Synthesis of chiral-at-metal rhodium complexes from achiral tripodal tetradentate ligands: resolution and application to enantioselective Diels–Alder and 1,3-dipolar cycloadditions†

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An improved synthesis of the racemic rhodium compound \([\text{RhCl}_2(\kappa^6C,N,N',P,L1)]\) (1) containing an achiral tripodal tetradentate ligand is reported. Their derived solvate complexes \([\text{Rh}(\kappa^6C,N,N',P,L1)(\text{Solv})_2][\text{SbF}_6]_2\) (Solv = NCMe, 2; H2O, 3) are resolved into their two enantiomers. Complexes 2 and 3 catalyze the Diels–Alder (DA) reaction between methacrolein and cyclopentadiene and the 1,3-dipolar cycloaddition reaction between methacrolein and the nitroene N-benzylidenephénylamine-N-oxide. When enantipure \((A_{R_3}R_1)-2\) was employed as the catalyst, enantiomeric ratios >99/1, in the R at C2 adduct, and up to 94/6, in the 3,5-endo isomer, were achieved in the DA reaction and in the 1,3-dipolar cycloaddition reaction, respectively. A plausible catalytic cycle that accounts for the origin of the observed enantioselectivity is proposed.

Introduction

Typically, a homogeneous chiral metal catalyst consists of a metal surrounded by a chiral ligand and accompanied by a range of achiral ligands. The source of chirality is the chiral ligand which configures a “chiral pocket” in the vicinity of the metal where the reaction between the catalytic substrates takes place.1

As early as 1979, Brunner argued: “It therefore seemed promising to move the inducing chirality to the metal atom where the catalysis is occurring”.2 However, in 2000, 21 years later, Muñiz and Bolm recognized that asymmetric catalysis with enantiomeric chiral-at-metal complexes was still unknown and that the only approach to this type of catalysts was diastereomeric complexes in which the configuration of the metal centre is predetermined by the coordination of a chiral ligand.3

It is interesting to point out that enantioselectively pure coordination compounds have been known since 1911 when Werner reported the resolution of the complex \([\text{Co}(\text{en})_3(\text{-NH}_3)]^{2+}\) [en = ethylenediamine, X = Cl or Br] into the individual mirror-imaged Δ- and Δ-enantiomers.4 However, it should be noted that the design and synthesis of efficient enantiopure catalysts in which the only stereogenic centre is the metal is not exempt from difficulties. On the one hand, given that different structural isomers are usually possible, the most suitable isomer must be selectively prepared or separated from the obtained mixture. In addition, this isomer will be obtained as a racemate that must be resolved following protocols that generally require a great deal of time with no guarantee of success.5 On the other, for these catalysts to achieve high efficiency, it is very convenient that they meet two requirements that seem contradictory a priori: configurational stability and labile coordination positions. Most likely for these reasons, the systematic development of chiral-only-at-metal catalysts has occurred only very recently. Indeed, although in 2003, Fontecave et al. demonstrated for the first time that chiral-only-at-metal bis-diimine Ru(n) complexes catalyze the oxidation of sulfides to sulfoxides by hydrogen peroxide with up to 59/41 er,6 only since 2013 the synthesis, resolution and catalytic applications of stereogenic-only-at-metal chiral catalysts have been systematically developed. From this date, Meggers group and others have been developing a family of bis-cyclometalated octahedral rhodium(m), iridium(m), ruthenium(n) and iron(n) complexes that efficiently catalyze a variety of organic transformations following Lewis acid7 and photoredox8 catalytic protocols.

Also in this line, Gladysz’s group reported that octahedral enantiopure salts of the tris-chelate ethylenediamine complexes \([\text{M}(\text{en})_3][\text{Barf}]_n\) [M = Co, Cr, Rh, Ir, n = 3; M = Pt, n = 4; Barf = B(3,5-C6H3(CF3)2)3] catalyze the addition of malonates to...
cyclopentenone or trans-β-nitrostyrene and that of the methyl 2-oxocyclopentanecarboxylate to di-tert-butyldiazodicarboxylate following a second coordination sphere mechanism involving substrate activation by hydrogen bonding to the coordinate NH₂ units. Modest enantioselectivities were achieved in all cases. Also, pseudo-octahedral geometry presents an enantiopure chiral-at-ruthenium complex containing a cyclometalated chelating NHC ligand that efficiently catalyzes the asymmetric cross metathesis, asymmetric ring closing metathesis and asymmetric ring opening cross metathesis of olefins.

