Figure 1(c)). The key point of this strategy is to generate an intermediate with entropic or enthalpic driving force to promote the selective dealkenylative C-C bond cleavage. Herein, we unlock a novel and efficient C-C nitrogeneration or hydroxylation reaction of styrenes for the preparation of high-value arylamines and phenols (Figure 1(d)). The significance of this chemistry is trifold: (1) this chemistry provides a new approach to arylamines or phenols under metal-free and simple operation conditions, which are of

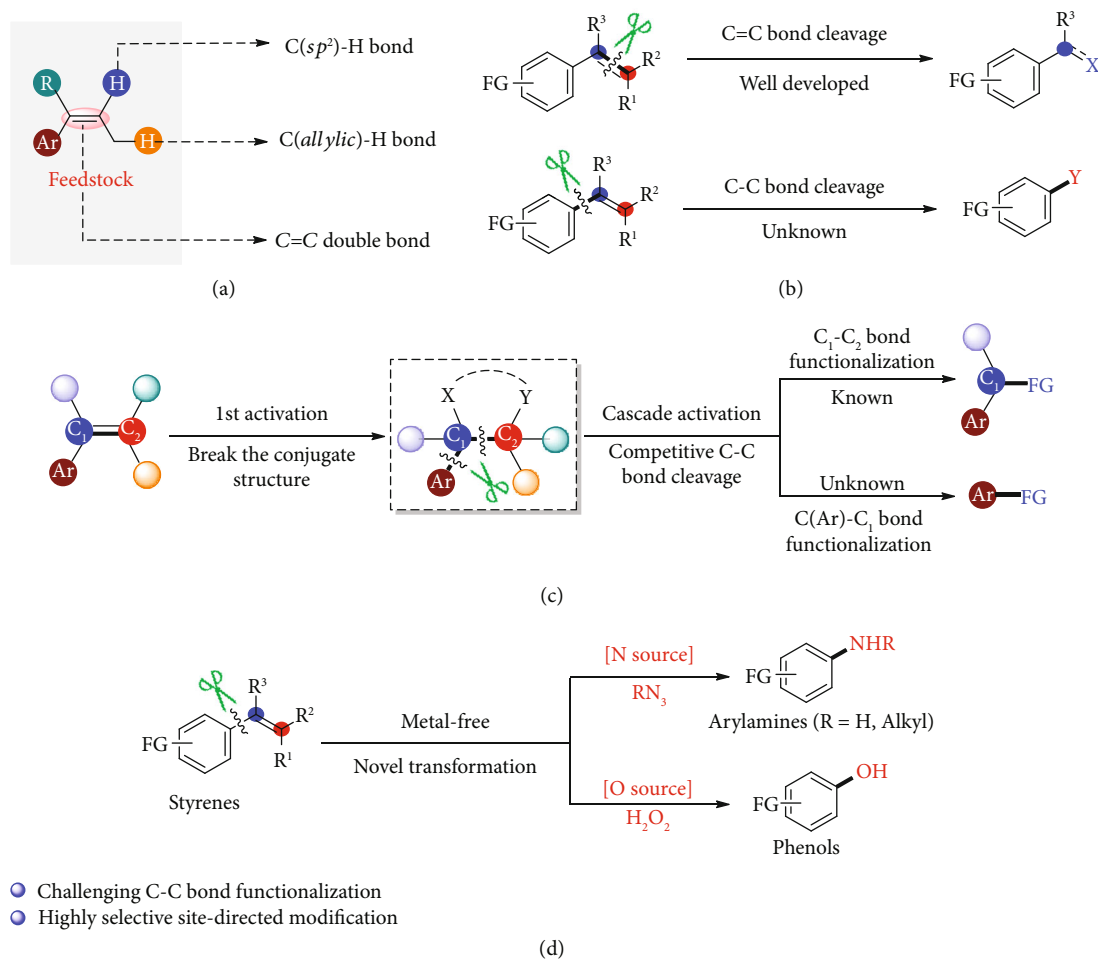


FIGURE 1: Functionalization based on styrenes. (a) Typical representative transformation patterns of styrene derivatives: C(sp^2)-H activation (for example, Heck-type reaction), C(allylic)-H activation, and C=C double bond activation (for example, wacker oxidation, difunctionalization, polymerization, and metathesis). However, other activation mode is still undeveloped yet very desired to synthetic chemistry. (b) Long-standing unmet challenges in the field of carbon-carbon bond cleavage chemistry of styrenes: although the C=C bond cleavage has been well studied, the dealkenylative C(Ar)-C(alkenyl) single bond cleavage is still unknown. (c) The proposed cascade activation strategy whereby the initial C=C double bond preactivation and the consecutive C(Ar)-C₁ single bond cleavage sequence may provide a chance to address the above unsolved dealkenylative transformation. However, the C₁-C₂ single bond cleavage in conventional styrene chemistry would be a challenging competitive path of the desired process. (d) This work: dealkenylative C-C bond amination and hydroxylation. FG: functional group.

considerable interest as synthons for the preparation of fine chemicals, pharmaceuticals, agrochemicals, and polymers [47, 48]; (2) compared to the aromatic C-H functionalization approach for the synthesis of arylamines and phenols which suffers from limited substrate scope, harsh conditions, and poor regioselectivity [49–53], the C(Ar)-C(alkenyl) single bond cleavage of styrene contributes a novel site-specific pathway for substituted arylamines and phenols synthesis; and (3) to the best of our knowledge, this chemistry is the first transformation of styrenes via the dealkenylative C-C single bond cleavage, which may inspire further methodology development based on olefins.

