

## From Imide to Lactam Metallo-pyridocarbazoles: Distinct Scaffolds for the Design of Selective Protein Kinase Inhibitors

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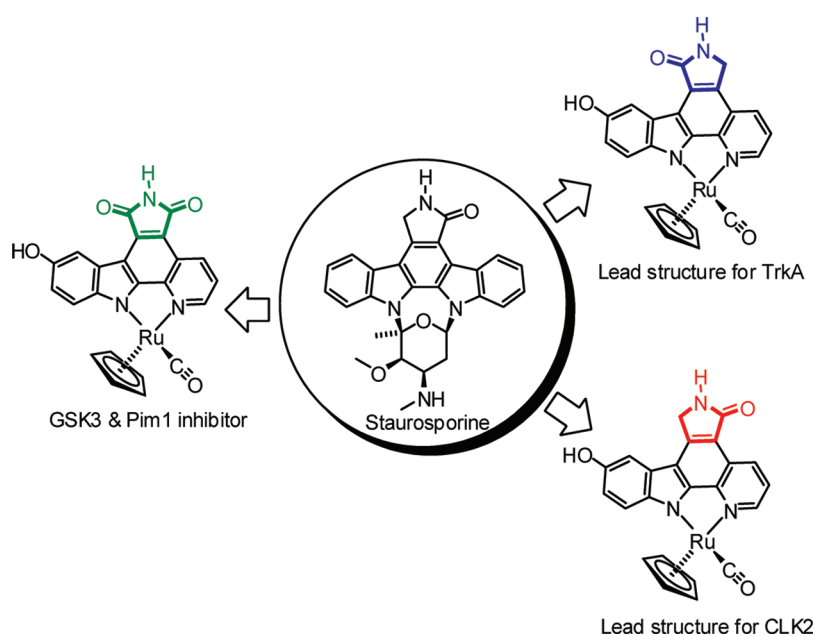
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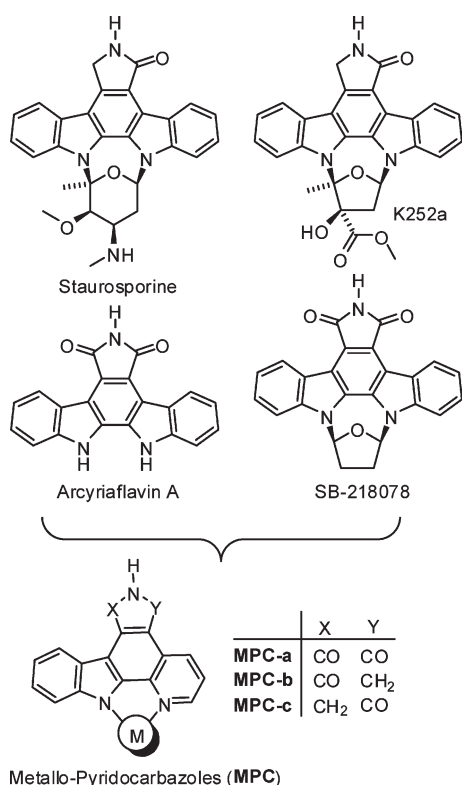


Organometallic pyridocarbazole scaffolds are investigated as protein kinase inhibitors. Whereas our previous designs employed solely a maleimide pharmacophore for achieving the two crucial canonical hydrogen bonds to the hinge region of the ATP binding site, we have now extended our investigations to include the related lactam metallo-pyridocarbazoles. The synthetic access of the two regioisomeric lactam pyridocarbazoles is described, and the distinct biological properties of the two lactam scaffolds are revealed by employing a ruthenium half sandwich complex as a model system, resulting in organometallic lead structures for the inhibition of the protein kinases TrkA and CLK2. These new lactam metallo-pyridocarbazoles expand our existing molecular toolbox and assist toward the generation of metal complex scaffolds as lead structures for the design of selective inhibitors for numerous kinases of the human kinome.

### Introduction

Recently, our group has initiated a research program that aims at exploring the versatility of organometallic complexes as structural scaffolds for the design of enzyme inhibitors. Unlike traditional applications for metals in medicine, we

envisioned that substitutionally inert transition metals could act as a scaffold to organize organic ligands in three-dimensional space. As a proof-of-principle, we chose kinases as our main targets and started by morphing the indolocarbazole natural product class and synthetic derivatives thereof into



**FIGURE 1.** Indolocarbazole natural products and derivatives as an inspiration for metallo-pyridocarbazoles as protein kinase inhibitors.

chemically inert metallo-pyridocarbazoles (**MPC**, Figure 1).<sup>1</sup> On the basis of this scaffold, we have reported over the past 5 years organometallic inhibitors for the kinases GSK3,<sup>2–7</sup> Pim1,<sup>4,6,8</sup> MSK1,<sup>4</sup> PAK1,<sup>9</sup> and PI3K.<sup>10</sup> Multiple co-crystal structures of metal complexes bound to the ATP binding sites of kinases are deposited in the Protein Databank (e.g., 2BZH, 2BZI, 2OI4, 2BZJ, 2IWI, 3CST, 3BWF, 3FXZ), all of them confirming that the metal exerts a purely structural role and is not in direct contact with any residue in the active site.<sup>7–11</sup> Some of the published inhibitors belong to the most potent and selective compounds known for their respective kinases, and we hypothesize that this is caused at least in part by

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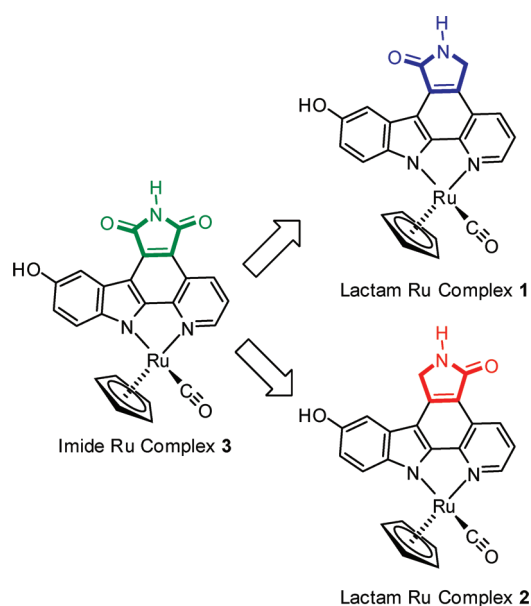
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**FIGURE 2.** Ruthenium half sandwich complexes as a model system to compare the protein kinase inhibition properties of imide and lactam metallo-pyridocarbazoles. Compounds **1–3** are racemic, but only one enantiomer is shown for clarity.

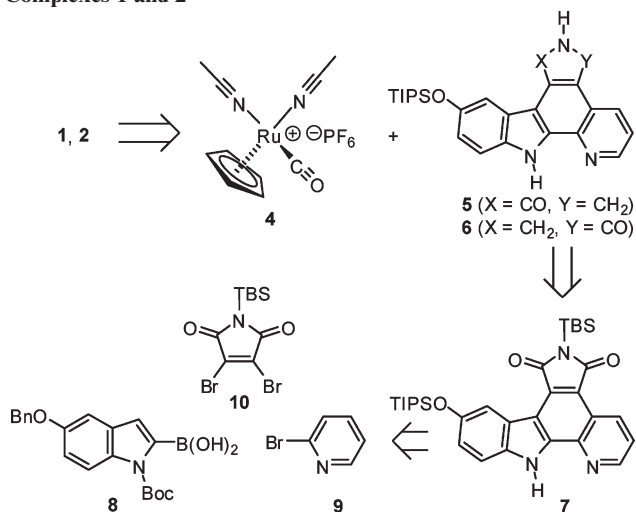
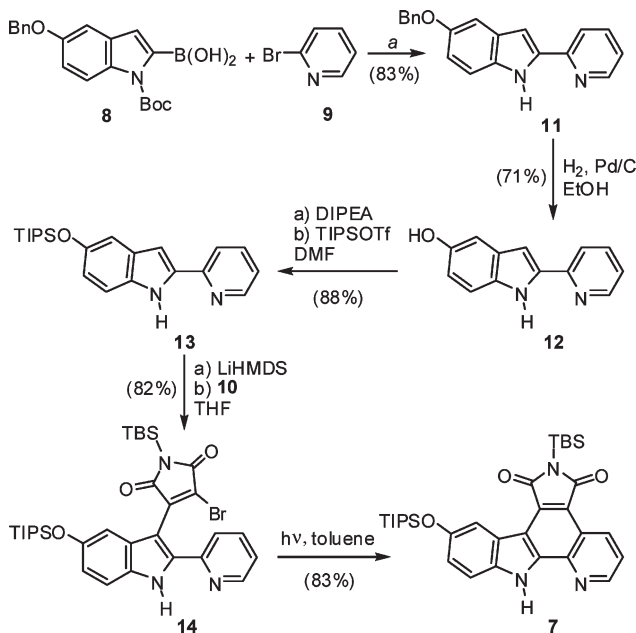
the combination of the globular shape and rigidity of these scaffolds.

To date, all published **MPC** complexes contain the malimide pharmacophore **MPC-a**, whereas the indolocarbazole alkaloids staurosporine and K252a, among others, possess a lactam moiety instead (Figure 1).<sup>12</sup> The lactam or imide is essential for the binding to most kinases, as they both can undergo two key hydrogen bonds with the hinge region of the ATP binding site. However, at the same time it is known that certain kinases prefer the one pharmacophore over the other. Thus, in order to expand the versatility and generality of our **MPC** scaffold for the design of selective kinase inhibitors, we were seeking synthetic access of the two regioisomeric lactam pyridocarbazoles complexes **MPC-b** and **MPC-c** (Figure 1). We here now report our progress into this direction and disclose the synthesis of lactam ruthenium complexes **1** and **2** (Figure 2). In fact, it turns out that lactam complexes **1** and **2** and the previously reported maleimide complex **3** differ significantly in their kinase selectivity profiles. For example, whereas maleimide **3** is a promising lead structure for developing selective inhibitors for GSK3 and Pim1,<sup>3,5–8</sup> lactam **1** constitutes a lead structure for TrkA, and lactam **2** is a potential starting point for CLK2 inhibitor design.

## Results and Discussion

**Synthesis.** A retrosynthetic analysis of the ruthenium complexes **1** and **2** is shown in Scheme 1. The complexes **1** and **2** are accessed from the ruthenium half sandwich complex **4** and the triisopropylsilyl (TIPS)-protected lactam pyridocarbazoles **5** and **6**, respectively. Both lactams **5** and **6** can be synthesized in two consecutive reduction steps starting with the imide pyridocarbazole **7**. Heterocycle **7**

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**SCHEME 1. Retrosynthetic Analysis of Lactam Pyridocarbazole Complexes 1 and 2**

**SCHEME 2. Synthesis of Maleimide Pyridocarbazole 7<sup>a</sup>**


<sup>a</sup>Reaction conditions: first, 1.1 equiv of **8**, 1.0 equiv of **9**, 0.1 equiv of Pd(0) catalyst, 2.75 equiv of  $\text{Na}_2\text{CO}_3$ , DME/ $\text{H}_2\text{O}$ , reflux, overnight; then 10:1 mass ratio silica gel/indole, high vacuum, 80 °C, overnight.

itself can be assembled from three easily accessible building blocks (**8–10**) in a series of six linear steps that include three C–C bond formation reactions.<sup>13,14</sup>

The synthesis of imide pyridocarbazole **7** is outlined in Scheme 2. Suzuki–Miyaura cross-coupling of boronic acid **8** with  $\alpha$ -halogenated pyridine **9** furnished the desired 2-pyridin-2-yl-1*H*-indole with the Boc protection group partially removed. Unable to completely separate the two compounds, the purified mixture was adsorbed onto silica gel

and heated under high vacuum to completely remove the Boc group to afford the 2-pyridin-2-yl-1*H*-indole **11** in 83% over two steps. Alternatively, the potassium trifluoroborate analogue of indole **8** could also be used for this transformation.<sup>13,15,16</sup> Before conducting the next two C–C bond formation reactions, the benzyl-protection group was first exchanged for a TIPS group in order to render the scaffold more soluble in organic solvents. This was accomplished by hydrogenolysis with  $\text{H}_2$ , Pd/C furnishing the phenol derivative **12** (71%), followed by treatment with TIPS triflate in the presence of the base di(isopropyl)ethylamine to afford the TIPS-protected pyridoindole **13** (88%). 2-Pyridin-2-yl-1*H*-indole **13** was deprotonated with LiHMDS and reacted with dibromomaleimide **10** to afford monobromide **14** in 82% yield. The final C–C bond could then be installed by a photocyclization of **14** in toluene with a medium pressure Hg lamp using an uranium filter (50% transmission at 350 nm) to provide pyridocarbazole **7** (83%). In summary, the *bis*-silyl protected ligand **7** was prepared in a total of six steps in an overall yield of 35%.

The first of two reduction steps conducted on the imide pyridocarbazole **7** is shown in Scheme 3. Two equivalents of  $\text{NaBH}_4$  were added at 0 °C each hour for a total of eight hours, followed by warming to room temperature to furnish regioisomeric products **15** and **16** in a ratio of 3:1 in a combined yield of 84%. Slow addition of the reducing agent at low temperature is critical, preventing the loss of the *tert*-butyldimethylsilyl (TBS) protection group. Migration of the TBS group to the newly formed  $sp^3$  oxygen atom retains the TBS group required for the solubility of the compounds in organic solvents and the successful separation of the two regioisomers using column chromatography.