On the other hand, tetrahedral complexes with stereogenic metal centres are rarely used as chiral catalysts mainly because they are too labile to ensure the absolute configuration of the metal centre. However, Shionoya et al. have reported the preparation and resolution of the tetrahedral chiral zinc complex 1 with high configurational stability using the achiral tetradentate ligand H₂L depicted in Scheme 1. When enantiopure S-1 was applied to the asymmetric oxo-Diels–Alder reaction of Danishefsky’s diene with 1-naphthaldehyde the Diels–Alder R-adduct was obtained in 98% yield with 93.5/6.5 er.

We have recently reported the application of achiral tripodal tetratdentate ligands to the synthesis of octahedral chiral-at-metal rhodium, iridium and ruthenium complexes. In particular, enantiopure rhodium complexes efficiently catalyze the enantioselective Diels–Alder reaction between methacrolein and cyclopentadiene with enantiomeric ratios of up to >99/1 (ref. 13c) and, on the basis of experimental NMR studies and DFT calculations, a detailed mechanistic pathway for the Friedel–Crafts reaction of trans-β-nitrostyrene and N-methyl-2-methylindole reaction is proposed, including an explanation of the origin of the obtained enantioselectivity.

Herein, we report on the synthesis and stereochemical resolution of the rhodium complex 2 containing the tripodal tetratdentate L₁ (Scheme 2) and its application to enantioselective Diels–Alder and 1,3-dipolar cycloadditions.

**Results and discussion**

**Synthesis of neutral dichloride complex [RhCl₂(k⁴C,N,N’,P-L₁)] (1)**

The achiral tripodal tetratdentate ligand L₁ has been prepared as reported elsewhere. The synthesis of the dichlorido complex [RhCl₂(k⁴C,N,N’,P-L₁)] (1) has been previously accomplished following the three steps protocol A depicted in Scheme 3. Reaction of RhCl₃ with the ligand L₁ in refluxing ethanol affords a mixture of fac-OC-6-43 (ref. 14) and mer-OC-6-41 trichlorido isomers that can be completely isomerized to the fac diastereomer by refluxing the mixture overnight in the same solvent. The cyclometalated isomer OC-6-54 of the dichlorido complex 1 could be obtained by refluxing in methanol pure trichlorido fac-OC-6-43 in the presence of NaOAc.

We have developed an improved synthesis of the dichlorido complex 1 following a one-pot, two steps reaction. First, RhCl₃ reacts with L₁ in refluxing EtOH; then, the resulting mixture is heated in the same solvent in the presence of NaOAc for 3 h (protocol B, Scheme 3). Only metalation at C6 carbon of the trifluoro aryl substituent was observed in both preparative routes.

**Synthesis of the solvate complexes [Rh(k⁴C,N,N’,P-L₁)(NCMe)₂][SbF₆]₂ (2) and [Rh(k⁴C,N,N’,P-L₁)(OH₂)₂][SbF₆]₂ (3)**

Treatment of the dichlorido complex 1 with AgSbF₆ in acetonitrile or in wet dichloromethane affords the solvate complexes [Rh(k⁴C,N,N’,P-L₁)(NCMe)₂][SbF₆]₂ (2) or [Rh(k⁴C,N,N’,P-L₁)(OH₂)₂][SbF₆]₂ (3), respectively (eqn (1)).

![Scheme 2 Chiral rhodium complex 2 bearing the achiral ligand L₁.](image)

The compounds were characterized by analytical and spectroscopic methods (see Experimental) and by the X-ray determination of the crystal structure of the acetonitrile compound 2. The metal and the sp³ nitrogen atoms of the ligand are stereochemically centres. However, the H₁, H₃P[¹H] and ¹²C[¹H] NMR spectra of complexes 2 and 3 consists of only one set of sharp resonance signals indicating that only one of the three possible isomers has been isolated. NMR measurements allow us to ascertain the geometry of the isolated isomer. As the spectroscopy features of both compounds are comparable, we will focus our discussion on the acetonitrile compound 2 as the model. The value of the J(PH) coupling constants together with COSY, HSQC and HMBC experiments permit the assignment of the three pairs of diastereotopic protons which have been labelled...
as CH$_2$(Py), CH$_2$(Ph) and CH$_2$(P) (see Scheme 4A). Scheme 4B shows the NOE pattern measured for these six methylene protons. Additionally, the methyl protons of the acetonitrile ligand trans to the sp$^3$ nitrogen atom exhibit NOE relationship with the 6-CH(Py) and 6-CH(Ph) protons of the tetradentate ligand. Accordingly, a $J$(PH) coupling constant of 4.0 Hz was measured for the proton-6 of the pyridine moiety (6-CH(Py)). At this point, it should be noted that when the ligand L1 coordinates with a metal in an octahedral geometry with a $\kappa^4$C,N,N'$,P$ coordination mode, the relative configuration of the sp$^3$

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Scheme 3  Synthesis of the dichlorido complex [RhCl$_2$(k$^4$C,N,N'$,P$-L1)] as described in the bibliography.\textsuperscript{15e} (A) and as reported in this work (B).

