

Cu(I)-Catalyzed Diamination of Disubstituted Terminal Olefins: An Approach to Potent NK₁ Antagonist

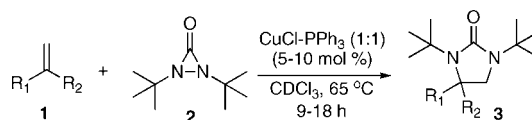
Yuehong Wen, Baoguo Zhao, and Yian Shi*

Department of Chemistry, Colorado State University, Fort Collins, Colorado 80523

yian@lamar.colostate.edu

Received April 13, 2009

ABSTRACT



This paper describes a diamination process using di-*tert*-butyldiaziridinone as nitrogen source and CuCl as catalyst. A wide variety of disubstituted terminal olefins can be efficiently diaminated in good yields under mild condition. This diamination process was used to synthesize potent NK₁ antagonist Sch 425078.

Diamination of olefins provides an effective approach to vicinal diamines, which are biologically and chemically important functional moieties.¹ Various metal-mediated^{2,3} and metal-catalyzed⁴⁻⁶ diaminations have been developed. Recently, we have reported the Pd(0)- and Cu(I)-catalyzed

regioselective diamination of conjugated dienes and trienes⁷⁻⁹ as well as the dehydrogenative diamination of terminal olefins^{10,11} using di-*tert*-butyldiaziridinone,^{13,14} di-aziridinimine,¹⁵ or di-*tert*-butylthiadiaziridine 1,1-dioxide¹⁶

(1) For leading reviews, see: (a) Lucet, D.; Le Gall, T.; Mioskowski, C. *Angew. Chem., Int. Ed.* **1998**, 37, 2580. (b) Mortensen, M. S.; O'Doherty, G. A. *Chemtracts: Org. Chem.* **2005**, 18, 555. (c) Kotti, S. R. S. S.; Timmons, C.; Li, G. *Chem. Biol. Drug Des.* **2006**, 67, 101. (d) de Figueiredo, R. M. *Angew. Chem., Int. Ed.* **2009**, 48, 1190.

(2) For examples of metal-mediated diaminations, see the following. Ti: (a) Aranda, V. G.; Barluenga, J.; Aznar, F. *Synthesis* **1974**, 504. Os: (b) Chong, A. O.; Oshima, K.; Sharpless, K. B. *J. Am. Chem. Soc.* **1977**, 99, 3420. (c) Muñiz, K. *Eur. J. Org. Chem.* **2004**, 2243. Pd: (d) Bäckvall, J.-E. *Tetrahedron Lett.* **1978**, 163. Hg: (e) Barluenga, J.; Alonso-Cires, L.; Asensio, G. *Synthesis* **1979**, 962. Co: (f) Becker, P. N.; White, M. A.; Bergman, R. G. *J. Am. Chem. Soc.* **1980**, 102, 5676. Mn: (g) Fristad, W. E.; Brandvold, T. A.; Peterson, J. R.; Thompson, S. R. *J. Org. Chem.* **1985**, 50, 3647.

(3) For a recent Cu(II)-mediated intramolecular diamination, see: (a) Zabawa, T. P.; Kasi, D.; Chemler, S. R. *J. Am. Chem. Soc.* **2005**, 127, 11250. (b) Zabawa, T. P.; Chemler, S. R. *Org. Lett.* **2007**, 9, 2035.

(4) For recent Pd(II)- and Ni(II)-catalyzed intramolecular diamination of olefins, see: (a) Streuff, J.; Hövelmann, C. H.; Nieger, M.; Muñiz, K. *J. Am. Chem. Soc.* **2005**, 127, 14586. (b) Muñiz, K.; Streuff, J.; Hövelmann, C. H.; Núñez, A. *Angew. Chem., Int. Ed.* **2007**, 46, 7125. (c) Muñiz, K. *J. Am. Chem. Soc.* **2007**, 129, 14542. (d) Muñiz, K.; Hövelmann, C. H.; Streuff, J. *J. Am. Chem. Soc.* **2008**, 130, 763. (e) Sibbald, P. A.; Michael, F. E. *Org. Lett.* **2009**, 11, 1147.

