tolyl), 391 ($M^+ - C_6H_4CO_2Et$), 300 ($M^+ -$ tolyl $- C_6H_4CO_2Et$), 209, 149. IR (neat): 1715 (C═O), 1584, 1387, 1281, 1103, 795, 752 cm^{-1}. Anal. Calcd for $C_{23}H_{23}BiO_2$: C, 51.12; H, 4.29. Found: C, 51.82; H, 4.33.

(4-Acetylphenyl)bis(4-methylphenyl)bismuthane (6f). Colorless crystals. Mp: 101−104 °C. ^{1}H NMR: δ 2.32 (s, 6H), 2.56 (s, 3H), 7.21 (d, J = 7.6 Hz, 4H), 7.61 (d, J = 7.6 Hz, 4H), 7.83 (d, J = 6.0 Hz, 2 H), 7.92 (d, J = 6.0 Hz, 2H). FABMS (m/z): 511 ($M^+ + 1$), 419 ($M^+ -$ tolyl), 391 ($M^+ - C_6H_4COMe$), 300 ($M^+ -$ tolyl $- C_6H_4COMe$), 209. IR (KBr): 1684 (C═O), 1576, 1485, 1385, 1269, 1184, 1007, 953, 797 cm^{-1}. Anal. Calcd for $C_{22}H_{21}BiO$: C, 51.77; H, 4.15. Found: C, 51.70; H, 4.15.

Reaction of Arylbis(4-methylphenyl)bismuthanes 6b,c with Sulfuryl Chloride. Typical Procedure. To a dichloromethane (10 mL) solution of bismuthane **6b** (0.30 g, 0.44 mmol) was added sulfuryl chloride (0.47 μL, 0.48 mmol) at −78 °C. As the mixture was warmed gradually to room temperature, the color of the reaction mixture turned yellow. After 1 h, the solvent was removed under reduced pressure to leave dichloride **11** as a yellow solid, which was purified by recrystallization from a mixture of dichloromethane and hexane.

Bis(4-methylphenyl)(1-phenanthryl)bismuth Dichloride (11). Yield: 90%. Mp: 145−147 °C. ^{1}H NMR: δ 2.49 (s, 6H), 7.57 (d, J = 8.2 Hz, 4H), 7.6−7.8 (m, 5H), 8.0−8.1 (m, 1H), 8.25 (s, 1H), 8.65 (d, J = 8.2 Hz, 4H), 8.7−8.8 (m, 2H). FABMS (m/z): 603 ($M^+ -$ Cl), 391 ($M^+ - 2$Cl − phenanthryl), 443, 268, 209. Anal. Calcd for $C_{28}H_{23}BiCl_2$: C, 52.60; H, 3.63. Found: C, 52.23; H, 3.53.

(9-Anthryl)bis(4-methylphenyl)bismuth Dichloride (12). Yield: 88%. Mp: 170−173 °C. ^{1}H NMR: δ 2.49 (s, 6H), 7.4−7.5 (m, 4H), 7.54 (d, J = 8.5 Hz, 4H), 8.0−8.1 (2H, m), 8.3−8.4 (m, 2H), 8.62 (s, 1H), 8.65 (d, J = 8.5 Hz, 4H). FABMS (m/z): 603 ($M^+ -$ Cl), 476 ($M^+ -$ tolyl − H), 391 ($M^+ -$ anthryl), 268, 212, 209. IR (KBr): 1522, 1473, 1446, 1385, 1266, 1248, 1183, 997, 899, 795, 727 cm^{-1}. Anal. Calcd for $C_{28}H_{23}BiCl_2$: C, 52.60; H, 3.63. Found: C, 52.43; H, 3.50.

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Additions and Corrections

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Tetsuo Ohta, Yoichi Tonomura, Kyoko Nozaki, Hidemasa Takaya, and Kazushi Mashima*: An Anionic Dinuclear BINAP−Ruthenium(II) Complex: Crystal Structure of $[NH_2Et_2][\{RuCl((R)$-*p*-MeO-BINAP$)\}_2(\mu$-Cl$)_3]$ and Its Use in Asymmetric Hydrogenation.

Pages 1522−1523. An improper computer operation caused wrong calculations of bond distances and angles of complex (*R*)-**2**. Selected bond lengths (Å) and angles (deg) in the caption to Figure 1 are corrected as follows: Ru−Cl(1) = 2.437(3), Ru−Cl(2) = 2.423(3), Ru−Cl(3) = 2.493(3), Ru−Cl(3*) = 2.518(3), Ru−P(1) = 2.271(3), Ru−P(2) = 2.269(3); Cl(1)−Ru−Cl(2) = 163.9(1), Cl(1)−Ru−Cl(3) = 79.8(1), Cl(1)−Ru−Cl(3*) = 79.3(1), Cl(2)−Ru−Cl(3) = 87.6(1), Cl(2)−Ru−Cl(3*) = 88.7(1), Cl(3)−Ru−Cl(3*) = 80.5(1), Cl(1)−Ru−P(1) = 96.0(1), Cl(1)−Ru−P(2) = 103.5(1), Cl(2)−Ru−P(1) = 94.8(1), Cl(2)−Ru−P(2) = 88.0(1), Cl(3)−Ru−P(1) = 94.0(1), Cl(3)−Ru−P(2) = 173.0(1), Cl(3*)−Ru−P(1) = 173.3(1), Cl(3*)−Ru−P(2) = 94.0(1), P(1)−Ru−P(2) = 91.8(1). The dihedral angle between the two naphthyl planes of **2** was 70.7°.

Supporting Information Available: Corrected tables for the crystallographic study of compound (*R*)-**2** (21 pages). Ordering information is given on any current masthead page.

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