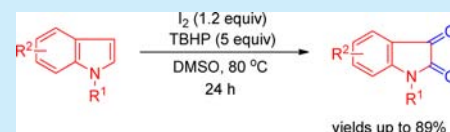


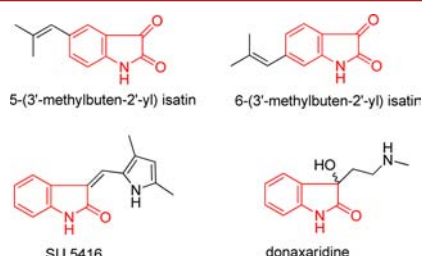
Synthesis of Isatins by I₂/TBHP Mediated Oxidation of IndolesYou Zi,[†] Zhong-Jian Cai,[†] Shun-Yi Wang,* and Shun-Jun Ji*

Key Laboratory of Organic Synthesis of Jiangsu Province, College of Chemistry, Chemical Engineering and Materials Science & Collaborative Innovation Center of Suzhou Nano Science and Technology, Soochow University, Suzhou 215123, China

S Supporting Information

ABSTRACT: An I₂/TBHP mediated oxidation of commercially available indoles has been developed, which affords isatins in moderate to good yields.

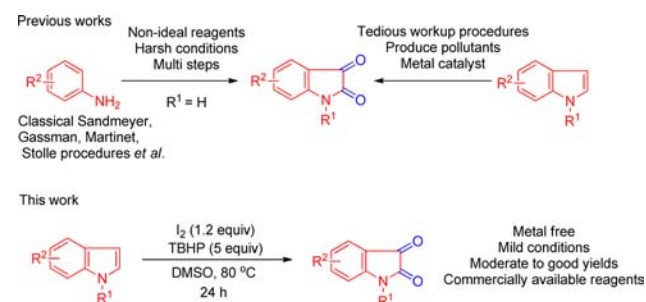
Isatins widely exist in natural products and pharmaceuticals. They are important heterocycles with diverse biological properties such as antimalarial,¹ anticancer,² anticonvulsant,³ antifungal,⁴ and anti-inflammatory agents.⁵ Moreover, they are also important intermediates for the construction of valuable molecules such as the synthesis of SU5416 (VEGFR-2 inhibitor)⁶ and donaxaridine⁷ (Figure 1). Sustained attention has been paid to developing general and direct methods for the preparation of isatins.

**Figure 1.** Pharmacologically active indoline-2,3-diones (isatins) and their derivatives.

Classical methods for the preparation of isatin derivatives include the Sandmeyer,⁸ Gassman,⁹ Martinet,¹⁰ and Stollé¹¹ procedures. However, they typically suffer from harsh conditions or reagents, elevated temperatures, and nonideal solvents. Recently, several improved protocols for the construction of isatins have been reported such as aryne-based methods,¹² Sandmeyer modifications,¹³ metal catalyzed oxidations,¹⁴ sulfur ylide mediated carbonyl homologation,¹⁵ and C–H amination.¹⁶ Although few reports are available for the direct synthesis of isatins from commercially available indoles,¹⁷ a mild and metal-free strategy for the direct synthesis of isatins from indoles is still desirable. More recently, we have reported an I₂/TBHP mediated amination of indoles with anilines.¹⁸ In our continuous efforts to construct heterocycles under oxidative conditions,¹⁹ herein, we demonstrate an I₂/TBHP mediated oxidation of commercially available indoles to give isatins in moderate to good yields (Scheme 1).

Our studies were initiated by the reaction of *N*-methylindole 1a in DMSO at 80 °C in the presence of 1.2 equiv of I₂ and 2 equiv of TBHP. The desired product *N*-methyl isatin 2a could

Scheme 1. Preparation of Isatins



be obtained in 38% LC-yield (Table 1, entry 1). To our surprise, the yield of 2a could be increased to 83% LC-yield (82% isolated yield) (Table 1, entry 2). When the reaction was mediated by other iodine sources such as KI and TBAI instead of I₂, the yield of 2a was decreased (Table 1, entries 3 and 4). Further screenings of the oxidant and reaction temperature (more details see Supporting Information) revealed no better results (Table 1, entries 5 to 12).