2. Results

Although the traditional C=C double bond cleavage leading to the corresponding aldehyde or ketone derivatives [18] pro-

vides a great challenge for the desired dealkenylative carbon-carbon functionalization, we investigated the hypothesis by a nucleophilic addition process to initially break the conjugate structure of the substrates. When 4-vinyl-1,1'-biphenyl (1a) was treated with azido nucleophile in the solvents such as DCE and CH₃CN, unfortunately, the substrate consumed but we did not detect any obvious products except some polymers (see Supplementary Table S1). To our delight, the aniline product 2a was obtained in the solvent of *n*-Hexane or CCl₄ under acidic conditions (see Supplementary Table S1), which indicated that the two-phase reaction condition generated by the combination of polar acid and the nonpolar solvent was vital to this process. Under the polar acidic conditions, the polymerization of the styrene is a very challenging inherent side reaction, so the choice of the nonpolar solvent such as CCl₄ is of importance for this dealkenylative transformation due to the formation of the

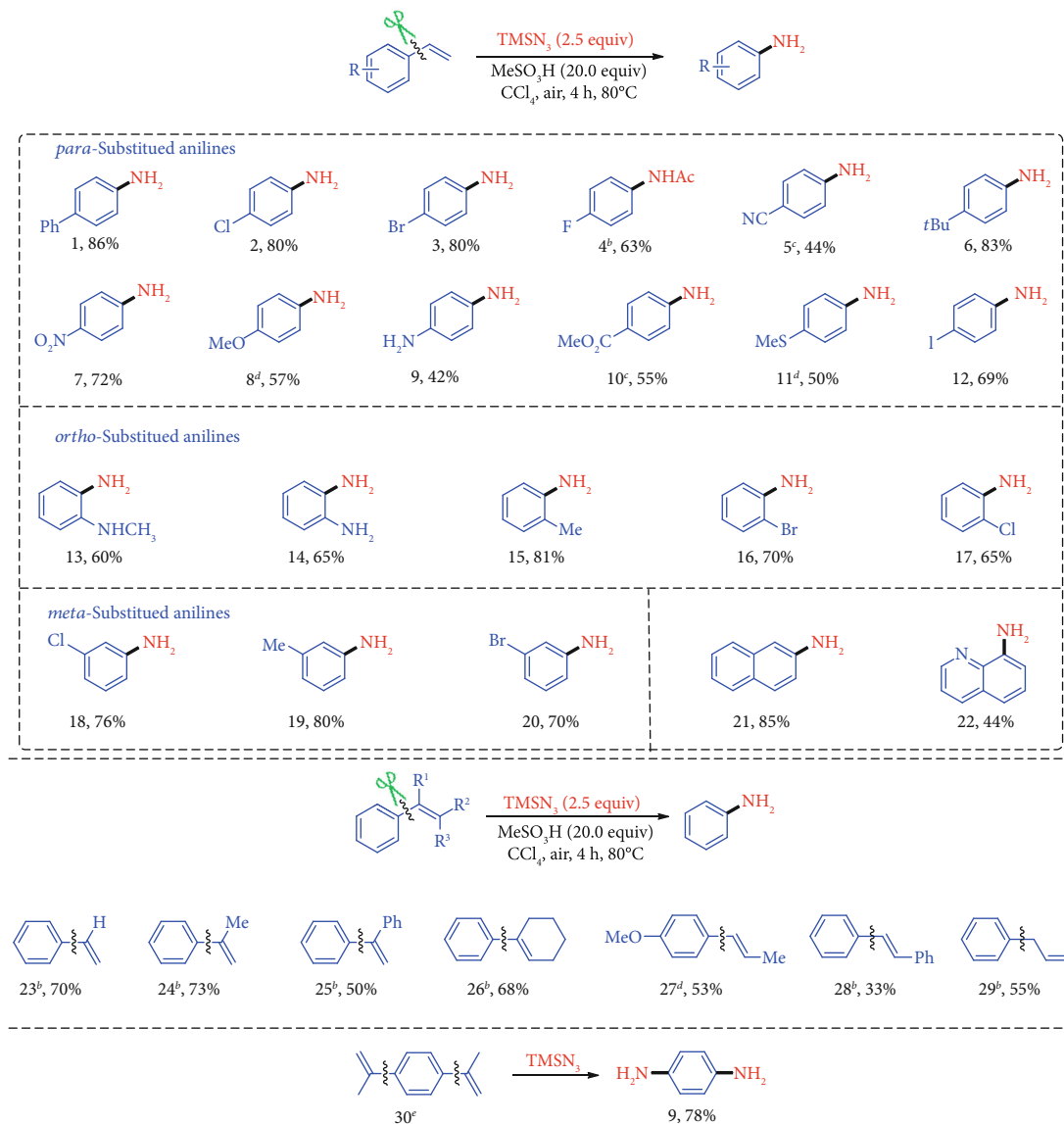


FIGURE 2: Substrate scope for the aniline synthesis from styrenes. ^aStandard conditions: reactions were performed with styrene (0.3 mmol), TMSN₃ (0.75 mmol), and MeSO₃H (6.0 mmol) in CCl₄ (1.0 mL) at 80°C for 4 h under atmosphere and isolated yields. ^bThe crude product was acetylated by acetyl chloride. ^cThe reaction was conducted at 40°C instead. ^dMeSO₃H (1.5 mmol) was used as the acid. ^eTMSN₃ (1.5 mmol) was used instead. CCl₄: tetrachloromethane.

two-phase reaction system with the polar acid to attenuate the side reaction. After the further screening of the acid additives, nitrogenation reagents, and other parameters (see Supplementary Table S3-4), this C-C amination reaction with the conditions of MeSO₃H (6.0 mmol) and TMSN₃ (0.75 mmol) in CCl₄ (1.0 mL) afforded the desired aniline 2a in 86% yield (Figure 2). The subsequent control experiment demonstrates that this chemistry is redox neutral with the acid additive as an essential player.

