A quadruple cascade protocol for the one-pot synthesis of fully-substituted hexahydroisoindolinones from simple substrates

Hong-Bo Zhang, Yong-Chun Luo, Xiu-Qin Hu, Yong-Min Liang and Peng-Fei Xu*

Abstract
A new and efficient synthetic method to obtain fully-substituted hexahydroisoindolinones was developed by using bifunctional tertiary amine-thioureas as powerful catalysts. As far as we know, there is no efficient synthetic method developed toward fully-substituted hexahydroisoindolinones. The products were obtained in good yield and diastereoselectivity. The one-pot cascade quadruple protocol features readily available starting materials, simple manipulation, mild conditions and good atom economy.

Introduction
Isoindolines and their congeners are one kind of the most widespread compounds in nature. They feature not only high biological activity, but also diverse chemical properties [1-16]. Therefore, it is highly desirable to develop efficient methods toward the synthesis of isoindoline derivatives, which is a frontier in organic synthesis.

However, compared with the synthesis of their congeners, the synthesis of fully-substituted hexahydroisoindolinones is much more difficult due to the steric hindrance and the high strain of the molecular architectures [17]. Three methods to synthesize 3-substituted isoindolinones have been developed. The first method was the synthesis of 3-substituted isoindolinones from the corresponding N-methylmaleimides by the Diels–Alder reaction with 1,3-butadiene followed by hydrogenation. The second and the third methods employed the corresponding dicarboxylic acids and the carboxylic acid anhydrides, respectively [17]. To the best of our knowledge, no efficient method toward the synthesis of fully-substituted hexahydroisoindolinones has been developed so far.
The synthesis of complicated molecular structures can now be achieved by organocatalytic cascade reactions [18-33]. By simplifying the experimental procedures and reducing the usage of both solvents and reagents, one-pot reactions can improve the synthesis efficiency and both save time and reduce cost [34]. Although a few types of complicated molecules were generated through multicomponent quadruple cascade reactions, there is no report about the cascade synthesis of isoidolines in the past few decades [35-46], not mention the quadruple cascade synthesis of difficult fully-substituted hexahydroisoindolinones. Previously, we established organocatalytic domino reactions to construct very useful molecular architectures [47-60]. Based on this past experience, we decided to develop a one-pot quadruple protocol to construct this difficult molecular architecture using easily accessible substrates.

Results and Discussion

We initiated this study by using 2-benzylidenemalononitrile (1a) and 2-oxo-N,3-diphenylpropanamide (2a) [61-64] in 0.5 mL of CH₃CN in the presence of 10 mol % of DABCO. After 12 h at room temperature, the reaction afforded the expected product rac-3a in 59% yield (Table 1, entry 1). We then tested different catalysts to optimize the reaction. When Et₃N was used, the reaction afforded the product with 41% yield (Table 1, entry 2). However, a complex mixture was observed when DBU was used (Table 1, entry 3), while no reaction was observed when K₂CO₃ was used as the catalyst (Table 1, entry 4). When thioureas were used as the catalysts, we also did not get the expected product (Table 1, entries 5 and 6). Since bifunctional tertiary amine-thioureas have been proved as powerful catalysts that can catalyze a variety of organocascade

---

**Table 1: Screening the reaction conditions.**

| entry | cat. | solvent | dr | yield [%] |
|-------|------|---------|----|-----------|
| 1     | DABCO | CH₃CN   | 4:1 | 59        |
| 2     | Et₃N  | CH₃CN   | 4:1 | 41        |
| 3     | DBU   | CH₃CN   | n.d. | complex  |
| 4     | K₂CO₃ | CH₃CN   | n.d. | n.r.      |
| 5     | cat-1 | CH₃CN   | n.d. | n.r.      |
| 6     | cat-2 | CH₃CN   | n.d. | n.r.      |
| 7     | DABCOᵈ | CH₃CN   | 4:1 | 62        |
| 8     | Et₃Nᵈ | CH₃CN   | 5:1 | 52        |
| 9     | cat-3 | CH₃CN   | 9:1 | 87        |
| 10    | cat-3 | DCM     | 4:1 | 33        |
| 11    | cat-3 | THF     | 4:1 | 34        |
| 12    | cat-3 | toluene | n.d. | trace    |
| 13    | cat-3 | CH₃OH   | n.d. | trace    |
| 14ʰ   | cat-3 | CH₃CN   | 6:1 | 87        |

ᵃUnless otherwise noted, the reactions were carried out with 1a (0.25 mmol, 38.5 mg), 2a (0.1 mmol, 23.9 mg), catalyst (0.01 mmol, 10 mol %) in the indicated solvent (0.5 mL) at rt for 12 h.ᵇDetermined by ¹H NMR analysis of the crude products.ᶜColumn chromatography yields.ᵈ10 mol % cat-2 was added.ʰThe reaction was carried out at 35 °C.