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Scheme 4  Assignment of the methylene descriptors (A) and NOE pattern (B) on OC-6-35 isomers of complex 2.
The crystal structure of racemic cation of 2. The \((S)_N\)-OC-6-35-C enantiomer of the racemate is depicted. For clarity, hydrogen atoms have been omitted, and only ipso-carbon atoms of phenyl groups are depicted.

Table 1 Selected bond lengths (Å) and angles (°) of complex 2

| Bond/ Angle | Value 1 | Value 2 | Value 3 |
|-------------|---------|---------|---------|
| Rh-P        | 2.2968(12) | N(1)-Rh-N(2) | 80.10(15) |
| Rh-N(1)     | 2.069(4)    | N(1)-Rh-N(3) | 174.12(15) |
| Rh-N(2)     | 2.102(4)    | N(1)-Rh-N(4) | 90.59(14)  |
| Rh-N(3)     | 2.028(4)    | N(1)-Rh-C(28) | 84.30(16)  |
| Rh-N(4)     | 2.170(4)    | N(2)-Rh-N(3) | 94.02(15)  |
| Rh-C(28)    | 2.007(4)    | N(2)-Rh-N(4) | 88.84(14)  |
| P-Rh-N(1)   | 95.17(11)   | N(2)-Rh-C(28) | 86.84(16)  |
| P-Rh-N(2)   | 174.91(11)  | N(3)-Rh-N(4) | 89.56(14)  |
| P-Rh-N(3)   | 90.69(11)   | N(3)-Rh-C(28) | 95.16(16)  |
| P-Rh-N(4)   | 93.15(11)   | N(4)-Rh-C(28) | 173.81(16) |
| P-Rh-C(28)  | 90.81(13)   | N(4)-Rh-C(28) | 173.81(16) |

Resolution of the \(A_{Rh,R_N}/C_{Rh,S_N}\) racemate of the solvate complex 2

As above mentioned, complex 2 was obtained as an equimolar mixture of the \(C_{Rh,S_N}\) and \(A_{Rh,R_N}\) diastereomers. In order to apply this complex as an asymmetric catalyst, it is necessary to resolve the obtained racemate. To this end, we developed a diastereomerization protocol of resolution using \((S)\)-phenylglycine as a chiral auxiliary.

Reaction of complexes 2 with \((S)\)-phenylglycine. Racemic 2 reacted with enantiopure \((S)\)-phenylglycine in refluxing ethanol in the presence of NaHCO₃ to afford 1/1 diastereomeric mixtures of \([A_{Rh,S_N,S_C}]\)-4 and \([C_{Rh,R_N,S_C}]\)-4 (eqn (2)).

The new complexes were characterized by spectroscopic and analytical methods (see Experimental). In particular, for both diastereomers, a NOE relationship between the NH₂ protons of the amino carboxylate ligand and one of the protons of each one of the CH₂(Py) and CH₂(P) methylene groups, indicates that the isolated isomers are those in which the oxygen atom of the amino carboxylato group is trans to the \(sp^3\) nitrogen of the diphosphino group, the other face is unencumbered. Therefore, when, in complex 2, prochiral substrates coordinate the metal displacing the labile acetonitrile ligands, the attack of a putative reagent would take preferably through the less hindered enantioface of the substrate originating enantioselectivity in the involved process.
tetradeutate ligand (eqn (2)). Probably, as it was found by DFT calculations carried out on a phenylglycinato rhodium complex containing a related tripodal tetradeutate ligand,183 diastereomers with the oxygen atom trans to the sp² nitrogen of the tetradeutate ligand are more stable than the corresponding isomers in which the oxygen atom is trans to the metalated carbon atom. Assuming a similar behaviour for complex 2, stereoselection in the formation of the isolated glycinato complexes 4 would mainly rely on thermodynamics.