With the optimal reaction conditions in hand, we investigated the scope of the reaction utilizing indoles. As shown in Figure 2, all the reactions proceeded well to furnish the desired isatins. Isatin 2b could be obtained in 65% yield by direct oxidation of indole 1b under the optimized conditions. It should be noted that the reaction of 4-methylindole 1c could lead to 2c in 89% yield. The reactions of 5-methylindole 1d, 6-methylindole 1e, and 7-methylindole 1f could also afford the desired isatins 2d, 2e, and 2f in 64%, 87%, and 73% yields, respectively. The reactions of indole bearing electron-donating groups such as 4-(benzyloxy)-1*H*-indole 1g and 5-methoxyindole 1h proceeded well to give 2g and 2h in 57% and 59% yields, respectively. When an indole bearing electron-withdrawing group such as 5-bromo-1*H*-indole 1i was subjected to the reaction, the yield of the corresponding product 2i is decreased to 11% for its lower reactivity. Notably, *N*-methylindoles bearing either electron-donating or -withdrawing groups increased the yields of the desired products significantly.

Received: April 27, 2014

Published: May 21, 2014

Table 1. Reaction Optimization^a

entry	additive (equiv)	oxidant (equiv)	yield (%) ^b
1	I ₂ (1.2)	TBHP (2)	38
2	I ₂ (1.2)	TBHP (5)	83 (82 ^c)
3	KI (1.2)	TBHP (5)	14
4	TBAI (1.2)	TBHP (5)	12
5	I ₂ (0.1)	TBHP (5)	trace
6	I ₂ (1.5)	TBHP (5)	65
7	I ₂ (1.2)	DTPB (5)	9
8	I ₂ (1.2)	TBPB (5)	20
9	I ₂ (1.2)	CPO (5)	20
10	I ₂ (1.2)	PAA ^d (5)	11
11	I ₂ (1.2)	TBHP (5)	57 ^e
12	I ₂ (1.2)	TBHP (5)	67 ^f

^aReaction conditions: **1a** (0.5 mmol), I₂ (0.6 mmol), TBHP (2.5 mmol) in DMSO (6 mL), 80 °C, 24 h. ^bDetermined by LC-MS analysis with biphenyl as the internal standard. ^cIsolated yield. ^dPAA = peracetic acid. ^eThis reaction proceeded at room temperature. ^fThis reaction proceeded at 110 °C.

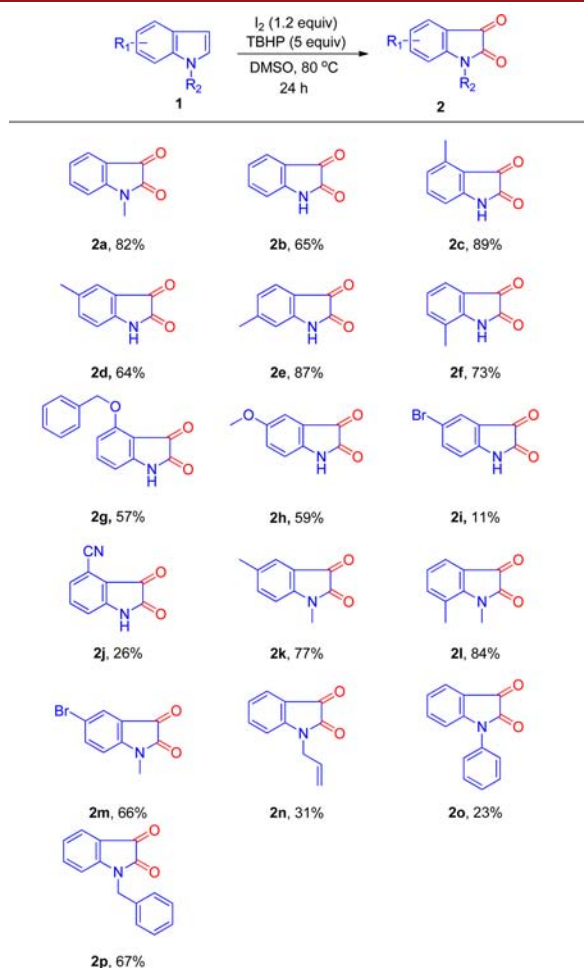
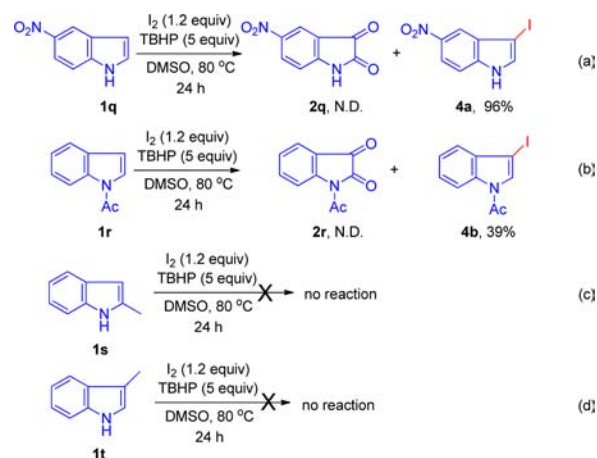


Figure 2. Substrate scope. Reaction conditions: **1** (0.5 mmol), I₂ (0.6 mmol), and TBHP (2.5 mmol) in DMSO (6 mL), 80 °C, 24 h. Isolated yields are shown.

