

Synthesis of *N*-aryl-1-aminoindoles via intermolecular redox amination

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Supporting information

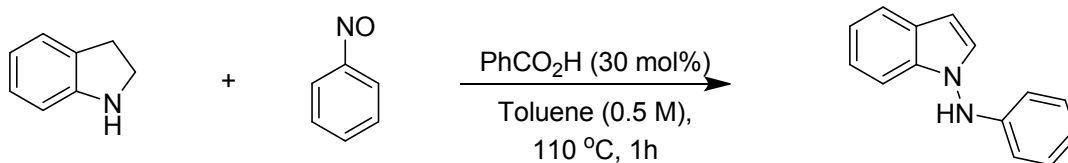
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General Information:

All reactions were run in flame-dried glassware under argon atmosphere using standard Schlenk techniques or in an inert atmosphere glove box and 1-dram sealed pressure vials. Commercially available reagents and solvents were used without additional purification unless otherwise stated. Indolines¹ and nitrosobenzenes² were prepared according to literature procedures. Compound purification was effected by flash chromatography using 230 × 400 mesh, 60 Å porosity silica. ¹H NMR and ¹³C NMR spectra were obtained on a Bruker Avance 400 or a Bruker Avance 500 DRX spectrometer and referenced to residual protio solvent signals (most spectra were taken using a QNP Cryoprobe). Structural assignments are based on ¹H, ¹³C, DEPT-135, COSY, HSQC and IR spectroscopies. Mass Spectrometry was run using EI or ESI techniques.

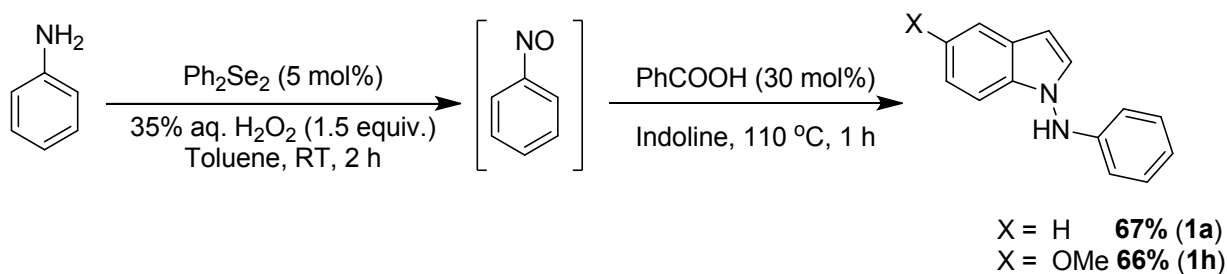
Procedure for the intermolecular redox amination reactions:



A: To a mixture of indoline (59.58 mg, 0.5 mmol) and 30 mol% benzoic acid (18.32 mg, 0.15 mmol) in dry toluene (0.2 mL), nitrosobenzene (53.56 mg, 0.5 mmol) in 0.8 mL dry toluene was added dropwise over 1 hour at 110 °C via a septum-sealed pressure vial. After heating in an aluminum block for 1 hour, completion of the reaction was indicated via TLC analysis. At this point, the reaction mixture was passed through a pad of silica gel and the product was isolated by flash column chromatography using silica gel as the stationary phase and 95:5 hexane:ethyl acetate as the eluent. 83 mg (80%) of the product *N*-phenyl-1*H*-indol-1-amine (**1a**) was obtained as an oil.

B: To 5-nitroindoline (164.16 mg, 1.0 mmol) and 30 mol% of benzoic acid (18.32 mg, 0.15 mmol) nitrosobenzene (53.56 mg, 0.5 mmol) in 1.0 mL of dry toluene was added to a pressure vial equipped with a septum. The mixture was heated in an aluminum block for 1 hour, at which point completion of reaction was indicated via TLC. The reaction mixture was then passed through a short pad of silica gel and the product was isolated by flash column chromatography using silica gel as the stationary phase and 95:5 hexane:ethyl acetate as the eluent. 63 mg (50%) of the product 5-nitro-*N*-phenyl-1*H*-indol-1-amine (**1k**) was obtained as a yellow oil.

