Electronic Supplementary Information for

Donor-acceptor interactions between cyclic trinuclear pyridinate gold(I)-complexes and electron-poor guests: Nature, energetics and templating of guest-binding on graphite

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1 Materials and methods

Chemicals and solvents were purchased from Sigma-Aldrich, Acros-Organics, Alfa-Aesar, TCI, Merck, or ChemPur and used directly unless otherwise noted. Reactions with dry solvents were carried out under an argon atmosphere in oven-dried glassware. Anhydrous solvents (THF, CH₂Cl₂) were withdrawn from an MB-SPS-800 unit from M-Braun and stored over molecular sieves (4 Å). Solvents for column chromatography were purified by distillation. 2-(Dimethylamino)-ethanol was distilled under argon and stored over molecular sieves (4 Å).

Analytical Thin Layer Chromatography (TLC) was carried out using TLC Silica gel 60 F₂₅₄ aluminum sheets from Merck KGaA or POLYGRAM® Alox N/UV₂₅₄ polyester sheets from Macherey-Nagel GmbH & Co. KG. Detection was carried out using short wave UV light (254 nm and 366 nm). Column chromatography was accomplished using silica gel S (0.040-0.062 mm) or aluminum oxide purchased from Merck KGaA and deactivated with 15% w/w H₂O. ¹H and ¹³C NMR spectra were recorded at room temperature on the following spectrometers: Bruker Avance DPX 500 (¹H = 500 MHz, ¹³C = 126 MHz), Bruker Avance DPX 400 (¹H = 400 MHz, ¹³C = 101 MHz), Bruker Avance DPX 300 (¹H = 300 MHz, ¹³C = 76 MHz) (all University of Bonn), Bruker Avance III HD 500 (¹H = 500 MHz, ¹³C = 126 MHz) and Bruker Avance II 400 (¹H = 400 MHz, ¹³C = 101 MHz) (University of Freiburg, 300 K). Unless stated otherwise all spectra were recorded in deuterchloroform or benzene-d₆, which were purchased from Sigma-Aldrich, Acros-Organics or Deutero, and all chemical shifts are given in δ units relative to tetramethyldisilane (singlet: δ = 0.00). Calibration of ¹H spectra was achieved using the residual solvent signal of chloroform at δ_H = 7.26 ppm or benzene at δ_H = 7.16 ppm. ¹³C spectra were calibrated to the central signal line of the triplet of chloroform at δ_C = 77.16 ppm or the singlet of benzene at δ_C = 128.06 ppm.[¹] Analyses followed first order, and the following abbreviations were used throughout: s = singlet, d = doublet, t = triplet, dd = doublet of doublets, m = multiplet. Coupling constants (J) are given in Hertz [Hz]. Mass spectra (MS) were recorded on a Bruker Daltonics ESI microTOF-Q time of flight, a Thermo Finnigan MAT 95 XL sector mass, a Bruker Daltonics autoflex II TOF/TOF spectrometer (University of Bonn), a Thermo Exactive with Orbitrap-analysers and electrospray ionization (ESI) or a MALDI-TOF autoflex™ by Bruker (University of Freiburg). Melting points were determined using a BÜCHI 510 melting point (University of Bonn). UV/Vis spectra were recorded on a Perkin-Elmer Lambda 950 UV/VIS spectrometer.

The titration of alky magnesium halide or alkyl magnesium halide-lithium chloride suspensions was performed following a procedure by Peters et al.[²] A stock solution containing N-phenyl-4-phenylazoaniline (10 mg, 37 μmol) in dry and degassed toluene (8.0 mL) and 2-butanol (0.80 mL) was prepared and stored under argon over 3 Å molecular sieves (stable over month). The Grignard suspension was added dropwise under inert conditions to the stock solution (0.2-0.4 mL). The equivalence point was indicated by a color change from orange to red. The procedure was performed twice, carried out before every
use, and the calculated concentration of 2-butanol was used to determine the concentration of the Grignard suspension.