With the developed optimal reaction conditions, we next investigated the scope of this C-C amination with a series of commercially available or readily prepared styrenes as substrates (Figure 2). As expected, various *para*-substituted styrenes derivatives were compatible with this reaction system, and the corresponding anilines with different electronic

properties could be efficiently synthesized. For example, the styrenes bearing electron-donating groups (6, 8, 9, 11, R=OMe, *t*Bu, MeS, NH₂) underwent the amination process successfully to produce the *para*-substituted anilines in high efficiencies. Substrates containing halogen substituent (2, 3, 12) also performed well to give the corresponding products in good yields, leaving halogens available for the subsequent synthetic transformations. It is noteworthy that substrates with a strong electron-withdrawing group (4, 5, 7, 10, R=F, CN, NO₂, CO₂Me) could also deliver the corresponding anilines efficiently using this newly developed method, which is difficult to be prepared through the traditional nitration/reduction sequence or C-H amination pathway. The unprotected amino group is tolerant under these conditions and provides a novel pathway for the synthesis of aryl diamines

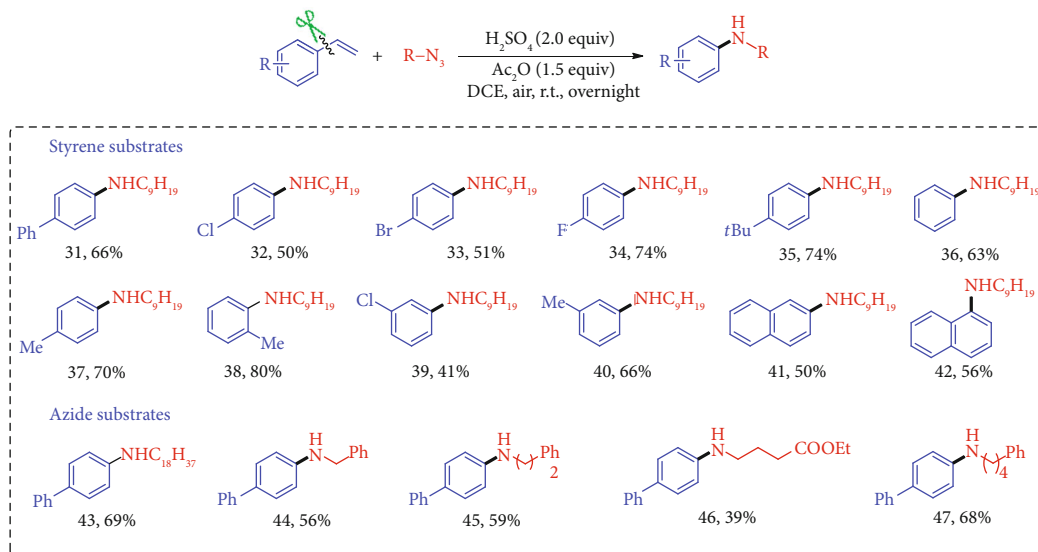


FIGURE 3: Substrate scope for the arylamine synthesis from styrenes. ^aStandard conditions: reactions were performed with styrene (0.3 mmol), alkyl azide (0.6 mmol), Ac₂O (0.45 mmol), and H₂SO₄ (0.6 mmol) in DCE (1.0 mL) at room temperature under air atmosphere overnight. Isolated yields. DCE: 1,2-dichloroethane.

(9) in moderate yields. The sulfide group which is relatively sensitive to oxidative condition or harsh nitration conditions was not destroyed in this protocol and afforded 4-(methylthio)aniline 11 in 50% yield.

Compared with the traditional nitration/reduction procedure in which the regioselectivity control is very challenging with the substituted arene substrates, the *ortho*-substituted anilines could be synthesized efficiently and selectively using the alkenyl group as a traceless site-directed group, thus without the extra complex isolation of the mixed *ortho*- and *para*-products (13–17). Notably, previously inaccessible *meta*-substituted anilines by the nitration/reduction process could also be prepared in good yields through the present C-C amination process (18–20). In addition, naphthyl and quinoline heterocyclic rings were also compatible, providing the expected product 21 and 22 in 85% and 44% yields, respectively.

To explore the effect of the alkenyl group on the styrenes, 1,1-disubstituted styrenes (24, 25 Figure 2) were first surveyed which produced the aniline products in good yields under the optimized conditions. Besides the terminal styrenes derivatives, the internal styrenes with bulky steric hindrance were also investigated. A natural bioactive molecule 1,2-disubstituted styrene (*trans*-anethole, 27) and (*E*-) stilbene (28) proceeded smoothly to form the target product. Moreover, 1,1,2-trisubstituted styrene, which bears bulkier hindrance, was also tolerated affording the aniline in 68% yield (26). To our delight, allylbenzene 29 could also furnish this C-C single bond cleavage due to the isomerization of the allyl group under acidic conditions. When styrene 30 bearing two alkenyls groups was employed as the substrate, two alkenyl groups on the aryl ring were cleaved simultaneously affording benzene-1,4-diamine 9 in 78% yield.

Interestingly, this dealkenylative C-C bond nitrogenation chemistry could also be successfully expanded to synthesize alkyl-substituted arylamines with alkyl azides as the *N*-source

under the conditions when employed H₂SO₄ (2.0 equiv) and Ac₂O (1.5 equiv) as the additives in DCE (for the results in different conditions, see Supplementary Table S5). During the reaction screening and optimization for the arylamine synthesis, many additives had been tried for the transformation and found that the anhydride had promoted the reaction, but it was not indispensable for the process. Definitely, its actual role in this synthetic route was still not completely clear yet. As shown in Figure 3, a series of styrenes containing substituents at the *para*-, *ortho*-, and *meta*-positions of the aromatic ring worked well and afforded the corresponding arylamines in moderate to good yields. Moreover, other alkyl azide reagents were tolerated in this transformation leading to various *N*-alkyl-substituted aniline products (43–47).

Although the epoxidation of styrenes was a known and favored process under oxidative conditions, inspired by the dealkenylative C-C bond amination results, we further investigated the C-C hydroxylation process with commercially available aqueous hydrogen peroxide as the oxygen source. Through the careful screening (see Supplementary Table S6), we optimized the conditions as MeSO₃H (2.0 equiv) and H₂O₂ (30%, 5.0 equiv) in MeNO₂/HFIP (4.5/1.5 mL, 0.05 M) stirring at 60°C. The reaction of styrenes under these conditions could afford the designed phenols by the novel dealkenylative C-C bond oxygenation process. The low reaction concentrations, the type of solvent, and acid were crucial to suppress the undesired by-products such as polymerization and epoxidation. As shown in Figure 4, a series of alkenyl groups on the styrenes (23–26, 28, 55, 56) were successfully replaced by the hydroxyl group to give the phenol products in moderate to good yields. Notably, the very active chalcones (57, 58) and cinnamyl alcohol (59) also worked albeit in low efficiency.

