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Citation	Tetrahedron : Asymmetry, 22(8), 907-915 https://doi.org/10.1016/j.tetasy.2011.05.011
Issue Date	2011-04-30
Doc URL	http://hdl.handle.net/2115/46945
Type	article (author version)
File Information	TetA22-8_907-915.pdf



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Catalytic enantioselective C–H functionalization of indoles with α -diazopropionates using chiral dirhodium(II) carboxylates: asymmetric synthesis of the (+)- α -methyl-3-indolylacetic acid fragment of acremoauxin A

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ARTICLE INFO

Article history:

Received

Received in revised form

Accepted

Available online

ABSTRACT

An enantioselective C–H functionalization of *N*-methoxymethyl (MOM)-protected 2,3-unsubstituted indoles with α -diazopropionates has been effected under catalysis by dirhodium(II) tetrakis[*N*-phthaloyl-(*S*)-triethylalaninate], Rh₂(*S*-PTTEA)₄, providing α -methyl-3-indolylacetates in high yields and with enantioselectivities of up to 86% ee. The effectiveness of this protocol was demonstrated by the first catalytic asymmetric synthesis of the (+)- α -methyl-3-indolylacetic acid fragment of acremoauxin A, a potent plant-growth inhibitor. Furthermore, the Fujioka protocol using a combination of TMSOTf and 2,2'-bipyridyl was shown to be superior for removal of the *N*-MOM group.

1. Introduction

Functionalized indoles are prevalent structural motifs in a large number of biologically active alkaloids and pharmaceuticals.¹ It is therefore not surprising that an enormous amount of effort has recently been devoted to the development of efficient methods for direct catalytic functionalization of indoles.² The majority of these studies have been focused on the Friedel–Crafts reaction and its enantioselective variants using a range of electrophilic reagents.³ In this context, C–H functionalization of indoles with metal carbenes generated from α -diazocarbonyl compounds under catalysis by metal complexes is a potentially powerful means for installing functionalized alkyl groups on the pyrrole portion of the indole ring system.^{4,5} Since publication of the seminal work of Davies and Lian on catalysis by Rh₂(*S*-DOSP)₄ **4**⁶ (Figure 1),⁷ the development of an enantioselective version of this catalytic process has been a challenging objective. Very recently, Fox and co-workers made a major breakthrough in this field when they demonstrated that enantioselective C–H functionalization of indoles with ethyl α -alkyl- α -diazooacetates under catalysis by Rh₂(*S*-NTTL)₄ **3**^{8–10} provides α -alkyl-substituted 3-indolylacetates in high yields and enantioselectivities (82–96% yields, 79–99% ee, Scheme 1).^{11,12} While the scope of the reaction was found to accommodate a wide array of substrates, an exceptionally high performance was achieved with diazoesters bearing longer-chained alkyl groups than α -methyl substituent when *N*-aryl or *N*-alkyl indole

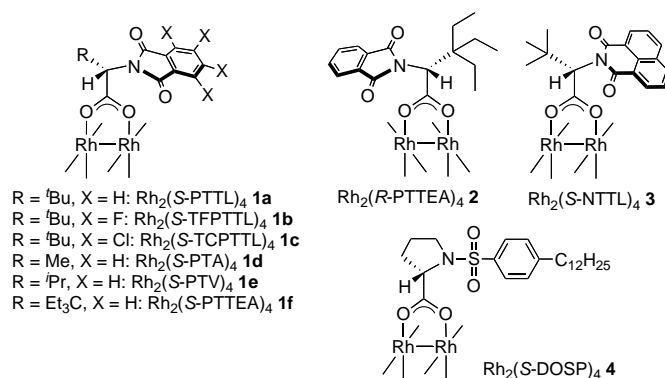
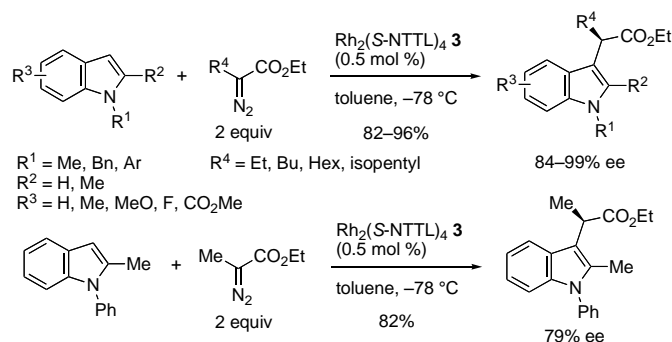


Figure 1. Chiral dirhodium(II) catalysts.



Scheme 1. The enantioselective C–H functionalization of indoles with ethyl α -alkyl- α -diazooacetates under catalysis by Rh₂(*S*-NTTL)₄ **3** reported by Fox.¹¹

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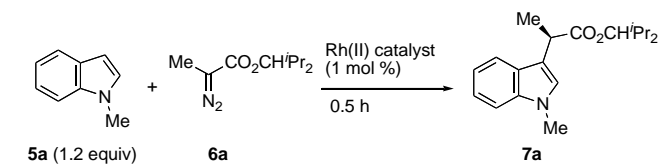
derivatives with small 2-substituents ($R^2 = \text{H, Me}$) were used. We have recently been engaged in the enantioselective C–H functionalization of indoles with α -diazopropionates catalyzed by chiral dirhodium(II) carboxylates. Although Fox and co-workers described the catalytic enantioselective reaction of 2-methyl-1-phenylindole with ethyl α -diazopropionate (79% ee, Scheme 1),¹¹ no examples of the reaction of 2,3-unsubstituted indoles with α -diazopropionates have been reported. Herein, we report the first example of a catalytic enantioselective C–H functionalization of 2,3-unsubstituted indoles with α -diazopropionates.

2. Results and discussion

At the outset, we explored the reaction of 2,3-unsubstituted *N*-methylindole **5a** (1.2 equiv) with 2,4-dimethyl-3-pentyl α -diazopropionate **6a** using 1 mol % of $\text{Rh}_2(\text{S-PTTL})_4$ **1a**^{13–15} (Table 1, entry 1). The reaction in dichloromethane at room temperature proceeded smoothly to give α -methyl-3-indolylacetate **7a** in 82% yield with no signs of α,β -unsaturated ester formation via a 1,2-hydride shift.¹⁶ The enantiomeric excess of **7a** was determined to be 52% by HPLC using a Daicel Chiralcel OD-H column. To further enhance the enantioselectivity, we evaluated the performance of other chiral dirhodium(II) carboxylate catalysts. Although $\text{Rh}_2(\text{S-TFP TTL})_4$ **1b**¹⁷ and $\text{Rh}_2(\text{S-TCPTTL})_4$ **1c**,^{18–20} fluorinated and chlorinated analogues of **1a**, provided **7a** in high yields, poor enantioselectivities were obtained (43% and 22% ee, respectively, entries 2 and 3). Catalysis with $\text{Rh}_2(\text{S-PTA})_4$ **1d** and $\text{Rh}_2(\text{S-PTV})_4$ **1e** resulted in lower product yields and enantioselectivities than those obtained with **1a** (21% and 32% ee, respectively, entries 4 and 5). We recently developed dirhodium(II) tetrakis[*N*-phthaloyl-(*S*)-triethylalaninate], $\text{Rh}_2(\text{S-PTTEA})_4$ **1f**, characterized by an exceptionally bulky triethylmethyl group, for the enantioselective intramolecular C–H insertion of aryldiazooacetates.²¹ Gratifyingly, the use of **1f** was found to increase the enantioselectivity to 67% ee (entry 6). A survey of solvents with **1f** revealed that dichloromethane was the optimal solvent for this transformation in terms of both product yield and enantioselectivity (entry 6 vs entries 7–10).

Table 1

Enantioselective C–H functionalization of *N*-methylindole **5a** with 2,4-dimethyl-3-pentyl α -diazopropionate **6a** catalyzed by chiral dirhodium(II) carboxylates^a



Entry	Rh(II) catalyst	Solvent	Yield ^b (%)	ee ^c (%)
1	$\text{Rh}_2(\text{S-PTTL})_4$ 1a	CH_2Cl_2	82	52
2	$\text{Rh}_2(\text{S-TFP TTL})_4$ 1b	CH_2Cl_2	84	43
3	$\text{Rh}_2(\text{S-TCPTTL})_4$ 1c	CH_2Cl_2	90	22
4	$\text{Rh}_2(\text{S-PTA})_4$ 1d	CH_2Cl_2	64	21
5	$\text{Rh}_2(\text{S-PTV})_4$ 1e	CH_2Cl_2	72	32
6	$\text{Rh}_2(\text{S-PTTEA})_4$ 1f	CH_2Cl_2	83	67
7	$\text{Rh}_2(\text{S-PTTEA})_4$ 1f	toluene	87	59
8	$\text{Rh}_2(\text{S-PTTEA})_4$ 1f	hexane	92	58
9	$\text{Rh}_2(\text{S-PTTEA})_4$ 1f	AcOEt	76	65
10	$\text{Rh}_2(\text{S-PTTEA})_4$ 1f	acetone	40	54

^a All reactions were carried out as follows: Rh(II) catalyst (1 mol %) was added to a solution of **5a** (0.030 mL, 0.24 mmol, 1.2 equiv) and **6a** (39.7 mg, 0.20 mmol) in the indicated solvent (1 mL) at room temperature.