The constitutions of the two regioisomeric lactams **15** and **16** were assigned by rotational frame nuclear Overhauser effect spectroscopy (ROESY) analysis of both regioisomers **15** and **16** (Figure 3). We were initially surprised by the observed regioselectivity because we expected the reaction to occur primarily at the more electrophilic carbonyl group **a** of heterocycle **7** (Scheme 3).<sup>17,18</sup> This carbonyl group is directly conjugated to the electron poor pyridine, unlike the carbonyl **b**, which is directly conjugated to the electron rich indole. Additionally, we suspected that nucleophilic attack of carbon center **b** would be more hindered as a result of the hydride approaching from the same side as the bulky TIPS group.<sup>19–21</sup> However, a likely explanation for this observed result could be the difference in the transition state energies, with migration of the TBS group preferring the more electron rich oxygen atom conjugated to the indole.

Transformation of heterocycles **15** and **16** to their corresponding lactams is outlined in Scheme 3. First, the TBS group of **15** was replaced by an ethyl group through treatment with 0.1% ethanol in the presence of 10% TFA to

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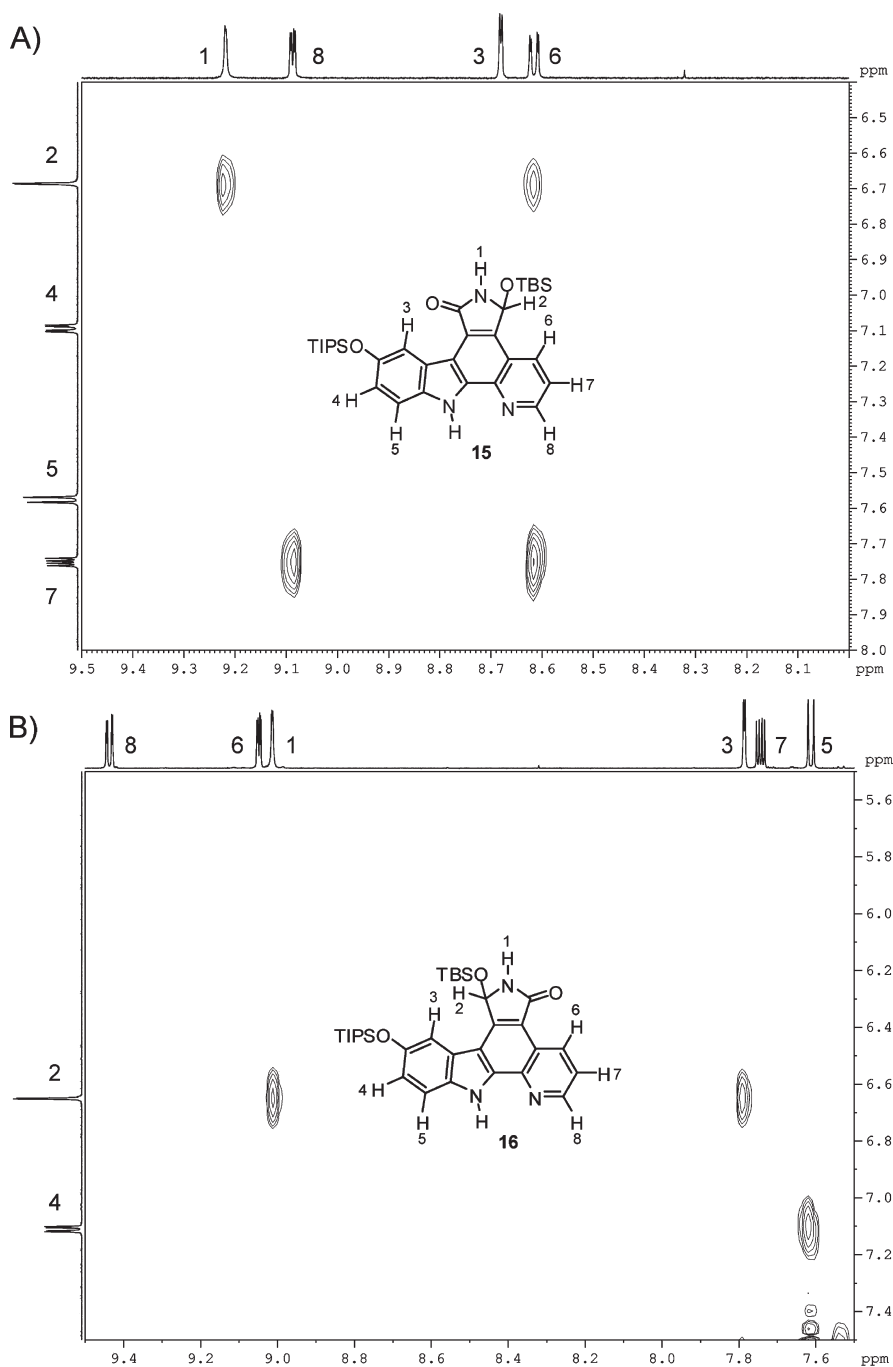
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**FIGURE 3.**  $^1\text{H}$ – $^1\text{H}$ -ROESY spectra ( $\text{DMSO-}d_6$ , 600 MHz) of compounds **15** and **16**. (A) Compound **15**: through-space correlation between H2 and H6. (B) Compound **16**: through-space correlation between H2 and H3.

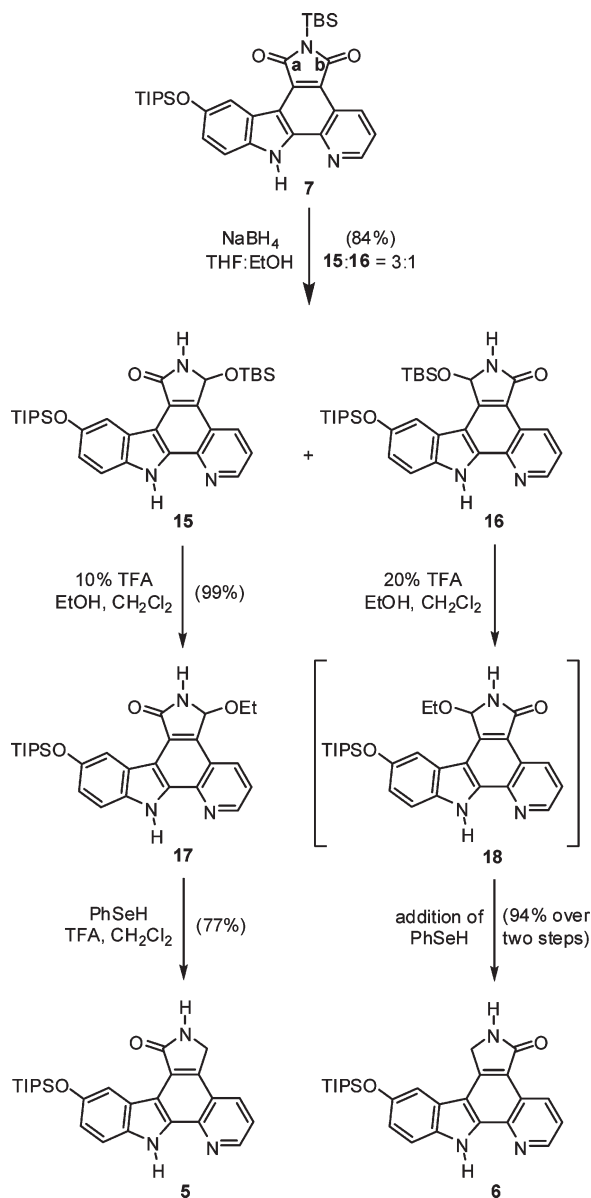
afford heterocycle **17** in near quantitative yield. This ether derivative could then smoothly be reduced to lactam **5** in 77% yield using phenylselenol and a slight excess of TFA in refluxing methylene chloride.<sup>22,23</sup> It is noteworthy, that initial attempts to directly convert TBS-protected ligand **15** to lactam **5** failed and we attributed this to the size of the TBS ether, precluding the necessary nucleophilic attack of

phenylselenol. In analogy, treatment of TBS-protected ligand **16** with 0.1% ethanol in now 20% TFA afforded intermediate **18**, which due to its poor solubility was immediately reacted with phenylselenol in refluxing methylene chloride to furnish lactam **6** in excellent yield (94%) over two steps.

Introduction of the ruthenium half sandwich fragment using both lactam ligands **5** and **6** is shown in Scheme 4. Unable to optimize the coordination step using regular heating with an oil bath, we were pleased to discover that the complexation proceeds very smoothly under microwave irradiation. Lactams **5** and **6** were each reacted with

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**SCHEME 3. Synthesis of Lactam Pyridocarbazoles 5 and 6 from Maleimide 7**


ruthenium precursor 4 in the presence of  $\text{K}_2\text{CO}_3$  in anhydrous DMF at 80 °C. After only 10 min, both reactions were complete as judged by thin layer chromatography. After column chromatography, TIPS-protected ruthenium complexes 19 and 20 were isolated in high yields of 79% and 87%, respectively. Lactam inhibitor complexes 1 and 2 were then prepared by careful treatment with tetrabutylammonium fluoride (TBAF) (76% and 71%, respectively). It should be noted that lactam ruthenium complexes 20 and 2, in contrast to 19 and 1, are slightly air-sensitive, slowly oxidizing to their corresponding imide analogues if left in solution for long periods of time. Also, this process is accelerated in the presence of base. Therefore, it is critical to thoroughly purge both reactions 6  $\rightarrow$  20 and 20  $\rightarrow$  2 with inert gas. Furthermore, as a precaution, DMSO stock solutions of ruthenium complex 2 were purged with nitrogen and stored frozen at  $-20$  °C until assayed for activity.

**Kinase Inhibition.** To identify similarities and differences between the new lactam half sandwich complexes 1 and 2 compared to the imide compound 3, racemic mixtures of complexes 1–3 were screened at one concentration against a panel of 253 human kinases (Millipore KinaseProfiler).<sup>24</sup> As expected, the racemic mixture of maleimide 3 displayed high potencies for GSK3 ( $\alpha$ -isoform: 0% activity at 30 nM, 10  $\mu\text{M}$  ATP) and Pim1 and Pim2 (1% and  $-1\%$  activity at 30 nM, respectively, 10  $\mu\text{M}$  ATP), whereas most other protein kinases were not inhibited to a significant degree (see Supporting Information). In contrast, lactams 1 and 2 displayed only very moderate inhibition of GSK3 with activities above 40% at concentrations of 100 nM (10  $\mu\text{M}$  ATP). Furthermore, under the conditions of the screening (100 nM compound, 10  $\mu\text{M}$  ATP), lactam 1 demonstrated a surprising selectivity for just very few kinases (Flt3(D835Y), MLCK, Pim1, and TrkA), whereas lactam 2 showed highest inhibitory activity against the protein kinases CLK2, PKG1 $\beta$ , and Pim1 (1%, 2%, and 2% remaining activities at 100 nM of 2, 10  $\mu\text{M}$  ATP, respectively; see Supporting Information).

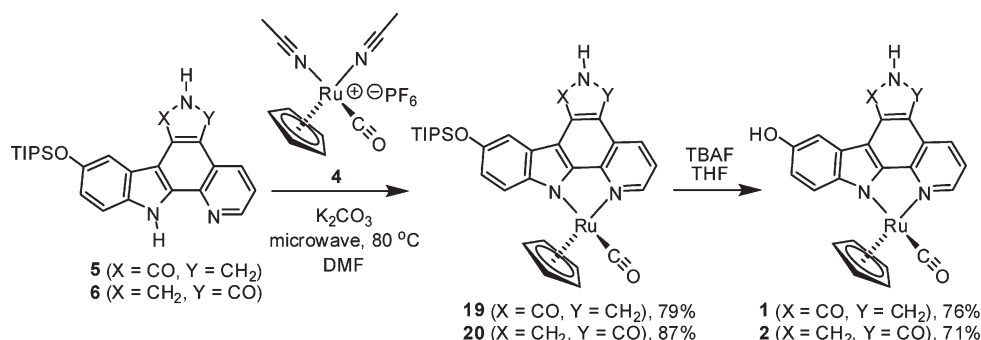
To investigate the inhibition of TrkA and CLK2 in more detail, the racemic half sandwich complexes 1–3 were first resolved into their individual enantiomers. The chiral separation of imide 3 has been reported recently,<sup>8</sup> and the chiral separation of lactam ruthenium complexes 1 and 2 was accomplished by resolving the individual enantiomers of their TIPS-protected half sandwich analogues 19 and 20, with the TIPS group serving as a handle to increase solubility in organic solvents. The resolution of 10 mg of each enantiomer of TIPS-protected complexes 19 and 20 was achieved using an analytical HPLC chiral column, and subsequently all four stereoisomers were treated separately with TBAF and purified over column chromatography. Assignment of the absolute configuration for each enantiomer for both complexes was achieved by comparing individual CD spectra with known reference compounds (see Supporting Information).