---

*Beilstein J. Org. Chem. 2016, 12, 253–259.*
reactions, we also tested thiourea catalysts, cat-1 to cat-3. Interestingly, the thioureas cat-1 and cat-2 were able to promote the reaction (Table 1, entries 7 and 8), but we obtained an even better yield when the tertiary amine-thiourea cat-3 was used as the catalyst (Table 1, entry 9). All products were racemic even when chiral catalysts were used (see Supporting Information File 1 for details). Next, we performed a solvent screening. As shown in Table 1, when DCM and THF were used as the solvent, the yield of the desired product was 33% and 34%, respectively (Table 1, entries 10 and 11). Only traces of the product were seen when toluene or methanol was used as the solvent (Table 1, entries 12 and 13). Furthermore, raising the reaction temperature was not beneficial for the diastereoselectivity of the reaction (Table 1, entry 14).

With the optimal conditions in hand, we next examined the reaction scope (Table 2). All reactions afforded the corresponding products 3a–t with medium to good yield and diastereoselectivity using the simple protocol at room temperature. To our delight, with our optimized reaction system, various types of substrates 1 showed very good reaction activities. Different types of substrates 1, bearing either electron withdrawing or donating groups in para-, meta- and ortho-positions, gave the desired products in good yield and diastereoselectivity (Table 2, entries 12 and 13).

Table 2: Substrates scope.

| entry | R¹ | R² | R³ | dr | yield [%] |
|-------|----|----|----|----|-----------|
| 1     | C₂H₅ | C₂H₅ | C₂H₅ | 9:1 | 87 (3a)   |
| 2     | 2-MeC₆H₄ | C₂H₅ | C₂H₅ | >20:1 | 89 (3b) |
| 3     | 3-MeC₆H₄ | C₂H₅ | C₂H₅ | 10:1 | 69 (3c) |
| 4     | 4-OMeC₆H₄ | C₂H₅ | C₂H₅ | 10:1 | 66 (3d) |
| 5     | 2-BrC₆H₄ | C₂H₅ | C₂H₅ | >20:1 | 84 (3e) |
| 6     | 3-ClC₆H₄ | C₂H₅ | C₂H₅ | 4:1  | 72 (3f) |
| 7     | 4-FC₆H₄ | C₂H₅ | C₂H₅ | >20:1 | 82 (3g) |
| 8     | 4-CF₃C₆H₄ | C₂H₅ | C₂H₅ | >20:1 | 86 (3h) |
| 9     | 2-NO₂C₆H₄ | C₂H₅ | C₂H₅ | >20:1 | 89 (3i) |
| 10    | 3-NO₂C₆H₄ | C₂H₅ | C₂H₅ | >20:1 | 91 (3j) |
| 11    | 4-NO₂C₆H₄ | C₂H₅ | C₂H₅ | 3:1  | 42 (3k) |
| 12    | 2-naphthalene | C₂H₅ | C₂H₅ | >20:1 | 90 (3l) |
| 13    | 2-thiophene | C₂H₅ | C₂H₅ | 3:1  | 51 (3m) |
| 14    | 3,4-diClC₆H₃ | C₂H₅ | C₂H₅ | >20:1 | 84 (3n) |
| 15    | 3,5-diOMeC₆H₃ | C₂H₅ | C₂H₅ | 15:1 | 55 (3o) |
| 16    | C₆H₅ | 4-OMeC₆H₄ | C₂H₅ | 4:1  | 56 (3p) |
| 17    | C₆H₅ | 4-ClC₆H₄ | C₂H₅ | >20:1 | 89 (3q) |
| 18    | 2-naphthalene | 4-OMeC₆H₄ | C₂H₅ | 14:1 | 88 (3r) |
| 19    | C₆H₅ | C₂H₅ | 4-MeC₆H₄ | 8:1 | 61 (3s) |
| 20    | C₆H₅ | C₂H₅ | 4-FC₆H₄ | 8:1 | 61 (3t) |
| 21    | C₂H₅(CH₂)₂ | C₂H₅ | C₂H₅ | n.d. | n.r. |
| 22    | CH₃(CH₂)₆ | C₂H₅ | C₂H₅ | n.d. | n.r. |
| 23    | C₂H₅ | CH₃(CH₂)₃ | C₂H₅ | n.d. | n.r. |
| 24    | C₂H₅ | CH₃CH₂ | C₂H₅ | n.d. | n.r. |
| 25    | C₂H₅ | C₂H₅ | H | n.d. | n.r. |
| 26    | C₂H₅ | C₂H₅ | CH₃ | n.d. | n.r. |

