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**Title:** Enantioselective Synthesis of Indolines, Benzodihydrothiophenes, and Indanes by C–H Insertion of Donor/Donor Carbenes

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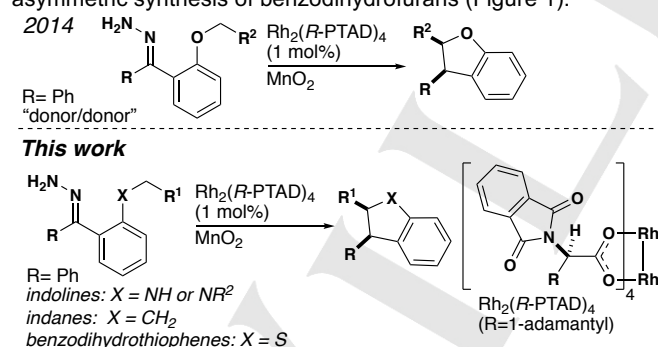
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# Enantioselective Synthesis of Indolines, Benzodihydrothiophenes, and Indanes by C–H Insertion of Donor/Donor Carbenes

Lucas W. Souza,<sup>†</sup> Richard A. Squitieri,<sup>†</sup> Christine A. Dimirjian, Blanka M. Hodur, Leslie A. Nickerson, Corinne N. Penrod, Jesus Cordova, James C. Fettingner, Jared T. Shaw<sup>\*[a]</sup>

**Abstract:** We employ a single catalyst/oxidant system to enable the asymmetric syntheses of indolines, benzodihydrothiophenes, and indanes by C–H insertion of donor/donor carbenes. This methodology enables the rapid construction of densely substituted five-membered rings that form the core of many drug targets and natural products. Furthermore, oxidation of hydrazones to the corresponding diazo compounds proceeds *in situ*, enabling a relatively facile one- or two-pot protocol in which isolation of potentially explosive diazo alkanes is avoided. Regioselectivity studies were performed to determine the impact of sterics and electronics in donor/donor metal carbene C–H insertions to form indolines. This methodology was applied to a variety of substrates in high yield, diastereomeric, and enantiomeric ratios and to the synthesis of a patented indane estrogen receptor agonist with anti-cancer activity.

The insertion of metal carbenes into otherwise unreactive C–H bonds is a powerful method for the synthesis of complex molecules. Chiral metal complexes, specifically rhodium tetracarboxylates, can induce high levels of enantioselectivity for both intra- and intermolecular reactions.<sup>[1]</sup> The asymmetric C–H insertion reactions of metal carbenes flanked by electron withdrawing groups have been well-documented since Teyssie's seminal publication in 1982.<sup>[2]</sup> Recently, our group has demonstrated the utility of metal carbenes with two pendant electron donating groups, or "donor/donor" metal carbenes, in the asymmetric synthesis of benzodihydrofurans (Figure 1).<sup>[3]</sup>