Table 1 displays the  $\text{IC}_{50}$  values of the individual enantiomers of 1–3 against Pim1, GSK3 $\beta$ , TrkA, and CLK2. The data demonstrate that by simply removing one carbonyl group of imide 3 a profound influence on the potency and selectivity of the half sandwich scaffold is achieved. As reported recently, ( $R_{\text{Ru}}$ )-3 and some derivatives show a high selectivity for GSK3 and Pim1, whereas the enantiomer ( $S_{\text{Ru}}$ )-3 is a very selective inhibitor of Pim1.<sup>5–8</sup> However, replacing one carbonyl group by a methylene group of the Pim1 inhibitor lead structure ( $S_{\text{Ru}}$ )-3 ( $\text{IC}_{50}$  = 0.2 nM) reduces the binding affinities for Pim1 by the factors of 200 and 30 for the lactam complexes ( $S_{\text{Ru}}$ )-1 and ( $S_{\text{Ru}}$ )-2, respectively. Even more pronounced, replacing the imide group for a lactam in the GSK3 inhibitor lead structure ( $R_{\text{Ru}}$ )-3 ( $\text{IC}_{50}$  = 1.0 nM) reduces the binding affinities for GSK3 $\beta$  by factors of 330 and 1000, for the complexes ( $R_{\text{Ru}}$ )-1 and ( $R_{\text{Ru}}$ )-2, respectively.

This dramatic loss in binding affinities of ( $R_{\text{Ru}}$ )-3 for GSK3 by removing a single carbonyl group can be rationalized with the help of a recently solved crystal structure of a related ruthenium half sandwich complex with GSK3 $\beta$ .<sup>7</sup>

(24) Manning, G.; Whyte, D. B.; Martinez, R.; Hunter, T.; Sudarsanam, S. *Science* 2002, 298, 1912–1934.



SCHEME 4. Synthesis of Lactam Complexes 1 and 2<sup>a</sup>

<sup>a</sup>Formed as racemates; only one enantiomer is shown for clarity.

TABLE 1. IC<sub>50</sub> Values of Imide and Lactam Half Sandwich Complexes 1–3 and (R<sub>Ru</sub>)-21 against Pim1, GSK3β, TrkA, and CLK2<sup>a</sup>

	Pim1	GSK3β	TrkA	CLK2
(R <sub>Ru</sub> )-3	0.7	1.0	65	<sup>b</sup>
(S <sub>Ru</sub> )-3	0.2	15	40	<sup>b</sup>
(R <sub>Ru</sub> )-1	80	330	20	<sup>c</sup>
(S <sub>Ru</sub> )-1	40	1000	50	<sup>c</sup>
(R <sub>Ru</sub> )-2	25	1000	75	30
(S <sub>Ru</sub> )-2	6	370	15	5
(R <sub>Ru</sub> )-21	390	1200	6	<sup>d</sup>

<sup>a</sup>IC<sub>50</sub> data were obtained by phosphorylation of substrate peptides with [<sup>γ</sup>-<sup>32</sup>P]ATP or [<sup>γ</sup>-<sup>33</sup>P]ATP and 100 μM ATP in the presence of organometallic kinase inhibitors. Each IC<sub>50</sub> value was determined from at least two independent measurements and the average was taken. See Experimental Section for more details.

<sup>b</sup>Racemic 3: IC<sub>50</sub> = 35 nM at 100 μM ATP. <sup>c</sup>Racemic 1: 26% CLK2 activity at 100 nM 1 and 10 μM ATP. <sup>d</sup>50% CLK2 activity at 100 nM of (R<sub>Ru</sub>)-21 at 10 μM ATP.

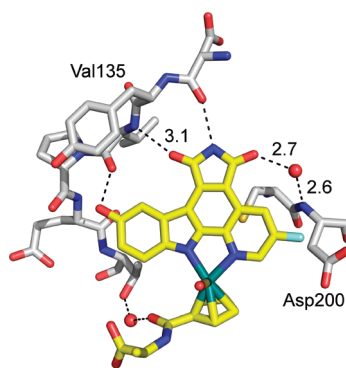


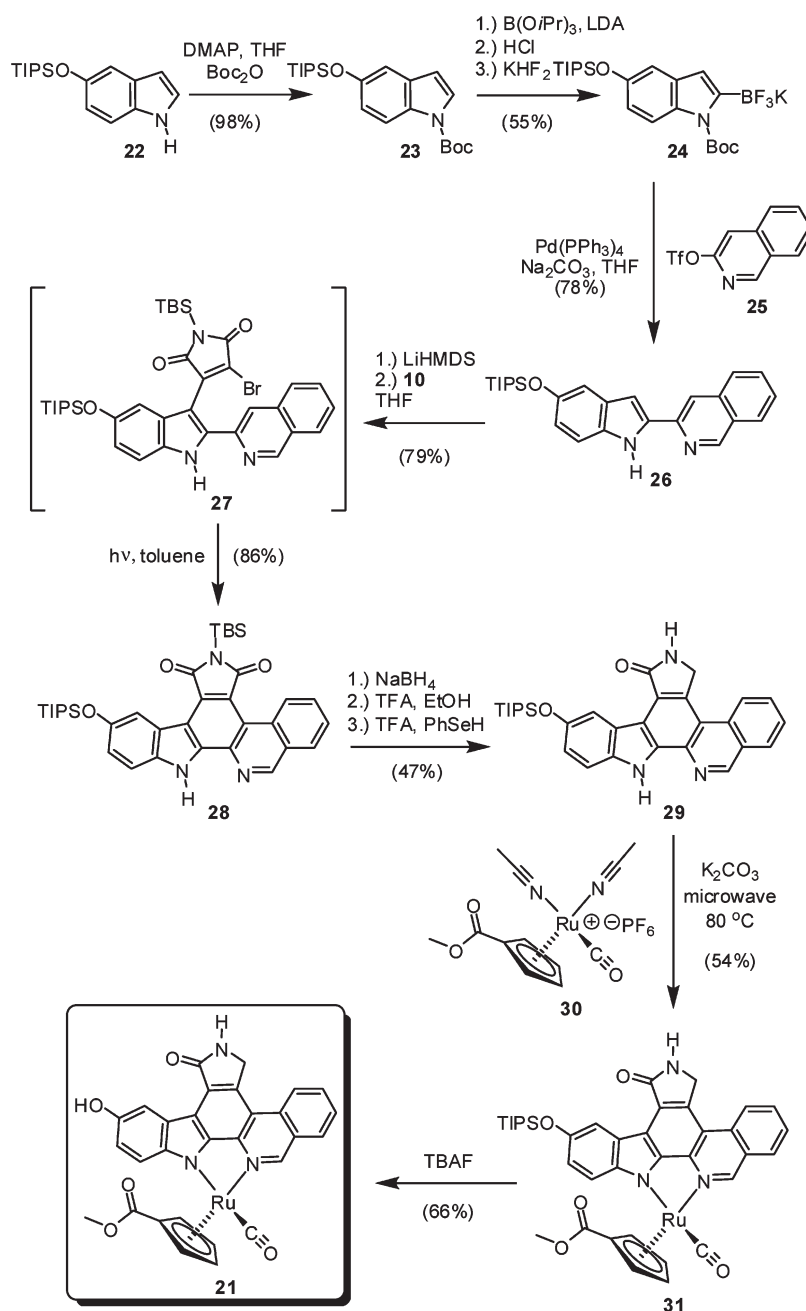
FIGURE 4. Crystal structure of a maleimide-containing ruthenium pyridocarbazole half sandwich complex with GSK3β (PDB code 2JLD) showing the hydrogen-bonding interactions of the maleimide carbonyl groups in the ATP binding site of GSK3β.

Figure 4 shows the hydrogen bond interactions of the maleimide pyridocarbazole in the ATP binding site of GSK3β and reveals that both carbonyl groups are involved in hydrogen bonds, one with the hinge residue valine 135 and a second water-mediated contact with aspartate 200. This explains the preference of GSK3 for the maleimide over the lactam, which can provide only one of the two hydrogen-bonding contacts.<sup>25</sup>

(25) For the binding modes of small molecules to the ATP binding sites of protein kinases, see for example: Ghose, A. K.; Herbertz, T.; Pippin, D. A.; Salvino, J. M.; Mallamo, J. P. *J. Med. Chem.* **2008**, *51*, 5149–5171.

Thus, having eliminated the intrinsic preference of the half sandwich scaffold for GSK3 and Pim1 by transforming the imide group of 3 into lactam moieties, the individual enantiomers of complexes 1 and 2 can now be used as lead structures for other kinases. In fact, Table 1 demonstrates that the enantiomer (R<sub>Ru</sub>)-1 displays a significant selectivity for TrkA over Pim1 (4-fold) and GSK3β (16.5-fold) and is therefore a promising lead structure for the development of further improved inhibitors of protein kinase TrkA, whereas (S<sub>Ru</sub>)-2 displays an IC<sub>50</sub> of just 5 nM (100 μM ATP) against CLK2, thus serving as a lead structure for the development of CLK2 inhibitors.

**Improved Inhibitor for Protein Kinase TrkA.** To verify that the obtained lactam complexes are indeed promising lead structures for the design of organometallic complexes as inhibitors for protein kinases different from Pim1 or GSK3, we decided to use (R<sub>Ru</sub>)-1 as a starting point for the proof-of-principle design of a more potent and selective inhibitor for TrkA. From previous structure activity relationships with organoruthenium half sandwich complexes,<sup>5,13,14</sup> we envisioned that the complex (R<sub>Ru</sub>)-21, differing from complex (R<sub>Ru</sub>)-1 by a fused benzene ring at the pyridine moiety and a methylester at the cyclopentadienyl ring, might be an improved inhibitor for TrkA with a diminished affinity for Pim1. The synthesis of racemic 21 is summarized in Scheme 5. Starting with 5-(triisopropylsilyloxy)indole 22, Boc-protection afforded 23 (98%), which was subsequently converted to the stable trifluoroborate 24 (55%). Suzuki–Miyaura cross-coupling of 24 with triflate 25 in refluxing anhydrous THF furnished 2-isoquinolin-3-yl-1*H*-indole 26 (78%) with complete loss of the Boc group. Next, isoquinolinyl-indole 26 was deprotonated with LiHMDS and condensed with dibromomaleimide 10 to afford the rather unstable monobromide 27 (79%). The final C–C bond could then be installed by a photocyclization in toluene with a medium pressure Hg lamp using a uranium filter to provide heterocycle 28 in a yield of 86%. Next, 28 was converted to the lactam 29 in a yield of 47% by first treatment with NaBH<sub>4</sub>, followed by EtOH with 0.1% TFA, and subsequently PhSeH in the presence of 2 equiv of TFA. The lactam ligand 29 was reacted with ruthenium half sandwich complex 30 in the presence of K<sub>2</sub>CO<sub>3</sub> in anhydrous DMF at 80 °C under microwave irradiation. After 10 min and column chromatography, TIPS-protected

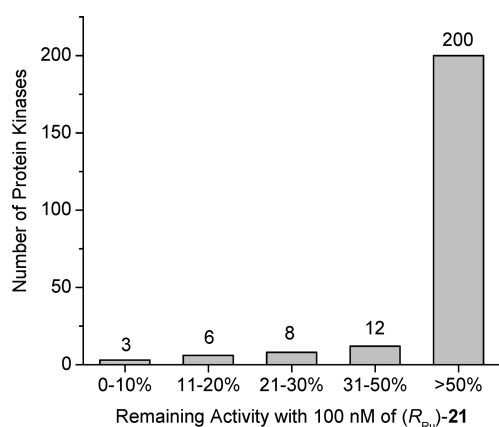
SCHEME 5. Synthesis of Lactam Complex **21**<sup>a</sup>

<sup>a</sup>The half sandwich complexes were formed as racemates, but only the (*R*<sub>Ru</sub>)-enantiomer is shown for clarity.

ruthenium complex **31** was isolated in a yield of 54%. Finally, treatment of **31** with TBAF at 0 °C afforded the ruthenium complex **21** (66%) and the racemic mixture of **21** was resolved into its enantiomers at this final stage using an analytical chiral HPLC column. The absolute configuration for both enantiomers was assigned by comparison of CD spectra with known reference compounds (see Supporting Information).

IC<sub>50</sub> values of (*R*<sub>Ru</sub>)-**21** at 100 μM ATP against TrkA, GSK3β, and Pim1 are presented in Table 1. Gratifyingly, (*R*<sub>Ru</sub>)-**21** shows a significantly improved potency and selectivity compared to the initial lead structure (*R*<sub>Ru</sub>)-**1**. With an IC<sub>50</sub> value of 6 nM for TrkA, (*R*<sub>Ru</sub>)-**21** prefers binding to

TrkA over Pim1 by a factor of 65 (IC<sub>50</sub> = 390 nM) and even a factor of 200 over GSK3β (IC<sub>50</sub> = 1200 nM). Thus, (*R*<sub>Ru</sub>)-**21** displays high selectivity over Pim1 and GSK3 and probably most other protein kinases. In fact, a screening of complex (*R*<sub>Ru</sub>)-**21** against a panel of human wild-type protein kinases demonstrates the excellent selectivity profile of (*R*<sub>Ru</sub>)-**21** (Figure 5). From 229 tested protein kinases, 200 remained more than 50% active at a concentration of 100 nM of (*R*<sub>Ru</sub>)-**21** (10 μM ATP), and only three kinases, including TrkA and TrkB, displayed activities below 10% under these conditions. Thus, it can be concluded that (*R*<sub>Ru</sub>)-**21** constitutes a highly selective molecular probe for the protein kinase TrkA.