The determination of the absolute configuration of each of the complexes has been carried out by NMR from a mixed solution of the two diastereomers (see ESI†). In particular, the 6-CH(Py) proton (δ = 8.32 ppm) of one diastereomer exhibits a nuclear Overhauser effect (NOE) with the asymmetric C≡H proton (δ = 3.98 ppm) determining the configuration for this diastereomer as (A[Rh,S,N,S]C)⁴. In the other diastereomer, determined as (C[Rh,R,S,N,S]C)⁴, the C≡H proton (δ = 3.10 ppm) correlates by NOE with the phenyl protons of the PPh₂ fragment (δ = 7.74 ppm). The ⁴¹P{¹H} NMR spectrum of the mixture of diastereomers consists of two doublets centred at 29.67 ppm (J(RHP) = 130.8 Hz), assigned to the (A[Rh,S,N,S]C)⁴ isomer, and at 28.83 ppm (J(RHP) = 130.1 Hz), attributed to the (C[Rh,R,S,N,S]C)⁴ isomer (see ESI†). Crystallization of the 1/1 mixture of isomers from MeOH/Et₂O affords a 57/43 molar ratio mixture from which individual diastereomers were separated by chromatography.

Separation of the diastereomeric mixture of glycinato complexes. Diastereomeric mixtures of (A[Rh,S,N,S,C])⁴ and (C[Rh,R,S,N,S,C])⁴ were separated by flash chromatography on a silica column using a CH₂Cl₂/MeOH mixture as eluent (see Experimental). Three fractions A, B and C were consecutively collected. Fraction A (ca. 21% yield) was diastereopure (A[Rh,S,N,S,C])⁴. Fraction B (ca. 48% yield) was a mixture of the two diastereomers (A[Rh,S,N,S,C])⁴ and (C[Rh,R,S,N,S,C])⁴ that was recycled. Fraction C (ca. 18% yield) was diastereopure (C[Rh,R,S,N,S,C])⁴.

The ⁴¹P{¹H} NMR spectrum of the starting mixture and those of each diastereomer, once separated by chromatography, are shown in Fig. 3.

Replacement of the coordinated auxiliary. The chiral auxiliary was replaced with two chlorido ligands, under retention of the configuration, by addition of excess of concentrated HCl(aq). From diastereopure (A[Rh,S,N,S,C])⁴ and (C[Rh,R,S,N,S,C])⁴, enantiopure dichlorido (A[Rh,R,S,C])¹ and (C[Rh,S,N,S,C])¹ were obtained, respectively, in more than 99.5/0.5 enantiomeric ratio in both cases.¹⁷ Fig. 4 shows the HPLC traces of a racemic mixture of the dichlorido complex 1 as well as those of the pure enantiomers (C[Rh,R,S,C])¹ and (A[Rh,R,S,C])¹. The circular dichroism spectra of both enantiomers are shown in Fig. 4, too. No significant changes of the HPLC traces were observed when enantiopure 1 was heated at 80 °C in methanol, in a high-pressure NMR tube, for 48 h. Thus, it can be argued that the configuration at the metal is thermally stable.

From the enantiopure chlorido complex (A[Rh,R,S,C])¹, enantiopure solvate complex (A[Rh,R,S,C])² was obtained following the treatment above reported for the preparation of racemic solvate complexes from racemic chlorido complex 1 (eqn (1)).

Catalytic Diels–Alder reaction between methacrolein and cyclopentadiene

Complexes 2 and 3 catalyze the Diels–Alder reaction between methacrolein (MA) and cyclopentadiene (HCp) in
dichloromethane. To determine the optimal conditions, we first tested the activity of rac-2 and rac-3. As shown in Table 2, at 298 K, to get good conversions with a catalyst loading of 5 or 10 mol%, several hours are needed (entries 1, 2, 5 and 6). Decreasing temperature increases the reaction time and diminishes the obtained exo/endo ratio of the adduct (entries 3 and 4). Complex 2 is more active than complex 3 (compare entries 1 and 2 with entries 5 and 6, respectively). At 5 mol% of catalyst loading, better exo/endo ratios than at 10 mol% catalyst loading were achieved but, to reach similar conversions, it is necessary to increase importantly the reaction time (entries 1, 2 and 5, 6). Taking into account all these data, we carried out the reaction between methacrolein and cyclopentadiene using 10 mol% of enantiopure (A$_{Rh}$,R$_N$)-2, at 298 K. After 8 h of reaction, 97% of conversion was reached with an exo/endo ratio of 93/7. Notably, enantioselectivity in the R at C2 isomer almost perfect was obtained (er > 99/1, entry 7). Comparing the catalytic results using the CF$_3$-substituted (A$_{Rh}$,R$_N$)-2 complex with those of the unsubstituted one$^{14}$ (after 24 h at 298 K, conv. 93%; exo/endo ratio: 88/12; er: 91/9), the CF$_3$-substituted complex has about 3 times higher catalytic activity, better diastereoselectivity and a remarkable increase in enantioselectivity, from 91/9 to >99/1 of er. These catalytic differences are understood by a large difference in the trans effect favourable to the CF$_3$ group which enhances the catalytic activity by favouring the de-and coordination of the substrates. On the other hand, enantioselectivity increases because the non-catalytic route to products becomes less competitive versus the catalytic route.