**Pyrazolate CTCs 2** were synthesized following procedures of Omary *et al.* (2a)[3] and Dias *et al.* (2b,c).[4]

**Chloro(triphenylarsine)gold(I)** was synthesized following a procedure of Nieto-Oberhuber *et al.*[5]
2 Synthetic manipulations

2.1 Preparation of Grignard reagents

2.1.1 Preparation of alkyl magnesium halide suspensions

The preparation of alkyl magnesium halide solutions was performed following a modified procedure by Peters et al.\textsuperscript{[2]} A dry three-necked round-bottom flask was charged with magnesium turnings (1.2 eq.), equipped with a reflux condenser and heated \textit{in vacuo} for 7–10 min. After cooling to room temperature, 1/3 of a total amount of dry and degassed THF (1.7 mmol/mL) was added, and the resulting suspension was sonicated for 10 min. The respective alkyl halide (1.0 eq.) and a small crystal of iodine were added. The reaction mixture was heated to 85 °C without stirring to initiate activation of the Grignard formation. As soon as the formation of the Grignard reagent had started, the remaining THF was added, and the reaction mixture was heated under reflux for 1 h. The suspension was allowed to cool to room temperature and stirred overnight. The resulting Grignard suspension was used without purification and titrated as described above.

2.1.2 Preparation of alkyl magnesium halide-lithium chloride suspensions

The preparation of the alkyl magnesium halide-lithium chloride suspensions was performed following a modified procedure by Peters et al.\textsuperscript{[2]} To ground and vacuum-dried (120 °C, 8 h, $10^{-3}$ mbar) LiCl (1.0 eq.) was added the respective alkyl magnesium halide suspension in THF (1.0 eq.) under inert conditions. The reaction mixture was stirred at room temperature for 20 h until all solids had dissolved. It was then titrated and used without purification.

2.2 Synthesis of hexylpyridines 6b and 6c

\begin{center}
\begin{tikzpicture}
\node (a) at (0,0) {\textbf{4}};
\node (b) at (1,0) {\textbf{5a (58\%)}};
\node (c) at (2,0) {\textbf{6b (55\%)}};
\node (d) at (3,0) {\textbf{6c (27\%)}};
\node (e) at (0,-1) {\textbf{5a}};
\path (a) edge [above, bend left = 15] node {THF \(-50 \ ^\circ \text{C} \text{ to } \text{rt}\)} (b);
\path (b) edge [above, bend left = 15] node {THF \text{rt, 16 h}} (c);
\path (b) edge [above, bend left = 15] node {Ni(dppe)Cl\textsubscript{2}} (d);
\end{tikzpicture}
\end{center}

2.2.1 3-Bromo-4-hexylpyridine (5a)

The synthesis of 5a was performed in analogy to a procedure by Knochel and coworkers.\textsuperscript{[6]} To a stirred solution of 3-bromopyridine (4, 0.60 mL, 970 mg, 6.1 mmol, 1.0 eq) in dry THF (31 mL) was added BF\textsubscript{3} • OEt\textsubscript{2} (0.86 mL, 960 mg, 6.8 mmol, 1.1 eq) at 0 °C. The solution was stirred at 0 °C for 20–30 min and cooled to –30 to –50 °C. LiCl • BrMgC\textsubscript{6}H\textsubscript{13} (1.4 M suspension in THF, 8.5 mL, 12 mmol, 2.0 eq.) was added over a period of 10–20 min. The orange suspension was
stirred at ambient temperature for 3 h. Chloranil (3.0 g, 12 mmol, 2.0 eq.) was suspended in dry THF (10 mL) and added to the reaction mixture at −30 °C. The resulting green suspension was allowed to warm to room temperature and stirred overnight. The resulting yellow suspension was quenched with NH₃ (9 mL, 25% w/w in H₂O) and brine (90 mL) and extracted with Et₂O (4 × 60 mL). The combined organic extracts were concentrated in vacuo. The residue was dissolved in CH₂Cl₂ (40 mL), washed with brine (20 mL) and dried over Na₂SO₄. Two consecutive chromatography steps (silica gel, CH₂Cl₂ and silica gel, cyclohexane/EtOAc: 15/1) yielded 5a (870 mg, 3.6 mmol, 58%) as a yellow liquid. Rₓ 0.50 (cyclohexane/EtOAc: 4/1); ¹H NMR (400 MHz, CDCl₃): δ = 8.64 (s, 1H), 8.39 (d, J = 4.9 Hz, 1H), 7.14 (d, J = 4.9 Hz, 1H), 2.71 (t, J = 7.8 Hz, 2H), 1.65–1.58 (m, 2H), 1.44–1.25 (m, 6H), 0.90–0.86 (m, 3H); ¹³C NMR (101 MHz, CDCl₃): δ = 152.0, 151.0, 148.3, 125.2, 123.3, 35.5, 31.7, 29.1, 29.0, 22.7, 14.2; HRMS (ESI⁺): m/z calcd for C₂₃H₄₀BrN 242.0539 [M+H]⁺, found 242.0537.