**FIGURE 5.** Selectivity profile of ( $R_{Ru}$ )-21 tested against a panel of 229 human wild-type protein kinases at a single concentration of 100 nM and an ATP concentration of 10  $\mu$ M. Duplicate measurements were performed and the average was taken.

## Conclusions

We have here reported synthetic access to the class of lactam metallo-pyridocarbazoles **MPC-b** and **MPC-c**, thus complementing the previously developed imide metallo-pyridocarbazole scaffold **MPC-a** and therefore expanding our toolbox of using metallo-pyridocarbazoles as general scaffolds for the design of selective kinase inhibitors. The distinct properties of the lactam scaffolds have been demonstrated by using the half sandwich complex **3** as a model system. By replacing the imide pharmacophore of **3** for the lactam moieties in **1** and **2**, we have changed the selectivity profile from GSK3/Pim1 selectivity of ( $R_{Ru}$ )-**3** and Pim1 selectivity of ( $S_{Ru}$ )-**3** to TrkA selectivity of ( $R_{Ru}$ )-**1** and CLK2 selectivity of ( $S_{Ru}$ )-**2**. These lactam metallo-pyridocarbazoles will serve as scaffolds for the design of protein kinase inhibitors with new selectivity profiles as has been demonstrated with the TrkA-selective organoruthenium complex ( $R_{Ru}$ )-**21**.

## Experimental Section

All reactions were carried out using oven-dried glassware and conducted under a positive pressure of nitrogen unless otherwise specified. NMR spectra were recorded on an Avance 300 (300 MHz), DRX-400 (400 MHz), or DRX-500 (500 MHz) spectrometer. ROESY experiments were recorded on a Bruker Avance 600 MHz spectrometer. Infrared spectra were recorded on either a Bruker Alpha FTIR or Nicolet 510 FTIR spectrometer. High resolution mass spectra were obtained with a Finnigan LTQ-FT instrument using either APCI or ESI. Reagents and solvents were used as received from standard suppliers unless otherwise specified. CD spectra were recorded on a JASCO J-810 spectropolarimeter. Ruthenium complexes **4** and **30** were prepared as previously reported.<sup>5,26</sup>

**Compound 11.** A biphasic suspension of pyridine **9** (0.87 mL, 9.1 mmol), indole **8** (3.67 g, 10.0 mmol), tetrakis(triphenylphosphine)palladium(0) (1.05 g, 0.91 mmol), and  $\text{Na}_2\text{CO}_3$  (2.65 g, 25.0 mmol) in 1,2-dimethoxyethane (39 mL) and water (11 mL) was purged with nitrogen for 15 min and then refluxed overnight. The resulting orange reaction mixture was cooled to room temperature, diluted with water, and extracted with EtOAc. The combined organic layers were washed with brine, dried using  $\text{Na}_2\text{SO}_4$ , filtered, and concentrated to dryness

in vacuo. The crude material was adsorbed onto silica gel and chromatographed using hexanes/EtOAc (8:1 to 1:1). The combined product eluents were isolated as a mixture of a Boc-protected 2-pyridin-2-yl-1H-indole and compound **11**. The resulting orange solid (2.82 g) was dissolved into EtOAc and adsorbed onto silica gel (28.2 g) using rotary evaporation. The powder was heated to 80 °C overnight under high vacuum. The silica gel was cooled to room temperature, filtered through cotton with  $\text{CH}_2\text{Cl}_2/\text{MeOH}$  (15:1) and the filtrate dried in vacuo to provide 2-pyridin-2-yl-1H-indole **11** (2.27 g, 83% over 2 steps) as a yellow solid.  $^1\text{H}$  NMR (300 MHz,  $\text{CDCl}_3$ ):  $\delta$  (ppm) 9.54 (s, 1H), 8.57 (ddd,  $J = 4.9, 1.7, 1.0$  Hz, 1H), 7.79–7.67 (m, 2H), 7.51–7.48 (m, 2H), 7.43–7.29 (m, 4H), 7.18–7.14 (m, 2H), 6.98 (dd,  $J = 8.8, 2.4$  Hz, 1H), 6.94 (dd,  $J = 2.1, 0.8$  Hz, 1H), 5.12 (s, 2H).  $^{13}\text{C}$  NMR (75 MHz,  $\text{CDCl}_3$ ):  $\delta$  (ppm) 153.9, 150.6, 149.4, 137.9, 137.6, 136.8, 132.2, 129.7, 128.7, 128.0, 127.8, 122.1, 119.9, 114.8, 112.3, 104.3, 100.5, 71.1. IR  $\nu$  ( $\text{cm}^{-1}$ ) 3192, 1622, 1594, 1562, 1543, 1469, 1443, 1299, 1257, 1214, 1150, 1119, 785, 775, 741. HRMS calcd for  $\text{C}_{20}\text{H}_{16}\text{N}_2\text{ONa}$  ( $M + \text{Na}$ )<sup>+</sup> 323.1155, found ( $M + \text{Na}$ )<sup>+</sup> 323.1152.

**Compound 12.** A suspension of 2-pyridin-2-yl-1H-indole **11** (2.14 g, 7.13 mmol) in ethanol (98 mL) was purged with nitrogen for 15 min. Following thorough nitrogen purge, palladium (10 wt % on activated carbon, dry) (2.14 g) was added, and the reaction mixture was carefully purged with hydrogen gas for approximately 10 min. The reaction mixture was then stirred at room temperature under an atmosphere of hydrogen, monitoring by thin layer chromatography. After 2 h, the resulting dark reaction mixture was filtered through Celite with  $\text{CH}_2\text{Cl}_2/\text{MeOH}$  (10:1), and the filtrate was concentrated to dryness in vacuo. The crude material was adsorbed onto silica gel and subjected to silica gel chromatography with hexanes/EtOAc (1:1). The combined product eluents were dried in vacuo to provide the 2-pyridin-2-yl-1H-indole **12** (1.07 g, 71%) as an off-white solid.  $^1\text{H}$  NMR consistent with literature reported spectrum.<sup>3</sup>

**Compound 13.** A solution of 2-pyridin-2-yl-1H-indole **12** (1.07 g, 5.10 mmol) in DMF (18.3 mL) was cooled to 0 °C, and  $N,N$ -diisopropylethylamine (4.50 mL, 25.5 mmol) was added dropwise. After stirring for 10 min at 0 °C, triisopropylsilyl trifluoromethanesulfonate (1.44 mL, 5.36 mmol) was added dropwise over 15 min, and the reaction mixture was stirred at room temperature for an additional 1 h. The resulting yellow solution was cooled to 0 °C, and excess 1 M  $\text{NH}_4\text{OAc}$  was carefully added. The mixture was then further diluted with water and extracted with  $\text{CH}_2\text{Cl}_2$ . The combined organic layers were washed with brine, dried using  $\text{Na}_2\text{SO}_4$ , filtered, and concentrated to dryness in vacuo. The crude material was adsorbed onto silica gel and subjected to silica gel chromatography with  $\text{CH}_2\text{Cl}_2/\text{MeOH}$  (100:1). The combined product eluents were dried in vacuo to provide the 2-pyridin-2-yl-1H-indole **13** (1.65 g, 88%) as a white solid.  $^1\text{H}$  NMR (300 MHz,  $\text{CDCl}_3$ ):  $\delta$  (ppm) 10.30 (s, 1H), 8.59 (ddd,  $J = 4.9, 1.7, 1.0$  Hz, 1H), 7.80 (td,  $J = 8.0, 1.1$  Hz, 1H), 7.70 (ddd,  $J = 8.0, 7.4, 1.7$  Hz, 1H), 7.17–7.13 (m, 3H), 6.95 (dd,  $J = 2.1, 0.8$  Hz, 1H), 6.86 (dd,  $J = 8.8, 2.3$  Hz, 1H), 1.39–1.27 (m, 3H), 1.17 (d,  $J = 6.9$  Hz, 18H).  $^{13}\text{C}$  NMR (75 MHz,  $\text{CDCl}_3$ ):  $\delta$  (ppm) 150.7, 150.1, 149.1, 137.5, 136.7, 132.5, 129.8, 121.9, 119.9, 117.8, 111.8, 109.9, 100.4, 18.1, 12.8. IR  $\nu$  ( $\text{cm}^{-1}$ ) 3150, 2940, 2861, 1595, 1545, 1442, 1292, 1280, 1222, 1150, 963, 879, 804, 776, 679. HRMS calcd for  $\text{C}_{22}\text{H}_{31}\text{N}_2\text{OSi}$  ( $M + \text{H}$ )<sup>+</sup> 367.2200, found ( $M + \text{H}$ )<sup>+</sup> 367.2198.

**Compound 14.** A solution of high-vacuum-dried 2-pyridin-2-yl-1H-indole **13** (1.55 g, 4.23 mmol) in freshly distilled THF (11 mL) was cooled to –15 °C. Lithium bis(trimethylsilyl)amide (1 M solution in hexanes) (12.7 mL, 12.7 mmol) was added dropwise over 15 min, and the resulting orange reaction mixture was stirred at –15 °C for an additional 45 min. A solution of vacuum-dried **10** (1.64 g, 4.44 mmol) in freshly distilled THF

(26) Gill, T. P.; Mann, K. R. *Organometallics* **1982**, *1*, 485–488.



(11 mL) cooled to 0 °C was added, and the reaction mixture was stirred at -15 °C for 20 min and room temperature overnight. The resulting dark purple reaction mixture was carefully poured into stirring ice cold water (300 mL) and extracted with EtOAc. The combined organic layers were washed with brine, dried using Na<sub>2</sub>SO<sub>4</sub>, filtered, and concentrated to dryness in vacuo. The crude material was adsorbed onto silica gel and subjected to silica gel chromatography with hexanes/EtOAc (6:1 to 3:1). The combined product eluents were dried in vacuo to provide monobromide **14** (2.25 g, 82%) as an orange solid. <sup>1</sup>H NMR (300 MHz, CDCl<sub>3</sub>): δ (ppm) 10.22 (s, 1H), 8.58 (d, *J* = 4.2 Hz, 1H), 7.61 (ddd, *J* = 7.8, 7.8, 1.8 Hz, 1H), 7.31 (d, *J* = 8.1 Hz, 1H), 7.25 (d, *J* = 8.7 Hz, 1H), 7.18 (ddd, *J* = 7.5, 4.9, 1.2 Hz, 1H), 6.99 (d, *J* = 2.3 Hz, 1H), 6.88 (dd, *J* = 8.7, 2.3 Hz, 1H), 1.34–1.23 (m, 3H), 1.12 (d, *J* = 7.0 Hz, 18H), 1.00 (s, 9H), 0.49 (s, 6H). <sup>13</sup>C NMR (75 MHz, CDCl<sub>3</sub>): δ (ppm) 173.4, 171.1, 151.1, 149.8, 149.4, 141.7, 137.2, 136.8, 131.5, 128.2, 125.6, 122.8, 122.7, 118.9, 112.5, 110.4, 101.7, 26.5, 19.1, 18.3, 12.9, -4.2. IR ν (cm<sup>-1</sup>) 3337, 2947, 2865, 1703, 1635, 1473, 1313, 1269, 1213, 1186, 1086, 971, 892, 826, 735. HRMS calcd for C<sub>32</sub>H<sub>45</sub>BrN<sub>3</sub>O<sub>3</sub>Si<sub>2</sub> (M + H)<sup>+</sup> 654.2177, found (M + H)<sup>+</sup> 654.2182.

**Compound 7.** An orange suspension of monobromide **14** (1.10 g, 1.68 mmol) in toluene (250 mL) was irradiated with a medium pressure Hg lamp using a uranium filter (50% transmission at 350 nm) for 6 h with constant nitrogen flow through the reaction mixture. The resulting red solution was concentrated, and the reaction was repeated using an additional 1.10 g (1.68 mmol) of **14**. Once combined, the red solution was concentrated to dryness in vacuo, adsorbed onto silica gel, and subjected to silica gel chromatography with hexanes/EtOAc (6:1 to 1:1). The combined product eluents were dried in vacuo to provide pyridocarbazole **7** (1.60 g, 83%) as a yellow solid. <sup>1</sup>H NMR (300 MHz, CDCl<sub>3</sub>): δ (ppm) 10.48 (s, 1H), 9.42 (dd, *J* = 8.5, 1.7 Hz, 1H), 8.99 (dd, *J* = 4.3, 1.7 Hz, 1H), 8.69 (d, *J* = 2.5 Hz, 1H), 7.63 (dd, *J* = 8.5, 4.3 Hz, 1H), 7.41 (d, *J* = 8.7 Hz, 1H), 7.18 (dd, *J* = 8.7, 2.5 Hz, 1H), 1.46–1.34 (m, 3H), 1.18 (d, *J* = 7.3 Hz, 18H), 1.07 (s, 9H), 0.63 (s, 6H). <sup>13</sup>C NMR (75 MHz, CDCl<sub>3</sub>): δ (ppm) 175.7, 174.1, 151.5, 150.4, 140.8, 138.7, 134.9, 134.6, 131.0, 123.3, 122.9, 122.1, 121.5, 120.4, 115.4, 114.6, 111.9, 26.7, 19.3, 18.3, 12.9, -3.9. IR ν (cm<sup>-1</sup>) 3456, 2946, 2866, 1751, 1695, 1527, 1473, 1338, 1314, 1281, 1216, 1182, 905, 827, 795. HRMS calcd for C<sub>32</sub>H<sub>44</sub>N<sub>3</sub>O<sub>3</sub>Si<sub>2</sub> (M + H)<sup>+</sup> 574.2916, found (M + H)<sup>+</sup> 574.2926.