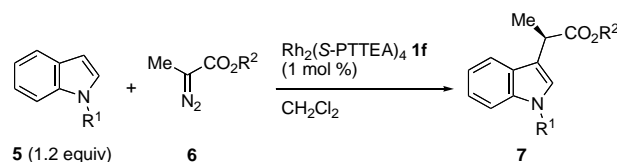
^b Isolated yield.

^c Determined by HPLC (Daicel Chiralcel OD-H).

Using $\text{Rh}_2(\text{S-PTTEA})_4$ **1f** as a catalyst, we next examined the effects of the *N*-substitution of the indole ring (Table 2). As might be expected from precedents,^{4,5g,5i,5l,11,22} the reaction of **6a** with indoles **5b** and **5c** carrying strongly electron-withdrawing Boc or Ts groups failed to give the corresponding indole products **7b** and **7c** (entries 2 and 3); instead, azine derived from **6a** was obtained as a major product. While the reaction of *N*-benzylindole **5d** proceeded smoothly to give **7d** in 89% yield with 74% ee (entry 4), *N*-methoxymethyl (MOM)-protected indole **5e** provided **7e** in 83% yield and with a greatly improved enantioselectivity of 82% ee (entry 5). An examination of the temperature profile demonstrated that lowering the reaction temperature to -40 or -60 °C had only a marginal effect on enantioselectivity (78% and 75% ee), although a sharp drop in product yield was observed (70% and 64%, respectively, entries 6 and 7). The effects of the ester moiety were also evaluated. Reactions of **5e** with *tert*-butyl and ethyl esters **6b** and **6c** resulted in lower enantioselectivities than that observed with **6a** (45% and 41% ee, respectively, entries 8 and 9). These results demonstrate that the use of the 2,4-dimethyl-3-pentyl ester moiety²³ is crucial for a high degree of enantioselection.

Table 2

Enantioselective C–H functionalization of indoles **5a–e** with α -diazopropionates **6a–c** catalyzed by $\text{Rh}_2(\text{S-PTTEA})_4$ **1f**



Entry	Indole 5		Diazoester 6		<i>T</i> (°C)	<i>t</i> (h)	Product 7		
	<i>R</i> ¹	<i>R</i> ²	<i>R</i> ¹	<i>R</i> ²			Yield ^a (%)	ee ^b (%)	
1	5a	Me	6a	^t Pr ₂ CH	23	0.5	7a	83	67
2	5b	Boc	6a	^t Pr ₂ CH	23	0.5	7b	ND ^c	–
3	5c	Ts	6a	^t Pr ₂ CH	23	0.5	7c	ND ^c	–
4	5d	Bn	6a	^t Pr ₂ CH	23	0.5	7d	89	74
5	5e	MOM	6a	^t Pr ₂ CH	23	0.5	7e	83	82
6	5e	MOM	6a	^t Pr ₂ CH	-40	2	7e	70	78
7	5e	MOM	6a	^t Pr ₂ CH	-60	15	7e	64	75
8	5e	MOM	6b	^t Bu	23	0.5	7f	72	45
9	5e	MOM	6c	Et	23	0.5	7g	69	32

^a Isolated yield.

^b Determined by HPLC.

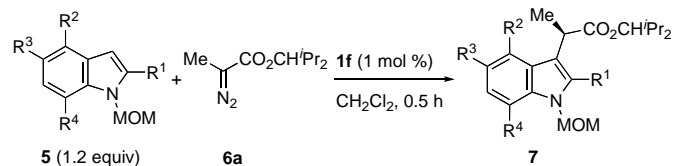
^c Not detected.

Under optimized reaction conditions, we then explored the scope of the reaction with respect to the indole component (Table 3). The reaction with 2-methylindole derivative **5f** resulted in only a trace amount of product **7h** (entry 1), which was markedly different from Fox's result with ethyl α -diazopropionate.²⁴ The reaction with 4-methylindole **5g** afforded **7i** in 72% yield and 69% ee (entry 2). The reaction was also tolerant of methoxy, bromo, and nitro groups at the 5-position of the *N*-MOM-protected indole nucleus (69–83% yields, 58–72% ee, entries 3–5). The highest enantioselectivity (86% ee) was obtained in the reaction with 7-methylindole **5k** (entry 6).

To determine the preferred absolute configuration of **7e**, we explored removal of the MOM protecting group (Scheme 2). Hydrolysis of **7e** with 2 M HCl in THF followed by treatment with 15% NaOH²⁵ provided indole **9** in 51% yield. Treatment with BBr_3 ,²⁶ aq. formic acid²⁷ or TMSI²⁸ gave a complex mixture of products. After considerable experimentation, the Fujioka protocol using a combination of TMSOTf and 2,2'-bipyridyl²⁹ proved to be the method of choice. Treatment of **7e** with TMSOTf (2 equiv) and 2,2'-bipyridyl (3 equiv) in CH_2Cl_2 at 0 °C led to the exclusive cleavage of the methyl ether linkage of

Table 3

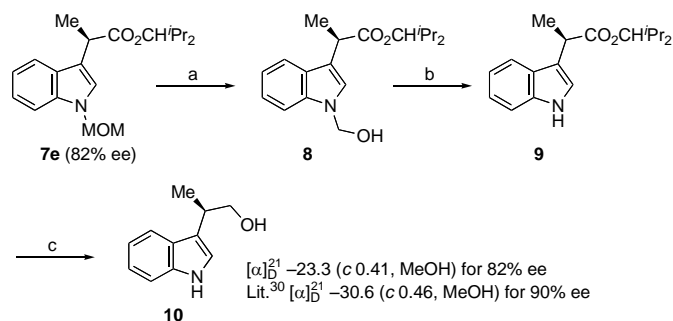
Enantioselective C–H functionalization of indoles **5f–k** with 2,4-dimethyl-3-pentyl α -diazopropionate **6a** catalyzed by $\text{Rh}_2(\text{S-PTTEA})_4$ **1f**



Entry	Indole 5				Product 7	
	R ¹	R ²	R ³	R ⁴	Yield ^a (%)	ee ^b (%)
1	5f	Me	H	H	7h	trace
2	5g	H	Me	H	7i	72
3	5h	H	H	MeO	7j	77
4	5i	H	H	Br	7k	83
5	5j	H	H	NO ₂	7l	69
6	5k	H	H	Me	7m	83

^a Isolated yield.

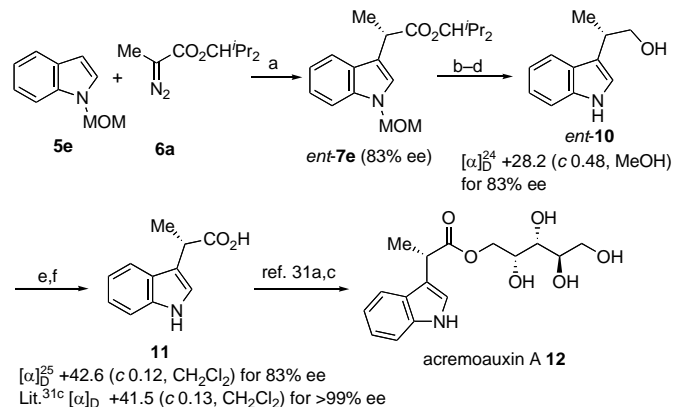
^b Determined by HPLC.



Scheme 2. Reagents and conditions: (a) TMSOTf, 2,2'-bipyridyl, CH_2Cl_2 , 0 °C, 0.5 h; (b) NaOH, THF/ H_2O (9:1), 24 h, 98% (two steps); (c) LiAlH_4 , THF, 4 h, 87%.

the MOM protecting group to give **8**, which, upon exposure to NaOH in THF/ H_2O (9:1), afforded **9** in 98% yield. Reduction of 2,4-dimethyl-3-pentyl ester **9** with LiAlH_4 provided the known alcohol **10** $\{[\alpha]_{\text{D}}^{21} -23.3$ (c 0.41, MeOH) for 82% ee; lit.,³⁰ $[\alpha]_{\text{D}}^{21} -30.6$ (c 0.46, MeOH) for 90% ee of (*R*)-**10** $\}$ in 87% yield. Thus, the preferred absolute configuration of **7e** was established as *R*.