The reaction of 1,7-dimethyl-1H-indole **1l** and 5-bromo-1-methyl-1H-indole **1m** lead to the desired isatins **2l** and **2m** in 84% and 66% yields, respectively. In addition, the reaction of 1-benzyl-1H-indole **1p** also proceeded well and could furnish the desired product **2p** in 67% yield. However, 1-allylindoline-2,3-dione **2n** and 1-phenylindoline-2,3-dione **2o** could only be obtained in 31% and 23% yields by oxidation of the corresponding indoles **1n** and **1o**, respectively.

When 5-nitroindole **1q** was subjected to the reaction under the optimal conditions, 3-iodo-5-nitroindole **4a** was obtained instead of 5-nitroisatin **2q** (Scheme 2, reaction a). Despite this,

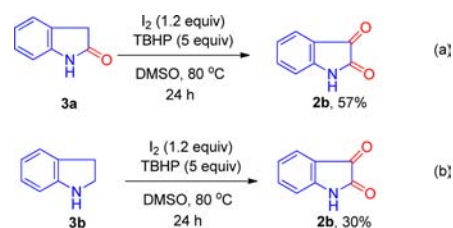
Scheme 2. Oxidation Reaction of Other Indoles



the 5-nitroisatin **2q** could be synthesized in excellent yield via a simple nitration of isatin **2b**.²⁰ However, the reaction of 1-acetylindole **1r** led to 1-acetyl-3-iodoindole **4b** in 39% yield instead of the isatin derivative **2r** under the identical conditions (Scheme 2, reaction b).

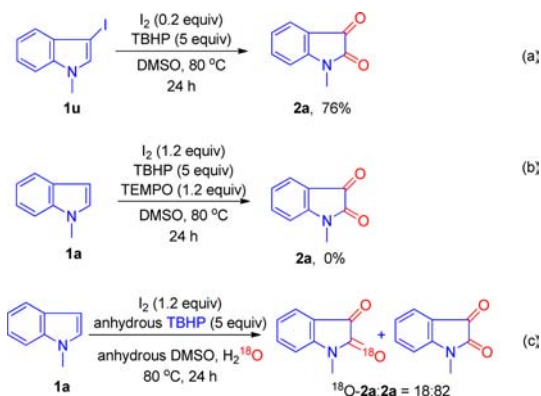
As expected, no desired product was obtained when 2-methyl-indole **1s** and 3-methyl-indole **1t** were subjected to the reaction under the optimized conditions (Scheme 2, reactions c and d).

To our delight, 2-indolone **3a** and indoline **3b** could also be oxidized under identical conditions to give the isatin **2b** in 57% and 30% yields, respectively (Scheme 3).

Scheme 3. Oxidation Reactions of 2-Indolone **3a** and Indoline **3b**

In order to better understand the mechanism of this reaction, we attempted several control experiments. When 3-iodo-1-methyl-indole **1u** was subjected to the reaction under similar conditions, **2a** could be obtained in 76% yield (Scheme 4, reaction a). This result indicated that 3-iodo-1-methyl-indole **1u** was the possible intermediate or its corresponding tautomer of the model reaction. As anticipated, when TEMPO (1.2 equiv) was added to the system, the reaction was completely