To further demonstrate the utility of this transformation, we carried out gram-scale reactions with styrene (23) as the

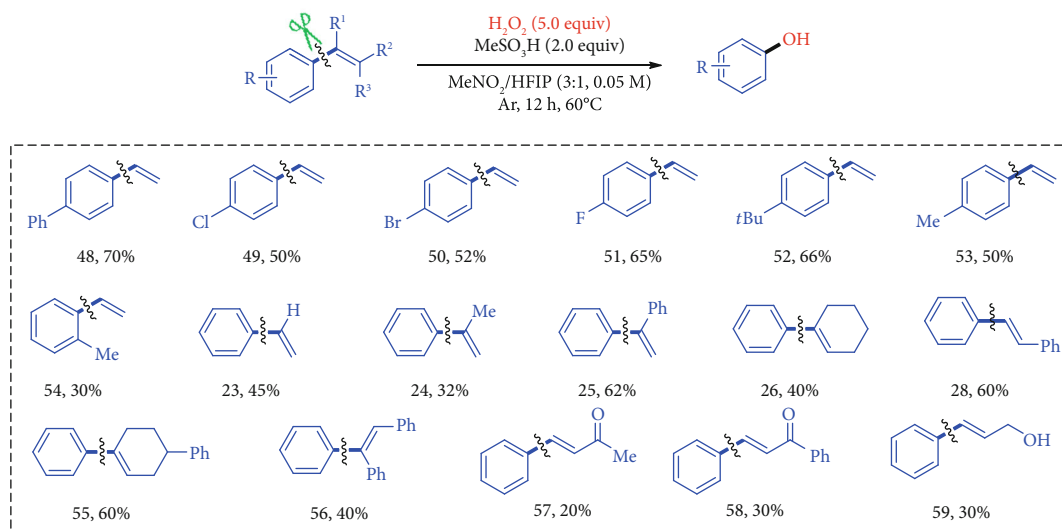


FIGURE 4: Substrate scope for the phenol synthesis from styrene. ^aStandard conditions: reactions were performed with styrene (0.3 mmol), H₂O₂ (1.5 mmol), and MeSO₃H (0.6 mmol) in MeNO₂/HFIP (4.5:0.5 mL, 0.05 M) at 60°C for 12 h under Ar atmosphere and isolated yields. MeNO₂: nitromethane; HFIP: hexafluoroisopropanol.

substrate which is a bulk chemical from natural sources and coal/petroleum products. The reaction offered the aniline in good yield, indicating its potential industrial application possibility (Figure 5(a)). In addition, the late-stage functionalization of complex bioactive molecules was further evaluated. Interestingly, 61 derived from (+)- δ -tocopherol was proven to be tolerated in this carbon-carbon amination process, affording the corresponding 62 in 47% yield. Additionally, the alkene-containing tyrosine derivative (63) and estrone derivative (65) could also furnish this transformation in good efficiency, giving 64 and 66 in 64% and 70% yield, respectively (Figure 5(b)).

Moreover, in order to testify the intermediates of this process and trace the alkenyl group, we first conducted an in situ reduction reaction with regard to the carbon-carbon amination procedure with NaBH₄ as the hydrogenative reagent, and arylamine 67 and 69 were produced in 55% and 41% yields, respectively, which indicates that the protonated imine 68 and 70 might be the key intermediates of this transformation. The result of the benzyl alcohol 71 under this C-C hydroxylation conditions quantitatively yielding the corresponding phenol (Figure 5(c)) suggests that the benzylic cation is probably involved in this oxygenation process. To explore the regiochemistry for the dealkenylative transformation, substrates of 73 and 74 have been conducted under the standard conditions (Figure 5(d)). High regioselectivities were obtained in these cases which was controlled by the stability of the generated benzylic carbon cation intermediate during the hydroazidation of alkene.

On the basis of the above results and previous reports [46, 54–59], the mechanism of this transformation was described in Figure 5(e). Initially, the acid-assisted hydroazidation of the C=C double-bond of styrenes occurs to generate the intermediate A with Markovnikov's rule [54, 55], which undergoes the subsequent Schmidt-type rearrangement process to afford the imine intermediate B through the cleavage of the C(Ar)-C(alkenyl) single bond [46, 56–59]. The final

hydrolysis of species B produces the desired anilines and aldehyde side products. Alternatively, a similar process occurs for the styrene substrate to generate the intermediate C in situ, which undergoes the traditional Hock process [60] to produce the phenol product.

3. Conclusions

This chemistry has described a novel carbon-carbon amination and hydroxylation of styrenes for the efficient and site-specific synthesis of arylamines and phenols. Significantly, this protocol provides a highly selective dealkenylative C-C bond activation mode of styrenes under transition-metal free and redox-neutral conditions with azide reagents as the nitrogenation reagents or aqueous hydrogen peroxide as the oxygen source. Compared to the poor regioselectivity and limited substrate scope in the typical aromatic C-H amination and hydroxylation process, this chemistry features site-directed selectivity and broad substrate scope. The simple and mild conditions make it applicable to the late-stage modification of some bioactive molecules. This strategy may open new avenues for the development of other novel transformations of alkenes through the C-C bond cleavage.

4. Methods

4.1. General C-C Amination Procedure. The substrate alkenes (0.3 mmol, 1.0 equiv), TMSN₃ (0.75 mmol, 2.5 equiv), and CCl₄ (1.0 mL), were added into a 20 mL vial equipped with a stir bar. Then, MeSO₃H (6.0 mmol, 20.0 equiv) was added. The reaction was refluxed under air at 80°C for 4 h. After cooling down to room temperature, the reaction mixture was quenched by 2 M NaOH (5 mL) and extracted by EA (5 × 2 mL), and the combined organic phase was washed with brine and dried over Na₂SO₄. Then, the mixture was concentrated and purified by flash chromatography on a short silica gel (eluent: PE/EA = 10/1) to afford the desired anilines.