2.2.2 4-Hexylpyridine (6b) and 3,4-dihexylpyridine (6c)

The synthesis of 6b and 6c was performed following a modified procedure by Kumada and coworkers.⁷ To a stirred solution of 3-bromo-4-hexylpyridine (5a, 460 mg, 1.9 mmol, 1.0 eq.) and Ni(dppp)Cl₂ (52 mg, 130 µmol, 0.07 eq.) in dry THF (23 mL) was added C₆H₁₃MgBr (1.44 M suspension in THF, 1.7 mL, 2.5 mmol, 1.3 eq.) at room temperature. After stirring overnight, another portion of C₆H₁₃MgBr (1.44 M solution in THF, 0.20 mL, 0.29 mmol, 0.15 eq.) was added to the dark suspension. The reaction mixture was stirred for 2.5 h and quenched with HCl (8.0 mL, 10% w/w in H₂O), after TLC indicated complete consumption of the starting material 5a. A saturated aqueous solution of NaHCO₃ (20 mL) was slowly added, and the mixture was extracted with Et₂O (50 mL) and EtOAc (3 × 15 mL). The combined organic extracts were washed with brine and dried over Na₂SO₄. Purification by column chromatography (silica gel, cyclohexane/EtOAc: 10/1) yielded 6b (170 mg, 1.0 mmol, 55%) and 6c (130 mg, 0.52 mmol, 27%) as colorless liquids.

6b: Rₓ 0.23 (cyclohexane/EtOAc: 5/2); ¹H NMR (400 MHz, CDCl₃): δ = 8.47 (dd, J = 4.5, 1.5 Hz, 2H), 7.09 (dd, J = 4.5, 1.5 Hz, 2H), 2.59 (t, J = 7.7 Hz, 2H), 1.65–1.58 (m, 2H), 1.37–1.25 (m, 6H), 0.89–0.86 (m, 3H); ¹³C NMR (76 MHz, CDCl₃): δ = 151.9, 149.8, 124.0, 35.4, 31.7, 30.4, 29.0, 22.7, 14.2; HRMS (ESI⁺): m/z calcd for C₁₁H₁₂N 164.1434 [M+H]⁺, found 164.1431.

6c: Rₓ 0.38 (cyclohexane/EtOAc: 5/2); ¹H NMR (400 MHz, CDCl₃): δ = 8.33 (s, 1H), 8.31 (d, J = 5.0 Hz, 1H), 7.03 (d, J = 5.0 Hz, 1H), 2.63–2.56 (m, 4H), 1.62–1.52 (m, 4H), 1.42–1.26 (m, 12H), 0.91–0.88 (m, 6H); ¹³C NMR (76 MHz, CDCl₃): δ = 150.5, 149.4, 147.3, 136.2, 123.9, 32.0, 31.8, 31.1, 30.3, 30.2, 29.4, 22.7, 22.7, 14.2; HRMS (ESI⁺): m/z calcd for C₁₇H₂₉N 248.2373 [M+H]⁺, found 248.2368.
2.3 **Synthesis of 3,4-dioctadecanlypyridine (6e)**

![Chemical structure](image)