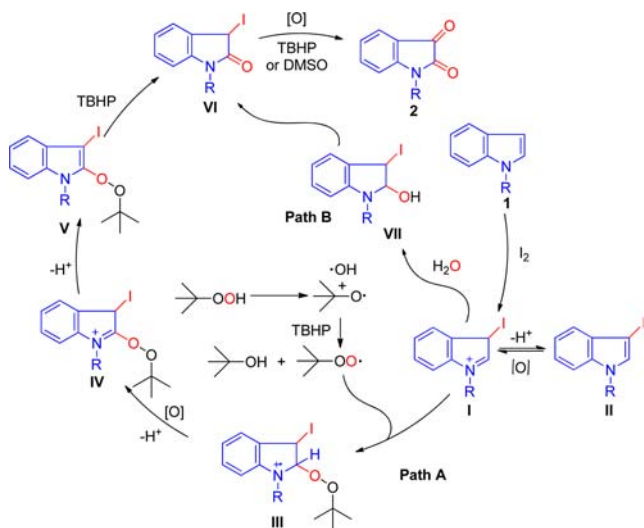
Scheme 4. Investigation into the Reaction Mechanism



suppressed (Scheme 4, reaction b), which indicated that the reaction involved radical processes. When the reaction was carried out with anhydrous DMSO and TBHP in the presence of H_2^{18}O , **2a** and ^{18}O labeled **2a** with the ratio of 82:18 could be obtained (Scheme 4, reaction c). This result meant that one of the oxygen atoms of the isatin comes from the water or TBHP.

Based on the above results and the literature,^{21–23} a plausible mechanism for this reaction via two possible pathways is proposed in Scheme 5. First, indole reacts with I_2 to give the

Scheme 5. Plausible Reaction Mechanism



iminium intermediate **I**. Then, **I** reacts with a *tert*-butylperoxy free radical^{22,22b} to furnish intermediate **III**,^{22,22c,23} which can be oxidized by TBHP to give intermediate **IV**. After the isomerization of **IV** and oxidation by TBHP, 3-iodo-indolin-2-one **VI** can be generated (path A). On the other hand, water attacks the intermediate **I** to give intermediate **VII**. After simple oxidation of **VII** by TBHP, **VI** can also be performed (path B). After further oxidation of **VI** by TBHP or DMSO, isatin **2** is formed.^{21,21e,f}

In conclusion, we have developed a convenient, simple, and metal-free protocol for the direct synthesis of isatins in moderate to good yields by the direct oxidation of commercially available indoles under mild conditions.

■ ASSOCIATED CONTENT

§ Supporting Information

Experimental procedures and full spectroscopic data for all new compounds. This material is available free of charge via the Internet at <http://pubs.acs.org>.

■ AUTHOR INFORMATION

Corresponding Authors

*E-mail: shunyi@suda.edu.cn.

*E-mail: shunjun@suda.edu.cn.

Author Contributions

[†]Y.Z. and Z.-J.C. contributed equally.

Notes

The authors declare no competing financial interest.

■ ACKNOWLEDGMENTS

We gratefully acknowledge the Natural Science Foundation of China (Nos. 21172162, 21372174), the Ph.D. Programs Foundation of Ministry of Education of China (2013201130004), the Young National Natural Science Foundation of China (No. 21202111), the Young Natural Science Foundation of Jiangsu Province (BK2012174), Key Laboratory of Organic Synthesis of Jiangsu Province (KJS1211), PAPD, the Project of Scientific and Technologic Infrastructure of Suzhou (SZS201207), and Soochow University for financial support.