Procedure for the one-pot synthesis of *N*-amino indolines:



To aniline (93.13 mg, 1.0 mmol), Ph₂Se₂ (15.6 mg, 0.05 mmol) was added followed by 35% aq. hydrogen peroxide (146 μL, 1.5 mmol) in 1.0 mL of dry toluene. The solution was stirred in a

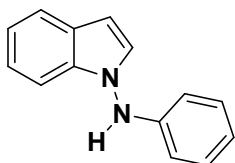
pressure vial at room temperature for 2 hours. After 2 hours, indoline (59.58 mg, 0.5 mmol) and benzoic acid (18.32 mg, 0.15 mmol) were added and the resulting mixture was heated in an aluminum block for 1 hour or until completion of the reaction was indicated by TLC. The reaction mixture was passed through a short pad of silica gel and the product was isolated by flash column chromatography using silica gel as the stationary phase and 95:5 hexane:ethyl acetate as the eluent. 70 mg (67%) of the product *N*-phenyl-1*H*-indol-1-amine (**1a**) was obtained as a slightly red oil.

References:

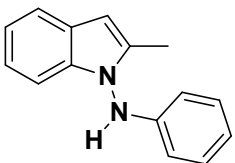
1. Vu, A. T.; Cohn, S. T.; Zhang, P.; Kim, C. Y.; Mahaney, P. E.; Bray, J. A.; Johnston, G. H.; Koury, E. J.; Cosmi, S. A.; Deecher, D. C.; Smith, V. A.; Harrison, J. E.; Leventhal, L.; Whiteside, G. T.; Kennedy, J. D.; Trybulski, E. J. *J. Med. Chem.* **2010**, *53*, 2051–2062.
2. Zhao, D.; Johansson, M.; Bäckvall, J. E. *Eur. J. Org. Chem.* **2007**, 4431–4436.

Spectroscopic Data of compounds for 1a-1l & 2a-2e:

***N*-Phenyl-1*H*-indol-1-amine (1a):** Prepared following the procedure A and purified by column chromatography using EtOAc/hexane and isolated as a reddish liquid; IR (neat): $\nu_{\max}$ 3327, 3053, 1601, 1495, 1473, 1458, 1327, 1090, 1215, 742, 692 cm^{-1} ; ^1H NMR (500 MHz, CDCl_3) δ 7.68 (d, J = 7.0 Hz, 1H), 7.29 (d, J = 8.2 Hz, 1H), 7.22–7.16 (m, 5H), 6.92 (t, J = 7.5 Hz, 1H), 6.60 (s, 1H), 6.56 (dd, J = 3.3, 0.7 Hz, 1H), 6.52 (dd, J = 8.6, 0.9 Hz, 2H). ^{13}C NMR (126 MHz, CDCl_3) δ 147.3, 135.8, 129.4, 128.6, 126.6, 122.4, 121.2, 121.1, 120.3, 112.6, 109.4, 100.7. HRMS m/z 207.0906 ($M - \text{H}^+$), calcd for $\text{C}_{14}\text{H}_{11}\text{N}_2$ 207.0922.

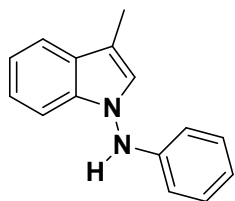


2-Methyl-*N*-phenyl-1*H*-indol-1-amine (1b): Prepared following the procedure A and purified by column chromatography using EtOAc/hexane and isolated as a white solid; IR (neat): $\nu_{\max}$ 3333, 2918, 2361, 1601, 1497, 1456, 1325, 1242, 746, 692 cm^{-1} ; ^1H NMR (500 MHz, CDCl_3) δ 7.56 – 7.54 (m, 1H), 7.18 (dt, J = 11.8, 5.9 Hz, 3H), 7.11 – 7.09 (m, 2H), 6.88 (t, J = 7.5 Hz, 1H), 6.47 (dd, J = 9.0, 1.0 Hz, 2H), 6.44 (s, 1H), 6.30 (t, J = 1.0 Hz, 1H), 2.34 (s, 3H). ^{13}C NMR (126 MHz,



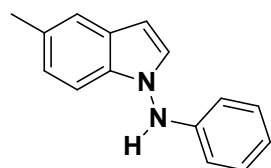
CDCl₃) δ 146.8, 137.6, 135.5, 129.6, 126.0, 121.2, 120.8, 120.3, 119.9, 112.5, 108.8, 98.7, 11.4. HRMS m/z 221.1053 ($M - H^+$), calcd for C₁₅H₁₃N₂ 221.1079.