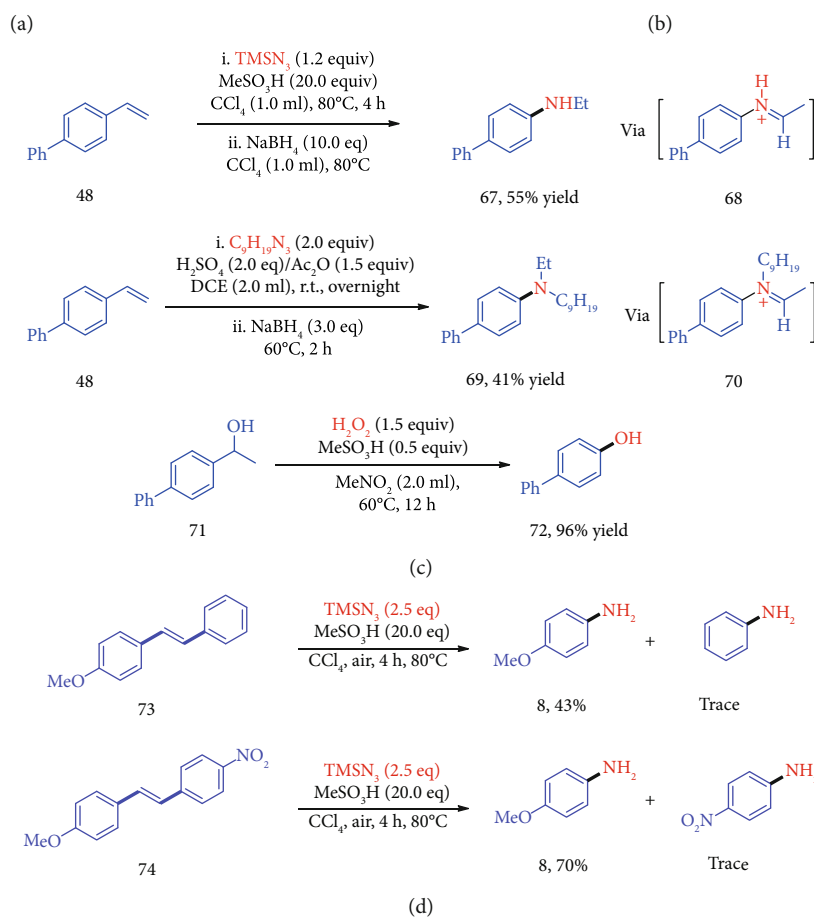
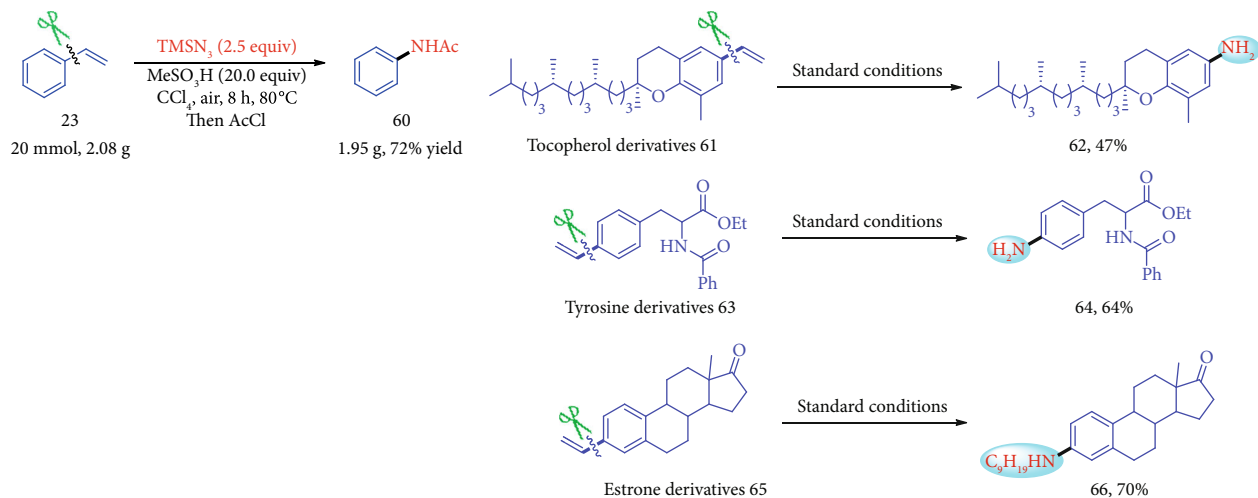


FIGURE 5: Continued.

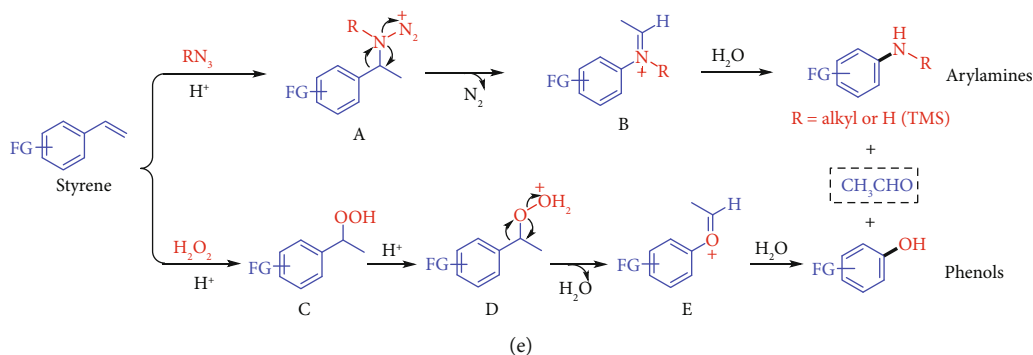


FIGURE 5: Synthetic applications and mechanism study of the dealkenylative C-C bond functionalization. (a) A gram-scale reaction with substrate 23 under standard conditions. (b) Modification of some pharmaceutical derivatives by this C-C amination protocol. (c) Mechanistic studies for the determination of the reaction intermediate. The formation of 67 and 69 suggests that the imine 68 and 70 are involved as the intermediates. The transformation from 71 to 72 shows the possible benzylic cation intermediate. (d) Regiochemistry exploration for this process. (e) The proposed mechanism of this dealkenylative C-C functionalization.