**Catalytic 1,3-dipolar cycloaddition reaction between methacrolein and N-benzylidenphenylamine-N-oxide**

Complex 2 also catalyzes the 1,3-dipolar cycloaddition reaction between methacrolein and the nitrone N-
benzylidenphenylamine-N-oxide. Table 3 lists a selection of the results obtained together with the reaction conditions employed. First, we tested rac-2 as the catalyst (entries 1–4). To achieve moderate to good conversions, 24 h of reaction are necessary even operating at 298 K (entry 1). Only endo isomers were detected and good regioselectivity in favour of the 3,5-endo isomer was obtained under all conditions explored. At 298 K, using (Arh,RN)-2 as the catalyst, an enantiomeric ratio of 94/6 was measured for the major adduct of the 3,5-endo diastereomer (entry 5). Lowering temperature to 273 K diminishes the er from 94/6 to 84/16 (entry 6). As long reaction times are necessary to reach acceptable conversions, a greater contribution of the thermal reaction is, probably, at the origin of the observed lowering of the er.

**Proposed mechanism and origin of the enantioselectivity**

The proposed catalytic cycle is depicted in Fig. 5A. One molecule of methacrolein (MA) replaces the more labile molecule of acetonitrile trans to the aromatic carbon atom of the enantiopure catalyst (Arh,RN)-2. Assuming that the methacrolein coordinates as a planar molecule with an s-trans conformation for the double carbon–oxygen and carbon–carbon bonds and with an E-configuration around the carbon–oxygen double bond,20 its Si-face becomes much more accessible to the cyclopentadiene or to the nitrone than its Re-face, because, in the chiral pocket generated in the diastereopure catalyst, the Re-face is shielded by the phenyl rings of the PPh2 group (Fig. 5B). An attack of the cyclopentadiene substrate through the Si-face of the coordinated methacrolein renders the R at C2 enantiomer of the DA adduct as it was experimentally observed. On the other hand, an endo approach of the nitrone through the Si-face of the coordinated methacrolein would render the 3R,5R isomer of the corresponding isoxazolidine.

**Conclusions**

Judiciously designed achiral tripodal tetradentate ligands are able to generate octahedral rhodium complexes in which the metal is a stereogenic centre. This is the case of the dichlorido complex 1. The frame generated by the polydentate ligand confers configurational stability to the octahedral molecule even when the two remaining positions are occupied by labile solvent molecules as in complexes 2 and 3. Notably, the compounds 2 and 3 meet two important requirements that make them good candidates for asymmetric catalysts: configurational stability and labile coordination positions. Indeed, after stereochemical resolution the chiral complex (Arh,RN)-2 catalyzes the Diels–Alder reaction between methacrolein and cyclopentadiene with a >99/1 enantiomeric ratio. Furthermore, the same catalyst mediates the 1,3-dipolar cycloaddition reaction between methacrolein and the nitrone.

**Table 3**

Selected data for the reaction between MA and N-benzylidenphenylamine-N-oxide catalyzed by complex 2a

| Entry | Cat. | T (K) | t (h) | Conv. (%) | (3,4)/(3,5) | erd |
|-------|------|-------|-------|-----------|-------------|-----|
| 1     | (rac)-2 | 298   | 24    | 64        | 5/95        | —   |
| 2     | (rac)-2 | 281   | 24    | 17        | 2/98        | —   |
| 3     | (rac)-2 | 273   | 24    | 8         | 6/94        | —   |
| 4     | (rac)-2 | 263   | 24    | 0         | —           | —   |
| 5     | (Arh,RN)-2 | 298 | 24    | 78        | 4/96        | 94/6|
| 6     | (Arh,RN)-2 | 273 | 164   | 68        | 12/88       | 84/16|