Using the present catalytic protocol, we conducted an asymmetric synthesis of (+)- α -methyl-3-indolylacetic acid **11**,³¹ a phytohormone of the auxin class³² and a constituent of a potent plant-growth inhibitor acremoauxin A **12** (Scheme 3).^{33,34} Alcohol *ent*-**10** was prepared using $\text{Rh}_2(\text{R-PTTEA})_4$ **2**. Fortunately, Overman and Govek reported the chemoselective



Scheme 3. Reagents and conditions: (a) $\text{Rh}_2(\text{R-PTTEA})_4$ **2** (1 mol %), CH_2Cl_2 , 0.5 h, 83%; (b) TMSOTf, 2,2'-bipyridyl, CH_2Cl_2 , 0 °C, 0.5 h; (c) NaOH, THF/ H_2O (9:1), 24 h, 98% (two steps); (d) LiAlH_4 , THF, 4 h, 97%; (e) SO_3 ·pyridine, Et_3N , DMSO; (f) NaClO_2 , NaH_2PO_4 · H_2O , 2-methyl-2-butene, THF/ H_2O , 61% (two steps).

oxidation of a primary alcohol tethered to an unmasked indole substituent to carboxylic acid in the total synthesis of (+)-asperazine.³⁵ Following this precedent, oxidation of alcohol *ent*-**10** under standard Parikh–Doering conditions³⁶ and subsequent Lindgren–Kraus oxidation³⁷ of the crude aldehyde provided carboxylic acid **11** in 61% yield. The spectroscopic data (¹H and ¹³C NMR, and IR) of synthetic material **11** were identical to those reported by Baran and co-workers.^{31c} The sign of optical rotation of synthetic product **11** $\{[\alpha]_{\text{D}}^{25} +42.6$ (c 0.12, CH_2Cl_2) for 83% ee $\}$ was the same as that reported in the literature $\{$ lit.^{31c} $[\alpha]_{\text{D}} +41.5$ (c 0.13, CH_2Cl_2) for >99% ee $\}$.³⁸ This is the first catalytic asymmetric synthesis of the (+)- α -methyl-3-indolylacetic acid fragment of **12**.

3. Conclusion

We have developed a catalytic enantioselective C–H functionalization protocol for *N*-methoxymethyl (MOM)-protected 2,3-unsubstituted indoles using 2,4-dimethyl-3-pentyl α -diazopropionates. In this process, $\text{Rh}_2(\text{S-PTTEA})_4$, which is characterized by an exceptionally bulky triethylmethyl group, has emerged as the catalyst of choice for providing α -methyl-3-indolylacetates in high yields and with enantioselectivities of up to 86% ee. This represents the first example of a catalytic enantioselective C–H functionalization of 2,3-unsubstituted indoles with α -diazopropionates. Using this methodology, we achieved the first catalytic asymmetric synthesis of the (+)- α -methyl-3-indolylacetic acid fragment of acremoauxin A. In our approach, the use of a MOM group as a protecting group on the indole nitrogen was not only crucial for high levels of enantioselection but also synthetically advantageous since the removal of the *N*-MOM group could be conducted efficiently under the Fujioka conditions. Further applications of this methodology to catalytic asymmetric synthesis of indole alkaloids as well as stereochemical studies are currently in progress.

4. Experimental

4.1. General

Optical rotations were measured on a JASCO P-1030 digital polarimeter at the sodium D line (589 nm). IR spectra were recorded on a JASCO FT/IR-5300 spectrometer and absorbance bands are reported in wavenumber (cm^{-1}). ¹H NMR spectra were recorded on JEOL JNM-ECX 400P (400 MHz) spectrometer or JNM-ECA 500 (500 MHz) spectrometer. Chemical shifts are reported relative to internal standard (tetramethylsilane at δ_{H} 0.00 or CDCl_3 at δ_{H} 7.26). Data are presented as follows: chemical shift (δ , ppm), multiplicity (s = singlet, d = doublet, t = triplet, q = quartet, m = multiplet, br = broad), coupling constants, integration and assignment. ¹³C NMR spectra were recorded on JEOL JNM-ECX 400P (100 MHz) spectrometer or JEOL JNM-ECA 500 (125 MHz) spectrometer. The following internal references were used (CDCl_3 at δ 77.0). EI mass spectra were recorded on a JEOL JMS-FABmate spectrometer, operating with ionization energy of 70 eV. ESI mass spectra were recorded on a JEOL JMS-T100LCP spectrometer. Column chromatography was carried out on Kanto silica gel 60 N (63–210 mesh). Analytical thin layer chromatography (TLC) was carried out on Merck Kieselgel 60 F₂₅₄ plates. Visualization was accomplished with UV light, anisaldehyde stain solution or phosphomolybdic acid stain solution followed by heating. Analytical high performance liquid chromatography (HPLC) was performed on a JASCO PU-1580 intelligent HPLC pump with JASCO UV-1575

intelligent UV/VIS detector. Detection was performed at 254 nm. Chiralcel OD-H, Chiralpak AD-H, Chiralpak IC columns (0.46 cm × 25 cm) and Chiralpak IC-3 column (0.46 cm × 15 cm) from Daicel were used. Retention times (t_R) and peak ratios were determined with JASCO-Borwin analysis system.

All non-aqueous reactions were carried out in flame-dried glassware under Ar atmosphere unless otherwise noted. Reagents and solvents were purified by standard means. Dehydrated CH_2Cl_2 , THF and DMF were purchased from Kanto Chemical Co., Inc. Chiral dirhodium(II) carboxylates **1a-f**^{14a,17a,18a,21} and **2**,²¹ 1-methoxymethylindole **5e**,³⁹ 1-methoxymethyl-5-nitroindole **5j**⁴⁰ and α -diazopropionates **6a**,^{23c} **6b**⁴¹ and **6c**⁴¹ were prepared according to literature procedures.

4.2. Preparation of *N*-MOM-protected indoles

4.2.1. 1-Methoxymethyl-2-methylindole **5f**

A solution of 2-methylindole (1.00 g, 6.16 mmol) in DMF (5 mL) was added to a solution of NaH (60% in oil, 0.34 g, 8.38 mmol) in DMF (10 mL) at 0 °C. After stirring for 0.5 h at room temperature, MOMCl (0.70 mL, 9.22 mmol) was added to the reaction mixture. After stirring for 1 h, the reaction was quenched by addition of water (30 mL). The mixture was extracted with EtOAc (2 × 20 mL) and the combined organic layers were washed with water (30 mL) and brine (30 mL), and dried over anhydrous Na_2SO_4 . Filtration and evaporation in vacuo furnished the crude product (1.65 g), which was purified by column chromatography (silica gel, 20:1 hexane/EtOAc) to provide **5f** (1.20 g, 90%) as a colorless oil: TLC R_f = 0.38 (20:1 hexane/EtOAc); IR (neat) ν 2948, 2252, 1460, 1308 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 2.48 (s, 3H, CH_3Ar), 3.24 (s, 3H, CH_3OCH_2), 5.43 (s, 2H, CH_3OCH_2), 6.29 (s, 1H, Ar), 7.09 (ddd, J = 0.9, 7.7, 8.2 Hz, 1H, Ar), 7.16 (ddd, J = 1.4, 7.2, 8.2 Hz, 1H, Ar), 7.40 (d, J = 8.2 Hz, 1H, Ar), 7.51 (d, J = 7.2 Hz, 1H, Ar); ^{13}C NMR (100 MHz, CDCl_3) δ 12.4 (CH_3), 55.5 (CH_3), 73.6 (CH_2), 101.7 (CH), 109.0 (CH), 119.7 (CH), 120.0 (CH), 121.1 (CH), 128.2 (C), 136.7 (CH), 137.5 (C); HRMS (ESI) calcd for $\text{C}_{11}\text{H}_{14}\text{NO}$ ($\text{M} + \text{H}$)⁺ 176.10699, found 176.10708.

4.2.2. 1-Methoxymethyl-4-methylindole **5g**

According to the procedure for the preparation of **5f**, **5g** was prepared from 4-methylindole (0.500 g, 3.08 mmol). The crude product was purified by column chromatography (silica gel, 20:1 hexane/EtOAc) to provide the title compound (0.639 g, 96%) as a colorless oil: TLC R_f = 0.31 (20:1 hexane/EtOAc); IR (neat) ν 2928, 2251, 1519, 1493, 1303, 1236 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 2.56 (s, 3H, CH_3Ar), 3.23 (s, 3H, CH_3OCH_2), 5.45 (s, 2H, CH_3OCH_2), 6.56 (dd, J = 0.9, 3.2 Hz, 1H, Ar), 6.95 (ddd, J = 0.9, 0.9, 8.2 Hz, 1H, Ar), 7.15 (d, J = 8.2 Hz, 1H, Ar), 7.17 (d, J = 3.2 Hz, 1H, Ar), 7.33 (d, J = 8.2 Hz, 1H, Ar); ^{13}C NMR (100 MHz, CDCl_3) δ 18.7 (CH_3), 55.9 (CH_3), 77.5 (CH_2), 101.1 (CH), 107.5 (CH), 120.5 (CH), 122.3 (CH), 127.5 (CH), 129.0 (C), 130.5 (CH), 136.1 (C); HRMS (ESI) calcd for $\text{C}_{11}\text{H}_{14}\text{NO}$ ($\text{M} + \text{H}$)⁺ 176.10699, found 176.10704.