3-Methyl-*N*-phenyl-1*H*-indol-1-amine (1c): Prepared following the procedure **A** and purified



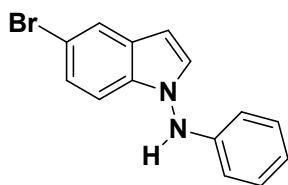
by column chromatography using EtOAc/hexane and isolated as a white solid; IR (neat): $\nu_{\max}$ 3327, 3053, 2916, 1601, 1497, 1458, 1306, 1227, 1111, 1009, 743, 692 cm⁻¹; ¹H NMR (500 MHz, CDCl₃) δ 7.62 (d, J = 7.9 Hz, 1H), 7.26 (d, J = 8.2 Hz, 1H), 7.18 (dt, J = 15.9, 7.2 Hz, 4H), 6.96 (d, J = 1.0 Hz, 1H), 6.90 (t, J = 7.4 Hz, 1H), 6.52 (dd, J = 8.6, 1.0 Hz, 3H). 2.36 (s, 3H). ¹³C NMR (126 MHz, CDCl₃) δ 147.4, 136.1, 129.1, 126.9, 126.0, 122.3, 121.0, 119.7, 119.0, 112.6, 110.1, 109.2, 9.6. HRMS m/z 221.1052 ($M - H^+$), calcd for C₁₅H₁₃N₂ 221.1079.

5-Methyl-*N*-phenyl-1*H*-indol-1-amine (1d): Prepared following the procedure **A** and purified



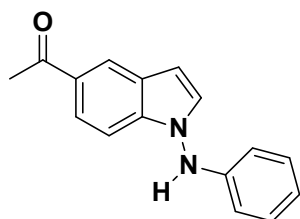
by column chromatography using EtOAc/hexane and isolated as a white solid; IR (neat): $\nu_{\max}$ 3323, 2901, 1601, 1495, 1472, 1331, 1246, 1209, 1150, 1090, 1045, 796, 752, 692 cm⁻¹; ¹H NMR (500 MHz, CDCl₃) δ 7.46 (s, 1H), 7.20 (dd, J = 8.5, 7.4 Hz, 2H), 7.16 (dd, J = 5.8, 2.5 Hz, 2H), 7.02 (d, J = 8.3 Hz, 1H), 6.91 (t, J = 7.8 Hz, 1H), 6.57 (s, 1H), 6.51 (d, J = 9.4 Hz, 2H), 6.47 (d, J = 4.0 Hz, 1H), 2.47 (s, 3H). ¹³C NMR (126 MHz, CDCl₃) δ 147.4, 134.1, 129.6, 129.3, 128.7, 126.9, 124.0, 121.1, 120.8, 112.6, 109.1, 100.2, 21.4. HRMS m/z 221.1052 ($M - H^+$), calcd for C₁₅H₁₃N₂ 221.1079.

5-Bromo-*N*-phenyl-1*H*-indol-1-amine (1e): Prepared following the procedure **A** and purified



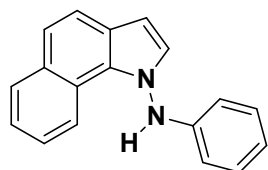
by column chromatography using EtOAc/hexane and isolated as a white solid; IR (neat): $\nu_{\max}$ 3329, 1601, 1495, 1458, 1325, 1207, 1092, 1043, 895, 798, 750, 692 cm⁻¹; ¹H NMR (500 MHz, CDCl₃) δ 7.68 (d, J = 1.6 Hz, 1H), 7.16 (dd, J = 8.7, 1.8 Hz, 1H), 7.12 – 7.07 (m, 3H), 7.04 (d, J = 8.6 Hz, 1H), 6.82 (t, J = 7.4 Hz, 1H), 6.46 (s, 1H), 6.38 – 6.37 (m, 3H). ¹³C NMR (126 MHz, CDCl₃) δ 146.9, 134.5, 129.8, 129.4, 128.1, 125.3, 123.5, 121.4, 113.6, 112.6, 110.9, 100.4. HRMS m/z 285.0017 ($M - H^+$), calcd for C₁₄H₁₀N₂Br 285.0027.