■ REFERENCES

- (1) Rottmann, M.; McNamara, C.; Yeung, B. K. S.; Lee, M. C. S.; Zou, B.; Russell, B.; Seitz, P.; Pluoffe, D. M.; Dharia, N. V.; Tan, J.; Cohen, S. B.; Spencer, K. R.; Gonzalez-Paez, G. E.; Lakshminarayana, S. B.; Goh, A.; Suwanarusk, R.; Jegla, T.; Schmitt, E. K.; Beck, H. P.; Brun, R.; Nosten, F.; Renia, L.; Dartois, V.; Keller, T. H.; Fidock, D. A.; Winzeler, E. A.; Diagana, T. T. *Science* **2010**, 329, 1175.
- (2) Vine, K. L.; Locke, J. M.; Ranson, M.; Pyne, S. G.; Bremner, J. B. *J. Med. Chem.* **2007**, 50, 5109.
- (3) (a) Verma, M.; Pandeya, S. N.; Singh, K. N.; Stables, J. P. *Acta Pharm.* **2004**, 54, 49. (b) Bhattacharya, S. K.; Chakrabarti, A. *Indian J. Exp. Biol.* **1998**, 36, 118. (c) Pajouhesh, H.; Parson, R.; Popp, F. D. *J. Pharm. Sci.* **1983**, 72, 318.
- (4) Zahid, H. C.; Humayun, P.; Rauf, A.; Khalid, M. K.; Claudiu, T. S. *J. Enzyme Inhib. Med. Chem.* **2004**, 19, 417.
- (5) Prakash, C. R.; Raja, S.; Saravanan, G. *Int. J. Pharm. Sci. Biotechnol.* **2010**, 1, 105.
- (6) Sassatelli, M.; Debiton, E.; Aboab, B.; Prudhomme, M.; Moreau, P. *Eur. J. Med. Chem.* **2006**, 41, 709.
- (7) Kawasaki, T.; Nagaoka, M.; Satoh, T.; Okamoto, A.; Ukon, T. R.; Ogawa, A. *Tetrahedron* **2004**, 60, 3493.
- (8) (a) Sandmeyer, T. *Helv. Chim. Acta* **1919**, 2, 234. (b) Marvel, C. S.; Hiers, G. S. *Org. Synth.* **1941**, 1, 327. (c) Garden, S. J.; Torres, J. C.; Ferreira, A. A.; Silva, R. B.; Pinto, A. C. *Tetrahedron Lett.* **1997**, 38, 1501. (d) Rewcastle, G. W.; Sutherland, H. S.; Weir, C. A.; Blackburn, A. G.; Denny, W. A. *Tetrahedron Lett.* **2005**, 46, 8719. (e) Zhao, P. B.; Zhao, B. B.; Li, Y.; Gao, W. T. *Chin. J. Org. Chem.* **2014**, 34, 126.
- (9) Gassman, P. G.; Cue, B. W., Jr.; Luh, T. Y. *J. Org. Chem.* **1977**, 42, 1344.
- (10) (a) Guyot, A.; Martinet, J. *Compt. Rend.* **1913**, 166, 1625. (b) Bonnefoy, J.; Martinet, J. *Compt. Rend.* **1921**, 172, 220. (c) Rice, K. C.; Boone, B. J.; Rubin, A. B.; Rauls, T. J. *J. Med. Chem.* **1976**, 19, 887. (d) Hewawasam, P.; Meanwell, N. A. *Tetrahedron Lett.* **1994**, 35, 7303.
- (11) (a) Stollé, R. *J. Prakt. Chem.* **1923**, 105, 137. (b) Stollé, R. *Ber. Dtsch. Chem. Ges.* **1913**, 46, 3915. (c) Stollé, R.; Bergdoll, R.; Luther, M.; Auerhahn, A.; Wacker, W. *J. Prakt. Chem.* **1930**, 128, 1. (d) Welstead, W. J.; Moran, H. W.; Stauffer, H. F.; Turnbull, L. B.;