4.2.3. 5-Methoxy-1-methoxymethylindole **5h**

According to the procedure for the preparation of **5f**, **5h** was prepared from 5-methoxyindole (0.500 g, 3.40 mmol). The crude product was purified by column chromatography (silica gel, 10:1 hexane/EtOAc) to provide the title compound (0.583 g, 90%) as a colorless oil: TLC R_f = 0.29 (10:1 hexane/EtOAc); IR (neat) ν 2938, 1486, 1239, 1151, 1101 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 3.23 (s, 3H, CH_3OCH_2), 3.86 (s, 3H, CH_3OAr), 5.42 (s, 2H, CH_3OCH_2), 6.46 (d, J = 2.7 Hz, 1H, Ar), 6.90 (dd, J = 2.3, 9.1 Hz, 1H, Ar), 7.10 (d, J = 2.3 Hz, 1H, Ar), 7.15 (d, J = 3.2 Hz, 1H, Ar), 7.38 (d, J = 8.6 Hz, 1H, Ar); ^{13}C NMR (125 MHz, CDCl_3) δ 55.7

(CH_3), 55.8 (CH_3), 77.6 (CH_2), 102.1 (CH), 102.6 (CH), 110.6 (CH), 112.3 (CH), 128.7 (CH), 129.6 (C), 131.5 (C), 154.5 (C); HRMS (ESI) calcd for $\text{C}_{11}\text{H}_{14}\text{NO}_2$ ($\text{M} + \text{H}$)⁺ 192.10191, found 192.10193.

4.2.4. 5-Bromo-1-methoxymethylindole **5i**

According to the procedure for the preparation of **5f**, **5i** was prepared from 5-bromoindole (0.500 g, 2.55 mmol). The crude product was purified by column chromatography (silica gel, 10:1 hexane/EtOAc) to provide the title compound (0.596 g, 97%) as a colorless oil: TLC R_f = 0.26 (10:1 hexane/EtOAc); IR (neat) ν 2936, 2360, 1452, 1328, 1183, 1089 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 3.22 (s, 3H, CH_3OCH_2), 5.43 (s, 2H, CH_3OCH_2), 6.48 (d, J = 3.2 Hz, 1H, Ar), 7.18 (d, J = 3.2 Hz, 1H, Ar), 7.32 (dd, J = 1.8, 8.6 Hz, 1H, Ar), 7.37 (d, J = 8.6 Hz, 1H, Ar), 7.76 (d, J = 1.8 Hz, 1H, Ar); ^{13}C NMR (125 MHz, CDCl_3) δ 55.9 (CH_3), 77.5 (CH_2), 102.0 (CH), 111.4 (CH), 113.5 (C), 123.4 (CH), 125.0 (CH), 129.2 (CH), 130.8 (C), 134.9 (C); HRMS (ESI) calcd for $\text{C}_{10}\text{H}_{11}\text{BrNO}$ ($\text{M} + \text{H}$)⁺ 240.00185, found 240.00228.

4.2.5. 1-Methoxymethyl-7-methylindole **5k**

According to the procedure for the preparation of **5f**, **5k** was prepared from 7-methylindole (0.500 g, 3.08 mmol). The crude product was purified by column chromatography (silica gel, 20:1 hexane/EtOAc) to provide the title compound (0.658 g, 99%) as a colorless oil: TLC R_f = 0.25 (20:1 hexane/EtOAc); IR (neat) ν 2930, 1462, 1305 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 2.74 (s, 3H, CH_3Ar), 3.23 (s, 3H, CH_3OCH_2), 5.53 (s, 2H, CH_3OCH_2), 6.49 (d, J = 3.2 Hz, 1H, Ar), 6.98 (d, J = 7.2 Hz, 1H, Ar), 7.04 (dd, J = 7.2, 7.7 Hz, 1H, Ar), 7.10 (d, J = 3.2 Hz, 1H, Ar), 7.47 (d, J = 7.2 Hz, 1H, Ar); ^{13}C NMR (100 MHz, CDCl_3) δ 19.2 (CH_3), 55.2 (CH_3), 79.3 (CH_2), 102.3 (CH), 118.9 (CH), 120.5 (CH), 121.9 (C), 124.9 (CH), 130.1 (CH), 130.4 (C); HRMS (ESI) calcd for $\text{C}_{11}\text{H}_{14}\text{NO}$ ($\text{M} + \text{H}$)⁺ 176.10699, found 176.10714.

4.3. Enantioselective C–H functionalization of indoles with α -diazopropionates

4.3.1. Typical procedure for the C–H functionalization of indoles: (*R**)-2,4-Dimethyl-3-pentyl 2-[(1-methyl)-1*H*-indol-3-yl]propionate **7a** (Table 1, entry 6)

$\text{Rh}_2(\text{S-PTTEA})_4 \cdot 2\text{EtOAc}$ **1f** (3.18 mg, 0.002 mmol, 1 mol %) was added in one portion to a solution of 2,4-dimethyl-3-pentyl α -diazopropionate **6a** (39.7 mg, 0.200 mmol) and 1-methylindole **5a** (0.030 mL, 0.235 mmol) in CH_2Cl_2 (1.0 mL) at room temperature. After stirring for 0.5 h at room temperature, the reaction mixture was concentrated in vacuo, and the residue was purified by column chromatography (silica gel, 20:1 hexane/EtOAc) to provide **7a** (50.2 mg, 83%) as a colorless oil: TLC R_f = 0.43 (10:1 hexane/EtOAc); $[\alpha]_D^{24}$ –24.0 (c 0.99, CHCl_3) for 67% ee; IR (neat) ν 2966, 1730, 1471, 1371, 1330, 1242, 1177 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 0.68 (d, J = 6.8 Hz, 3H, $\text{CH}(\text{CH}_3)_2$), 0.72 (d, J = 6.8 Hz, 3H, $\text{CH}(\text{CH}_3)_2$), 0.82 (d, J = 6.8 Hz, 3H, $\text{CH}(\text{CH}_3)_2$), 0.84 (d, J = 7.2 Hz, 3H, $\text{CH}(\text{CH}_3)_2$), 1.63 (d, J = 7.2 Hz, 3H, CH_3CHAr), 1.77–1.92 (m, 2H, $\text{CH}(\text{CH}_3)_2$), 3.76 (s, 3H, CH_3N), 4.06 (q, J = 7.2 Hz, 1H, CH_3CHAr), 4.58 (t, J = 6.3 Hz, 1H, CO_2CH), 7.03 (s, 1H, Ar), 7.10 (m, 1H, Ar), 7.21 (m, 1H, Ar), 7.28 (d, J = 8.2 Hz, 1H, Ar), 7.68 (d, J = 8.2 Hz, 1H, Ar); ^{13}C NMR (100 MHz, CDCl_3) δ 17.0 (CH_3), 17.2 (CH_3), 18.3 (CH_3), 19.4 (CH_3), 19.5 (CH_3), 29.4 (CH), 32.7 (CH_3), 37.2 (CH), 82.6 (CH), 109.1 (CH), 114.3 (C), 118.9 (CH), 119.4 (CH), 121.6 (CH), 126.3 (CH), 127.0 (C), 136.9 (C), 175.3 (CO); HRMS (EI^+) calcd for $\text{C}_{19}\text{H}_{27}\text{NO}_2$ (M)⁺ 301.20418, found 301.20346. The enantiomeric excess of **7a** was determined to be 67% by HPLC with a Daicel Chiralpak OD-H

column (eluent: 100:1 hexane/2-propanol; flow: 1.0 mL/min): $t_R = 8.5$ min for minor enantiomer; $t_R = 9.5$ min for major enantiomer. The preferred absolute configuration of **7a** was not determined.