*Unless otherwise noted, the reactions were carried out with 1 (0.25 mmol), 2 (0.1 mmol), cat-3 (3.6 mg, 0.01 mmol, 10 mol %) in CH₃CN (0.5 mL) at rt for 12 h. †Determined by ¹H NMR analysis of the crude products. ‡Column chromatography yields.
entries 1–10 and 12), although 4-NO\textsubscript{2}C\textsubscript{6}H\textsubscript{4} gave the product in medium yield due to its poor solubility (Table 2, entry 11). A heteroaromatic substrate such as thiophene could also be successfully employed to afford rac-3 with medium yield and diastereoselectivity (Table 2, entry 13). 3,4-Dichloro-substituted and 3,5-dimethoxy-substituted substrates produced the desired products in 84% and 55% yield with 20:1 and 15:1 diastereoselectivity, respectively (Table 2, entries 14 and 15). When substrates with different R\textsuperscript{2} and R\textsuperscript{3} were used in this reaction, the corresponding products were obtained in medium yield and diastereoselectivity (Table 2, entries 16–20). The structure of 3p was determined by X-ray analysis [65]. However, substrates with aliphatic R\textsuperscript{1}, R\textsuperscript{2} or R\textsuperscript{3} did not produce the desired products (Table 2, entries 21–26).

This bifunctional catalysis cascade reaction was also amenable to scale-up. When the reaction was carried out on a 3 mmol scale, the desired product was obtained in 84% yield. Therefore, this method is fast and easy to implement, and it is suitable for large-scale synthesis (Scheme 1).

Many isoindolinone skeletons show high biological potential as antihypertensives, anesthetics, etc. [66-68]. The useful hydrolyzed product rac-4a was obtained in 80% yield by treating rac-3a with trifluoroacetic anhydride in DCM (Scheme 2).

Finally, we propose a mechanism for the reaction. Initially, substrate 1 is activated by catalyst (I), which reacts with substrate 2 via two Michael addition reactions to sequentially produce II and III. Then, IV is generated from III by an aldol reaction. Finally, the product is produced after the nucleophilic reaction, and the catalyst is regenerated (Scheme 3).

### Conclusion

In summary, we have developed a one-pot quadruple cascade protocol to obtain fully-substituted hexahydroisoindolinones. This new, synthetic method is simple, efficient and atom-economic. This reaction can be widely used in organic synthesis due to its advantages such as simple operation, availability of raw materials, mild conditions and high efficiency.

### Experimental

#### General procedure for the synthesis of fully-substituted hexahydroisoindolinones

Benzylidenemalononitrile (0.1 mmol), 2-oxo-N,3-diphenyl-propanamide (0.25 mmol) and cat-3 (0.01 mmol) were added to a test tube, then CH\textsubscript{3}CN (0.5 mL) was added to the mixture. The reaction mixture was stirred at 300 rpm at 21 °C in a stoppered carousel tube for 12 h. The solvent was removed in vacuo and the product was purified by silica gel flash column chromatography to give the corresponding product 3.

![Scheme 1: An example of scalable synthesis.](image1)

![Scheme 2: Hydrolysis reaction to produce a useful product.](image2)
Supporting Information

Supporting Information File 1
Experimental procedures, characterization data for all new compounds and X-ray analysis of compound 3.
[http://www.beilstein-journals.org/bjoc/content/supplementary/1860-5397-12-27-S1.pdf]

Acknowledgements
We are grateful to the NSFC (21172097, 21202070, 21302075 and 21372105), the International S&T Cooperation Program of China (2013DFR70580), the National Natural Science Foundation from Gansu Province of China (no. 1204WCGA015), and the “111” program from MOE of P. R. China.