(5) For Rh(II)- and Fe(III)-catalyzed diamination with TsNCl₂, see: (a) Li, G.; Wei, H.-X.; Kim, S. H.; Carducci, M. D. *Angew. Chem., Int. Ed.* **2001**, 40, 4277. (b) Wei, H.-X.; Kim, S. H.; Li, G. *J. Org. Chem.* **2002**, 67, 4777.

(6) For a recent Pd(II)-catalyzed intermolecular diamination of conjugated dienes, see: Bar, G. L. J.; Lloyd-Jones, G. C.; Booker-Milburn, K. I. *J. Am. Chem. Soc.* **2005**, 127, 7308.

(7) For Pd(0)-catalyzed intermolecular diamination of conjugated dienes and trienes using di-*tert*-butyldiaziridinone, see: (a) Du, H.; Zhao, B.; Shi, Y. *J. Am. Chem. Soc.* **2007**, 129, 762. (b) Xu, L.; Du, H.; Shi, Y. *J. Org. Chem.* **2007**, 72, 7038. (c) Du, H.; Yuan, W.; Zhao, B.; Shi, Y. *J. Am. Chem. Soc.* **2007**, 129, 11688.

(8) For Cu(I)-catalyzed intermolecular diamination of conjugated dienes and trienes using di-*tert*-butyldiaziridinone, see: (a) Yuan, W.; Du, H.; Zhao, B.; Shi, Y. *Org. Lett.* **2007**, 9, 2589. (b) Du, H.; Zhao, B.; Yuan, W.; Shi, Y. *Org. Lett.* **2008**, 10, 4231.

(9) For Cu(I)-catalyzed intermolecular diamination of conjugated dienes, trienes, and activated terminal olefins using diaziridinimine, see: Zhao, B.; Du, H.; Shi, Y. *Org. Lett.* **2008**, 10, 1087.

(10) For Pd(0)-catalyzed intermolecular allylic and homoallylic C-H diamination of terminal olefins using di-*tert*-butyldiaziridinone, see: (a) Du, H.; Yuan, W.; Zhao, B.; Shi, Y. *J. Am. Chem. Soc.* **2007**, 129, 7496. (b) Du, H.; Zhao, B.; Shi, Y. *J. Am. Chem. Soc.* **2008**, 130, 8590.

(11) For Pd(0)-catalyzed dehydrogenative diamination of terminal olefins using di-*tert*-butylthiadiaziridine 1,1-dioxide, see: Wang, B.; Du, H.; Shi, Y. *Angew. Chem., Int. Ed.* **2008**, 47, 8224.

(12) For Cu(I)-catalyzed intermolecular diamination of activated terminal olefins using di-*tert*-butylthiadiaziridine 1,1-dioxide, see: Zhao, B.; Yuan, W.; Du, H.; Shi, Y. *Org. Lett.* **2007**, 9, 4943.

(13) Greene, F. D.; Stowell, J. C.; Bergmark, W. R. *J. Org. Chem.* **1969**, 34, 2254.

(14) For a leading review on diaziridinones, see: Heine, H. W. In *The Chemistry of Heterocyclic Compounds*; Hassner, A., Ed.; John Wiley & Sons, Inc.: Hoboken, NJ, 1983; p 547.

as the nitrogen sources. The Cu(I)-catalyzed diamination process has also been extended to activated monosubstituted terminal olefins such as styrenes, enynes, enol ether, etc.^{9,12} Considering the fact that 4,4-disubstituted 2-imidazolidinones (Figure 1) have been shown to be potent NK₁ antagonists,¹⁷

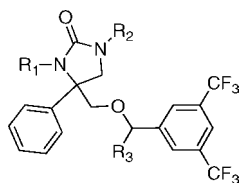
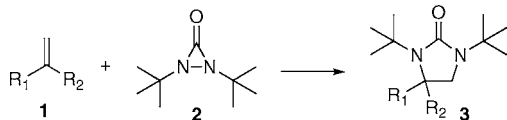


Figure 1. NK₁ antagonists.

and in conjunction with our efforts to expand the diamination scope, we have investigated the diamination of disubstituted simple terminal olefins using di-*tert*-butyldiaziridinone (**2**) as the nitrogen source (Scheme 1).⁸ Herein, we wish to report our studies on this subject.