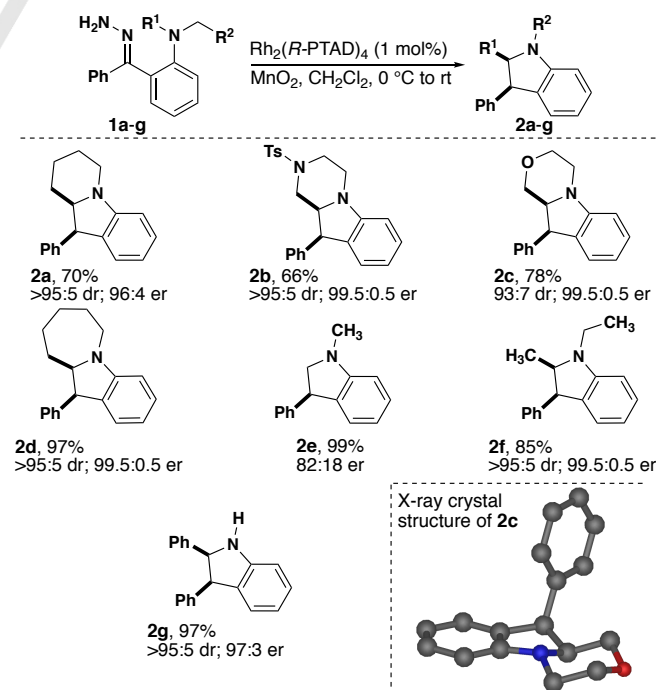


**Figure 1.** Additional cores accessible by C–H insertion reactions of donor/donor metal carbenes.

Moreover, the requisite diazo precursor for the donor/donor carbene can be produced *in situ* from the parent hydrazone. Herein, we apply our methodology to the synthesis of three new classes of products, i.e. indolines, benzodihydrothiophenes and indanes.

Indolines represent a common core structure among natural products and drug discovery candidates.<sup>[4]</sup> Although these heterocycles can be prepared by a variety of strategies, formation of the C2–C3 bond by C–H insertion offers the opportunity to create two stereogenic centers.<sup>[5]</sup> Intramolecular C–H insertion has been used previously for the synthesis of indolines, however few examples of substrates containing basic nitrogen have been shown to proceed with high levels of enantioselectivity.<sup>[6]</sup> We documented the first enantioselective synthesis of an indoline from a donor/donor carbene, albeit with modest enantioselectivity on a substrate lacking a basic nitrogen atom.<sup>[3a]</sup> Recently, Zhu et. al. reported the enantioselective synthesis of indolines from donor/donor carbenes via a non-diazo approach.<sup>[5b]</sup>

A variety of indolines were synthesized with high levels of diastereo- and enantioselectivity (Figure 2). Fused indolines derived from cyclic anilines (**2a–d**) were formed in good to excellent er, dr, and yield. C–H Insertion into primary centers is known to be electronically unfavorable.<sup>[2, 7]</sup> However, indoline **2e** was produced in high yield and modest er. Changing from a



**Figure 2.** Indolines synthesized by the C–H insertion reaction of donor/donor metal carbenes.

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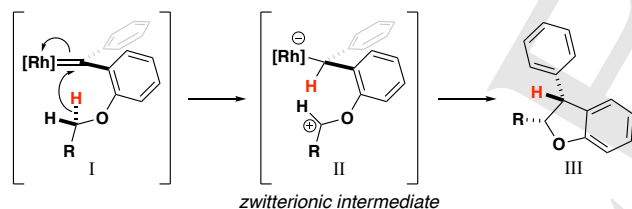
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methyl to an ethyl substituent has a drastic effect on the enantioselectivity of the reaction, as indoline **2f** was produced as primarily one enantiomer. This observation suggests that increased steric bulk at the insertion center plays a role in the high degree of stereoselectivity observed. An indoline derived from an *N*-tosyl protected amine was also tolerated (**2b**) allowing for further modification after deprotection. Additionally, indoline **2g** derived from a free amine was also synthesized in excellent er, dr and yield. X-ray crystallography of **2c** indicated that the relative and absolute configurations of the product were consistent with the sense of induction observed for the formation of benzodihydrofurans.<sup>[3a, 8]</sup>

Over the course of our synthesis of indolines, it became apparent that unsymmetrical aniline substrates presented the opportunity to insert into two chemically different C–H bonds. X-ray crystal structures by Fürstner show that the  $\pi$  system of the pendant phenyl group is in conjugation with the carbene center.<sup>[9]</sup> Additionally, computational experiments in our previous work with benzodihydrofurans have demonstrated that the electron donation afforded by the phenyl ring reduces the electrophilicity of the resultant metal carbene and stabilizes its formation.<sup>[3b]</sup> These factors motivate a stepwise reaction mechanism unique to our donor/donor system. In contrast, the mechanism of donor/acceptor substituted carbenes is known to be concerted asynchronous.<sup>[10]</sup> With this mechanistic framework in mind we set out to test the effect of sterics and electronics on the regiochemistry of C–H insertion reactions to form indolines. The insertion begins with a hydride transfer followed by rapid collapse of the resultant zwitterionic intermediate II.<sup>[3b]</sup> As such, insertion is favored by substituents that stabilize cations at the insertion carbon (Scheme 1). However, steric demand near the insertion carbon can be unfavorable, especially with crowded chiral catalysts.