* Reaction conditions: catalyst 1.16 × 10−3 mmol (5.0 mol%); MA 7.00 × 10−3 mmol, 10 mg of 4 Å MS, and nitrone 3.50 × 10−3 mmol in 0.5 mL of CD2Cl2. Based on nitrone. Determined by 1H NMR. Only endo isomers were detected. Determined for the 3,5-endo isomer by integration of the 1H NMR signals of the diastereomeric (S)-methylbenzylimine derivatives.21
N-benzylidenphenylamine-N-oxide affording only endo-adducts with enantiomeric ratios of up to 94/6 for the 3,5-endo adduct. The molecular structure of complex 2, determined by X-ray diffraction analysis, reveals the presence of a chiral pocket around the labile solvent molecules coordination sites. When a prochiral substrate coordinates the metal inside this chiral pocket, the two phenyl groups of the tetradequate ligand shield one of its enantiopairs. Consequently, an incoming reagent preferentially approaches the coordinated substrate through its clearer enantiopair resulting in enantioselection.

Finally, the findings reported in this work open the door to the design and use of new polydentate ligands capable of generating stereogenic metal centres both in octahedral metal geometries and in metal complexes with other coordination numbers. The studies can be extended to other metals and, in particular, to cheap, non-toxic and earth-abundant metals of the first transition series, with inherent economic and environmental benefits. Further work in this area is in progress in our laboratory and will be reported in due course.

Experimental

All preparations have been carried out under argon, unless otherwise stated. All solvents were treated in a PS-400-6 Innovative Technologies Solvent Purification System (SPS). Infrared spectra were recorded on PerkinElmer Spectrum-100 (ATR mode) FT-IR spectrometer. 1H, 13C, 31P spectra were obtained using standard 1H, 13C, 31P) correlation spectra were obtained using standard 1H, 13C, 31P) correlation spectra were obtained using standard procedures. Flash chromatography was performed on an Interchim (PuriFlash 450) instrument using a PF-50SILHP column (25 g, 30 μm).

Improved synthesis of the complex 1

At RT, to a suspension of RhCl₃ × H₂O (1.00 g, 3.85 mmol) in 20 mL of ethanol, 2.07 g of L₁ were added. The resulting suspension was stirred under reflux overnight. During this time, the colour of the suspension gradually changes from pink-red to yellow. After the reaction time, the suspension was cooled to room temperature and 2.21 g [8.31 mmol] of NaAcO was added. The resulting suspension was stirred for 3 h under reflux. During this time, a yellow precipitate was formed. The solid was filtered off and extracted with CH₂Cl₂ (3 × 15 mL). The resulting yellow solution was vacuum-dried to give analytically pure compound 1. Yield: 2.01 g (73%). The spectroscopic data for complex 1 were in agreement with the previously reported in the literature.¹

Preparation and characterization of the complex 2

To a solution of [RhCl₂(k⁺CN,N,N,P-L₁)] (1) (1.00 g, 1.40 mmol) in 40 mL of acetonitrile, 963.4 mg (2.81 mmol) of AgSbF₆ were added. The resulting suspension was placed under reflux for 40 h and then filtered to remove any insoluble material. The resulting light yellow solution was evaporated to ca. 1 mL. The slow addition of Et₂O (25 mL) led to the precipitation of a pale yellow solid that was filtered off, washed with Et₂O (3 × 5 mL) and vacuum-dried. Yield: 1.54 g (92%) Anal. calcd C₃₇H₃₃F₁₅N₂O₁₉RhSb₂: C, 34.30; H, 2.73; N, 4.10. Found: C, 34.13; H, 2.67; N, 3.97. IR (solid, cm⁻¹): ν(CF₃) 2172, 1376, 1204, 1099. CD spectra were determined in CH₂Cl₂/acetone-d₆, RT, ppm: δ = 5.0 (broad pseudo-triplet, J = 4.6 Hz, 1H, 6-CH₂(P)), 8.01 (ptd, J = 7.8, 1.4 Hz, 1H, CH(Ar)), 7.80–7.42 (m, 14H, H(Ar)), 6.97 (d, J = 9.6 Hz, 1H, 5-CH(P)), 6.84–6.78 (m, 4H, 2×CH(Ar)), 6-CH(P), 3-CH(P)), 5.22 (d, J = 15.8 Hz, 1H, pro-S-CH₂(P)), 4.75 (d, J = 16.0 Hz, 1H, pro-R-CH₂(P)), 4.74 (d, J = 14.0 Hz, 1H, pro-CH(P)), 4.64 (d, J = 17.9 Hz, 1H, pro-R-CH₂(P)), 4.52 (d, J = 14.1, 5.0, 1.8 Hz, 1H, pro-S-CH₂(P)), 4.07 (d, J = 17.9 Hz, 1H, pro-S-CH₂(P)), 2.69 (s, 3H, NCMe cis to C), and 2.42 (s, 3H, NCMe trans to C). ¹³C¹H NMR (125.77 MHz, CD₂Cl₂/acetone-d₆, RT, ppm): δ = 155.79 (d, J = 1.9 Hz, 2-C(Py)), 149.30 (brdd, J_Rh-C = 29.1 Hz, J_P-C = 8.7 Hz, Crh), 148.30 (s, 6-CH(P)), 148.16 (s, 2-C(Py)), 141.05–128.45 (m, 18C(H)(Ar)), 127.30 (q, J_Rh-C = 32.4 Hz, 4-C(P)), 127.08 (d, J_Rh-C = 5.8 Hz, NCMe cis to C), 123.52 (q, J_P-C = 271.40 Hz, CFS), 125.98–120.26 (m, 6C(H)(Ar)), 124.13 (brs, NCMe trans to C), 120.26 (m, 3C(H)(Ar)), 119.30 (m, 3C(H)(Ar)), 114.1 Hz).
Preparation and characterization of the complex 3