1-(1-(Phenylamino)-1*H*-indol-5-yl)ethanone (1f): Prepared following the procedure **A** and purified by column chromatography using EtOAc/hexane and isolated



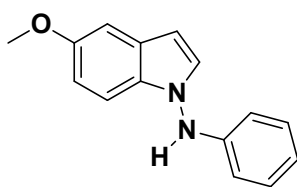
as a yellow liquid; IR (neat): $\nu_{\max}$ 3325, 3128, 2959, 1603, 1560, 1495, 1302, 1229, 1063, 970, 822, 753 cm^{-1} ; ^1H NMR (500 MHz, CDCl_3) δ 8.20 (d, J = 1.5 Hz, 1H), 7.72 (d, J = 8.6 Hz, 1H), 7.20 (d, J = 8.6 Hz, 1H), 7.15 – 7.14 (m, 1H), 7.10 (dd, J = 8.5, 7.5 Hz, 2H), 6.82 (t, J = 7.4 Hz, 2H), 6.54 (d, J = 3.3 Hz, 1H), 6.41 (d, J = 7.7 Hz, 2H), 2.54 (s, 3H). ^{13}C NMR (126 MHz, CDCl_3) δ 198.3, 146.9, 138.6, 130.4, 130.2, 129.4, 126.0, 123.2, 122.7, 121.4, 112.7, 109.4, 102.6, 26.6. HRMS m/z 251.1182 ($\text{M} + \text{H}^+$), calcd for $\text{C}_{16}\text{H}_{15}\text{N}_2\text{O}$ 251.1184.

***N*-Phenyl-1*H*-benzo[*g*]indol-1-amine (1g):** Prepared following the procedure **A** and purified by column chromatography using EtOAc/hexane and isolated as a white



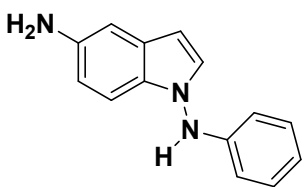
solid; IR (neat): $\nu_{\max}$ 3335, 1603, 1495, 1485, 1402, 1354, 1236, 806, 750, 687 cm^{-1} ; ^1H NMR (500 MHz, CDCl_3) δ 8.73 (d, J = 8.3 Hz, 1H), 7.98 (d, J = 7.6 Hz, 1H), 7.76 (d, J = 8.6 Hz, 1H), 7.62 (d, J = 8.6 Hz, 1H), 7.46 – 7.42 (m, 2H), 7.22 (t, J = 7.5 Hz, 2H), 7.07 (d, J = 3.2 Hz, 1H), 6.94 (t, J = 7.4 Hz, 1H), 6.71 (d, J = 3.2 Hz, 1H), 6.64 (s, 1H), 6.52 (d, J = 7.9 Hz, 2H). ^{13}C NMR (126 MHz, CDCl_3) δ 147.1, 131.3, 129.5, 129.0, 128.5, 127.2, 125.5, 124.0, 122.7, 122.1, 121.6, 121.3, 121.2, 120.8, 112.9, 102.8. HRMS m/z 259.1208 ($\text{M} + \text{H}^+$), calcd for $\text{C}_{18}\text{H}_{15}\text{N}_2$ 259.1235.

5-Methoxy-*N*-phenyl-1*H*-indol-1-amine (1h): Prepared following the procedure **A** and purified



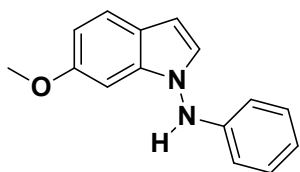
by column chromatography using EtOAc/hexane and isolated as a white solid; IR (neat): $\nu_{\max}$ 3319, 1603, 1497, 1473, 1252, 1232, 1148, 1028, 752, 692 cm^{-1} ; ^1H NMR (500 MHz, CDCl_3) δ 7.18 (ddd, J = 24.5, 18.9, 10.2 Hz, 5H), 6.91 (t, J = 7.4 Hz, 1H), 6.85 (dd, J = 8.8, 2.3 Hz, 1H), 6.59 (s, 1H), 6.51 (d, J = 8.5 Hz, 2H), 6.47 (d, J = 3.2 Hz, 1H), 3.87 (s, 3H). ^{13}C NMR (126 MHz, CDCl_3) δ 154.6, 147.4, 130.9, 129.3, 129.2, 127.0, 121.1, 112.6, 110.2, 102.8, 100.3, 55.8. HRMS m/z 237.1004 ($\text{M} - \text{H}^+$), calcd for $\text{C}_{15}\text{H}_{13}\text{N}_2\text{O}$ 237.1028.