2.3.1 **3-Bromo-4-octadecanlypyridine (5b)**

The synthesis of 5a was performed in analogy to a procedure by Knochel and coworkers.\(^6\) To a stirred solution of 3-bromopyridine (4, 0.76 mL, 1.2 g, 7.8 mmol, 1.0 eq.) in dry THF (39 mL) was added BF\(_3\) • Et\(_2\)O (1.1 mL, 1.2 g, 8.6 mmol, 1.1 eq.) at 0 °C. The solution was stirred at 0 °C for 20–30 min and cooled to −30 to −50 °C. LiCl • ClMgC\(_{18}\)H\(_{37}\) (1.1 M suspension in THF, 14 mL, 16 mmol, 2.0 eq.) was added over a period of 10 to 20 min. The orange suspension was stirred at ambient temperature for 3 h. Chloranil (3.8 g, 16 mmol, 2.0 eq.) was suspended in dry THF (15 mL) and added to the reaction mixture at −30 °C. The resulting green suspension was allowed to warm to room temperature and stirred overnight. The yellow suspension was quenched with NH\(_3\) (13 mL, 25% w/w in H\(_2\)O). Brine (80 mL) was added, and the resulting dark suspension was extracted with Et\(_2\)O (70 mL) and ethyl acetate (3 × 100 mL). The combined organic extracts were concentrated in vacuo, dissolved in CH\(_2\)Cl\(_2\) and filtered through a column of silica gel. The filtrate was concentrated in vacuo, diluted in CH\(_2\)Cl\(_2\) (50 mL), washed with brine (50 mL) and dried over Na\(_2\)SO\(_4\). Two consecutive chromatography steps (silica gel, cyclohexane/EtOAc: 20/1 and silica gel, CH\(_2\)Cl\(_2\)) yielded 5b (820 mg, 2.0 mmol, 26%) as a white solid. R\(_f\) 0.70 (cyclohexane/EtOAc: 5/2); Mp 26 °C; \(^1\)H NMR (500 MHz, CDCl\(_3\)): \(\delta = 8.64\) (s, 1H), 8.39 (d, \(J = 5.0\) Hz, 1H), 7.14 (d, \(J = 5.0\) Hz, 1H), 2.72–2.69 (m, 2H), 1.65–1.58 (m, 2H), 1.39–1.25 (m, 30H), 0.88 (t, \(J = 6.8\) Hz, 3H); \(^{13}\)C NMR (126 MHz, CDCl\(_3\)): \(\delta = 152.0, 151.0, 148.3, 125.2, 123.3, 35.5, 32.1, 29.9, 29.8, 29.8, 29.7, 29.5, 29.4, 29.0, 22.9, 14.3\); HRMS (ESI+): \(m/z\) calcd for C\(_{23}\)H\(_{46}\)BrN 432.2236 [M+Na]\(^+\), found 432.2235.

2.3.2 **3,4-Dioctadecanlypyridine (6e)**

![Chemical structure](image)

The synthesis was performed following a modified procedure by Kumada and coworkers.\(^7\) To a stirred solution of 3-bromo-4-octadecanlypyridine (5b, 620 mg, 1.5 mmol, 1.0 eq.) and Ni(dpdp)Cl\(_2\) (44 mg, 106 \(\mu\)mol, 0.07 eq.) in dry THF (6 mL) was added C\(_{18}\)H\(_{37}\)MgCl (0.92 M in
THF, 2.2 mL, 2.0 mmol, 1.3 eq.) at room temperature. After stirring at room temperature for 15 min, another portion of C_{18}H_{37}MgCl (0.92 m suspension in THF, 0.4 mL, 0.4 mmol, 0.25 eq.) was added to the dark suspension in two equal portions within 30 min. The reaction mixture was stirred for an additional 30 min and quenched with HCl (4 mL, 10% w/w in H_2O), after TLC indicated complete consumption of the starting material 5b. A saturated aqueous solution of NaHCO_3 (8 mL) was slowly added, and the mixture was extracted with Et_2O (25 mL) and EtOAc (3 × 20 mL). The combined organic extracts were washed with brine and dried over Na_2SO_4. Purification by column chromatography (silica gel, cyclohexane/EtOAc: 20/1) yielded 6e (230 mg, 400 μmol, 26%) as a white solid. Rf 0.55 (cyclohexane/EtOAc: 4/1); Mp 31 °C; ^1H NMR (400 MHz, CDCl_3): δ = 8.34 (s, 1H), 8.32 (d, J = 5.1 Hz, 1H), 7.09 (d, J = 5.1 Hz, 1H), 2.63–2.58 (m, 4H), 1.62–1.53 (m, 4H), 1.40–1.25 (m, 60H), 0.88 (t, J = 6.8 Hz, 6H); ^13C NMR (101 MHz, CDCl_3): δ = 150.5, 149.5, 147.3, 136.2, 123.9, 32.1, 32.0, 31.2, 30.4, 30.2, 29.9, 29.8, 29.7, 29.7, 29.6, 29.5, 22.8, 14.3; HRMS (ESI+): m/z calcd for C_{41}H_{77}N 584.6129 [M+H]^+, found 584.6134.