At room temperature, to a solution of [RhCl₂(κ⁴C,N,N′,P-L1)] (1) (500.0 mg, 0.701 mmol) in 30 mL of wet dichloromethane, 1.55 mmol of AgSbF₆ were added. The pale yellow suspension was stirred for 72 h and then was filtered to remove the AgCl formed. The solution was evaporated to ca. 5 mL. Addition of 10 mL Et₂O and 10 mL of n-pentane led to the precipitation of a light yellow solid that was filtered off and vacuum-dried to afford complex 3 as a microcrystalline solid. Yield: 406.2 mg (79%). Anal. calc. C₅₁H₃₅F₉N₃O₂PRhSb: C, 49.77; H, 3.56; N, 4.24. Found: C, 49.38; H, 3.47; N, 4.00. IR (solid, cm⁻¹): ν(NH₂) 2959 (br), ν(C=O) 1640 (s), ν(SbF₆) 656 (s).

Preparation y characterization of the diastereomers (C₈Rh₄N₅S₄C₄Rh₄N₅S₄C₄)-4

To a suspension of 72.1 mg (0.477 mmol) of (S)-phenylglycine in 40 mL of EtOH, 60.1 mg (0.715 mmol) of NaHCO₃ were added. A light yellow solid that was filtered off and vacuum-dried to afford complex 3 as a microcrystalline solid. Yield: 406.2 mg (79%). Molar ratio (C₈Rh₄N₅S₄C₄):(A₈Rh₄N₅S₄C₄)-4, 1/1. Crystallization of

{(C₈Rh₄N₅S₄C₄)-4:(A₈Rh₄N₅S₄C₄)-4, 57/43 molar ratio mixture of diastereomers. Anal. calc. C₄₁H₃₅F₉N₃O₂PRhSb: C, 49.77; H, 3.56; N, 4.24. Found: C, 49.38; H, 3.47; N, 4.00. IR (solid, cm⁻¹): ν(NH₂) 2959 (br), ν(C=O) 1640 (s), ν(SbF₆) 656 (s).}
(d, J = 2.0 Hz, 2-C(Py)), 147.42 (s, 6-CH(Py)), 145.10−115.88 (m, 34C(H)(Ar)), 74.84 (s, CH3(Py)), 68.52 (d, J = 5.6 Hz, CH3(P)), 67.69 (s, CH2(Ph)), 58.61 (s, C*). 19F[1H] NMR (282.33 MHz, CD2Cl2, RT, ppm): δ = −62.54 (s). 31P[1H] NMR (202.46 MHz, CD2Cl2, RT, ppm): δ = 28.83 (d, J = 130.1 Hz).

Chromatographic separation of the diastereomers (C$_{Rh}$R$_N$S$_C$/A$_{Rh}$S$_N$S$_C$)-4

An equimolar mixture of diastereomers (C$_{Rh}$R$_N$S$_C$/A$_{Rh}$S$_N$S$_C$): [Rh(k$_5$C$_{5}$N$_2$N$_2$F$_2$LL)](S-phenylglycinato)[SbF$_6$] (200.0 mg, 0.194 mmol) was eluted with a CH$_2$Cl$_2$/MeOH (98.75/1.25, v/v, 30 mL min$^{-1}$) mixture on a silica column (SI-HP, 25 g, 30 µm). Three fractions A, B and C, were consecutively collected. Fraction A (43.0 mg, 0.042 mmol, 21%) was diastereopure (C$_{Rh}$R$_N$S$_C$)-4. Fraction B (96.0 mg, 0.093 mmol, 48%) was an equimolar mixture of the diastereomers (C$_{Rh}$R$_N$S$_C$/A$_{Rh}$S$_N$S$_C$)-4. Fraction C (36.0 mg, 0.035 mmol, 18%) was diastereopure (C$_{Rh}$R$_N$S$_C$)-4.