4.3.2. (*R**)-2,4-Dimethyl-3-pentyl 2-[(1-benzyl)-1*H*-indol-3-yl]propionate **7d** (Table 2, entry 4)

According to the typical procedure for C–H functionalization of indole, **7d** was prepared from **1f** (3.18 mg, 0.002 mmol, 1 mol %), **6a** (39.7 mg, 0.200 mmol) and 1-benzylindole **5d** (49.7 mg, 0.24 mmol). The crude product was purified by column chromatography (silica gel, 30:1 hexane/EtOAc) to provide **7d** (67.4 mg, 89%) as a colorless oil: TLC $R_f = 0.43$ (10:1 hexane/EtOAc); $[\alpha]_D^{21} -16.4$ (c 1.00, CHCl₃) for 74% ee; IR (neat) ν 2966, 1728, 1468, 1173 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ 0.64 (d, $J = 6.8$ Hz, 3H, CH(CH₃)₂), 0.70 (d, $J = 6.8$ Hz, 3H, CH(CH₃)₂), 0.78 (d, $J = 6.8$ Hz, 3H, CH(CH₃)₂), 0.81 (d, $J = 6.8$ Hz, 3H, CH(CH₃)₂), 1.63 (d, $J = 7.2$ Hz, 3H, CH₃CHAr), 1.76–1.87 (m, 2H, CH(CH₃)₂), 4.07 (q, $J = 7.2$ Hz, 1H, CH₃CHAr), 4.57 (t, $J = 6.3$ Hz, 1H, CO₂CH), 5.27 (d, $J = 16.3$ Hz, 1H, ArCHH), 5.32 (d, $J = 16.3$ Hz, 1H, ArCHH), 7.06–7.17 (m, 5H, Ar), 7.22–7.29 (m, 4H, Ar), 7.72 (d, $J = 7.7$ Hz, 1H, Ar); ¹³C NMR (100 MHz, CDCl₃) δ 16.9 (CH₃), 17.2 (CH₃), 18.1 (CH₃), 19.4 (CH₃), 19.4 (CH₃), 29.4 (CH), 29.4 (CH), 37.2 (CH), 49.9 (CH₂), 82.5 (CH), 109.6 (CH), 115.0 (C), 119.1 (CH), 119.6 (CH), 121.8 (CH), 125.6 (CH), 126.6 (CH), 127.3 (C), 127.5 (CH), 128.6 (CH), 136.4 (C), 137.5 (C), 175.0 (CO); HRMS (ESI) calcd for C₂₅H₃₁NO₂Na (M + Na)⁺ 400.22470, found 400.22451. The enantiomeric excess of **7d** was determined to be 74% by HPLC with a Daicel Chiralpak OD-H column (eluent: 100:1 hexane/2-propanol; flow: 1.0 mL/min): $t_R = 16.0$ min for minor enantiomer; $t_R = 17.7$ min for major enantiomer. The preferred absolute configuration of **7d** was not determined.

4.3.3. (*R*)-2,4-Dimethyl-3-pentyl 2-[(1-methoxymethyl)-1*H*-indol-3-yl]propionate **7e** (Table 2, entry 5)

According to the typical procedure for C–H functionalization of indole, **7e** was prepared from **1f** (3.18 mg, 0.002 mmol, 1 mol %), **6a** (79.3 mg, 0.400 mmol) and 1-methoxymethylindole **5e** (77.4 mg, 0.480 mmol). The crude product was purified by column chromatography (silica gel, 5:1 hexane/Et₂O) to provide **7e** (111 mg, 83%) as a colorless oil: TLC $R_f = 0.30$ (10:1 hexane/EtOAc); $[\alpha]_D^{20} -20.3$ (c 1.04, CHCl₃) for 82% ee; IR (neat) ν 2966, 1730, 1466, 1181, 1090 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ 0.65 (d, $J = 6.8$ Hz, 3H, CH(CH₃)₂), 0.71 (d, $J = 6.8$ Hz, 3H, CH(CH₃)₂), 0.80 (d, $J = 6.3$ Hz, 3H, CH(CH₃)₂), 0.82 (d, $J = 6.8$ Hz, 3H, CH(CH₃)₂), 1.64 (d, $J = 7.2$ Hz, 3H, CH₃CHAr), 1.75–1.90 (m, 2H, CH(CH₃)₂), 3.19 (s, 3H, CH₃OCH₂), 4.05 (q, $J = 7.2$ Hz, 1H, CH₃CHAr), 4.58 (t, $J = 6.3$ Hz, 1H, CO₂CH), 5.39 (d, $J = 10.9$ Hz, 1H, CH₃OCHH), 5.44 (d, $J = 11.3$ Hz, 1H, CH₃OCHH), 7.15 (m, 1H, Ar), 7.15 (s, 1H, Ar), 7.23 (m, 1H, Ar), 7.45 (d, $J = 8.2$ Hz, 1H, Ar), 7.70 (d, $J = 7.7$ Hz, 1H, Ar); ¹³C NMR (100 MHz, CDCl₃) δ 16.9 (CH₃), 17.2 (CH₃), 18.0 (CH₃), 19.3 (CH₃), 19.5 (CH₃), 29.4 (CH), 37.1 (CH), 55.7 (CH₃), 77.3 (CH₂), 82.6 (CH), 109.8 (CH), 115.9 (C), 119.6 (CH), 119.9 (CH), 122.3 (CH), 125.4 (CH), 127.8 (C), 136.5 (C), 174.9 (CO); HRMS (ESI) calcd for C₂₀H₂₉NO₃Na (M + Na)⁺ 354.20451, found 354.20480. The enantiomeric excess of **7e** was determined to be 82% by HPLC analysis with a Daicel Chiralpak IC column (eluent: 100:1 hexane/2-propanol; flow: 1.0 mL/min): t_R (minor) = 9.8 min for (*S*)-**7e**; t_R (major) = 11.0 min for (*R*)-**7e**.

4.3.4. (*R**)-*tert*-Butyl 2-[(1-methoxymethyl)-1*H*-indol-3-yl]propionate **7f** (Table 2, entry 8)

According to the typical procedure for C–H functionalization of indole, **7f** was prepared from **1f** (3.18 mg, 0.002 mmol, 1

mol %), *tert*-butyl α -diazopropionate **6b** (31.2 mg, 0.200 mmol) and **5e** (38.7 mg, 0.24 mmol). The crude product was purified by column chromatography (silica gel, 10:1 hexane/EtOAc) to provide **7f** (41.6 mg, 72%) as a colorless oil: TLC $R_f = 0.13$ (10:1 hexane/EtOAc); $[\alpha]_D^{22} -24.4$ (c 1.06, CHCl₃) for 45% ee; IR (neat) ν 2978, 1726, 1466, 1367, 1149, 1089 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ 1.41 (s, 9H, (CH₃)₃C), 1.56 (d, $J = 7.2$ Hz, 3H, CH₃CHAr), 3.23 (s, 3H, CH₃OCH₂), 3.90 (q, $J = 7.2$ Hz, 1H, CH₃CHAr), 5.42 (s, 2H, CH₃OCH₂), 7.11 (s, 1H, Ar), 7.15 (dd, $J = 7.2, 8.0$ Hz, 1H, Ar), 7.24 (dd, $J = 8.0, 8.0$ Hz, 1H, Ar), 7.45 (d, $J = 8.4$ Hz, 1H, Ar), 7.69 (d, $J = 7.6$ Hz, 1H, Ar); ¹³C NMR (100 MHz, CDCl₃) δ 17.8 (CH₃), 28.0 (CH₃), 37.8 (CH), 55.8 (CH₃), 77.3 (CH₂), 80.4 (C), 109.8 (CH), 116.0 (C), 119.6 (CH), 119.8 (CH), 122.3 (CH), 125.3 (CH), 127.8 (C), 136.6 (C), 174.3 (CO); HRMS (ESI) calcd for C₁₇H₂₃NO₃Na (M + Na)⁺ 312.15701, found 312.15655. The enantiomeric excess of **7f** was determined to be 45% by HPLC with a Daicel Chiralpak OD-H column (eluent: 100:1 hexane/2-propanol; flow: 1.0 mL/min): $t_R = 10.6$ min for major enantiomer; $t_R = 11.4$ min for minor enantiomer. The preferred absolute configuration of **7f** was not determined.

4.3.5. (*R**)-Ethyl 2-[(1-methoxymethyl)-1*H*-indol-3-yl]propionate **7g** (Table 2, entry 9)

According to the typical procedure for C–H functionalization of indole, **7g** was prepared from **1f** (3.18 mg, 0.002 mmol, 1 mol %), ethyl α -diazopropionate **6c** (25.6 mg, 0.200 mmol) and **5e** (38.7 mg, 0.24 mmol). The crude product was purified by column chromatography (silica gel, 10:1→5:1 hexane/EtOAc) to provide **7g** (36.0 mg, 69%) as a colorless oil: TLC $R_f = 0.10$ (10:1 hexane/EtOAc); $[\alpha]_D^{22} -19.5$ (c 1.00, CHCl₃) for 32% ee; IR (neat) ν 2980, 1731, 1466, 1330, 1181, 1089 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ 1.22 (t, $J = 6.8$ Hz, 3H, CH₃CH₂O), 1.60 (d, $J = 7.2$ Hz, 3H, CH₃CHAr), 3.25 (s, 3H, CH₃OCH₂), 4.00 (q, $J = 7.2$ Hz, 1H, CH₃CHAr), 4.07–4.20 (m, 2H, CH₃CH₂O), 5.42 (s, 2H, CH₃OCH₂), 7.13 (s, 1H, Ar), 7.16 (dd, $J = 7.2, 7.7$ Hz, 1H, Ar), 7.25 (d, $J = 7.2, 8.2$ Hz, 1H, Ar), 7.46 (d, $J = 8.2$ Hz, 1H, Ar), 7.69 (d, $J = 7.7$ Hz, 1H, Ar); ¹³C NMR (100 MHz, CDCl₃) δ 14.2 (CH₃), 17.9 (CH₃), 36.9 (CH), 55.9 (CH₃), 60.7 (CH₂), 77.4 (CH₂), 109.9 (CH), 115.7 (C), 119.5 (CH), 120.0 (CH), 122.4 (CH), 125.4 (CH), 127.7 (C), 136.7 (C), 175.0 (CO); HRMS (ESI) calcd for C₁₅H₁₉NO₃Na (M + Na)⁺ 284.12571, found 284.12538. The enantiomeric excess of **7g** was determined to be 32% by HPLC with a Daicel Chiralpak OD-H column (eluent: 9:1 hexane/2-propanol; flow: 1.0 mL/min): $t_R = 8.0$ min for major enantiomer; $t_R = 10.7$ min for minor enantiomer. The preferred absolute configuration of **7g** was not determined.