The substrate alkenes (0.2 mmol, 1.0 equiv), alkyl azide (0.4 mmol, 2.0 equiv), acetic anhydride (0.3 mmol, 1.5 equiv), and DCE (2.0 mL), were added into a 20 mL vial equipped with a stir bar. The mixture was stirred at 25°C. Then, *conc.* H₂SO₄ (0.4 mmol, 2.0 equiv) was added to the mixture in 5 seconds. The mixture was stirred at 25°C overnight. The reaction was quenched with 20% NaOH and was extracted with EA, purified by flash chromatography on a short silica gel (eluent: PE/EA = 50/1) to afford the desired arylamines.

4.2. General C-C hydroxylation procedure. The substrate alkenes (0.3 mmol, 1.0 equiv), MeNO₂ (4.5 mL)/HFIP (1.5 mL), were added into a 20 mL vial equipped with a stir bar. Then, 30% aqueous hydrogen peroxide solution (1.5 mmol, 5.0 equiv) and MeSO₃H (0.6 mmol, 2.0 equiv) were added in order. The reaction was heated under Ar at 60°C for 12 h. After cooling down to room temperature, the reaction mixture was quenched by *sat.* NaHCO₃ (5 mL) and extracted by EA (5 × 2 mL), and the combined organic phase was washed with brine and dried over Na₂SO₄. Then, the mixture was concentrated and purified by flash chromatography on a short silica gel (eluent: PE/EA = 10/1) to afford the desired phenols.

Conflicts of Interest

The authors declare no competing interests.

Authors' Contributions

J.L., J.P., and N.J. conceived and designed the experiments. J.L., J.P., and X.L. carried out most of the experiments. J.L., J.P., X.L., X.Q., C.Z., and N.J. analysed the data. J.L. and N.J. prepared the manuscript. N.J. directed the project. Jianzhong Liu and Jun Pan contributed equally to this work.

Acknowledgments

This study was financially supported from the National Natural Science Foundation of China (Nos. 21632001, 21772002, 81821004), the Drug Innovation Major Project

(2018ZX09711-001), Peking University Health Science Center (No. BMU20160541), and the Open Research Fund of Shanghai Key Laboratory of Green Chemistry and Chemical Processes which are greatly appreciated.

Supplementary Materials

Table S1: the effects of solvents for the synthesis of anilines. Table S2: the effects of additives for the synthesis of anilines. Table S3: the effects of acid for the synthesis of anilines. Table S4: the effects of nitrogeneration reagents and temperature for the synthesis of anilines. Table S5: the reaction optimization for the synthesis of arylamines. Table S6: the reaction optimization for the synthesis of phenols. The synthesis of substrates. General procedures. ¹H NMR and ¹³C NMR spectra of product. (*Supplementary Materials*)

References

- [1] B. A. Vaughan, M. S. Webster-Gardiner, T. R. Cundari, and T. B. Gunnoe, "A rhodium catalyst for single-step styrene production from benzene and ethylene," *Science*, vol. 348, no. 6233, pp. 421–424, 2015.
- [2] D. Ozokwelu and M. Mukherjee, *New process for producing styrene cuts costs SE, and reduces greenhouse gas emissions by U.S. Department of Energy*, 2013.
- [3] G. A. Olah and Á. Molnár, *Hydrocarbon chemistry*, Wiley, Hoboken, NJ, 2 edition, 2003.
- [4] H. A. Wittcoff, B. G. Reuben, and J. S. Plotkin, *Industrial organic chemicals*, Wiley, Hoboken, NJ, 2004.
- [5] J. Smidt, W. Hafner, R. Jira, R. Sieber, J. Sedlmeier, and A. Sabel, "The oxidation of olefins with palladium chloride catalysts," *Angewandte Chemie, International Edition*, vol. 1, no. 2, pp. 80–88, 1962.
- [6] R. Jira, "Acetaldehyde from ethylene—a retrospective on the discovery of the Wacker process," *Angewandte Chemie, International Edition*, vol. 48, no. 48, pp. 9034–9037, 2009.
- [7] R. I. McDonald, G. Liu, and S. S. Stahl, "Palladium(II)-catalyzed alkene functionalization via nucleopalladation: stereochemical pathways and enantioselective catalytic applications," *Chemical Reviews*, vol. 111, no. 4, pp. 2981–3019, 2011.