References
1. Jiaang, W.-T.; Chen, Y.-S.; Hsu, T.; Wu, S.-H.; Chien, C.-H.; Chang, C.-N.; Chang, S.-P.; Lee, S.-J.; Chen, X. Bioorg. Med. Chem. Lett. 2005, 15, 687–691. doi:10.1016/j.bmcl.2004.11.023
2. Van Goethem, S.; Van der Veken, P.; Dubois, V.; Soroka, A.; Lambeir, A.-M.; Chen, X.; Haemers, A.; Scharpé, S.; De Meester, I.; Augustyns, K. Bioorg. Med. Chem. Lett. 2008, 18, 4159–4162. doi:10.1016/j.bmcl.2008.05.079
3. Portevin, B.; Tordjman, C.; Pastoureau, P.; Bonnet, J.; De Nanteuil, G. J. Med. Chem. 2000, 43, 4582–4593. doi:10.1021/jm990965x
4. Mancilla, T.; Correa-Basurto, J.; Alavés Carbajal, K. S.; Sanchez Escalante, E. J. T.; Ferrara, J. T. J. Mex. Chem. Soc. 2007, 51, 96–102.
5. Kukkola, P. J.; Bilci, N. A.; Ikeler, T. J. Tetrahedron Lett. 1996, 37, 5065–5068. doi:10.1016/0040-4039(96)01018-0
55. Wang, Y.; Luo, Y.-C.; Hu, X.-Q.; Xu, P.-F. Org. Lett. 2011, 13, 5346–5349. doi:10.1021/ol2022092
56. Zhao, Y.-L.; Wang, Y.; Cao, J.; Liang, Y.-M.; Xu, P.-F. Org. Lett. 2014, 16, 2438–2441. doi:10.1021/ol5008185
57. Zhao, S.; Lin, J.-B.; Zhao, Y.-Y.; Liang, Y.-M.; Xu, P.-F. Org. Lett. 2014, 16, 1802–1805. doi:10.1021/ol500547e
58. Lu, H.; Lin, J.-B.; Liu, J.-Y.; Xu, P.-F. Chem. – Eur. J. 2014, 20, 11659–11663. doi:10.1002/chem.201402947
59. Gao, T.-P.; Lin, J.-B.; Hu, X.-Q.; Xu, P.-F. Chem. Commun. 2014, 50, 8934–8936. doi:10.1039/C4CC03896G
60. Wang, Y.; Lu, H.; Xu, P.-F. Acc. Chem. Res. 2015, 48, 1832–1844. doi:10.1021/acs.accounts.5b00217
61. Goudedranche, S.; Pierrot, D.; Constantieux, T.; Bonne, D.; Rodriguez, J. Chem. Commun. 2014, 50, 15605–15608. doi:10.1039/C4CC07731H
62. Wang, L.; Ni, Q.; Bürmel, M.; Shu, T.; Raabe, G.; Enders, D. Chem. – Eur. J. 2015, 21, 8033–8037. doi:10.1002/chem.201500661
63. Joie, C.; Deckers, K.; Enders, D. Synthesis 2014, 46, 799–808. doi:10.1055/s-0033-1340565
64. Joie, C.; Deckers, K.; Raabe, G.; Enders, D. Synthesis 2014, 46, 1539–1546. doi:10.1055/s-0033-1340982
65. CCDC 1404942 (3p, minor) and CCDC 1409146 (3p, major) contains the supplementary crystallographic data for this paper. These data can be obtained free of charge from The Cambridge Crystallographic Data Centre via http://www.ccdc.cam.ac.uk/data_request/cif.
66. Comins, D. L.; Schilling, S.; Zhang, Y. Org. Lett. 2005, 7, 95–98. doi:10.1021/ol047824w
67. Comins, D. L.; Hiebel, A.-C. Tetrahedron Lett. 2005, 46, 5639–5642. doi:10.1016/j.tetlet.2005.06.105
68. Chitanda, J. M.; Prokopchuk, D. E.; Quail, J. W.; Foley, S. R. Organometallics 2008, 27, 2337–2345. doi:10.1021/om800080e

License and Terms

This is an Open Access article under the terms of the Creative Commons Attribution License (http://creativecommons.org/licenses/by/2.0), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

The license is subject to the Beilstein Journal of Organic Chemistry terms and conditions: (http://www.beilstein-journals.org/bjoc)

The definitive version of this article is the electronic one which can be found at: doi:10.3762/bjoc.12.27