Scheme 1



Initial studies were carried out using 2-phenylpropene as the substrate and di-*tert*-butyldiaziridinone (**2**) as the nitrogen source. When Pd(PPh₃)₄ was used as catalyst, only a trace amount of diamination product was observed. However, >95% conversion was obtained when the reaction was carried out with 5 mol % of CuCl–PPh₃ (1:1) in CDCl₃ at 65 °C.¹⁸ As shown in Table 1, various 2-phenylpropenes with different substituents on the phenyl ring can be successfully diami-

Table 1. Catalytic Diamination of Disubstituted Terminal Olefins^a

entry	substrate 1	product 3	yield (%) ^d
1	1a , X = H		91
2	1b , X = <i>o</i> -F		47
3 ^b	1c , X = <i>o</i> -OMe		50
4	1d , X = <i>m</i> -Br		86
5	1e , X = <i>m</i> -OMe		74
6	1f , X = <i>m</i> -Me		83
7	1g , X = <i>m</i> -CN		80
8	1h , X = <i>p</i> -Me		64
9	1i , X = <i>p</i> -Cl		90
10 ^c	1j		86
11	1k , R = Me		71
12	1l , R = Ph		66
13 ^b	1m , R = OMe		63
14	1n		55
15	1o , R = Ph		65
16	1p , R = Me		54

^a All reactions were carried out with olefin (**1**) (0.4 mmol), di-*tert*-butyldiaziridinone (**2**) (0.8 mmol) (added by syringe pump over 8 h), and CuCl–PPh₃ (1:1) (0.02 mmol) in CDCl₃ (0.3 mL) at 65 °C unless otherwise stated. Upon complete addition of **2**, the reaction mixture was stirred at 65 °C for an additional time period (1 h for entries 1, 4, 5, 6, 8, and 9; 2 h for entries 7, 10, and 15; 4 h for entry 11; 7 h for entries 3, 12, 14, and 16; 10 h for entries 2 and 13). ^b CuCl–PPh₃ (1:1) (0.04 mmol) was used. ^c The reaction was carried out with olefin (0.2 mmol), di-*tert*-butyldiaziridinone (**2**) (0.4 mmol), and CuCl–PPh₃ (1:1) (0.02 mmol) in CDCl₃ (0.3 mL). ^d Isolated yield based on olefin.

nated in moderate to good yield (Table 1, entries 1–9).¹⁹ 2-Isopropenylnaphthalene **1j** is also an effective substrate (Table 1, entry 10). Substrates with different alkyl substituents (such as ethyl, benzyl, and methoxymethyl groups) can also be diaminated in 55–71% yield (Table 1, entries 11–14). The diamination process can also be extended to α,β-unsaturated esters (Table 1, entries 15 and 16). However,

(19) **Representative diamination procedure** (Table 1, entry 1): To a 1.5 mL vial equipped with a stir bar were added CuCl (0.002 g, 0.02 mmol), triphenylphosphine (0.0052 g, 0.02 mmol), and CDCl₃ (0.3 mL). After the mixture was stirred at room temperature for 10 min, 2-phenylpropene (**1a**) (0.047 g, 0.4 mmol) was added. The reaction mixture was warmed to 65 °C using an oil bath with stirring, and di-*tert*-butyldiaziridinone (**2**) (0.136 g, 0.8 mmol) was added by syringe pump over 8 h. The reaction mixture was stirred at this temperature for an additional 1 h and purified by flash chromatography (silica gel, hexane/ether = 10/1) to give diamination product **3a** as a white solid (0.105 g, 91%).

(15) (a) Quast, H.; Schmitt, E. *Angew. Chem., Int. Ed.* **1969**, *8*, 448. (b) L'abbé, G.; Verbruggen, A.; Minami, T.; Toppet, S. *J. Org. Chem.* **1981**, *46*, 4478. (c) Mestres, R.; Palomo, C. *Synthesis* **1980**, 755.