- [8] J. Gui, C. M. Pan, Y. Jin et al., "Practical olefin hydroamination with nitroarenes," *Science*, vol. 348, no. 6237, pp. 886–891, 2015.
- [9] Y. Yang, S. L. Shi, D. W. Niu, P. Liu, and S. L. Buchwald, "Catalytic asymmetric hydroamination of unactivated internal olefins to aliphatic amines," *Science*, vol. 349, no. 6243, pp. 62–66, 2015.
- [10] N. K. Fu, G. S. Sauer, A. Saha, A. Loo, and S. Lin, "Metal-catalyzed electrochemical diazidation of alkenes," *Science*, vol. 357, no. 6351, pp. 575–579, 2017.
- [11] M. Gaydou, T. Moragas, F. Juliá-Hernández, and R. Martin, "Site-selective catalytic carboxylation of unsaturated hydrocarbons with CO₂ and water," *Journal of the American Chemical Society*, vol. 139, no. 35, pp. 12161–12164, 2017.
- [12] A. J. Musacchio, B. C. Lainhart, X. Zhang, S. G. Naguib, T. C. Sherwood, and R. R. Knowles, "Catalytic intermolecular hydroaminations of unactivated olefins with secondary alkyl amines," *Science*, vol. 355, no. 6326, pp. 727–730, 2017.
- [13] T. M. Monos, R. C. McAtee, and C. R. J. Stephenson, "Arylsulfonylacetamides as bifunctional reagents for alkene aminoarylation," *Science*, vol. 361, no. 6409, pp. 1369–1373, 2018.
- [14] L. Legnani, G. Prina-Cerai, T. Delcaillau, S. Willems, and B. Morandi, "Efficient access to unprotected primary amines by iron-catalyzed aminochlorination of alkenes," *Science*, vol. 362, no. 6413, pp. 434–439, 2018.
- [15] S. Aoshima and S. Kanaoka, "A renaissance in living cationic polymerization," *Chemical Reviews*, vol. 109, no. 11, pp. 5245–5287, 2009.
- [16] S. R. Chemler and P. H. Fuller, "Heterocycle synthesis by copper facilitated addition of heteroatoms to alkenes, alkynes and arenes," *Chemical Society Reviews*, vol. 36, no. 7, pp. 1153–1160, 2007.
- [17] A. Minatti and K. Muñoz, "Intramolecular aminopalladation of alkenes as a key step to pyrrolidines and related heterocycles," *Chemical Society Reviews*, vol. 36, no. 7, pp. 1142–1152, 2007.
- [18] G. Urgoitia, R. SanMartin, M. T. Herrero, and E. Domínguez, "Aerobic cleavage of alkenes and alkynes into carbonyl and carboxyl compounds," *ACS Catalysis*, vol. 7, no. 4, pp. 3050–3060, 2017.
- [19] T. Wang and N. Jiao, "TEMPO-catalyzed aerobic oxygenation and nitrogenation of olefins via C=C double-bond cleavage," *Journal of the American Chemical Society*, vol. 135, no. 32, pp. 11692–11695, 2013.
- [20] Y. Imada, Y. Okada, K. Noguchi, and K. Chiba, "Selective functionalization of styrenes with oxygen using different electrode materials: olefin cleavage and synthesis of tetrahydrofuran derivatives," *Angewandte Chemie, International Edition*, vol. 58, no. 1, pp. 125–129, 2019.
- [21] C. C. C. Johansson Seechurn, M. O. Kitching, T. J. Colacot, and V. Snieckus, "Palladium-catalyzed cross-coupling: a historical contextual perspective to the 2010 Nobel Prize," *Angewandte Chemie, International Edition*, vol. 51, no. 21, pp. 5062–5085, 2012.
- [22] E. W. Werner, T.-S. Mei, A. J. Burckle, and M. S. Sigman, "Enantioselective Heck arylations of acyclic Alkenyl alcohols using a redox-relay strategy," *Science*, vol. 338, no. 6113, pp. 1455–1458, 2012.
- [23] R. Grubbs and H. Olefin-Metathesis, "Olefin-metathesis catalysts for the preparation of molecules and materials (nobel lecture)," *Angewandte Chemie, International Edition*, vol. 45, no. 23, pp. 3760–3765, 2006.
- [24] J. R. Ludwig, P. M. Zimmerman, J. B. Gianino, and C. S. Schindler, "Iron(III)-catalysed carbonyl-olefin metathesis," *Nature*, vol. 533, no. 7603, pp. 374–379, 2016.
- [25] T. T. Nguyen, M. J. Koh, T. J. Mann, R. R. Schrock, and A. H. Hoveyda, "Synthesis of E- and Z-trisubstituted alkenes by catalytic cross-metathesis," *Nature*, vol. 552, no. 7685, pp. 347–354, 2017.
- [26] G. Coin, R. Patra, S. Rana et al., "Fe-catalyzed aziridination is governed by the electron affinity of the active imido-iron species," *ACS Catalysis*, vol. 10, no. 17, pp. 10010–10020, 2020.
- [27] S. Maity, S. Manna, S. Rana, T. Naveen, A. Mallick, and D. Maiti, "Efficient and stereoselective nitration of mono- and disubstituted olefins with AgNO₂ and TEMPO," *Journal of the American Chemical Society*, vol. 135, no. 9, pp. 3355–3358, 2013.
- [28] M. Bera, A. Maji, S. K. Sahoo, and D. Maiti, "Palladium(II)-catalyzed meta-C-H olefination: constructing multisubstituted arenes through homo-diolefination and sequential hetero-diolefination," *Angewandte Chemie International Edition*, vol. 54, no. 29, pp. 8515–8519, 2015.
- [29] U. Dutta, S. Maiti, S. Pimparkar et al., "Rhodium catalyzed template-assisted distalpara-C-H olefination," *Chemical science*, vol. 10, no. 31, pp. 7426–7432, 2019.
- [30] A. Deb, A. Hazra, Q. Peng, R. S. Paton, and D. Maiti, "Detailed mechanistic studies on palladium-catalyzed selective C-H olefination with aliphatic alkenes: a significant influence of proton shuttling," *Journal of the American Chemical Society*, vol. 139, no. 2, pp. 763–775, 2017.
- [31] S. Maity, P. Dolui, R. Kancherla, and D. Maiti, "Introducing unactivated acyclic internal aliphatic olefins into a cobalt catalyzed allylic selective dehydrogenative Heck reaction," *Chemical science*, vol. 8, no. 7, pp. 5181–5185, 2017.
- [32] T. K. Achar, X. Zhang, R. Mondal et al., "Palladium-catalyzed directed meta-selective C-H allylation of arenes: unactivated internal olefins as Allyl surrogates," *Angewandte Chemie, International Edition*, vol. 58, no. 30, pp. 10353–10360, 2019.
- [33] T. A. Ramirez, B. Zhao, and Y. Shi, "Recent advances in transition metal-catalyzed sp³ C-H amination adjacent to double bonds and carbonyl groups," *Chemical Society Reviews*, vol. 41, no. 2, pp. 931–942, 2012.
- [34] M. E. Harvey, D. G. Musaev, and J. Du Bois, "A diruthenium catalyst for selective, intramolecular allylic C-H amination: Reaction Development and Mechanistic Insight Gained through Experiment and Theory," *Journal of the American Chemical Society*, vol. 133, no. 43, pp. 17207–17216, 2011.
- [35] W. Liu, S. Z. Ali, S. E. Ammann, and M. C. White, "Asymmetric allylic C-H alkylation via palladium(II)/cis-ArSOX catalysis," *Journal of the American Chemical Society*, vol. 140, no. 34, pp. 10658–10662, 2018.
- [36] A. J. Smaligo, M. Swain, J. C. Quintana, M. F. Tan, D. A. Kim, and O. Kwon, "Hydrodealkenylative C(sp³)-C(sp²) bond fragmentation," *Science*, vol. 364, no. 6441, pp. 681–685, 2019.
- [37] A. J. Smaligo and O. Kwon, "Dealkenylative thiylation of C(sp³)-C(sp²) bonds," *Organic Letters*, vol. 21, no. 21, pp. 8592–8597, 2019.
- [38] A. Zavitsas and A. The, "The relation between bond lengths and dissociation energies of carbon-carbon bonds," *The Journal of Physical Chemistry. A*, vol. 107, no. 6, pp. 897–898, 2003.