4.3.6. (*R**)-2,4-Dimethyl-3-pentyl 2-[(1-methoxymethyl-4-methyl)-1*H*-indol-3-yl]propionate **7i** (Table 3, entry 2)

According to the typical procedure for C–H functionalization of indole, **7i** was prepared from **1f** (3.18 mg, 0.002 mmol, 1 mol %), **6a** (39.7 mg, 0.200 mmol) and 1-methoxymethyl-4-methylindole **5g** (42.1 mg, 0.24 mmol). The crude product was purified by column chromatography (silica gel, 5:1 hexane/Et₂O) to provide **7i** (49.5 mg, 72%) as a colorless oil: TLC $R_f = 0.18$ (5:1 hexane/Et₂O); $[\alpha]_D^{22} -28.1$ (c 0.89, CHCl₃) for 69% ee; IR (neat) ν 2966, 1730, 1465, 1181, 1096 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ 0.64 (d, $J = 6.8$ Hz, 3H, CH(CH₃)₂), 0.72 (d, $J = 6.8$ Hz, 3H, CH(CH₃)₂), 0.81 (d, $J = 7.2$ Hz, 3H, CH(CH₃)₂), 0.83 (d, $J = 7.7$ Hz, 3H, CH(CH₃)₂), 1.61 (d, $J = 6.8$ Hz, 3H, CH₃CHAr), 1.76–1.90 (m, 2H, CH(CH₃)₂), 2.76 (s, 3H, CH₃Ar), 3.18 (s, 3H, CH₃OCH₂), 4.40 (q, $J = 6.8$ Hz, 1H, CH₃CHAr), 4.58 (t, $J = 6.3$ Hz, 1H, CO₂CH), 5.36 (d, $J = 11.3$ Hz, 1H, CH₃OCHH), 5.44 (d, $J = 11.3$ Hz, 1H, CH₃OCHH), 6.89 (d, $J = 7.2$ Hz, 1H, Ar), 7.11 (dd, $J = 7.7, 7.7$ Hz, 1H, Ar), 7.18 (s, 1H, Ar), 7.31 (d, $J = 8.2$ Hz, 1H, Ar); ¹³C NMR (100 MHz, CDCl₃) δ 16.9 (CH₃), 17.2 (CH₃),

19.4 (CH₃), 19.5 (CH₃), 19.8 (CH₃), 20.5 (CH₃), 29.4 (CH), 29.4 (CH), 37.5 (CH), 55.7 (CH₃), 77.4 (CH₂), 82.6 (CH), 107.9 (CH), 116.9 (C), 122.2 (CH), 125.8 (CH), 126.4 (C), 130.8 (C), 136.6 (C), 175.5 (CO); HRMS (ESI) calcd for C₂₁H₃₁NO₃Na (M + Na)⁺ 368.21962, found 368.21912. The enantiomeric excess of **7i** was determined to be 69% by HPLC with a Daicel Chiralpak OD-H column (eluent: 200:1 hexane/2-propanol; flow: 1.0 mL/min): *t*_R = 15.5 min for minor enantiomer; *t*_R = 17.9 min for major enantiomer. The preferred absolute configuration of **7i** was not determined.

4.3.7. (*R**)-2,4-Dimethyl-3-pentyl 2-[(5-methoxy-1-methoxymethyl)-1*H*-indol-3-yl]propionate **7j** (Table 3, entry 3)

According to the typical procedure for C–H functionalization of indole, **7j** was prepared from **1f** (3.18 mg, 0.002 mmol, 1 mol %), **6a** (39.7 mg, 0.200 mmol) and 5-methoxy-1-methoxymethylindole **5h** (45.9 mg, 0.24 mmol). The crude product was purified by column chromatography (silica gel, 5:1 hexane/Et₂O) to provide **7j** (55.6 mg, 77%) as a colorless oil: TLC *R*_f = 0.13 (10:1 hexane/EtOAc); [α]_D²² –24.7 (c 1.12, CHCl₃) for 58% ee; IR (neat) ν 2966, 1730, 1488, 1222, 1172, 1088 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ 0.66 (d, *J* = 6.3 Hz, 3H, CH(CH₃)₂), 0.72 (d, *J* = 6.8 Hz, 3H, CH(CH₃)₂), 0.81 (d, *J* = 6.8 Hz, 3H, CH(CH₃)₂), 0.83 (d, *J* = 6.8 Hz, 3H, CH(CH₃)₂), 1.63 (d, *J* = 7.2 Hz, 3H, CH₃CHAR), 1.76–1.91 (m, 2H, CH(CH₃)₂), 3.18 (s, 3H, CH₃OCH₂), 3.86 (s, 3H, CH₃OAr), 4.00 (q, *J* = 7.2 Hz, 1H, CH₃CHAR), 4.58 (t, *J* = 6.3 Hz, 1H, CO₂CH), 5.35 (d, *J* = 11.3 Hz, 1H, CH₃OCHH), 5.40 (d, *J* = 11.3 Hz, 1H, CH₃OCHH), 6.89 (dd, *J* = 2.3, 8.6 Hz, 1H, Ar), 7.12 (s, 1H, Ar), 7.13 (d, *J* = 2.3 Hz, 1H, Ar), 7.34 (d, *J* = 8.6 Hz, 1H, Ar); ¹³C NMR (100 MHz, CDCl₃) δ 17.0 (CH₃), 17.2 (CH₃), 17.8 (CH₃), 19.4 (CH₃), 19.5 (CH₃), 29.4 (CH), 29.4 (CH), 37.1 (CH), 55.6 (CH₃), 55.8 (CH₃), 77.5 (CH₂), 82.6 (CH), 101.3 (CH), 110.7 (CH), 112.6 (CH), 115.4 (C), 126.0 (CH), 128.3 (C), 131.7 (C), 154.3 (C), 174.9 (CO); HRMS (ESI) calcd for C₂₁H₃₁NO₄Na (M + Na)⁺ 384.21453, found 384.21465. The enantiomeric excess of **7j** was determined to be 58% by HPLC with a Daicel Chiralpak AD-H column (eluent: 9:1 hexane/2-propanol; flow: 1.0 mL/min): *t*_R = 5.3 min for minor enantiomer; *t*_R = 6.2 min for major enantiomer. The preferred absolute configuration of **7j** was not determined.

4.3.8. (*R**)-2,4-Dimethyl-3-pentyl 2-[(5-bromo-1-methoxymethyl)-1*H*-indol-3-yl]propionate **7k** (Table 3, entry 4)

According to the typical procedure for C–H functionalization of indole, **7k** was prepared from **1f** (3.18 mg, 0.002 mmol, 1 mol %), **6a** (39.7 mg, 0.200 mmol) and 5-bromo-1-methoxymethylindole **5i** (57.6 mg, 0.24 mmol). The crude product was purified by column chromatography (silica gel, 5:1 hexane/Et₂O) to provide **7k** (67.9 mg, 83%) as a colorless oil: TLC *R*_f = 0.11 (10:1 hexane/EtOAc); [α]_D²² –36.0 (c 1.12, CHCl₃) for 64% ee; IR (neat) ν 2966, 1731, 1469, 1353, 1177, 1097 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ 0.66 (d, *J* = 6.8 Hz, 3H, CH(CH₃)₂), 0.71 (d, *J* = 6.8 Hz, 3H, CH(CH₃)₂), 0.82 (d, *J* = 6.8 Hz, 3H, CH(CH₃)₂), 0.84 (d, *J* = 6.8 Hz, 3H, CH(CH₃)₂), 1.62 (d, *J* = 7.2 Hz, 3H, CH₃CHAR), 1.78–1.90 (m, 2H, CH(CH₃)₂), 3.18 (s, 3H, CH₃OCH₂), 3.98 (q, *J* = 7.2 Hz, 1H, CH₃CHAR), 4.58 (t, *J* = 6.3 Hz, 1H, CO₂CH), 5.36 (d, *J* = 10.9 Hz, 1H, CH₃OCHH), 5.41 (d, *J* = 10.9 Hz, 1H, CH₃OCHH), 7.75 (s, 1H, Ar), 7.32 (d, *J* = 0.9 Hz, 1H, Ar), 7.32 (s, 1H, Ar), 7.85 (d, *J* = 0.9 Hz, 1H, Ar); ¹³C NMR (100 MHz, CDCl₃) δ 16.9 (CH₃), 17.2 (CH₃), 17.8 (CH₃), 19.3 (CH₃), 19.5 (CH₃), 29.4 (CH), 37.0 (CH), 55.8 (CH₃), 77.5 (CH₂), 82.9 (CH), 111.4 (CH), 113.4 (C), 115.5 (C), 122.4 (CH), 125.2 (CH), 126.5 (CH), 129.5 (C), 135.2 (C), 174.5 (CO); HRMS (ESI) calcd for C₂₀H₂₈BrNO₃Na (M + Na)⁺ 432.11448,

found 432.11508. The enantiomeric excess of **7k** was determined to be 64% by HPLC with a Daicel Chiralpak AD-H column (eluent: 9:1 hexane/2-propanol; flow: 1.0 mL/min): *t*_R = 4.5 min for minor enantiomer; *t*_R = 5.8 min for major enantiomer. The preferred absolute configuration of **7k** was not determined.