**Compounds 15 and 16.** A solution of pyridocarbazole **7** (510 mg, 0.890 mmol) in THF-EtOH (1:1) (51 mL) was cooled to 0 °C. Sodium borohydride (68 mg, 1.79 mmol) was added every hour for a total of 8 h, and the stirred reaction mixture was then allowed to naturally warm to room temperature overnight. The resulting yellow reaction mixture was again cooled to 0 °C, carefully quenched with addition of saturated aqueous NH<sub>4</sub>Cl, and extracted with EtOAc. The combined organic layers were washed with brine, dried using Na<sub>2</sub>SO<sub>4</sub>, filtered, and concentrated to dryness in vacuo. The crude material was adsorbed onto silica gel and subjected to silica gel chromatography with hexanes/EtOAc (6:1 to 3:1). The combined product eluents were dried in vacuo to individually provide pure **15** (320 mg, 63%) and **16** (110 mg, 21%). The constitutions of **15** and **16** were assigned by ROESY analysis (see Figure 3).

**Compound 15.** <sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>): δ (ppm) 12.55 (s, 1H), 9.21 (d, *J* = 1.7 Hz, 1H), 9.07 (dd, *J* = 4.3, 1.7 Hz, 1H), 8.66 (d, *J* = 2.6 Hz, 1H), 8.59 (dd, *J* = 8.4, 1.7 Hz, 1H), 7.73 (dd, *J* = 8.4, 4.3 Hz, 1H), 7.55 (d, *J* = 8.7 Hz, 1H), 7.07 (dd, *J* = 8.7, 2.6 Hz, 1H), 6.67 (d, *J* = 1.8 Hz, 1H), 1.39–1.27 (m, 3H), 1.11 (d, *J* = 7.4 Hz, 18H), 0.81 (s, 9H), 0.21 (s, 3H), -0.21 (s, 3H). <sup>13</sup>C NMR (100 MHz, DMSO-*d*<sub>6</sub>): δ (ppm) 170.3, 150.1, 149.0, 138.6, 137.3, 135.6, 134.7, 133.2, 125.6, 122.4, 121.1, 120.9, 119.5, 114.0, 113.4, 112.4, 79.1, 25.6, 17.9, 17.6, 12.1, -4.0. IR ν (cm<sup>-1</sup>) 3293, 3075, 2928, 2862, 1695, 1472, 1272,

1216, 1079, 1039, 981, 883, 832, 789, 668, 484. HRMS calcd for C<sub>32</sub>H<sub>45</sub>N<sub>3</sub>O<sub>3</sub>Si<sub>2</sub>Na (M + Na)<sup>+</sup> 598.2892, found (M + Na)<sup>+</sup> 598.2897.

**Compound 16.** <sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>): δ (ppm) 12.76 (s, 1H), 9.41 (dd, *J* = 8.4, 1.7 Hz, 1H), 9.03 (dd, *J* = 4.3, 1.7 Hz, 1H), 9.00 (d, *J* = 2.0 Hz, 1H), 7.76 (d, *J* = 2.4 Hz, 1H), 7.72 (dd, *J* = 8.4, 4.3 Hz, 1H), 7.59 (d, *J* = 8.8 Hz, 1H), 7.09 (dd, *J* = 8.7, 2.5 Hz, 1H), 6.63 (d, *J* = 2.0 Hz, 1H), 1.36–1.26 (m, 3H), 1.10 (d, *J* = 6.7 Hz, 18H), 0.73 (s, 9H), 0.03 (s, 3H), -0.50 (s, 3H). <sup>13</sup>C NMR (100 MHz, DMSO-*d*<sub>6</sub>): δ (ppm) 170.8, 149.4, 149.0, 143.6, 138.9, 137.2, 134.7, 131.9, 122.3, 122.1, 119.0, 115.4, 114.6, 112.7, 111.9, 78.7, 25.6, 17.8, 17.6, 12.1, -3.4, -3.9. IR ν (cm<sup>-1</sup>) 3156, 3071, 2945, 2927, 2865, 1686, 1670, 1530, 1471, 1280, 1251, 1213, 1094, 984, 957, 878, 831, 792, 780, 682, 665. HRMS calcd for C<sub>32</sub>H<sub>46</sub>N<sub>3</sub>O<sub>3</sub>Si<sub>2</sub> (M + H)<sup>+</sup> 576.3072, found (M + H)<sup>+</sup> 576.3080.

**Compound 17.** A yellow suspension of **15** (260 mg, 0.452 mmol) in CH<sub>2</sub>Cl<sub>2</sub> (23.4 mL) containing 0.1% ethanol was cooled to 0 °C. Trifluoroacetic acid (2.6 mL) was then carefully added, and the resulting orange solution was stirred at 0 °C for 10 min. The reaction mixture was then carefully quenched with addition of saturated aqueous NaHCO<sub>3</sub> and extracted with CH<sub>2</sub>Cl<sub>2</sub>. The combined organic layers were washed with brine, dried using Na<sub>2</sub>SO<sub>4</sub>, filtered, and concentrated to dryness in vacuo to provide **17** (220 mg, 99%) as a light yellow solid. See Supporting Information for a ROESY analysis. <sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>): δ (ppm) 12.57 (s, 1H), 9.28 (d, *J* = 1.3 Hz, 1H), 9.07 (dd, *J* = 4.3, 1.7 Hz, 1H), 8.65 (d, *J* = 2.6 Hz, 1H), 8.63 (dd, *J* = 8.3, 1.7 Hz, 1H), 7.73 (dd, *J* = 8.4, 4.3 Hz, 1H), 7.55 (d, *J* = 8.7 Hz, 1H), 7.07 (dd, *J* = 8.7, 2.6 Hz, 1H), 6.43 (d, *J* = 1.4 Hz, 1H), 3.67 (m, 1H), 3.39 (m, 1H), 1.40–1.27 (m, 3H), 1.15 (t, *J* = 7.0 Hz, 3H), 1.11 (d, *J* = 7.3 Hz, 18H). <sup>13</sup>C NMR (100 MHz, DMSO-*d*<sub>6</sub>): δ (ppm) 170.3, 150.1, 149.0, 138.3, 137.5, 134.7, 132.9, 132.7, 126.6, 122.4, 121.5, 121.1, 119.5, 114.1, 113.4, 112.4, 83.5, 60.7, 17.9, 15.3, 12.1. IR ν (cm<sup>-1</sup>) 3284, 3172, 3072, 2942, 2864, 1689, 1471, 1338, 1280, 1260, 1214, 1069, 1049, 881, 814, 792, 726, 682, 667, 621, 483. HRMS calcd for C<sub>28</sub>H<sub>35</sub>N<sub>3</sub>O<sub>3</sub>SiNa (M + Na)<sup>+</sup> 512.2340, found (M + Na)<sup>+</sup> 512.2338.

**Compound 5.** A suspension of **17** (220 mg, 0.450 mmol) in CH<sub>2</sub>Cl<sub>2</sub> was purged with nitrogen for approximately 15 min. Phenylselenol (191 μL, 1.80 mmol) followed by trifluoroacetic acid (52 μL, 0.675 mmol) were then added, and the resulting dark orange solution was heated to reflux for 5 h. The reaction mixture was then cooled to room temperature and carefully quenched with addition of saturated aqueous NaHCO<sub>3</sub>. The CH<sub>2</sub>Cl<sub>2</sub> was then removed in vacuo, and the resulting yellow precipitate was collected, washed with twice with water, followed by diethyl ether (2×) to provide lactam **5** (153 mg, 77%) as a light yellow solid. <sup>1</sup>H NMR (400 MHz, 100 °C, DMSO-*d*<sub>6</sub>): δ (ppm) 12.06 (s, 1H), 9.05 (dd, *J* = 4.0, 1.3 Hz, 1H), 8.77 (s, 1H), 8.53 (dd, *J* = 8.3, 1.1 Hz, 1H), 8.34 (s, 1H), 7.67 (dd, *J* = 8.1, 4.1 Hz, 1H), 7.57 (d, *J* = 8.7 Hz, 1H), 7.05 (d, *J* = 8.4 Hz, 1H), 4.85 (s, 2H), 1.38 (m, 3H), 1.16 (d, *J* = 7.3 Hz, 18H). <sup>13</sup>C NMR (100 MHz, 100 °C, DMSO-*d*<sub>6</sub>): δ (ppm) 171.1, 149.3, 148.4, 138.1, 135.9, 134.3, 134.1, 131.8, 126.2, 122.4, 120.7, 120.5, 118.6, 114.5, 113.4, 111.6, 44.2, 17.4, 11.7. IR ν (cm<sup>-1</sup>) 3283, 3163, 3060, 2940, 2865, 1680, 1661, 1469, 1261, 1214, 1171, 1058, 978, 914, 879, 671, 658, 623, 553, 457. HRMS calcd for C<sub>26</sub>H<sub>32</sub>N<sub>3</sub>O<sub>2</sub>Si (M + H)<sup>+</sup> 446.2258, found (M + H)<sup>+</sup> 446.2258.

**Compound 6.** A suspension of **16** (45 mg, 0.078 mmol) in CH<sub>2</sub>Cl<sub>2</sub> (3.6 mL) containing 0.1% ethanol was cooled to 0 °C. Trifluoroacetic acid (0.9 mL) was then carefully added, and the resulting orange solution was allowed to naturally warm to room temperature, stirring for a total of 3 h. Following thin layer chromatography analysis, the reaction mixture was purged with nitrogen for approximately 15 min. Phenylselenol (34 μL, 0.312 mmol) was then added, and the reaction was heated to

reflux overnight. The reaction mixture was then cooled to room temperature and carefully quenched with addition of saturated aqueous NaHCO<sub>3</sub>. The CH<sub>2</sub>Cl<sub>2</sub> was then removed in vacuo, and the resulting yellow precipitate was collected and washed twice with water, followed by diethyl ether (2×) to provide lactam **6** (32 mg, 94% over 2 steps) as a white solid. <sup>1</sup>H NMR (500 MHz, 100 °C, DMSO-*d*<sub>6</sub>): δ (ppm) 12.35 (s, 1H), 9.53 (d, *J* = 8.4 Hz, 1H), 8.99 (dd, *J* = 4.2, 1.5 Hz, 1H), 8.13 (s, 1H), 7.69–7.64 (m, 2H), 7.46 (d, *J* = 2.0 Hz, 1H), 7.10 (dd, *J* = 8.7, 2.2 Hz, 1H), 4.89 (s, 2H), 1.35 (septet, *J* = 7.4, 3H), 1.15 (d, *J* = 7.4 Hz, 18H). IR ν (cm<sup>-1</sup>) 3156, 3063, 2943, 2865, 1678, 1652, 1533, 1463, 1279, 1244, 1209, 1174, 964, 868, 809, 787, 672, 568, 434. HRMS calcd for C<sub>26</sub>H<sub>31</sub>N<sub>3</sub>O<sub>2</sub>SiNa (M + Na)<sup>+</sup> 468.2078, found (M + Na)<sup>+</sup> 468.2083.

**Compound 19.** To a vacuum-dried solid mixture of lactam **5** (20 mg, 0.045 mmol), ruthenium precursor **4**<sup>26</sup> (21 mg, 0.050 mmol), and K<sub>2</sub>CO<sub>3</sub> (7.0 mg, 0.050 mmol) was added freshly distilled DMF (2.0 mL), and the resulting suspension was purged with nitrogen for approximately 10 min. The reaction mixture was then heated to 80 °C for 10 min in a microwave reactor (CEM Discover). The resulting dark red solution was concentrated using high vacuum and coevaporated once with EtOAc. The crude material was subjected to silica gel chromatography with toluene:acetone (3:1). The combined product eluents were dried in vacuo to provide complex **19** (23 mg, 79%) as an orange film. <sup>1</sup>H NMR (300 MHz, CDCl<sub>3</sub>): δ (ppm) 8.86 (dd, *J* = 5.1, 1.1 Hz, 1H), 8.56 (d, *J* = 2.4 Hz, 1H), 8.17 (dd, *J* = 8.2, 1.1 Hz, 1H), 7.32–7.28 (m, 2H), 7.15 (dd, *J* = 8.8, 2.5 Hz, 1H), 6.92 (s, 1H), 5.20 (s, 5H), 4.76 (s, 2H), 1.48–1.38 (m, 3H), 1.21 (d, *J* = 7.2 Hz, 18H). <sup>13</sup>C NMR (100 MHz, CDCl<sub>3</sub>): δ (ppm) 200.6, 173.0, 154.1, 151.1, 148.7, 147.9, 146.1, 131.7, 129.3, 128.0, 124.5, 121.0, 119.6, 119.4, 115.8, 114.7, 114.0, 80.9, 45.1, 18.4, 13.0. IR ν (cm<sup>-1</sup>) 3183, 3077, 2942, 2925, 2865, 1951, 1731, 1686, 1607, 1544, 1462, 1405, 1267, 1244, 1210, 1189, 1127, 1056, 997, 977, 922, 896, 884, 835, 805, 791, 688. HRMS calcd for C<sub>32</sub>H<sub>36</sub>N<sub>3</sub>O<sub>3</sub>RuSi (M + H)<sup>+</sup> 640.1564, found (M + H)<sup>+</sup> 640.1561.