- Sancilio, L. F. *J. Med. Chem.* **1979**, *22*, 1074. (e) Bryant, W. M.; Huhn, G. F.; Jensen, J. H.; Pierce, M. E.; Stammbach, C. *Synth. Commun.* **1993**, *23*, 1617.
- (12) Rogness, D. C.; Larock, R. C. *J. Org. Chem.* **2011**, *76*, 4980.
- (13) (a) Klein, L. L.; Tufano, M. D. *Tetrahedron Lett.* **2013**, *54*, 1008. (b) Chouhan, M.; Senwar, K. R.; Sharma, R.; Grover, V.; Nair, V. A. *Green Chem.* **2011**, *13*, 2553.
- (14) (a) Tang, B. X.; Song, R. J.; Wu, C. Y.; Liu, Y.; Zhou, M. B.; Wei, W. T.; Deng, G. B.; Yin, D. L.; Li, J. H. *J. Am. Chem. Soc.* **2010**, *132*, 8900. (b) Sun, J.; Liu, B.; Xu, B. *RSC Adv.* **2013**, *3*, 5824. (c) Liu, T.; Yang, H.; Jiang, Y.; Fu, H. *Adv. Synth. Catal.* **2013**, *355*, 1169. (d) Liu, Y.; Chen, H.; Hu, X.; Zhou, W.; Deng, G. J. *Eur. J. Org. Chem.* **2013**, 4229.
- (15) Lollar, C. T.; Krennek, K. M.; Bruemmer, K. J.; Lippert, A. R. *Org. Biomol. Chem.* **2014**, *12*, 406.
- (16) Huang, P. C.; Gandeepan, P.; Cheng, C. H. *Chem. Commun.* **2013**, *49*, 8540.
- (17) (a) Chen, S.; Liu, Z.; Shi, E.; Chen, L.; Wei, W.; Li, H.; Cheng, Y.; Wan, X. *Org. Lett.* **2011**, *13*, 2274. (b) Yadav, J. S.; Reddy, B. V. S.; Reddy, C. S.; Krishna, A. D. *Synthesis* **2007**, 693. (c) Sriram, R.; Kumar, C. N. S. S. P.; Raghunandan, N.; Ramesh, V.; Sarangapani, M.; Rao, V. J. *Synth. Commun.* **2012**, *42*, 3419. (d) Kumar, P.; Devi, C. L.; Rao, V. J.; Palaniappan, S. *Synlett* **2008**, *13*, 2023. (e) Wei, W.; Shao, Y.; Hu, H.; Zhang, F.; Zhang, C.; Xu, Y.; Wan, X. *J. Org. Chem.* **2012**, *77*, 7157. (f) Sonawane, R. P.; Tripathi, R. R. *ILCPA* **2013**, *7*, 30. (g) Tatsugi, J.; Tong, Z. W.; Izawa, Y. *ARKIVOC* **2001**, 67. (h) Parrick, J.; Yahya, A.; Jin, Y. Z. *J. Chem. Soc., Perkin Trans.* **1989**, 2009. (i) Liu, X. L.; Chen, W. Z. *Organometallics* **2012**, *31*, 6614.
- (18) Cai, Z.-J.; Wang, S.-Y.; Ji, S.-J. *Org. Lett.* **2013**, *15*, 5226.
- (19) (a) Cai, Z.-J.; Wang, S.-Y.; Ji, S.-J. *Org. Lett.* **2012**, *14*, 6068. (b) Cai, Z.-J.; Wang, S.-Y.; Ji, S.-J. *Adv. Synth. Catal.* **2013**, *355*, 2686. (c) Zhu, T.-H.; Wang, S.-Y.; Wang, G.-N.; Ji, S.-J. *Chem.—Eur. J.* **2013**, *19*, 5850. (d) Hao, W.-J.; Wang, S.-Y.; Ji, S.-J. *ACS Catal.* **2013**, *3*, 2501. (e) Hao, W.-J.; Xu, X.-P.; Wang, S.-Y.; Ji, S.-J. *J. Org. Chem.* **2013**, *78*, 12362.
- (20) Magiatis, P.; Polychronopoulos, P.; Skaltsounis, A.-L.; Lozach, O.; Meijer, L.; Miller, D. B.; O'Callaghan, J. P. *Neurotoxicol. Teratol.* **2010**, *32*, 212.
- (21) (a) Zhang, C.; Xu, Z.; Zhang, L.; Jiao, N. *Angew. Chem., Int. Ed.* **2011**, *50*, 11088. (b) Zhang, C.; Jiao, N. *J. Am. Chem. Soc.* **2010**, *132*, 28. (c) Zhang, X.; Wang, L. *Green Chem.* **2012**, *14*, 2141. (d) Moorthy, J. N.; Senapati, K.; Singhal, N. *Tetrahedron Lett.* **2009**, *50*, 2493. (e) Gao, M.; Yang, Y.; Wu, Y.-D.; Deng, C.; Shu, W.-M.; Zhang, D.-X.; Cao, L.-P.; She, N.-F.; Wu, A.-X. *Org. Lett.* **2010**, *12*, 4026. (f) Zhu, Y. P.; Liu, M. C.; Jia, F. C.; Yuan, J. J.; Gao, Q. H.; Lian, M.; Wu, A. X. *Org. Lett.* **2012**, *14*, 3392. (g) Gao, M.; Yang, Y.; Wu, Y.-D.; Deng, C.; Cao, L.-P.; Meng, X.-G.; Wu, A.-X. *Org. Lett.* **2010**, *12*, 1856. (h) Yadav, J. S.; Reddy, B. V. S.; Singh, A. P.; Basak, A. K. *Tetrahedron Lett.* **2008**, *49*, 5880.
- (22) (a) He, Z.; Liu, W.; Li, Z. *Chem.—Asian J.* **2011**, *6*, 1340. (b) Boess, E.; Schmitz, C.; Klusmann, M. *J. Am. Chem. Soc.* **2012**, *134*, 5317. (c) Zhang, J.; Wang, Z.; Wang, Y.; Wan, C.; Zheng, X.; Wang, Z. *Green Chem.* **2009**, *11*, 1973. (d) Tasch, B. O. A.; Bensch, L.; Antovic, D.; Müller, T. J. J. *Org. Biomol. Chem.* **2013**, *11*, 6113.
- (23) Wu, W. B.; Huang, J. M. *Org. Lett.* **2012**, *14*, 5832.