**Scheme 1.** Mechanism of benzodihydrofuran insertion.

To study the effects of sterics and electronics on the regioselectivity of this chemistry, we synthesized two unsymmetrical hydrazones (**5a, b**) with both sterically and electronically favorable insertion sites (Table 1). In the presence of  $\text{Rh}_2(\text{OAc})_4$ , insertion at the benzylic carbon is favorable, to form **7a** (entry 1). Treatment with  $\text{Rh}_2(\text{Mes})_4$  and  $\text{Rh}_2(R\text{-PTAD})_4$ , shows a decrease in selectivity for the electronically favorable center as the bulk of the catalyst increases (entries 2, 3). When methyl is contrasted with isopropyl (**5b**), selectivity for the less substituted carbon is modest (entry 4) and use of the  $\text{Rh}_2(\text{Mes})_4$  catalyst heavily favors methyl insertion (entry 5). Finally, the reaction of **5b** with  $\text{Rh}_2(R\text{-PTAD})_4$  also favors methyl insertion to a lesser extent (entry 6). The results indicate that both steric and electronic factors compete for the most favorable insertion pathway.

Benzodihydrothiophenes form an interesting class of heterocycles for which there are currently few stereoselective synthetic methods.<sup>[11]</sup> Growing interest in sulfur-containing heterocycles as anti-tumor agents emphasizes the need for new method development in this area.<sup>[12]</sup> To date, there is only a single report of a benzodihydrothiophene synthesis employing rhodium

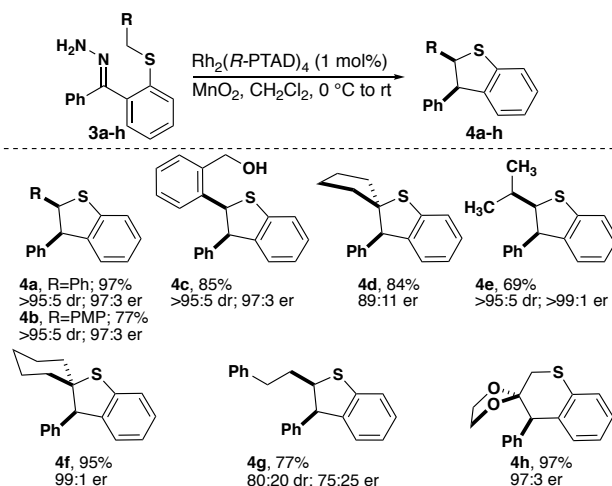
**Table 1.** Regioselectivity of indolines.

Entry	R	catalyst	6:7
1	Ph	$\text{Rh}_2(\text{OAc})_4$	7:93
2	Ph	$\text{Rh}_2(\text{Mes})_4$ <sup>[a]</sup>	25:75
3	Ph	$\text{Rh}_2(R\text{-PTAD})_4$	51:49
4	<i>i</i> -Pr	$\text{Rh}_2(\text{OAc})_4$	61:39
5	<i>i</i> -Pr	$\text{Rh}_2(\text{Mes})_4$ <sup>[a]</sup>	95:5
6	<i>i</i> -Pr	$\text{Rh}_2(R\text{-PTAD})_4$	82:18

<sup>[a]</sup>  $\text{Rh}_2(\text{Mes})_4 = \text{Rh}_2(2, 4, 6\text{-trimethylbenzoate})_4$

carbenes with acceptor-substituted diazo compounds.<sup>[13]</sup> The authors of this study hypothesize that the lack of development of this methodology is due to a propensity of the highly nucleophilic sulfur atom to attack the carbene directly to form an ylide.<sup>[14]</sup> Due to the reduced electrophilicity of our donor/donor carbenes we are able to synthesize a variety of benzodihydrothiophenes with excellent stereocontrol and high yields.