***N*¹-Phenyl-1*H*-indole-1,5-diamine (1i):** Prepared following the procedure **A** and purified by



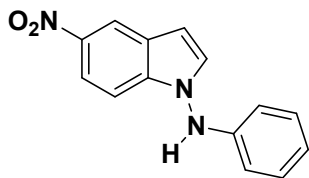
column chromatography using EtOAc/hexane and isolated as a yellow solid; IR (neat): $\nu_{\max}$, 3400, 3327, 3211, 2959, 2359, 1603, 1495, 1474, 1242, 1215, 1153, 800, 750, 619 cm^{-1} ; ^1H NMR (500 MHz, CDCl_3) δ 7.19 (dd, J = 8.5, 7.4 Hz, 2H), 7.12 (d, J = 3.3 Hz, 1H), 7.05 (d, J = 8.5 Hz, 1H), 6.94 (d, J = 2.1 Hz, 1H), 6.89 (t, J = 6.4 Hz, 1H), 6.63 (dd, J = 8.5, 2.1 Hz, 1H), 6.59 (s, 1H), 6.51 (dd, J = 8.6, 0.9 Hz, 2H), 6.35 (dd, J = 3.3, 0.8 Hz, 1H), 3.47 (s, 2H). ^{13}C NMR (126 MHz, CDCl_3) δ 147.5, 140.1, 130.5, 129.3, 129.1, 127.6, 121.0, 113.1, 112.6, 110.0, 105.9, 99.5. HRMS m/z 222.1056 ($M - \text{H}^+$), calcd for $\text{C}_{14}\text{H}_{12}\text{N}_3$ 222.1031.

6-Methoxy-*N*-phenyl-1*H*-indol-1-amine (1j): Prepared following the procedure **A** and purified



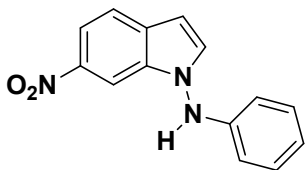
by column chromatography using EtOAc/hexane and isolated as a white solid; IR (neat): $\nu_{\max}$ 3323, 1601, 1493, 1286, 1231, 1084, 1043, 933, 752, 692, 635 cm^{-1} ; ^1H NMR (500 MHz, CDCl_3) δ 7.53 (d, J = 8.6 Hz, 1H), 7.22 (dd, J = 8.5, 7.5 Hz, 2H), 7.07 (d, J = 3.3 Hz, 1H), 6.92 (t, J = 7.4 Hz, 1H), 6.84 (dd, J = 8.6, 2.3 Hz, 1H), 6.79 (d, J = 2.2 Hz, 1H), 6.51 (dd, J = 15.8, 6.0 Hz, 4H), 3.79 (s, 3H). ^{13}C NMR (126 MHz, CDCl_3) δ 156.8, 147.2, 136.9, 129.4, 127.2, 121.7, 121.1, 120.4, 112.7, 110.5, 101.0, 92.6, 55.5. HRMS m/z 237.1006 ($M - \text{H}^+$), calcd for $\text{C}_{15}\text{H}_{13}\text{N}_2\text{O}$ 237.1028.

5-Nitro-*N*-phenyl-1*H*-indol-1-amine (1k): Prepared following the procedure **B** and purified by



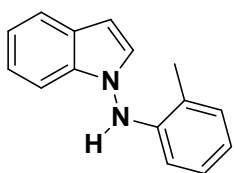
column chromatography using EtOAc/hexane and isolated as a yellow liquid; IR (neat): $\nu_{\max}$ 3323, 1603, 1578, 1514, 1497, 1205, 1327, 1067, 897, 743, 692, 602 cm^{-1} ; ^1H NMR (500 MHz, CDCl_3) δ 8.63 (s, 1H), 8.12 (d, J = 9.0 Hz, 1H), 7.39 (dd, J = 9.1, 6.2 Hz, 2H), 7.26 (dd, J = 8.5, 7.5 Hz, 2H), 6.99 (t, J = 7.4 Hz, 1H), 6.82 (s, 1H), 6.77 (d, J = 3.4 Hz, 1H), 6.55 (dd, J = 8.6, 0.9 Hz, 2H). ^{13}C NMR (126 MHz, CDCl_3) δ 146.4, 142.4, 139.1, 131.7, 129.6, 125.6, 122.0, 118.3, 118.2, 112.8, 109.6, 103.5. HRMS m/z 252.0757 ($M - \text{H}^+$), calcd for $\text{C}_{14}\text{H}_{10}\text{N}_3\text{O}_2$ 252.0773.