**Compound 20.** To a vacuum-dried solid mixture of lactam **6** (10 mg, 0.023 mmol), ruthenium precursor **4** (11 mg, 0.025 mmol) and K<sub>2</sub>CO<sub>3</sub> (3.5 mg, 0.025 mmol) was added freshly distilled DMF (1.0 mL) and the resulting suspension was purged with nitrogen for approximately 10 min. The reaction mixture was then heated to 80 °C for 10 min in a microwave reactor (CEM Discover). The resulting yellow-orange solution was concentrated using high vacuum and coevaporated once with EtOAc. The crude material was subjected to silica gel chromatography with toluene/acetone (3:1). The combined product eluents were dried in vacuo to provide complex **20** (13 mg, 87%) as a yellow solid. <sup>1</sup>H NMR (300 MHz, CDCl<sub>3</sub>): δ (ppm) 9.41 (dd, *J* = 8.3, 1.2 Hz, 1H), 8.81 (dd, *J* = 5.2, 1.2 Hz, 1H), 7.48 (d, *J* = 2.4 Hz, 1H), 7.40 (dd, *J* = 8.3, 5.2 Hz, 1H), 7.33 (d, *J* = 8.8 Hz, 1H), 7.13 (dd, *J* = 8.8, 2.4 Hz, 1H), 6.26 (s, 1H), 5.19 (s, 5H), 4.95 (s, 2H), 1.40–1.28 (m, 3H), 1.17 (d, *J* = 7.2 Hz, 18H). <sup>13</sup>C NMR (125 MHz, CDCl<sub>3</sub>): δ (ppm) 200.4, 173.7, 153.2, 153.1, 149.6, 147.8, 145.2, 145.0, 133.2, 124.6, 123.3, 121.0, 118.6, 115.7, 115.2, 111.9, 110.7, 80.8, 45.1, 18.3, 13.0. IR ν (cm<sup>-1</sup>) 2943, 2865, 1951, 1676, 1608, 1583, 1520, 1456, 1432, 1421, 1397, 1334, 1275, 1247, 1207, 1189, 923, 882, 685. HRMS calcd for C<sub>32</sub>H<sub>36</sub>N<sub>3</sub>O<sub>3</sub>RuSi (M + H)<sup>+</sup> 640.1564, found (M + H)<sup>+</sup> 640.1537.

**Compound 1.** To a vacuum-dried sample of complex **19** (23 mg, 0.036 mmol) was added freshly distilled THF (2.3 mL), and the resulting solution was purged with nitrogen while cooling to 0 °C. Tetrabutylammonium fluoride (1 M solution in THF, dried over 4 Å molecular sieves) (47 μL, 0.047 mmol) was then added, and the reaction mixture immediately became a dark red suspension. Within 1 min, the reaction was quenched with excess saturated aqueous NH<sub>4</sub>Cl (orange color returns) and extracted

with EtOAc. The combined organic layers were dried using Na<sub>2</sub>SO<sub>4</sub>, filtered, and concentrated to dryness in vacuo. The crude material was then subjected to silica gel chromatography with CH<sub>2</sub>Cl<sub>2</sub>/MeOH (20:1). The combined product eluents were dried in vacuo to provide complex **1** (13 mg, 76%) as a light red solid. <sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>): δ (ppm) 9.19 (dd, *J* = 5.1, 1.1 Hz, 1H), 8.79 (s, 1H), 8.72 (s, 1H), 8.56 (dd, *J* = 8.2, 1.1 Hz, 1H), 8.20 (d, *J* = 2.2 Hz, 1H), 7.57 (dd, *J* = 8.3, 5.2 Hz, 1H), 7.31 (d, *J* = 8.7 Hz, 1H), 6.93 (dd, *J* = 8.8, 2.6 Hz, 1H), 5.48 (s, 5H), 4.80 (s, 2H). <sup>13</sup>C NMR (125 MHz, DMSO-*d*<sub>6</sub>): δ (ppm) 201.7, 171.6, 155.3, 150.0, 149.4, 146.2, 144.9, 132.7, 128.9, 128.7, 124.0, 120.6, 120.2, 115.0, 114.9, 114.2, 108.4, 81.6, 44.6. IR ν (cm<sup>-1</sup>) 3263, 3108, 2930, 2856, 1940, 1680, 1608, 1473, 1406, 1223, 1186, 1048, 1024, 1004, 808, 793, 698. HRMS calcd for C<sub>23</sub>H<sub>15</sub>N<sub>3</sub>O<sub>3</sub>RuNa (M + Na)<sup>+</sup> 506.0049, found (M + Na)<sup>+</sup> 506.0055.

**Compound 2.** To a vacuum-dried sample of complex **20** (11 mg, 0.017 mmol) was added freshly distilled THF (1.1 mL), and the resulting solution was purged with nitrogen while cooling to -40 °C. Ice-cold nitrogen-purged tetrabutylammonium fluoride (1 M solution in THF, dried over 4 Å molecular sieves) (19 μL, 0.019 mmol) was then added, and the reaction mixture immediately became a dark suspension. Within 1 min, the reaction was quenched with excess glacial acetic acid, and the resulting yellow solution was concentrated to dryness in vacuo. The crude material was then subjected to silica gel chromatography with CH<sub>2</sub>Cl<sub>2</sub>/MeOH (20:1). The combined product eluents were dried in vacuo to provide complex **2** (5.8 mg, 71%) as a yellow orange solid. <sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>): δ (ppm) 9.25 (dd, *J* = 8.3, 1.2 Hz, 1H), 9.11 (dd, *J* = 5.2, 1.2 Hz, 1H), 8.95 (s, 1H), 8.31 (s, 1H), 7.57 (dd, *J* = 8.3, 5.2 Hz, 1H), 7.39–7.36 (m, 2H), 6.99 (dd, *J* = 8.8, 2.3 Hz, 1H), 5.46 (s, 5H), 4.84 (s, 2H). <sup>13</sup>C NMR (100 MHz, DMSO-*d*<sub>6</sub>): δ (ppm) 201.6, 172.0, 154.0, 151.5, 150.5, 146.1, 144.7, 143.8, 131.7, 124.3, 122.1, 121.1, 115.8, 114.6, 114.2, 111.7, 105.6, 81.6, 44.3. IR ν (cm<sup>-1</sup>) 3232, 3108, 2928, 2854, 1938, 1651, 1606, 1583, 1520, 1452, 1401, 1334, 1216, 1184, 1024, 1004, 814, 789, 715, 693, 559. HRMS calcd for C<sub>23</sub>H<sub>16</sub>N<sub>3</sub>O<sub>3</sub>Ru (M + H)<sup>+</sup> 484.0230, found (M + H)<sup>+</sup> 484.0233.

**Resolution of (*R*<sub>Ru</sub>)-1 and (*S*<sub>Ru</sub>)-1.** Baseline separation of the enantiomers of half sandwich complex **19** was achieved using the Chiral Pak 1B analytical HPLC column (Daicel/Chiral Technologies). First, **19** was dissolved in a minimal amount of methylene chloride and carefully filtered. Then, each injection was run using an ethanol gradient of 20% to 80% over 20 min in hexanes. A flow rate of 0.7 mL/min was used, and injection volumes of approximately 50 μL could be reached without compromising baseline separation. Each enantiomer was then treated separately with TBAF to provide both (*R*<sub>Ru</sub>)-1 and (*S*<sub>Ru</sub>)-1. Retention of enantiomeric purity was confirmed using the same Chiral Pak 1B analytical HPLC column. For this, 10 mM stock solutions (*R*<sub>Ru</sub>)-1 and (*S*<sub>Ru</sub>)-1 in DMSO were diluted with an equal volume of ethanol and carefully filtered. Then, analytical injections (5 μL) were run using an ethanol gradient of 50% to 95% over 20 min in hexanes with a flow rate of 0.7 mL/min. (*R*<sub>Ru</sub>)-1 and (*S*<sub>Ru</sub>)-1 were determined to be 99% and 98% enantiopure, respectively. Absolute configurations were assigned by CD-spectroscopy (see Supporting Information).

**Resolution of (*R*<sub>Ru</sub>)-2 and (*S*<sub>Ru</sub>)-2.** Baseline separation of the enantiomers of half sandwich complex **20** was achieved using the Chiral Pak 1B analytical HPLC column (Daicel/Chiral Technologies). First, **20** was dissolved in a minimal amount of methylene chloride and carefully filtered. Then, each injection was run using an acetone gradient of 25% to 75% over 20 min in hexanes. A flow rate of 0.7 mL/min was used and injection volumes of approximately 50 μL could be reached without compromising baseline separation. Additionally, a detection wavelength of 470 nm was used to avoid acetone absorption.

Each enantiomer was then treated separately with TBAF to provide both ( $R_{Ru}$ )-**2** and ( $S_{Ru}$ )-**2**. Retention of enantiomeric purity was confirmed using the same Chiral Pak 1B analytical HPLC column. First, 10 mM stock solutions ( $R_{Ru}$ )-**2** and ( $S_{Ru}$ )-**2** in DMSO were diluted with an equal volume of ethanol and carefully filtered. Then, analytical injections (5  $\mu$ L) were run using an acetone gradient of 40% to 75% over 2 h in hexanes with a flow rate of 0.7 mL/min and a detection wavelength of 470 nm. ( $R_{Ru}$ )-**2** and ( $S_{Ru}$ )-**2** were determined to be 99% and 98% enantiopure, respectively. Absolute configurations were assigned by CD-spectroscopy (see Supporting Information).

**5-(Triisopropylsilyloxy)indole (22).**<sup>27</sup> A solution of 5-benzoyloxyindole (2.50 g, 11.2 mmol) in EtOAc (100 mL) was purged with nitrogen for 15 min. Following thorough nitrogen purge, palladium (10 wt % on activated carbon, dry) (2.50 g) was added, and the reaction mixture was carefully purged with hydrogen gas for approximately 10 min. The reaction mixture was then stirred at room temperature under an atmosphere of hydrogen, monitoring by thin layer chromatography. After 45 min, the resulting dark reaction mixture was filtered through Celite with EtOAc, and the filtrate concentrated to dryness in vacuo. The resulting light yellow solid was then redissolved in DMF (40 mL) and purged with nitrogen while cooling to 0 °C. Then, *N,N*-diisopropylethylamine (9.70 mL, 56.0 mmol) was added dropwise. After stirring for 10 min at 0 °C, triisopropylsilyl trifluoromethanesulfonate (3.0 mL, 11.2 mmol) was added dropwise over 15 min, and the reaction mixture was stirred at room temperature for an additional 1 h. The reaction mixture was cooled to 0 °C, and excess 1 M  $NH_4OAc$  was carefully added. The mixture was then further diluted with water and extracted with EtOAc. The combined organic layers were washed with brine, dried using  $Na_2SO_4$ , filtered, and concentrated to dryness in vacuo. The crude material was subjected to a short flash column using only  $CH_2Cl_2$ . The combined product eluents were dried in vacuo to provide indole **22** (2.85 g, 87% over 2 steps) as a clear oil. <sup>1</sup>H NMR consistent with literature reported spectrum.<sup>27</sup>

**1-(tert-Butoxycarbonyl)-5-(triisopropylsilyloxy)indole (23).** A solution of indole **22** (2.85 g, 9.86 mmol) in THF (7.3 mL) was purged with nitrogen while cooling to 0 °C. To the solution was added di-*tert*-butyl dicarbonate (2.39 mL, 10.4 mmol) followed by the careful addition of solid (dimethylamino)pyridine (1.44 g, 10.4 mmol). The resulting yellow suspension was stirred overnight, naturally warming to room temperature. The reaction mixture was cooled to 0 °C, and 1 M HCl (7 mL) was carefully added. After stirring at room temperature for 5 min, the organic layer was separated, and the aqueous layer was extracted with EtOAc. The combined organic layers were washed with brine, dried using  $Na_2SO_4$ , filtered, and concentrated to dryness in vacuo. The crude material was adsorbed onto silica gel and subjected to silica gel chromatography with hexanes/EtOAc (20:1). The combined product eluents were dried in vacuo to provide **23** (3.76 g, 98%). <sup>1</sup>H NMR (300 MHz,  $CDCl_3$ ):  $\delta$  (ppm) 7.95 (d,  $J = 8.5$  Hz, 1H), 7.54 (d,  $J = 3.6$  Hz, 1H), 7.02 (d,  $J = 2.4$  Hz, 1H), 6.88 (dd,  $J = 8.9, 2.5$  Hz, 1H), 6.45 (d,  $J = 3.7$  Hz, 1H), 1.66 (s, 9H), 1.33–1.21 (m, 3H), 1.11 (d,  $J = 6.9$  Hz, 18H). <sup>13</sup>C NMR (75 MHz,  $CDCl_3$ ):  $\delta$  (ppm) 152.0, 150.0, 131.7, 130.4, 126.6, 117.7, 115.7, 110.8, 107.3, 83.6, 28.4, 18.2, 12.9. IR  $\nu$  ( $cm^{-1}$ ) 2941, 2866, 1723, 1458, 1376, 1327, 1280, 1257, 1217, 1183, 1150, 1122, 1078, 962, 877, 857, 811, 764, 725, 685, 637, 500, 478. HRMS calcd for  $C_{22}H_{36}NO_3Si$  ( $M + H$ )<sup>+</sup> 390.2459, found ( $M + H$ )<sup>+</sup> 390.2458.