2.4 Synthesis of 4-dodecylpyridine (6d)

The synthesis was performed following a modified procedure by Akiba et al.[^8] tert-Butyldimethylsilyl trifluoromethanesulfonate (12.4 g, 10.8 mL, 47.1 mmol, 1.0 eq.) was added dropwise to a stirred solution of dry pyridine (3.72 g, 3.80 mL, 47.1 mmol, 1.0 eq.) in dry CH_2Cl_2 (45 mL) at room temperature. After stirring at rt for 1 h, the solvent was removed in vacuo, and the residue was suspended in dry THF (45 mL). C_{12}H_{25}MgBr (1.0 m suspension in THF, 56 mL, 56 mmol, 1.2 eq.) was added to the white suspension, and the resulting yellow solution was stirred at room temperature for 5 h. Chloranil (12.7 g, 51.8 mmol, 1.1 eq.) was added, and the resulting suspension was stirred overnight. The reaction mixture was quenched with sat. aqueous NaHCO_3 (130 mL) and H_2O (100 mL) and extracted with Et_2O (200 mL) and EtOAc (2 × 100 mL). The combined organic layers were washed with brine (200 mL), concentrated in vacuo and dissolved in CH_2Cl_2 (80 mL). The organic layer was washed again with a saturated aqueous solution of NaHCO_3 (50 mL), brine (100 mL) and dried over Na_2SO_4. Column chromatography (silica gel, cyclohexane/EtOAc) yielded 4-dodecylpyridine 6d (2.40 g, 9.71 mmol, 21%) as a yellowish oil. Rf 0.28 (cyclohexane/EtOAc: 5/2); ^1H NMR (400 MHz, CDCl_3): δ = 8.48 (dd, J = 4.4 Hz, 1.7 Hz, 2H), 7.10 (dd, J = 4.4 Hz, 1.7 Hz, 2H), 2.60 (t, J = 7.8 Hz, 2H), 1.56–1.69 (m, 2H), 1.19–1.39 (m, 18H), 0.88 (t, J = 6.8 Hz, 3H); ^13C NMR (101 MHz, CDCl_3): δ = 151.9, 149.7, 124.0, 35.4, 30.4, 29.7, 29.7, 29.6, 29.5, 29.4, 29.3, 27.0, 22.8, 14.2; HRMS (ESI+): m/z calcd for C_{17}H_{30}N 248.2373 [M]^+, found 248.2371.
2.5 General procedure for synthesis of 2-bromopyridines 3

![Chemical structure of 3b-e](image)

The synthesis of 3 was performed following a modified procedure by Gros et al.\textsuperscript{[9]} To a stirred solution of 2-(dimethylamino)ethanol (2.0 eq.) in dry n-hexane (300 µmol/mL) was added n-BuLi (1.6 M solution in n-hexane, 4.0 eq.) dropwise at 0 °C. The colorless solution was stirred for 1–2 h at 0 °C, followed by addition of a solution of the respective alkylpyridine 6 (1.0 eq.) in dry n-hexane (700 µmol/mL). The resulting orange reaction mixture was stirred at 0 °C for 1–3 h, cooled to −90 °C, and treated with a solution of CBr\textsubscript{4} (2.5 eq.) in dry THF (1.0 mmol/mL). The resulting yellow suspension was stirred at −78 °C for 2 h and was then allowed to warm to room temperature and stirred for an additional 15 min. The resulting black suspension was cooled to 0 °C and quenched with H\textsubscript{2}O. Brine and Et\textsubscript{2}O were added, the organic layer was separated, and the aqueous layer was extracted with EtOAc. The combined organic extracts were dried over Na\textsubscript{2}SO\textsubscript{4}. Column chromatography yielded 2-bromopyridines 3 (36–51%) as brown solids or oils. Yields were not optimized.