6-Nitro-*N*-phenyl-1*H*-indol-1-amine (1l): Prepared following the procedure **B** and purified by



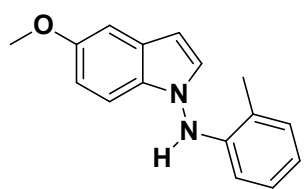
column chromatography using EtOAc/hexane and isolated as a yellow solid; IR (neat): $\nu_{\max}$ 3319, 1603, 1514, 1495, 1400, 1333, 1211, 1119, 1061, 752, 692, 625 cm^{-1} ; ^1H NMR (500 MHz, CDCl_3) δ 8.28 (d, $J = 2.0$ Hz, 1H), 8.02 (dd, $J = 8.7, 2.1$ Hz, 1H), 7.68 (d, $J = 8.8$ Hz, 1H), 7.49 (d, $J = 3.3$ Hz, 1H), 7.22 (dd, $J = 8.6, 7.5$ Hz, 2H), 6.95 (t, $J = 6.9$ Hz, 1H), 6.87 (s, 1H), 6.66 (dd, $J = 3.3, 0.9$ Hz, 1H), 6.52 (dd, $J = 8.6, 1.0$ Hz, 2H). ^{13}C NMR (126 MHz, CDCl_3) δ 146.6, 143.7, 134.8, 134.3, 131.0, 129.5, 121.9, 121.1, 115.8, 112.7, 106.4, 101.9. HRMS m/z 252.0773 ($\text{M} - \text{H}^+$), calcd for $\text{C}_{14}\text{H}_{10}\text{N}_3\text{O}_2$ 252.0773.

***N*-(*o*-Tolyl)-1*H*-indol-1-amine (2a):** Prepared following the procedure **A** and purified by



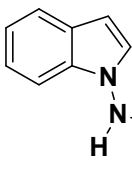
column chromatography using EtOAc/hexane and isolated as a white solid; IR (neat): $\nu_{\max}$ 3331, 3051, 1600, 1514, 1495, 1402, 1346, 1327, 1238, 1067, 808, 746, 688 cm^{-1} ; ^1H NMR (500 MHz, CDCl_3) δ 7.57 (dd, $J = 6.8, 1.4$ Hz, 1H), 7.14 (dd, $J = 7.8, 1.4$ Hz, 1H), 7.11 – 7.05 (m, 4H), 6.87 (t, $J = 7.0$ Hz, 1H), 6.74 (t, $J = 7.4$ Hz, 1H), 6.47 (dd, $J = 3.3, 0.8$ Hz, 1H), 6.43 (s, 1H), 5.94 (d, $J = 8.1$ Hz, 1H), 2.26 (s, 3H). ^{13}C NMR (126 MHz, CDCl_3) δ 145.0, 135.8, 130.5, 128.8, 127.3, 126.6, 122.4, 121.4, 121.1, 120.8, 120.3, 111.9, 109.3, 100.8, 17.1. HRMS m/z 221.1060 ($\text{M} - \text{H}^+$), calcd for $\text{C}_{15}\text{H}_{13}\text{N}_2$ 221.1079.

5-Methoxy-*N*-(*o*-tolyl)-1*H*-indol-1-amine (2b): Prepared following the procedure **A** and



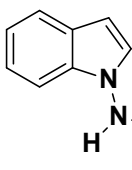
purified by column chromatography using EtOAc/hexane and isolated as a white solid; IR (neat): $\nu_{\max}$ 3342, 2935, 1622, 1589, 1472, 1283, 1236, 1148, 1030, 800, 752, 625 cm^{-1} ; ^1H NMR (500 MHz, CDCl_3) δ 7.18 (d, $J = 3.3$ Hz, 1H), 7.16 (d, $J = 7.4$ Hz, 1H), 7.12 (d, $J = 9.3$ Hz, 2H), 6.98 (t, $J = 7.7$ Hz, 1H), 6.85 (dd, $J = 14.0, 5.1$ Hz, 2H), 6.52 (s, 1H), 6.49 (d, $J = 3.2$ Hz, 1H), 6.04 (d, $J = 8.0$ Hz, 1H), 3.87 (s, 3H), 2.36 (s, 3H). ^{13}C NMR (126 MHz, CDCl_3) δ 154.6, 145.1, 130.9, 130.4, 129.3, 127.3, 126.9, 121.3, 120.7, 112.6, 111.8, 110.1, 102.7, 100.2, 55.8, 17.1. HRMS m/z 251.1154 ($\text{M} - \text{H}^+$), calcd for $\text{C}_{16}\text{H}_{15}\text{N}_2\text{O}$ 251.1184.

