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SUPPORTING INFORMATION

For

An Efficient Synthesis of Iminoquinones by Chemoseletive Domino *ortho*hydroxylation/oxidation/imidation sequence of 2- aminoaryl ketone

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Table of Contents

•	Synthesis of 2-amino benzophenone	2
•	Synthesis of secondary alcohol	2
•	References	3
•	¹ H and ¹³ C spectra for all compounds	4

Synthesis of 2-amino benzophenone derivatives¹ (1i-k)

Magnesium turnings were placed in a oven dried two neck 50 mL round bottom flask and the flask was further dried by hot air gun with applying vacuum. Then the setup was allowed to room temperature, refilled with nitrogen. Under nitrogen flow, a catalytic amount of iodine was added, followed by freshly distilled THF. Then followed by corresponding aryl halide was added by drop wise. After the disappearance of the iodine color, the reaction was allowed to stir at room temperature for 2 hours. Once all the magnesium turnings were dissolved. The Grignard reagent was added to solution of 2-amino benzonitrile dissolved in THF at 0 °C over 40 min. Then the reaction was allowed to warm to amibient temperature and allowed to stir at this temperature for 6h. The reaction was quenched by slow addition of 10 % HCl and made basic nature by the addition of NaOH at 0 °C. The organic layer was extracted with ethyl acetate. The combined organic layers were dried over Na₂SO₄ and the solvent was evaporated under vacuum. The residue was purified by column chromatography with ethyl acetate/hexanes solvents to provide pure products (**1i-1k**).



Synthesis of secondary alcohols¹ (5a-5c)

Grignard reagent was added to solution of 2-aminobenzaldehyde dissolved in THF at 0 °C over 40 min. Then the reaction was allowed to warm to amibient temperature and allowed to stir at this temperature for 6h. The reaction was quenched by slow addition of saturated NH_4Cl at 0 °C. The organic layer was extracted with ethyl acetate. The combined organic layers were dried over Na_2SO_4 and the solvent was evaporated under vacuum. The residue was purified by column chromatography with ethyl acetate/hexanes solvents to provide pure products (**5a-5c**).



Reference:

1) Z. Huang, Y. Yang, Q. Xiao, Y. Zhang, J. Wang, *Eur. J. Org. Chem.* **2012**, 6586-6593.



400 MHz ^{1H}NMR spectrum of **2a** in DMSO-d₆



100 MHz 13 C NMR spectrum of **2a** in DMSO-d₆





100 MHz 13 C NMR spectrum of **2b** in DMSO-d₆



400 MHz ¹H NMR spectrum of 2c in DMSO-d₆



100 MHz 13 C NMR spectrum of **2c** in DMSO-d₆



400 MHz ¹HNMR spectrum of **2d** in DMSO-d₆



100 MHz ^{13}C NMR spectrum of 2d in DMSO-d_6



400 MHz ¹H NMR spectrum of 2e in DMSO-d₆



100 MHz ¹³C NMR spectrum of **2e** in CDCl₃



400 MHz ¹H NMR spectrum of **2f** in CDCl₃



100 MHz ¹³C NMR spectrum of **2f** in CDCl₃



400 MHz ¹H NMR spectrum of **2g** in DMSO-d₆



100 MHz ¹³C NMR spectrum of **2g** in CDCl₃: DMSO-d₆



400 MHz ¹H NMR spectrum of 2h in DMSO-d₆



100 MHz 13 C NMR spectrum of **2h** in DMSO-d₆



400 MHz ¹³H NMR spectrum of **2i** in CDCl₃



100 MHz 13 C NMR spectrum of **2i** in CDCl₃



400 MHz ¹H NMR spectrum of 2j in CDCl₃



100 MHz ¹³C NMR spectrum of **2j** in CDCl₃



400 MHz ¹H NMR spectrum of **2k** in DMSO-d₆



100 MHz ^{13}C NMR spectrum of 2k in DMSO-d_6



400 MHz ¹³C NMR spectrum of **2l** in CDCl₃



100 MHz ¹³C NMR spectrum of **2l** in DMSO-d₆



400 MHz ¹H NMR spectrum of 2m in DMSO-d₆



100 MHz 13 C NMR spectrum of **2m** in DMSO-d₆



400 MHz ¹H NMR spectrum of **4a** in DMSO-d₆



100 MHz 13 C NMR spectrum of **4a** in DMSO-d₆



400 MHz ¹H NMR spectrum of **4b** in CDCl₃



100 MHz 13 C NMR spectrum of **4b** in CDCl₃



400 MHz ¹H NMR spectrum of 4c in DMSO-d₆



100 MHz ¹³CNMR spectrum of **4c** in DMSO-d₆



100 MHz ¹H NMR spectrum of **4d** in CDCl₃



100 MHz ¹³C NMR spectrum of **4d** in CDCl₃



400 MHz ¹H NMR spectrum of 6a in DMSO-d₆



100 MHz ¹³C NMR spectrum of **6a** in DMSO-d₆





100 MHz 13 C NMR spectrum of **6b** in CDCl₃



400 MHz ¹H NMR spectrum of 8a in CDCl₃



100 MHz ¹³C NMR spectrum of 8a in CDCl₃



400 MHz ¹H NMR spectrum of **8b** in CDCl₃



100 MHz ¹³C NMR spectrum of **8b** in CDCl₃



400 MHz ¹H NMR spectrum of 9 in DMSO-d₆



100 MHz 13 C NMR spectrum of **9** in DMSO-d₆