2.5.1 2-Bromo-4-hexylpyridine (3b)

![Chemical structure of 3b](image)

According to the general procedure, 2-(dimethylamino)ethanol (390 µL, 350 mg, 3.9 mmol, 2.0 eq.) in n-hexane (12.6 mL) was reacted with n-BuLi (1.6 M solution in n-hexane, 4.9 mL, 7.8 mmol, 4.0 eq.) and a solution of 4-hexylpyridine (6b, 316 mg, 1.94 mmol, 1.0 eq.) in n-hexane (2.5 mL). The reaction was treated with a solution of CBr\textsubscript{4} (1.61 g, 4.85 mmol, 2.5 eq.) in THF (5.0 mL). The resulting black suspension was quenched with H\textsubscript{2}O (20 mL). Brine (20 mL) and Et\textsubscript{2}O (25 mL) were added, separated and the aqueous layer was extracted with EtOAc (3 × 30 mL). Purification by column chromatography (silica gel, cyclohexane/CH\textsubscript{2}Cl\textsubscript{2}: 6/4) yielded 3b (221 mg, 913 µmol, 47%) as a light brown liquid. R\textsubscript{f} 0.20 (cyclohexane/CH\textsubscript{2}Cl\textsubscript{2}: 1/1); \textsuperscript{1}H NMR (400 MHz, CDCl\textsubscript{3}): \textit{δ} = 8.23 (d, J = 5.1 Hz, 1H), 7.31 (dd, J = 1.3, 0.6 Hz, 1H), 7.06, (dd, J = 5.1, 1.5 Hz, 1H), 2.57 (t, J = 7.8 Hz, 2H), 1.64–1.57 (m, 2H), 1.36–1.26 (m, 6H), 0.90–0.87 (m, 3H); \textsuperscript{13}C NMR (76 MHz, CDCl\textsubscript{3}): \textit{δ} = 155.2, 150.0, 142.5, 128.1, 123.2, 35.1, 31.7, 30.2, 28.9, 22.6, 14.2; MS (EI\textsuperscript{+}): m/z 171.0 [M–C\textsubscript{6}H\textsubscript{10}]\textsuperscript{+}; 162.2 [M–Br]\textsuperscript{+}, 92.1 [M–Br–C\textsubscript{6}H\textsubscript{10}]\textsuperscript{+}; HRMS (EI\textsuperscript{+}): m/z calcd for C\textsubscript{11}H\textsubscript{16}BrN 241.0461 [M]\textsuperscript{+}, found 241.0467.
2.5.2 2-Bromo-4,5-dihexylpyridine (3c)

![Structure of 3c]

According to the general procedure, 2-(dimethylamino)ethanol (80 µL, 71 mg, 0.80 mmol, 2.0 eq.) in n-hexane (2.6 mL) was reacted with n-BuLi (1.6 M in n-hexane, 1.0 mL, 1.6 mmol, 4.0 eq.) and a solution of 3,4-dihexylpyridine (6c, 99 mg, 400 µmol, 1.0 eq.) in n-hexane (0.60 mL). The reaction was treated with a solution of CBr₄ (332 mg, 1.00 mmol, 2.5 eq.) in THF (1.0 mL). The resulting black suspension was quenched with H₂O (4 mL). Brine (8 mL) and Et₂O (15 mL) were added, separated and the aqueous layer was extracted with EtOAc (3 × 15 mL). The combined organic extracts were dried over Na₂SO₄. Two consecutive chromatography steps (silica gel, cyclohexane/CH₂Cl₂, 7/3 and silica gel, cyclohexane/EtOAc: 70/1) yielded 3c (52.9 mg, 162 µmol, 41%) as a light yellow liquid. Rᵣ 0.65 (cyclohexane/EtOAc: 10/1); ¹H NMR (300 MHz, CDCl₃): δ = 8.07 (s, 1H), 7.23 (s, 1H), 2.57-2.52 (m, 4H), 1.62-1.48 (m, 4H), 1.41-1.25 (m, 12H), 0.93-0.85 (m, 6H); ¹³C NMR (76 MHz, CDCl₃): δ = 153.0, 150.5, 139.7, 135.7, 127.7, 31.9, 31.7, 31.7, 30.8, 30.1, 29.6, 29.4, 29.3, 22.7, 22.7, 14.2; HRMS (ESI+): m/z calcd for C₁₇H₂₈BrN 348.1297 [M+Na]⁺, found 348.1303.