***N*¹-(1*H*-Indol-1-yl)-*N*⁴,*N*⁴-dimethylbenzene-1,4-diamine (2c):** Prepared following the procedure **A** and purified by column chromatography using



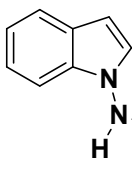
EtOAc/hexane and isolated as a yellow liquid; IR (neat): ν_{max} 3312, 2795, 1516, 1453, 1327, 1215, 1124, 1086, 945, 816, 743 cm^{-1} ; ^1H NMR (500 MHz, CDCl_3) δ 7.64 (d, J = 7.6 Hz, 1H), 7.33 (d, J = 7.3 Hz, 1H), 7.22 (d, J = 3.3 Hz, 1H), 7.19 – 7.16 (m, 1H), 7.13 (td, J = 7.5, 1.1 Hz, 1H), 6.67 – 6.64 (m, 2H), 6.53 – 6.50 (m, 3H), 6.42 (s, 1H), 2.84 (s, 6H). ^{13}C NMR (126 MHz, CDCl_3) δ 146.5, 138.6, 136.0, 128.7, 126.5, 122.2, 120.9, 120.1, 114.5, 109.5, 100.3, 41.5. HRMS m/z 250.1320 ($M - \text{H}^+$), calcd for $\text{C}_{16}\text{H}_{16}\text{N}_3$ 250.1344.

4-((1*H*-Indol-1-yl)amino)benzonitrile (2d): Prepared following the procedure **A** and purified by column chromatography using EtOAc/hexane and isolated as a



white solid; IR (neat): ν_{max} 3302, 2222, 1607, 1510, 1456, 1258, 1217, 1173, 829, 762, 744, 546 cm^{-1} ; ^1H NMR (500 MHz, CDCl_3) δ 7.67 (dd, J = 8.3, 1.4 Hz, 1H), 7.45 (d, J = 8.8 Hz, 2H), 7.22 – 7.17 (m, 3H), 7.14 (d, J = 3.4 Hz, 1H), 6.98 (s, 1H), 6.59 (d, J = 3.4 Hz, 1H), 6.50 (d, J = 8.8 Hz, 2H). ^{13}C NMR (126 MHz, CDCl_3) δ 150.7, 135.3, 133.9, 128.0, 126.6, 122.9, 121.4, 120.9, 119.3, 112.3, 109.0, 103.5, 101.9. HRMS m/z 232.0877 ($M - \text{H}^+$), calcd for $\text{C}_{15}\text{H}_{10}\text{N}_3$ 232.0875.

***N*-(4-Nitrophenyl)-1*H*-indol-1-amine (2e):** Prepared following the procedure **A** and purified by column chromatography using EtOAc/hexane and isolated as a



yellow solid; IR (neat): ν_{max} 3329, 2359, 2341, 1597, 1501, 1329, 1261, 1217, 1113, 839, 762, 743 cm^{-1} ; ^1H NMR (500 MHz, CDCl_3) δ 8.07 (d, J = 9.3 Hz, 2H), 7.70 – 7.68 (m, 1H), 7.24 – 7.18 (m, 3H), 7.15 (d, J = 3.4 Hz, 1H), 7.13 (s, 1H), 6.61 (d, J = 3.3 Hz, 1H), 6.48 (d, J = 9.1 Hz, 2H). ^{13}C NMR (126 MHz, CDCl_3) δ 152.5, 141.2, 135.3, 128.0, 126.6, 126.1, 123.0, 121.4, 121.0, 111.4, 109.0, 102.1. HRMS m/z 252.0754 ($M - \text{H}^+$), calcd for $\text{C}_{14}\text{H}_{10}\text{N}_3\text{O}_2$ 252.0773.