(16) Timberlake, J. W.; Alender, J.; Garner, A. W.; Hodges, M. L.; Özmeral, C.; Szilagy, S.; Jacobus, J. O. *J. Org. Chem.* **1981**, *46*, 2082.

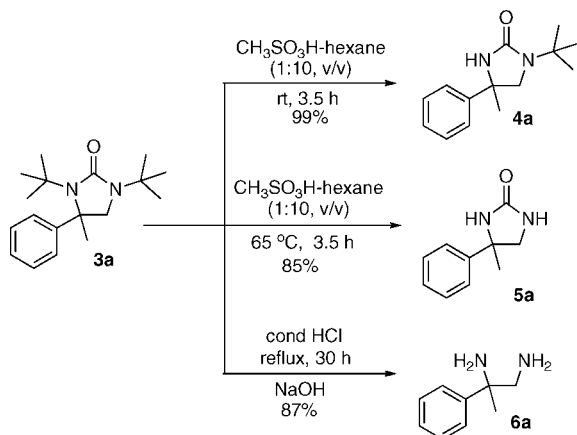
(17) For leading references, see: (a) Reichard, G. A.; Stengone, C.; Paliwal, S.; Mergelsberg, I.; Majmundar, S.; Wang, C.; Tiberi, R.; McPhail, A. T.; Piwinski, J. J.; Shih, N.-Y. *Org. Lett.* **2003**, *23*, 4249. (b) Shue, H.-J.; Chen, X.; Shih, N.-Y.; Blythin, D. J.; Paliwal, S.; Lin, L.; Gu, D.; Schwerdt, J. H.; Shah, S.; Reichard, G. A.; Piwinski, J. J.; Duffy, R. A.; Lachowicz, J. E.; Coffin, V. L.; Liu, F.; Nomeir, A. A.; Morgan, C. A.; Varty, G. B. *Bioorg. Med. Chem. Lett.* **2005**, *15*, 3896. (c) Shue, H.-J.; Chen, X.; Schwerdt, J. H.; Paliwal, S.; Blythin, D. J.; Lin, L.; Gu, D.; Wang, C.; Reichard, G. A.; Wang, H.; Piwinski, J. J.; Duffy, R. A.; Lachowicz, J. E.; Coffin, V. L.; Nomeir, A. A.; Morgan, C. A.; Varty, G. B.; Shih, N.-Y. *Bioorg. Med. Chem. Lett.* **2006**, *16*, 1065.

(18) When the CuCl-catalyzed diamination of 2-phenylpropene was carried out with di-*tert*-butylthiadiaziridine 1,1-dioxide or 1,2-di-*tert*-butyl-3-(cyanimino)diaziridine as the nitrogen source under the previously reported conditions (refs 9 and 12), only a small amount of diamination product or a messy mixture was obtained, respectively.

dialkyl terminal olefins, such as 2-methyl-3-phenyl-1-propene, are not effective substrates for this diamination.

The deprotection of the diamination products was investigated with compound **3a** (Scheme 2). Treating **3a** with

Scheme 2



$\text{CH}_3\text{SO}_3\text{H}$ in hexane (1:10, v/v) at room temperature gave monodeprotected compound **4a** in 99% yield.²⁰ The structure of compound **4a** was confirmed by NOE analysis and the X-ray structure of related compound **14** (Figure 2, Scheme 4).