4.3.9. (*R**)-2,4-Dimethyl-3-pentyl 2-[(1-methoxymethyl-5-nitro)-1*H*-indol-3-yl]propionate **7l** (Table 3, entry 5)

According to the typical procedure for C–H functionalization of indole, **7l** was prepared from **1f** (3.18 mg, 0.002 mmol, 1 mol %), **6a** (39.7 mg, 0.200 mmol) and 1-methoxymethyl-5-nitroindole **5j** (45.9 mg, 0.24 mmol). The crude product was purified by column chromatography (silica gel, 3:1 hexane/EtOAc) to provide **7l** (52.1 mg, 69%) as a colorless oil: TLC *R*_f = 0.50 (5:1 hexane/EtOAc); [α]_D²² –71.0 (c 0.50, CHCl₃) for 72% ee; IR (neat) ν 2967, 1731, 1519, 1336, 1179, 1102 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ 0.63 (d, *J* = 6.8 Hz, 3H, CH(CH₃)₂), 0.71 (d, *J* = 6.8 Hz, 3H, CH(CH₃)₂), 0.83 (d, *J* = 6.8 Hz, 3H, CH(CH₃)₂), 0.85 (d, *J* = 6.8 Hz, 3H, CH(CH₃)₂), 1.67 (d, *J* = 7.2 Hz, 3H, CH₃CHAR), 1.77–1.94 (m, 2H, CH(CH₃)₂), 3.23 (s, 3H, CH₃OCH₂), 4.09 (q, *J* = 7.2 Hz, 1H, CH₃CHAR), 4.59 (t, *J* = 6.3 Hz, 1H, CO₂CH), 5.43 (d, *J* = 10.9 Hz, 1H, CH₃OCHH), 5.47 (d, *J* = 10.9 Hz, 1H, CH₃OCHH), 7.31 (s, 1H, Ar), 7.50 (d, *J* = 9.1 Hz, 1H, Ar), 8.16 (d, *J* = 1.8, 9.1 Hz, 1H, Ar), 8.70 (d, *J* = 2.3 Hz, 1H, Ar); ¹³C NMR (100 MHz, CDCl₃) δ 16.9 (CH₃), 17.1 (CH₃), 17.8 (CH₃), 19.3 (CH₃), 19.5 (CH₃), 29.3 (CH), 36.8 (CH), 56.1 (CH₃), 77.8 (CH₂), 83.3 (CH), 110.1 (CH), 117.1 (CH), 118.1 (C), 118.3 (CH), 127.3 (C), 128.4 (CH), 139.4 (C), 142.0 (C), 174.2 (CO); HRMS (ESI) calcd for C₂₀H₂₈N₂O₅Na (M + Na)⁺ 399.18904, found 399.18926. The enantiomeric excess of **7l** was determined to be 72% by HPLC with a Daicel Chiralpak AD-H column (eluent: 9:1 hexane/2-propanol; flow: 1.0 mL/min): *t*_R = 7.2 min for minor enantiomer; *t*_R = 11.6 min for major enantiomer. The preferred absolute configuration of **7l** was not determined.

4.3.10. (*R**)-2,4-Dimethyl-3-pentyl 2-[(1-methoxymethyl-7-methyl)-1*H*-indol-3-yl]propionate **7m** (Table 3, entry 6)

According to the typical procedure for C–H functionalization of indole, **7m** was prepared from **1f** (3.18 mg, 0.002 mmol, 1 mol %), **6a** (39.7 mg, 0.200 mmol) and 1-methoxymethyl-7-methylindole **5k** (42.1 mg, 0.24 mmol). The crude product was purified by column chromatography (silica gel, 5:1 hexane/Et₂O) to provide **7m** (57.3 mg, 83%) as a colorless oil: TLC *R*_f = 0.17 (5:1 hexane/Et₂O); [α]_D²² –9.7 (c 0.95, CHCl₃) for 86% ee; IR (neat) ν 2968, 1725, 1463, 1183, 1106 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ 0.67 (d, *J* = 6.8 Hz, 3H, CH(CH₃)₂), 0.72 (d, *J* = 6.8 Hz, 3H, CH(CH₃)₂), 0.82 (d, *J* = 6.8 Hz, 3H, CH(CH₃)₂), 0.84 (d, *J* = 6.8 Hz, 3H, CH(CH₃)₂), 1.61 (d, *J* = 7.2 Hz, 3H, CH₃CHAR), 1.76–1.91 (m, 2H, CH(CH₃)₂), 2.72 (s, 3H, CH₃Ar), 3.19 (s, 3H, CH₃OCH₂), 4.03 (q, *J* = 7.2 Hz, 1H, CH₃CHAR), 4.58 (t, *J* = 6.3 Hz, 1H, CO₂CH), 5.46 (d, *J* = 10.9 Hz, 1H, CH₃OCHH), 5.54 (d, *J* = 11.3 Hz, 1H, CH₃OCHH), 6.98 (d, *J* = 7.2 Hz, 1H, Ar), 7.05 (dd, *J* = 7.2, 7.7 Hz, 1H, Ar), 7.09 (s, 1H, Ar), 7.52 (d, *J* = 7.7 Hz, 1H, Ar); ¹³C NMR (100 MHz, CDCl₃) δ 17.0 (CH₃), 17.2 (CH₃), 18.3 (CH₃), 19.2 (CH₃), 19.4 (CH₃), 19.5 (CH₃), 29.4 (CH), 36.9 (CH), 55.1 (CH₃), 79.2 (CH₂), 82.6 (CH), 115.3 (C), 117.2 (CH), 120.2 (CH), 121.9 (C), 125.1 (CH), 127.5 (CH), 129.1 (C), 134.8 (C), 175.0 (CO); HRMS (ESI) calcd for C₂₁H₃₁NO₃Na (M + Na)⁺ 368.21962, found 368.21957. The enantiomeric excess of **7m** was determined to be 86% by HPLC with a Daicel Chiralpak IC column (eluent: 200:1 hexane/2-propanol; flow: 1.0 mL/min): *t*_R = 17.5 min for minor enantiomer; *t*_R = 18.9 min for major enantiomer. The preferred absolute configuration of **7m** was not determined.

4.4. Determination of absolute configuration of (R)-2,4-dimethyl-3-pentyl 2-[(1-methoxymethyl)-1H-indol-3-yl]propionate **7e** (Scheme 2)