**Potassium-1-(tert-butoxycarbonyl)-5-(triisopropylsilyloxy)indol-2-yl-trifluoroborate (24).** Indole **23** (3.04 g, 7.81 mmol) was dried under high vacuum and dissolved in freshly distilled THF

(10.2 mL). Following the addition of triisopropyl borate (2.73 mL, 11.8 mmol), the reaction mixture was cooled to 0 °C, and lithium diisopropylamide (2.0 M solution in heptane/tetrahydrofuran/ethylbenzene) (5.9 mL, 11.8 mmol) was added dropwise over 1 h. After stirring for an additional 30 min at 0 °C, 2 M HCl (20 mL) was carefully added and allowed to stir at room temperature for 10 min. The resulting layers were separated, and the aqueous layer was extracted using EtOAc. The combined organic layers were dried using  $Na_2SO_4$ , filtered, and concentrated to dryness in vacuo to provide 1-(*tert*-butoxycarbonyl)-5-(triisopropylsilyloxy)indol-2-yl-boronic acid as an orange foam, which due to instability was converted to the trifluoroborate **24** by reacting with  $KHF_2$  (5.7 mL, 4.5 M in water) in MeOH. After 30 min, the solution was evaporated to dryness, and the crude solid was dissolved in acetone, sonicated, and then separated from the remaining insoluble material by filtration. The combined filtrates were concentrated in vacuo, the crude product was redissolved in a minimal amount of hot acetone, and the subsequent addition of hexane led to the precipitation of the product, which was filtered, collected, and dried in vacuo to provide compound **24** as a white solid (2.11 g, 55%). <sup>1</sup>H NMR (300 MHz,  $DMSO-d_6$ ):  $\delta$  (ppm) 7.85 (d,  $J = 8.8$  Hz, 1H), 6.83 (d,  $J = 2.4$  Hz, 1H), 6.64 (dd,  $J = 8.8, 2.4$  Hz, 1H), 6.30 (s, 1H), 1.55 (s, 9H), 1.27–1.19 (m, 3H), 1.06 (d,  $J = 7.1$  Hz, 18H). <sup>13</sup>C NMR (75 MHz,  $DMSO-d_6$ ):  $\delta$  (ppm) 151.2, 150.1, 132.6, 131.7, 114.9, 114.4, 111.6, 108.8, 81.1, 27.7, 17.8, 12.0 (C–B not observed). IR  $\nu$  ( $cm^{-1}$ ) 2968, 2943, 2867, 1699, 1458, 1366, 1303, 1256, 1188, 1127, 984, 969, 884, 873, 851, 689, 626, 478. HRMS calcd for  $C_{22}H_{34}BF_3NO_3Si$  ( $M - K$ )<sup>-</sup> 456.2359, found ( $M - K$ )<sup>-</sup> 456.2348.

**Compound 26.** To a vacuum-dried solid mixture of trifluoroborate **24** (2.23 g, 4.51 mmol), triflate **25** (1.00 g, 3.61 mmol), tetrakis(triphenylphosphine)palladium(0) (416 mg, 0.36 mmol), and  $Na_2CO_3$  (1.05 g, 9.93 mmol) was added freshly distilled anhydrous THF (20 mL). After purging with nitrogen for 15 min, the reaction mixture was heated to reflux overnight. The resulting orange reaction mixture was cooled to room temperature, diluted with water, and extracted with EtOAc. The combined organic layers were washed with brine, dried using  $Na_2SO_4$ , filtered, and concentrated to dryness in vacuo. The crude material was adsorbed onto silica gel and purified by chromatography using hexanes/EtOAc (10:1). The combined product eluents were dried in vacuo to provide compound **26** (1.17 g, 78%) as a white solid. <sup>1</sup>H NMR (300 MHz,  $CDCl_3$ ): 9.71 (s, 1H), 9.22 (s, 1H), 8.10 (s, 1H), 7.94 (d,  $J = 8.1$  Hz, 1H), 7.84 (d,  $J = 8.3$  Hz, 1H), 7.68 (ddd,  $J = 8.2, 6.9, 1.3$  Hz, 1H), 7.54 (ddd,  $J = 8.1, 6.9, 1.2$  Hz, 1H), 7.25 (d,  $J = 8.6$  Hz, 1H), 7.14 (d,  $J = 2.3$  Hz, 1H), 7.00 (dd,  $J = 2.1, 0.9$  Hz, 1H), 6.84 (dd,  $J = 8.7, 2.3$  Hz, 1H), 1.37–1.25 (m, 3H), 1.15 (d,  $J = 6.9$  Hz, 18H). <sup>13</sup>C NMR (75 MHz,  $CDCl_3$ ):  $\delta$  (ppm) 152.3, 150.3, 144.7, 138.0, 136.8, 132.3, 131.0, 130.3, 128.0, 128.0, 127.1, 126.9, 117.4, 115.5, 111.8, 109.9, 99.3, 18.3, 13.0. IR  $\nu$  ( $cm^{-1}$ ) 3409, 2944, 2866, 1621, 1575, 1470, 1447, 1407, 1258, 1211, 1182, 1143, 1123, 991, 960, 880, 853, 789, 752, 684, 641, 623, 575, 548, 502, 471. HRMS calcd for  $C_{26}H_{33}N_2OSi$  ( $M + H$ )<sup>+</sup> 417.2357, found ( $M + H$ )<sup>+</sup> 417.2355.

**Compound 27.** A solution of isoquinolinyl-indole **26** (750 mg, 1.80 mmol) in freshly distilled THF (4.7 mL) was prepared after drying **26** under high vacuum. After cooling the solution to –15 °C, lithium bis(trimethylsilyl)amide (1 M solution in hexanes) (5.4 mL, 5.4 mmol) was added dropwise over 15 min, and the resulting orange solution was stirred at –15 °C for an additional 45 min. A solution of vacuum-dried maleimide **10** (697 mg, 1.89 mmol) in freshly distilled THF (4.7 mL) cooled to 0 °C was added, and the reaction mixture was stirred at –15 °C for 20 min and room temperature overnight. The resulting dark purple reaction mixture was carefully poured into stirring ice-cold water (200 mL) and extracted with EtOAc. The combined organic layers were washed with brine, dried using  $Na_2SO_4$ ,

(27) Kondo, Y.; Kojima, S.; Sakamoto, T. *J. Org. Chem.* **1997**, *62*, 6507–6511.



filtered, and concentrated to dryness in vacuo. The crude material was adsorbed onto silica gel and subjected to silica gel chromatography with hexanes/EtOAc (10:1 to 8:1). The combined product eluents were dried in vacuo to provide the unstable monobromide **27** (1.0 g, 79%) as an orange solid.  $^1\text{H}$  NMR (300 MHz,  $\text{CDCl}_3$ ):  $\delta$  (ppm) 10.00 (s, 1H), 9.21 (s, 1H), 7.94 (d,  $J = 8.1$  Hz, 1H), 7.71–7.66 (m, 2H), 7.61–7.57 (m, 2H), 7.26 (d,  $J = 8.6$  Hz, 1H), 7.00 (d,  $J = 2.0$  Hz, 1H), 6.86 (d,  $J = 8.3$  Hz, 1H), 1.30 (septet,  $J = 7.4$  Hz, 3H), 1.14 (d,  $J = 7.4$  Hz, 18H), 0.98 (s, 9H), 0.47 (s, 6H). IR  $\nu$  ( $\text{cm}^{-1}$ ) 3335, 2942, 2892, 2863, 1697, 1648, 1625, 1580, 1466, 1431, 1312, 1266, 1217, 1182, 1076, 977, 897, 879, 844, 824, 795, 742, 709, 680, 619, 582, 556, 467, 405. HRMS calcd for  $\text{C}_{36}\text{H}_{47}\text{BrN}_3\text{O}_3\text{Si}_2$  ( $\text{M} + \text{H}$ ) $^+$  706.2313, found ( $\text{M} + \text{H}$ ) $^+$  706.2313.

**Compound 28.** An orange suspension of monobromide **27** (800 mg, 1.14 mmol) in toluene (250 mL) was irradiated with a medium pressure Hg lamp using an uranium filter (50% transmission at 350 nm) for 4 h with constant nitrogen flow through the reaction mixture. The resulting bright orange suspension was concentrated to dryness in vacuo, adsorbed onto silica gel, and subjected to silica gel chromatography with hexanes/EtOAc (6:1 to 3:1). The combined product eluents were dried in vacuo to provide heterocycle **28** (610 mg, 86%) as a yellow solid.  $^1\text{H}$  NMR (300 MHz,  $\text{CDCl}_3$ ):  $\delta$  (ppm) 10.23 (d,  $J = 8.6$  Hz, 1H), 9.95 (s, 1H), 9.31 (s, 1H), 8.87 (d,  $J = 2.5$  Hz, 1H), 8.09 (d,  $J = 8.0$  Hz, 1H), 8.00 (ddd,  $J = 8.6, 7.1, 1.5$  Hz, 1H), 7.98 (ddd,  $J = 8.0, 7.0, 0.9$  Hz, 1H), 7.46 (d,  $J = 8.7$  Hz, 1H), 7.21 (dd,  $J = 8.7, 2.5$  Hz, 1H), 1.49–1.39 (m, 3H), 1.20 (d,  $J = 7.3$  Hz, 18H), 1.09 (s, 9H), 0.68 (s, 6H).  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ ):  $\delta$  (ppm) 175.4, 173.9, 154.3, 151.2, 142.6, 135.5, 135.2, 132.2, 131.7, 129.9, 128.8, 128.5, 127.8, 123.1, 121.6, 120.4, 116.1, 115.6, 111.6, 26.8, 19.5, 18.3, 12.9, –3.5 (two hidden carbons). IR  $\nu$  ( $\text{cm}^{-1}$ ) 2939, 2862, 1686, 1482, 1463, 1362, 1333, 1301, 1276, 1248, 1212, 1178, 1156, 1080, 976, 884, 866, 846, 821, 787, 753, 682, 665, 607, 583, 478, 407. HRMS calcd for  $\text{C}_{36}\text{H}_{46}\text{N}_3\text{O}_3\text{Si}_2$  ( $\text{M} + \text{H}$ ) $^+$  624.3072, found ( $\text{M} + \text{H}$ ) $^+$  624.3072.