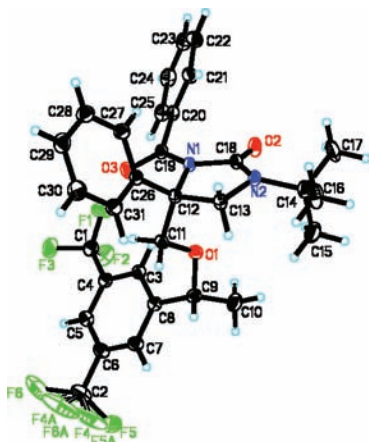
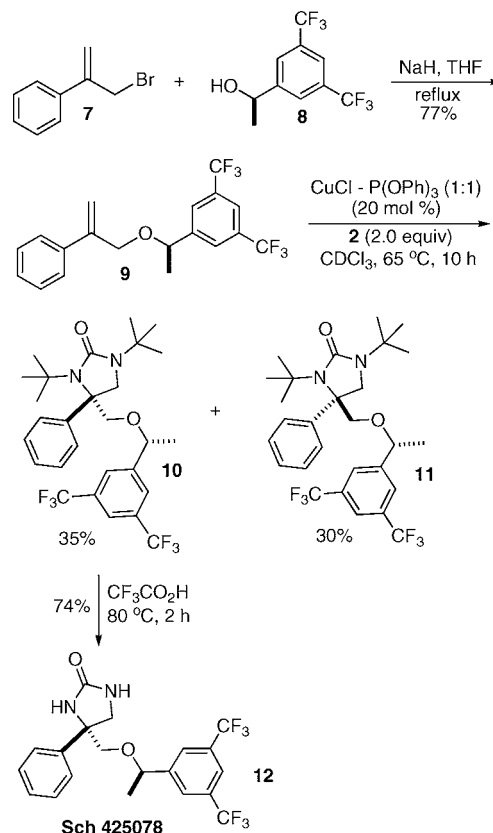


Figure 2. X-ray structure of compound **14**.

When the deprotection was carried out at 65 °C, both *tert*-butyl groups were smoothly removed in 85% yield. Free diamine **6a** can be obtained in 87% yield directly from **3a** by deprotection with concentrated HCl.^{7c,10}

The application of this catalytic diamination to the synthesis of potent NK₁ antagonist 4,4-disubstituted 2-im-

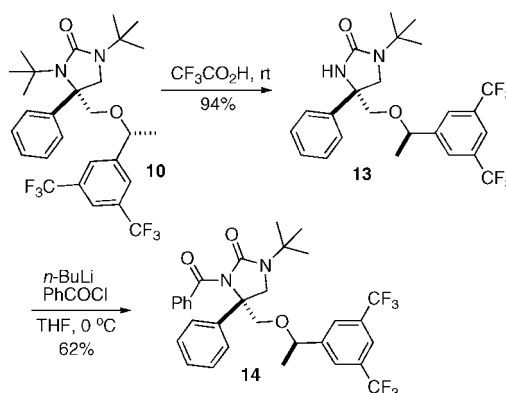
Scheme 3



Sch 425078

dazolidinone **Sch 425078**¹⁷ is outlined in Scheme 3. Disubstituted terminal olefin **9** was readily prepared in 77% yield by reaction between α -bromomethylstyrene (**7**) and commercially available (*R*)-1-(3,5-bis(trifluoromethyl)phenyl)ethanol using NaH as base. Diamination of olefin **9** with $\text{CuCl}-\text{P}(\text{OPh})_3$ and di-*tert*-butyldiaziridinone (**2**) gave 4,4-disubstituted 2-imidazolidinones **10** and **11** in 35% and 30% yield, respectively,²¹ after flash chromatography (the less polar spot on the TLC corresponds to compound **10**). Removal of both *tert*-butyl groups of compound **10** provided compound **12** (**Sch 425078**) in 74% yield.

Scheme 4



(20) Zhao, B.; Du, H.; Shi, Y. *J. Am. Chem. Soc.* **2008**, *130*, 7220.

(21) Diamination of olefin **9** with $\text{CuCl}-\text{PPh}_3$ (1:1) (10 mol %) in CDCl_3 at 65 °C for 10 h gave compound **10** in 30% yield and **11** in 33% yield. Diamination of **9** with $\text{CuCl}-(R)\text{-DTBM-SEGPHOS}$ (2:1) (10 mol %)⁸ in CDCl_3 at 65 °C for 10 h gave compound **10** in 32% yield and **11** in 29% yield.