Similarly to benzodihydrofurans,<sup>[3b]</sup> **4a, b** were synthesized with excellent stereoselectivity and yield (Figure 3). Hydrazone **3c** was selectively oxidized to the diazo compound with  $\text{MnO}_2$  in the presence of a benzylic alcohol. Furthermore, the corresponding diazo-alcohol was tolerated well under our reaction conditions, creating a useful chemical handle for further modification without the use of a protecting group (**4c**). Sterically crowded insertion centers were also tolerated allowing access to novel alkyl and spirocyclic compounds **4d-f**. In contrast, poor selectivity was observed with **4g**. This result supports our hypothesis that the degree of stereoselectivity observed corresponds to the presence of steric bulk near the insertion center as shown with indolines. Interestingly, with sufficient activation from the neighboring acetal, six-membered ring formation (**4h**) was achieved in high yield and er.



**Figure 3.** Insertion reactions for the synthesis of benzodihydrothiophenes.

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Indanes are rigid carbacycles that feature prominently in both natural products and in drug discovery. Unlike their heterocyclic counterparts, the electron-neutral benzene and benzylic carbon are less easily oxidized by enzymes responsible for drug clearance.<sup>[15]</sup> Enantioselective synthesis of indanes has been achieved primarily by asymmetric reduction of indenones.<sup>[16]</sup> Hashimoto and Lei have reported the enantioselective synthesis of indanes by C–H insertion of donor/acceptor carbenes.<sup>[17]</sup> In our seminal publication, we demonstrated that the formation of indanes by C–H insertion of donor/donor metal carbenes was possible in good yield and with a modest degree of stereoselectivity.<sup>[3a]</sup> In this work, we expand the scope of this chemistry to include a variety of indanes in excellent yield and selectivity.

Aliphatic spirocyclic five- and six-membered rings were particularly effective (Figure 4A, **9a, c**). Heteroatoms were also tolerated to generate the corresponding spirocycles **9d, e** including a Boc-protected amine as a chemical handle.<sup>[8]</sup> Disubstitution at the inserting carbon creates a high level of steric crowding, yet stabilizes carbocation formation at the insertion center more readily. As such, these substrates react to form quaternary carbons in high yield and with high levels of stereoselectivity (**9b**). The absolute configuration of the new stereogenic center was shown to be *S* upon treatment with  $\text{Rh}_2(\text{R-PTAD})_4$ , in agreement with the sense of induction for both benzodihydrofurans and indolines.

Substitution of the carbene pendant groups with halogens was also tolerated with excellent selectivity (**9f, g**). Furthermore, *J*-values of the relevant protons of these products were consistent

with the *cis* diastereomer observed both in this and in our previous work.<sup>[3a]</sup> We envisioned **9f** as the direct synthetic precursor to a class of indanes patented as an estrogen receptor agonist for the treatment of breast, prostate and ovarian cancers.<sup>[18]</sup> Treatment of substrate **9f** under modified Ullman coupling conditions allowed us to access the first enantioselective synthesis of drug lead **11a**<sup>[19]</sup> in 34% overall yield (Figure 4B).

In conclusion, we have expanded the C–H insertion methodology of donor/donor metal carbenes to the enantioselective synthesis of a variety of indolines, benzodihydrothiophenes, and indanes. Preliminary results suggest the regiochemistry of C–H insertions to form indolines is influenced by competition between electronics that favor carbocation stability and the steric demands of catalyst and substrate. Our methodology allows for the facile synthesis of complex and difficult-to-access benzodihydrothiophenes. Furthermore, insertion into deactivated C–H bonds was accomplished, yielding a variety of indanes. Finally, we demonstrated the utility of this chemistry by carrying out the first highly enantioselective synthesis of patented drug lead **11a** in good overall yield.<sup>[19]</sup>

## Experimental Section

Experimental procedures and compound characterization can be found in the Supporting Information (PDF). <sup>1</sup>H and <sup>13</sup>C NMR spectra for all new compounds and HPLC traces for all enantiomerically enriched insertion products (PDF). X-ray data for compound **2c** (CIF) and **9d** (CIF).