- [39] W. D. Jones, "The fall of the C–C bond," *Nature*, vol. 364, no. 6439, pp. 676–677, 1993.
- [40] M. Murakami, H. Amii, and Y. Ito, "Selective activation of carbon–carbon bonds next to a carbonyl group," *Nature*, vol. 370, no. 6490, pp. 540–541, 1994.
- [41] Y. J. Park, J.-W. Park, and C.-H. Jun, "Metal–organic cooperative catalysis in C–H and C–C bond activation and its concurrent recovery," *Accounts of Chemical Research*, vol. 41, no. 2, pp. 222–234, 2008.
- [42] A. Masarwa, D. Didier, T. Zabrodski, M. Schinkel, L. Ackermann, and I. Marek, "Merging allylic carbon–hydrogen and selective carbon–carbon bond activation," *Nature*, vol. 505, no. 7482, pp. 199–203, 2014.
- [43] L. Souillart and N. Cramer, "Catalytic C–C bond activations via oxidative addition to transition metals," *Chemical Reviews*, vol. 115, no. 17, pp. 9410–9464, 2015.
- [44] Y. Xia, G. Lu, P. Liu, and G. Dong, "Catalytic activation of carbon–carbon bonds in cyclopentanones," *Nature*, vol. 539, no. 7630, pp. 546–550, 2016.
- [45] G. Fumagalli, S. Stanton, and J. F. Bower, "Recent methodologies that exploit C–C single-bond cleavage of strained ring systems by transition metal complexes," *Chemical Reviews*, vol. 117, no. 13, pp. 9404–9432, 2017.
- [46] J. Liu, X. Qiu, X. Huang et al., "From alkylarenes to anilines via site-directed carbon–carbon amination," *Nature Chemistry*, vol. 11, no. 1, pp. 71–77, 2019.
- [47] S. A. Lawrence, Ed., *Amines: Synthesis, Properties and Applications*, Cambridge, Cambridge University Press, 2004.
- [48] M. Weber and M. Kleine-Boymann, *Phenol in Ullmann's Encyclopedia of Industrial Chemistry*, Wiley-VCH, Weinheim, 2004.
- [49] N. A. Romero, "Site-selective arene C–H amination via photoredox catalysis," *Science*, vol. 349, no. 6254, pp. 1326–1330, 2015.
- [50] Y.-W. Zheng, B. Chen, P. Ye et al., "Photocatalytic hydrogen-evolution cross-couplings: benzene C–H amination and hydroxylation," *Journal of the American Chemical Society*, vol. 138, no. 32, pp. 10080–10083, 2016.
- [51] C. Yuan, Y. Liang, T. Hernandez, A. Berriochoa, K. N. Houk, and D. Siegel, "Metal-free oxidation of aromatic carbon–hydrogen bonds through a reverse-rebound mechanism," *Nature*, vol. 499, no. 7457, pp. 192–196, 2013.
- [52] M. P. Paudyal, A. M. Adebesein, S. R. Burt et al., "Dirhodium-catalyzed C–H arene amination using hydroxylamines," *Science*, vol. 353, no. 6304, pp. 1144–1147, 2016.
- [53] A. Ruffoni, F. Juliá, T. D. Svejstrup, A. J. McMillan, J. J. Douglas, and D. Leonori, "Practical and regioselective amination of arenes using alkyl amines," *Nature Chemistry*, vol. 11, no. 5, pp. 426–433, 2019.
- [54] D. Balderman and A. Kalir, "Selective Reduction of azides. Improved preparation of α,α -disubstituted benzylamines," *Synthesis*, vol. 1978, no. 1, pp. 24–26, 2002.
- [55] J. Waser, H. Nambu, and E. M. Carreira, "Cobalt-catalyzed hydroazidation of olefins: convenient access to alkyl azides," *Journal of the American Chemical Society*, vol. 127, no. 23, pp. 8294–8295, 2005.
- [56] T. Sasaki, S. Eguchi, and N. Toi, "Synthesis of adamantane derivatives. 47. Photochemical synthesis of 4-azahomoadamant-4-enes and further studies on their reactivity in some cycloadditions," *The Journal of Organic Chemistry*, vol. 44, no. 21, pp. 3711–3715, 1979.
- [57] F. J. Lopez and D. Nitzan, "N-Methylanilines from benzylic azides," *Tetrahedron Letters*, vol. 40, no. 11, pp. 2071–2074, 1999.
- [58] W. H. Pearson, R. Walavalkar, J. M. Schkeryantz, W. K. Fang, and J. D. Blickendorf, "Intramolecular Schmidt reactions of azides with carbocations: synthesis of bridged-bicyclic and fused-bicyclic tertiary amines," *Journal of the American Chemical Society*, vol. 115, pp. 10183–10194, 1993.
- [59] A. Wroblewski, T. C. Coombs, C. W. Huh, S.-W. Li, and J. Aubé, "The Schmidt reaction," *Organic Reactions*, vol. 78, pp. 1–320, 2012.
- [60] H. Hock and S. Lang, "Autoxydation von Kohlenwasserstoffen, IX. Mitteil.: Über Peroxyde von Benzol-Derivaten," *Berichte der Deutschen Chemischen Gesellschaft*, vol. B77, pp. 257–264, 1944.