Elimination of the auxiliary. Crystallization of enantiopure (A$_{Rh}$R$_N$)-1 and (C$_{Rh}$S$_N$)-1

To a suspension of diastereopure (A$_{Rh}$R$_N$S$_C$)-4 (20.0 mg, 0.019 mmol) in 4 mL of EtOH, 0.20 mL of 12 M HCl(aq) were added. After 48 h at RT, a pale yellow microcrystalline solid precipitated from the bright yellow solution. It was collected by filtration and successively washed with EtOH (3 × 3 mL) and Et$_2$O (3 × 3 mL). The second fraction of the product was obtained from the filtrate (12 mg). Total yield: 89%. The isolated solid was characterized as (A$_{Rh}$R$_N$)-1. The diastereopure (C$_{Rh}$S$_N$)-1 was similarly obtained starting from (C$_{Rh}$R$_N$S$_C$)-4 (yield: 87%). The optical purity of the samples was determined by HPLC with a Chirapak IB column (0.46 × 25 cm) with IB guard (0.46 cm × 5 cm) (65/35, CH$_3$CN/H$_2$O, 0.1% CF$_3$COOH; 0.4 mL min$^{-1}$; t$_{R}$ 151.8 and 156.7 min for (A$_{Rh}$R$_N$)-1 and (C$_{Rh}$S$_N$)-1, respectively).

Preparation of the diastereopure solvate complex 2

Diastereopure (A$_{Rh}$R$_N$S$_C$)-2 was prepared starting from the corresponding diastereopure chlorido complex (A$_{Rh}$R$_N$)-1, as reported above for rac-2.

Catalytic procedure for the DA reaction between methacrolein and cyclopentadiene

At RT, in an NMR tube, to a solution of 1.16 × 10$^{-3}$ or 2.32 × 10$^{-3}$ mmol (5 or 10 mol%) of the corresponding catalyst in CD$_2$Cl$_2$ (0.5 mL), 2.0 µL [0.023 mmol] of methacrolein and 10 mg of 4 Å MS were added. After 5 min, 7.0 µL (0.081 mmol) of freshly distilled cyclopentadiene was added. After the indicated reaction times, the conversion and exo/endo ratio were determined by $^1$H NMR. The enantiomeric ratio for the exo isomer was determined by integration of the $^1$H NMR signals in presence of the chiral shift reagent (+)-Eu(hfc)$_3$. Major enantiomer (R), δ = 17.10 ppm; minor enantiomer (S), δ = 19.09 ppm.$^{18}$

Catalytic procedure for the 1,3-dipolar cycloaddition reaction between methacrolein and N-benzylidenephénylamine-N-oxide

In an NMR tube, complex 2 (1.16 × 10$^{-3}$ mmol (2.70 mg), 5 mol%) was dissolved in CD$_2$Cl$_2$ (0.2 mL) at −25 °C. Freshly distilled methacrolein (5 µL, 0.07 mmol) and 10.0 mg of activated 4 Å molecular sieves were added. After seeking the resulting suspension, a solution of N-benzylidenephénylamine-N-oxide (6.70 mg, 0.035 mmol) in CD$_2$Cl$_2$ (0.3 mL) was added. Regioselectivity was determined by the integration of the $^1$H NMR signals of the C3 proton on the crude mixture in CD$_2$Cl$_2$. Enantioselectivity was determined by the integration of the $^1$H NMR signals of the diastereomeric (S)-methylbenzylamine derivatives.

Conflicts of interest

The authors declare no competing financial interests.

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15 To simplify the notation of the stereochemical descriptors, we will label these diastereomers as $C_{\text{Rh}, \text{RN}}$ and $A_{\text{Rh}, \text{SN}}$. R and S refer to the configuration at the $sp^3$ nitrogen and C to the clockwise or anticlockwise arrangement of the donor atoms at the equatorial plane of the octahedral metal complex.

16 CCDC-2206804.

17 Due to the nomenclature rules, in this transformation, retention of the configuration implies the change of the metal descriptor from A to C, and vice versa.

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