4.4.1. (R)-2,4-Dimethyl-3-pentyl 2-(1H-indol-3-yl)propionate **9**

TMSOTf (0.061 mL, 0.33 mmol) was added to a solution of **7e** (55.0 mg, 0.166 mmol) and 2,2'-bipyridyl (77.8 mg, 0.498 mmol) in CH₂Cl₂ (1.0 mL) at 0 °C. After stirring for 0.5 h at 0 °C, the reaction was quenched by addition of water (3 mL). The mixture was extracted with EtOAc (2 × 5 mL), and the combined organic layers were washed with 1 M aq. HCl (2 × 3 mL), saturated aq. NaHCO₃ (3 mL) and brine (3 mL), and dried over anhydrous Na₂SO₄. Filtration and evaporation in vacuo furnished (R)-2,4-dimethyl-3-pentyl 2-[(1-hydroxymethyl)-1H-indol-3-yl]propionate **8** (60.5 mg), which was used without further purification. NaOH (66.4 mg, 1.66 mmol) was added to the solution of **8** (60.5 mg) in THF (3.6 mL) and water (0.4 mL). After stirring for 24 h at room temperature, the reaction was quenched with addition of saturated NH₄Cl (10 mL). The mixture was extracted with EtOAc (2 × 10 mL), and the combined organic layers were washed with brine (10 mL) and dried over anhydrous Na₂SO₄. Filtration and evaporation in vacuo furnished the crude product (62.0 mg), which was purified by column chromatography (silica gel, 6:1 hexane/EtOAc) to provide **9** (47.3 mg, 99%) as a yellow oil: [α]_D²² -40.4 (c 0.71, CHCl₃) for 82% ee; IR (neat) ν 3412, 2968, 1716, 1459, 1186 cm⁻¹; ¹H NMR (500 MHz, CDCl₃) δ 0.66 (d, J = 6.9 Hz, 3H, CH(CH₃)₂), 0.71 (d, J = 6.9 Hz, 3H, CH(CH₃)₂), 0.80 (d, J = 6.9 Hz, 3H, CH(CH₃)₂), 0.83 (d, J = 6.9 Hz, 3H, CH(CH₃)₂), 1.64 (d, J = 6.9 Hz, 3H, CH₃CHAR), 1.77–1.90 (m, 2H, CH(CH₃)₂), 4.07 (q, J = 6.8 Hz, 1H, CH₃CHAR), 4.58 (t, J = 6.3 Hz, 1H, CO₂CH), 7.10–7.13 (m, 1H, Ar), 7.17 (s, 1H, Ar), 7.19 (m, 1H, Ar), 7.35 (d, J = 8.0 Hz, 1H, Ar), 7.71 (d, J = 8.0 Hz, 1H, Ar), 8.03 (brs, 1H, NH); ¹³C NMR (125 MHz, CDCl₃) δ 16.9 (CH₃), 17.2 (CH₃), 18.0 (CH₃), 19.3 (CH₃), 19.5 (CH₃), 29.4 (CH), 29.4 (CH), 37.2 (CH), 77.2 (CH), 82.6 (CH), 111.1 (CH), 115.7 (C), 119.3 (CH), 121.5 (CH), 122.0 (CH), 126.5 (C), 136.1 (C), 175.2 (CO); HRMS (ESI) calcd for C₁₈H₂₅NO₂Na (M + Na)⁺ 310.17775, found 310.17797.

4.4.2. (R)-2-(1H-indol-3-yl)propane-1-ol **10**³⁰

LiAlH₄ (22.7 mg, 0.598 mmol) was added to a solution of **9** (43.0 mg, 0.150 mmol) in THF (1.0 mL) at 0 °C. After stirring for 4 h at room temperature, the reaction was quenched with addition of water (0.023 mL). 15% aq. NaOH (0.023 mL) and water (0.069 mL) were added to the reaction mixture. After stirring for 0.5 h at room temperature, filtration and evaporation in vacuo furnished the crude product (34.8 mg), which was purified by column chromatography (silica gel, 1:1 hexane/EtOAc) to provide **10** (22.7 mg, 87%) as a colorless oil: [α]_D²⁴ -23.3 (c 0.41, MeOH) for 82% ee [lit.,³⁰ [α]_D²⁵ +30.6 (c 0.46, MeOH) for (S)-**10** (90% ee)]; ¹H NMR (500 MHz, CDCl₃) δ 1.41 (d, J = 7.3 Hz, 3H, CH₃CHAR), 3.30–3.34 (m, 1H, CH₃CHAR), 3.80–3.86 (m, 2H, CH₂OH), 7.07 (s, 1H, Ar), 7.10 (m, 1H, Ar), 7.22 (m, 1H, Ar), 7.38 (d, J = 8.0 Hz, 1H, Ar), 7.67 (d, J = 8.0 Hz, 1H, Ar), 8.07 (brs, 1H, NH); ¹³C NMR (100 MHz, CDCl₃) δ 17.2 (CH₃), 33.9 (CH), 67.8 (CH₂), 111.3 (CH), 117.9 (C), 119.2 (CH), 119.3 (CH), 121.2 (CH), 122.1 (CH), 126.7 (C), 136.5 (C).

4.5. Synthesis of (S)-(+)-α-methyl-3-indolyacetic acid **11**, a constituent of acremoauxin A **12** (Scheme 3)

4.5.1. (S)-2,4-Dimethyl-3-pentyl 2-[(1-methoxymethyl)-1H-indol-3-yl]propionate *ent-7e*

According to the typical procedure for C–H functionalization of indole, *ent-7e* was prepared from Rh₂(R-PTTEA)₄·2EtOAc **2** (3.18 mg, 0.002 mmol, 1 mol %), **6a** (79.3 mg, 0.400 mmol) and **5e** (77.4 mg, 0.48 mmol). The crude product was purified by column chromatography (silica gel, 5:1 hexane/Et₂O) to provide *ent-7e* (110 mg, 83%) as a colorless oil: [α]_D²⁵ +20.6 (c 1.26, CHCl₃) for 83% ee.

4.5.2. (S)-2,4-Dimethyl-3-pentyl 2-(1H-indol-3-yl)propionate *ent-9*

According to the procedure for the preparation of **9**, *ent-9* was prepared from *ent-7e* (185.4 mg, 0.559 mmol). The crude product was purified by column chromatography (silica gel, 6:1 hexane/EtOAc) to provide the title compound (156.0 mg, 97%) as a colorless oil: [α]_D²⁵ +40.6 (c 0.98, CHCl₃) for 83% ee.

4.5.3. (S)-2-(1H-indol-3-yl)propane-1-ol *ent-10*³⁰

According to the procedure for the preparation of **10**, *ent-10* was prepared from *ent-9* (153.5 mg, 0.534 mmol). The crude product was purified by column chromatography (silica gel, 1:1 hexane/EtOAc) to provide the title compound (90.5 mg, 97%) as a colorless oil: [α]_D²⁴ +28.2 (c 0.48, MeOH) for 83% ee [lit.,³⁰ [α]_D²⁵ +30.6 (c 0.46, MeOH) for 90% ee].

4.5.3. (S)-(+)-α-methyl-3-indolyacetic acid **11**³¹

SO₃·pyridine complex (114 mg, 0.718 mmol) was added to a solution of *ent-10* (30.0 mg, 0.171 mmol) and Et₃N (0.11 mL, 0.789 mmol) in DMSO (1.0 mL) at room temperature. After stirring for 0.5 h at room temperature, the reaction mixture was poured into a mixture of saturated aq. NH₄Cl (5 mL) and saturated aq. NaHCO₃ (5 mL). The mixture was extracted with EtOAc (2 × 5 mL) and the combined organic layers were washed with brine (5 mL) and dried over anhydrous Na₂SO₄. Filtration and evaporation in vacuo gave the crude aldehyde, which was used without further purification. NaH₂PO₄·H₂O (534 mg, 3.42 mmol) and NaClO₂ (70.0 mg, 0.619 mmol) were added to the solution of the crude aldehyde in THF (1.0 mL), ^tBuOH (0.3 mL), 2-methyl-2-butene (0.3 mL) and water (1.0 mL). After stirring for 0.5 h, the reaction mixture was poured into saturated aq. NH₄Cl (5 mL), and whole was extracted with EtOAc (2 × 5 mL). The combined organic layers were washed with brine (5 mL) and dried over anhydrous Na₂SO₄. Filtration and evaporation in vacuo furnished the crude product (109 mg), which was purified by column chromatography (silica gel, 1:1 hexane/EtOAc) to provide **11** (19.8 mg, 61%) as a dark yellow oil: [α]_D²⁵ +42.6 (c 0.12, CH₂Cl₂) for 83% ee [lit.,^{31c} [α]_D²⁵ +41.5 (c 0.13, CH₂Cl₂) for >99% ee]; IR (neat) ν 3415, 2983, 1709, 1458, 1248 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ 1.63 (d, J = 7.3 Hz, 3H, CH₃CHAR), 4.05 (q, J = 7.3 Hz, 1H, CH₃CHAR), 7.13 (dd, J = 7.3, 8.0 Hz, 1H, Ar), 7.15 (s, 1H, Ar), 7.20 (dd, J = 7.3, 7.3 Hz, 1H, Ar), 7.36 (d, J = 8.0 Hz, 1H, Ar), 7.69 (d, J = 7.6 Hz, 1H, Ar), 8.07 (brs, 1H, NH); ¹³C NMR (100 MHz, CDCl₃) δ 17.5 (CH₃), 36.7 (CH), 111.2 (CH), 114.9 (C), 119.3 (CH), 119.7 (CH), 121.7 (CH), 122.4 (CH), 126.3 (C), 136.2 (C), 180.0 (CO). The enantiomeric excess of **11** was determined to be 83% by HPLC analysis with a Daicel Chiralpak IC-3 column [eluent: 9:1 hexane/2-propanol (contained 0.1% AcOH); flow: 0.5 mL/min; 40 °C]: t_R (major) = 12.2 min for (S)-**11**; t_R (minor) = 18.2 min for (R)-**11**.

Acknowledgments

This research was supported, in part, by a Grant-in-Aid for Scientific Research on Innovative Areas (Project No. 2105: Organic Synthesis Based on Reaction Integration) from the Ministry of Education, Culture, Sports, Science and Technology, Japan. We thank Ms. S. Oka, and M. Kiuchi of the Center for

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