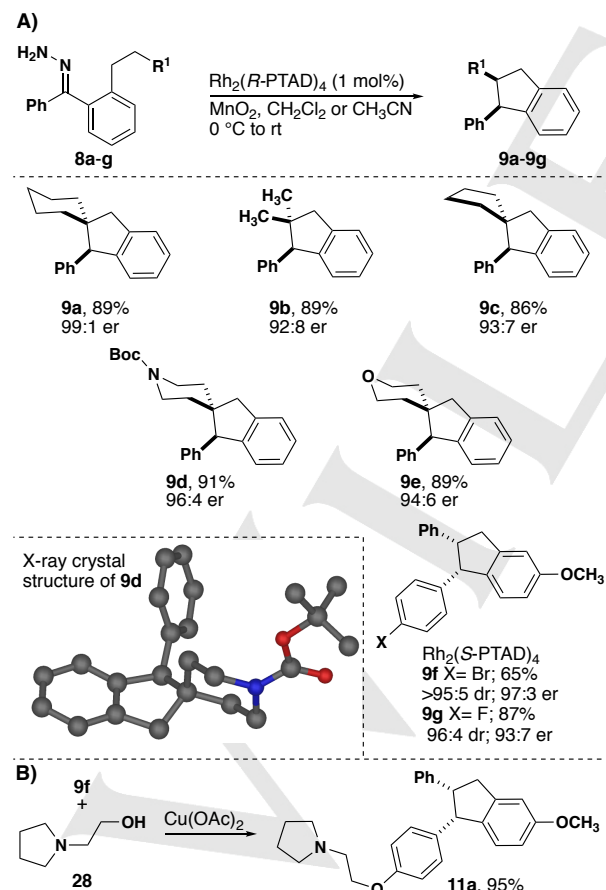
## Acknowledgements

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**Keywords:** asymmetric synthesis • C–H insertion • diastereoselectivity • enantioselectivity • rhodium carbenes

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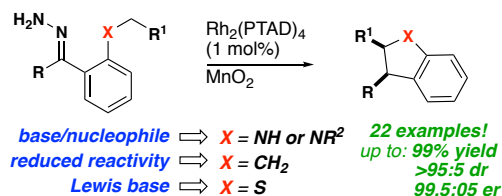
**Figure 4.** A) Insertion reactions for the synthesis of indanes. B) Synthesis of indane drug **11a**.



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## COMMUNICATION



Lucas W. Souza,<sup>†</sup> Richard A. Squitieri,<sup>†</sup> Christine A. Dimirjian, Blanka M. Hodur, Leslie A. Nickerson, Corinne N. Penrod, Jesus Cordova, James C. Fetting, Jared T. Shaw<sup>\*[a]</sup>

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**Enantioselective Synthesis of Indolines, Benzodihydrothiophenes, and Indanes by C–H Insertion of Donor/Donor Carbenes**

**The Carbene Abides.** Rhodium carbenes derived from substrates with no electron withdrawing groups enable the highly selective syntheses of indolines, benzodihydrothiophenes, and indanes in the presence of many previously incompatible functional groups, including strong bases/nucleophiles and free NH and OH groups. A single catalyst/oxidant system is employed for all substrates and in most cases C–H insertion proceeds as a one-pot process from the hydrazone. In the case of indoline formation, regioselectivity reflects a balance of both steric and electronic preferences. The reactions reported here represent the broadest level of functional group tolerance yet demonstrated for enantioselective C–H insertion.