To further confirm the configuration of the diamination product **10**, one *tert*-butyl group was selectively removed using CF₃CO₂H at room temperature to give compound **13**, which was converted to compound **14** with *n*-BuLi and benzoyl chloride²² (Scheme 4). The structure of compound **14** was determined by X-ray analysis (Figure 2). The determination of the structure of monodeprotected product **13** supports the structure assignment of monodeprotected compound **4a** in Scheme 2.

(22) Rahimizadeh, M.; Kam, K.; Jenkins, S. I.; McDonald, R. S.; Harrison, P. H. M. *Can. J. Chem.* **2002**, *80*, 517.

(23) For recent leading references on other approaches to 4,4-disubstituted-2-imidazolidinones, see: (a) Trost, B. M.; Fandrick, D. R. *J. Am. Chem. Soc.* **2003**, *125*, 11836. (b) Mehrotra, M. M.; Heath, J. A.; Smyth, M. S.; Pandey, A.; Rose, J. W.; Seroogy, J. M.; Volkots, D. L.; Nannizzi-Alaimo, L.; Park, G. L.; Lambing, J. L.; Hollenbach, S. J.; Scarborough, R. M. *J. Med. Chem.* **2004**, *47*, 2037. (c) Shen, D.; Zhang, F.; Brady, E. J.; Candelore, M. R.; Dallas-Yang, Q.; Ding, V. D.-H.; Dragovic, J.; Feeney, W. P.; Jiang, G.; McCann, P. E.; Mock, S.; Qureshi, S. A.; Saperstein, R.; Shen, X.; Tamvakopoulos, C.; Tong, X. C.; Tota, L. M.; Wright, M. J.; Yang, X.; Zheng, S.; Chapman, K. T.; Zhang, B.; Tata, J. R.; Parmee, E. R. *Bioorg. Med. Chem. Lett.* **2005**, *15*, 4564. (d) Brackmann, F.; Es-Sayed, M.; de Meijere, A. *Eur. J. Org. Chem.* **2005**, 2250. (e) Nadir, U. K.; Krishna, R. V.; Singh, A. *Tetrahedron Lett.* **2005**, *46*, 479. (f) Reference 6. (g) Reference 4a. (h) Fritz, J. A.; Nakhla, J. S.; Wolfe, J. P. *Org. Lett.* **2006**, *8*, 2531. (i) Kim, M.; Mulcahy, J. V.; Espino, C. G.; Du Bois, J. *Org. Lett.* **2006**, *8*, 1073. (j) Lanter, J. C.; Fiordeliso, J. J.; Allan, G. F.; Musto, A.; Hahn, D. W.; Sul, Z. *Bioorg. Med. Chem. Lett.* **2006**, *16*, 5646. (k) Reference 4d. (l) Fritz, J. A.; Wolfe, J. P. *Tetrahedron* **2008**, *64*, 6838. (m) Park, H.; Min, K.; Kwak, H.-S.; Koo, K. D.; Lim, D.; Seo, S.-W.; Choi, J.-U.; Platt, B.; Choi, D.-Y. *Bioorg. Med. Chem. Lett.* **2008**, *18*, 2900.

In summary, a variety of disubstituted terminal olefins have been effectively diaminated using CuCl as catalyst and di-*tert*-butyldiaziridinone as nitrogen source, which provides a rapid access to various 4,4-disubstituted 2-imidazolidinones.²³ In addition, the synthesis of 4,4-disubstituted 2-imidazolidinone **Sch 425078** (potent NK₁ antagonist) has been achieved in three steps using this diamination. The ability to selectively remove one or two protecting groups would provide opportunities to introduce different substituents on the nitrogens if desired. Future efforts will be devoted to the development of an asymmetric diamination process and its applications.

Acknowledgment. We are grateful for the generous financial support from the General Medical Sciences of the National Institutes of Health (GM083944-02).

Supporting Information Available: Experimental procedures, characterization of diamination products **3**, NOE studies of compound **4a**, and ¹H and ¹³C NMR spectra of compounds **3**, **4a**, **5a**, **6a**, and **9–14** along with the X-ray data of compound **14**. This material is available free of charge via the Internet at <http://pubs.acs.org>.

OL900808Z