**Compound 29.** A solution of imide **28** (135 mg, 0.217 mmol) in THF-MeOH (1:1) (14 mL) was cooled to 0 °C. Sodium borohydride (16 mg, 0.421 mmol) was added every hour for a total of 5 h, and the stirred reaction mixture was then allowed to naturally warm to room temperature overnight. The reaction mixture was again cooled to 0 °C, carefully quenched with addition of saturated aqueous  $\text{NH}_4\text{Cl}$ , and further diluted with water and extracted with  $\text{CH}_2\text{Cl}_2$ . The combined organic layers were washed with brine, dried using  $\text{Na}_2\text{SO}_4$ , filtered, and concentrated to dryness. The crude material was adsorbed onto silica gel and subjected to silica gel chromatography with hexanes/EtOAc (6:1 to 3:1) to provide a yellow solid. A yellow suspension of this solid (100 mg) in  $\text{CH}_2\text{Cl}_2$  (9.0 mL) containing 0.1% ethanol was cooled to 0 °C, trifluoroacetic acid (1.0 mL) was then carefully added, and the resulting orange solution was stirred at 0 °C for 15 min. The reaction mixture was then carefully quenched with saturated aqueous  $\text{NaHCO}_3$  and extracted with  $\text{CH}_2\text{Cl}_2$ . The combined organic layers were washed with brine, dried using  $\text{Na}_2\text{SO}_4$ , filtered, and concentrated to dryness in vacuo. A suspension of the crude material in  $\text{CH}_2\text{Cl}_2$  (10 mL) was purged with nitrogen for 15 min. Phenylselenol (68  $\mu\text{L}$ , 0.640 mmol) followed by trifluoroacetic acid (18.4  $\mu\text{L}$ , 0.240 mmol) were added, and the resulting dark orange solution was heated to reflux for 5 h. The reaction mixture was cooled to room temperature, carefully quenched with saturated aqueous  $\text{NaHCO}_3$ , and extracted with  $\text{CH}_2\text{Cl}_2$ . The combined organic layers were washed with brine, dried using  $\text{Na}_2\text{SO}_4$ , filtered, and concentrated to dryness in vacuo. The crude material was adsorbed onto silica gel and subjected to silica gel chromatography first with hexanes/EtOAc (3.5:1) and followed using  $\text{CH}_2\text{Cl}_2/\text{MeOH}$  (80:1). The combined product eluents were

dried in vacuo to provide lactam **29** (51 mg, 47% over 3 steps) as a light yellow solid. The constitution of **29** was confirmed by ROESY analysis.  $^1\text{H}$  NMR (300 MHz,  $\text{DMSO}-d_6$ ):  $\delta$  (ppm) 12.27 (s, 1H), 9.60 (s, 1H), 8.94 (s, 1H), 8.90 (d,  $J = 2.5$  Hz, 1H), 8.53 (d,  $J = 8.5$  Hz, 1H), 8.42 (d,  $J = 7.9$  Hz, 1H), 8.09 (t,  $J = 7.4$  Hz, 1H), 7.90 (t,  $J = 7.4$  Hz, 1H), 7.59 (d,  $J = 8.7$  Hz, 1H), 7.09 (dd,  $J = 8.7, 2.6$  Hz, 1H), 5.25 (s, 2H), 1.40–1.34 (m, 3H), 1.14 (d,  $J = 7.3$  Hz, 18H).  $^{13}\text{C}$  NMR (125 MHz,  $\text{DMSO}-d_6$ ):  $\delta$  (ppm) 171.0, 168.3, 153.5, 148.5, 137.9, 135.3, 132.6, 132.2, 132.1, 129.6, 127.3, 127.0, 126.7, 124.9, 122.3, 119.5, 116.5, 115.8, 114.5, 111.9, 48.6, 17.9, 12.1. IR  $\nu$  ( $\text{cm}^{-1}$ ) 3187, 3069, 2941, 2893, 2863, 1686, 1665, 1623, 1572, 1461, 1369, 1258, 1209, 1185, 1159, 1077, 1040, 1015, 998, 974, 881, 801, 745, 710, 679, 627, 600, 559, 503, 449. HRMS calcd for  $\text{C}_{30}\text{H}_{34}\text{N}_3\text{O}_2\text{Si}$  ( $\text{M} + \text{H}$ ) $^+$  496.2415, found ( $\text{M} + \text{H}$ ) $^+$  496.2427.

**Ruthenium Complex 31.** To a vacuum-dried solid mixture of lactam **29** (10 mg, 0.020 mmol), ruthenium precursor **30** $^5$  (11 mg, 0.022 mmol), and  $\text{K}_2\text{CO}_3$  (3.0 mg, 0.022 mmol) was added freshly distilled DMF (1.0 mL) and the resulting suspension was purged with nitrogen for approximately 10 min. The reaction mixture was then heated to 80 °C for 10 min in a microwave reactor (CEM Discover). The resulting dark red solution was concentrated using high vacuum and coevaporated once with EtOAc. The crude material was subjected to silica gel chromatography with toluene:acetone (3:1). The combined product eluents were dried in vacuo to provide ruthenium complex **31** (8.3 mg, 54%) as a red film.  $^1\text{H}$  NMR (300 MHz,  $\text{CDCl}_3$ ):  $\delta$  (ppm) 9.34 (s, 1H), 8.72 (d,  $J = 2.5$  Hz, 1H), 8.31 (d,  $J = 8.1$  Hz, 1H), 8.12 (d,  $J = 8.1$  Hz, 1H), 7.96 (t,  $J = 7.4$  Hz, 1H), 7.76 (t,  $J = 7.4$  Hz, 1H), 7.30 (d,  $J = 8.7$  Hz, 1H), 7.20 (dd,  $J = 8.7, 2.5$  Hz, 1H), 6.77 (s, 1H), 6.02 (m, 1H), 5.89 (m, 1H), 5.46 (m, 1H), 5.34 (m, 1H), 5.19 (d,  $J = 5.4$  Hz, 2H), 3.75 (s, 3H), 1.48–1.38 (m, 3H), 1.22 (d,  $J = 7.2$  Hz, 18H).  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ ):  $\delta$  (ppm) 199.1, 172.7, 167.2, 157.9, 151.7, 148.3, 147.8, 140.3, 133.3, 132.9, 129.4, 128.5, 127.3, 127.0, 126.6, 124.3, 124.1, 119.7, 117.3, 116.2, 114.6, 113.7, 93.8, 86.0, 85.3, 77.7, 75.3, 52.2, 48.5, 18.2, 12.8. IR  $\nu$  ( $\text{cm}^{-1}$ ) 2923, 2854, 1962, 1714, 1684, 1601, 1457, 1440, 1380, 1289, 1271, 1251, 1209, 1144, 884, 755. HRMS calcd for  $\text{C}_{37}\text{H}_{40}\text{N}_3\text{O}_4\text{RuSi}$  ( $\text{M} - \text{CO} + \text{H}$ ) $^+$  720.1826, found ( $\text{M} - \text{CO} + \text{H}$ ) $^+$  720.1823.

**Ruthenium Complex 21.** To a vacuum-dried sample of complex **31** (14.8 mg, 0.020 mmol) was added freshly distilled THF (2.1 mL), and the resulting solution was purged with nitrogen while cooling to 0 °C. Tetrabutylammonium fluoride (1 M solution in THF, dried over 4 Å molecular sieve) (25  $\mu\text{L}$ , 0.025 mmol) was then added, and the reaction mixture immediately became a dark red-brown suspension. Within 1 min, the reaction was quenched with excess saturated aqueous  $\text{NH}_4\text{Cl}$  and extracted with EtOAc. The combined organic layers were dried using  $\text{Na}_2\text{SO}_4$ , filtered, and concentrated to dryness in vacuo. The crude material was then subjected to silica gel chromatography first with  $\text{CH}_2\text{Cl}_2/\text{hexanes}$  (5:1) and followed by using  $\text{CH}_2\text{Cl}_2/\text{MeOH}$  (30:1). The combined product eluents were dried in vacuo to provide ruthenium complex **21** (7.8 mg, 66%) as a dark red solid.  $^1\text{H}$  NMR (300 MHz,  $\text{DMSO}-d_6$ ):  $\delta$  (ppm) 9.80 (s, 1H), 9.00 (s, 1H), 8.79 (s, 1H), 8.51 (d,  $J = 8.4$  Hz, 1H), 8.39–8.36 (m, 2H), 8.13 (ddd,  $J = 8.4, 7.2, 1.2$  Hz, 1H), 7.92 (t,  $J = 7.5$  Hz, 1H), 7.29 (d,  $J = 8.7$  Hz, 1H), 6.99 (dd,  $J = 8.7, 2.6$  Hz, 1H), 6.20 (m, 2H), 5.67 (m, 2H), 5.20 (m, 2H), 3.48 (s, 3H).  $^{13}\text{C}$  NMR (100 MHz,  $\text{DMSO}-d_6$ ):  $\delta$  (ppm) 200.8, 171.0, 166.0, 159.5, 150.9, 149.3, 146.4, 139.2, 133.4, 132.2, 129.7, 128.8, 127.8, 127.1, 126.8, 124.8, 123.7, 115.8, 115.7, 115.4, 114.2, 109.4, 89.9, 89.6, 81.4, 81.0, 79.2, 51.8, 47.9. IR  $\nu$  ( $\text{cm}^{-1}$ ) 2918, 2850, 2957, 1953, 1668, 1601, 1464, 1440, 1379, 1286, 1259, 1160, 1022, 798, 755, 718, 680, 559, 438, 402. HRMS calcd for  $\text{C}_{28}\text{H}_{20}\text{N}_3\text{O}_4\text{Ru}$  ( $\text{M} - \text{CO} + \text{H}$ ) $^+$  564.0492, found ( $\text{M} - \text{CO} + \text{H}$ ) $^+$  564.0491.



**Resolution of (*R*<sub>Ru</sub>)-**21** and (*S*<sub>Ru</sub>)-**21**.** Baseline separation of the enantiomers of racemic half sandwich complex **21** was achieved using a Chiral Pak 1B analytical HPLC column (Daicel/Chiral Technologies). First, **21** was dissolved in a minimal amount of CH<sub>2</sub>Cl<sub>2</sub> and carefully filtered. Then, each injection was run using an EtOH gradient of 50% to 90% over 20 min in hexanes. A flow rate of 0.7 mL/min was used, and injection volumes of approximately 25 μL could be reached without compromising baseline separation. Amounts of 2.8 mg of both (*R*<sub>Ru</sub>)-**21** and (*S*<sub>Ru</sub>)-**21** were isolated, and the absolute configuration was assigned by CD-spectroscopy (see Supporting Information).

**Protein Kinase Assays Using [ $\gamma$ -<sup>32</sup>P]ATP.** Protein kinases (human) and substrates were purchased from Upstate Biotechnology USA, Millipore. Various concentrations of (*R*<sub>Ru</sub>)-**1**, (*S*<sub>Ru</sub>)-**1**, (*R*<sub>Ru</sub>)-**2**, and (*S*<sub>Ru</sub>)-**2** were incubated at room temperature in 20 mM MOPS, 30 mM MgCl<sub>2</sub>, 0.8 μg/μL BSA, 5% DMSO (resulting from the inhibitor stock solution), pH 7.0, in the presence of substrate (GSK3β and CLK2: 20 μM and 100 μM phosphoglycogen synthase peptide-2, respectively; Pim1: 100 μM S6 kinase/Rsk2 substrate peptide 2; TrkA: 250 μM Trk peptide KKKSPGEYVNIEFG) and kinase (2.4 nM GSK3β, 0.8 nM Pim1, 8.0 nM TrkA, 0.6 nM CLK2). After 15 min, the reaction was initiated by adding ATP to a final concentration of 100 μM, including approximately 0.2 μCi/μL [ $\gamma$ -<sup>32</sup>P]ATP. Reactions were performed in a total volume of 25 μL and incubated at 30 °C. After 30 min (Pim1) or 45 min (GSK3β, TrkA, and CLK2), the reaction was terminated by spotting 17.5 μL on a circular P81 phosphocellulose paper (2.1 cm diameter, Whatman), followed by washing four times (5 min each wash) with 0.75% phosphoric acid and once with acetone. The dried P81 papers were transferred to a scintillation vial, 5 mL of scintillation cocktail was added, and the counts per minute (CPM) were determined with a Beckmann 6000 scintillation counter. IC<sub>50</sub> values were defined to be the concentration of inhibitor at which the CPM was 50% of the control sample, corrected by the background.

**Protein Kinase Assays Using [ $\gamma$ -<sup>33</sup>P]ATP.** Protein kinases (human) and substrates were purchased from Upstate Biotechnology USA, Millipore. Various concentrations of (*R*<sub>Ru</sub>)-**3**, (*S*<sub>Ru</sub>)-**3**, or (*R*<sub>Ru</sub>)-**21** were incubated at room temperature in 20 mM MOPS, 30 mM Mg(OAc)<sub>2</sub>, 0.8 μg/μL BSA, 5% DMSO (resulting from the inhibitor stock solution), pH 7.0, in the presence of substrate (GSK3β: 20 μM phosphoglycogen synthase peptide-2; Pim1: 50 μM S6 kinase/Rsk2 substrate peptide 2; TrkA: 150 μM Trk peptide KKKSPGEYVNIEFG) and kinase (0.4–2.0 nM GSK3β, 0.3–1.6 nM Pim1, 4.9–9.8 nM TrkA). After 15 min, the reaction was initiated by adding ATP to a final concentration of 100 μM, including approximately 0.05 μCi/μL [ $\gamma$ -<sup>33</sup>P]ATP (GSK3β and Pim1) or 0.1 μCi/μL [ $\gamma$ -<sup>33</sup>P]ATP (TrkA). Reactions were performed in a total volume of 25 μL. After 30 min (GSK3β and Pim1) or 2 h (TrkA), the reaction was terminated by spotting 15 μL (TrkA) or 17.5 μL (GSK3β, Pim1) on a circular P81 phosphocellulose paper (2.1 cm diameter, Whatman), followed by washing four times (5 min each wash) with 0.75% phosphoric acid and once with acetone. The dried P81 papers were transferred to a scintillation vial, 5 mL of scintillation cocktail was added, and the counts per minute (CPM) were determined with a Beckmann 6000 scintillation counter. IC<sub>50</sub> values were defined to be the concentration of inhibitor at which the CPM was 50% of the control sample, corrected by the background.

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**Supporting Information Available:** Screening results of compounds **1–3** and (*R*<sub>Ru</sub>)-**21** against a panel of human kinases; assignment of the absolute configurations of (*R*<sub>Ru</sub>)-**1**, (*S*<sub>Ru</sub>)-**1**, (*R*<sub>Ru</sub>)-**2**, (*S*<sub>Ru</sub>)-**2**, and (*R*<sub>Ru</sub>)-**21**; <sup>1</sup>H–<sup>1</sup>H-ROESY analysis of compound **17**; and NMR spectra for all new compounds. This material is available free of charge via the Internet at <http://pubs.acs.org>.