2.5.3 2-Bromo-4-dodecylpyridine (3d)

![Structure of 3d]

According to the general procedure, 2-(dimethylamino)ethanol (925 µL, 825 mg, 9.25 mmol, 2.0 eq.) in n-hexane (29 mL) was reacted with n-BuLi (1.6 M solution in n-hexane, 10.7 mL, 17.6 mmol, 4.0 eq.) and a solution of 4-dodecylpyridine (6d, 109 g, 4.41 mmol, 1.0 eq.) in n-hexane (7.0 mL). The reaction was treated with a solution of CBr₄ (3.66 g, 11.0 mmol, 2.5 eq.) in THF (11 mL). The resulting black suspension was quenched with H₂O (50 mL). Brine (100 mL) and Et₂O (50 mL) were added, separated and the aqueous layer was extracted with EtOAc (3 × 100 mL). Purification by column chromatography (silica gel, cyclohexane/EtOAc) yielded 3d (659 mg, 2.02 mmol, 46%) as a light brown liquid. Rᵣ 0.60 (cyclohexane/EtOAc: 5/1); ¹H NMR (400 MHz, CDCl₃): δ = 8.23 (dd, J = 5.1 Hz, 0.5 Hz, 1H), 7.31 (dd, J = 1.5 Hz, 0.5 Hz, 1H), 7.06, (dd, J = 5.1, 1.5 Hz, 1H), 2.57 (t, J = 7.8 Hz, 2H), 1.67-1.56 (m, 2H), 1.33-1.23 (m, 6H), 0.88 (t, J = 6.8 Hz, 3H); ¹³C NMR (101 MHz, CDCl₃): δ = 155.2, 149.8, 142.4, 128.1, 123.2, 35.0, 32.0, 30.2, 29.7, 29.7, 29.6, 29.5, 29.4, 29.2, 22.8, 14.2; HRMS (EI+): m/z calcd for C₁₇H₂₉NBr 326.1478 [M⁺], found 326.1479.
2.5.4 2-Bromo-4,5-dioctadecanlypyridine (3e)

According to the general procedure, 2-(dimethylamino)ethanol (120 µL, 107 mg, 1.20 mmol, 2.0 eq.) in n-hexane (3.8 mL) was reacted with n-BuLi (1.6 M solution in n-hexane, 1.5 mL, 2.4 mmol, 4.0 eq.) and a solution of 3,4-dioctadecanlypyridine (6e, 342 mg, 586 µmol, 1.0 eq.) in n-hexane (0.90 mL). The reaction was treated with a solution of CBr₄ (486 mg, 1.47 mmol, 2.5 eq.) in THF (1.5 mL). The resulting black suspension was quenched with H₂O (6 mL). Brine (10 mL) and Et₂O (15 mL) were added, separated and the aqueous layer was extracted with EtOAc (3 × 15 mL). Two consecutive chromatography steps (silica gel, cyclohexane/EtOAc: 200/1 and silica gel, cyclohexane/CH₂Cl₂, 10/1) yielded 3e (140 mg, 211 µmol, 36%) as a light yellow solid. Rf 0.25 (cyclohexane/EtOAc: 100/1); Mp 34 °C; ¹H NMR (400 MHz, CDCl₃): δ = 8.08 (s, 1H), 7.24 (s, 1H), 2.55 (t, J = 7.8 Hz, 2H), 2.55 (t, J = 7.8 Hz, 2H), 1.61–1.50 (m, 4H), 1.40–1.26 (m, 60H), 0.88 (t, J = 6.8 Hz, 6H); ¹³C NMR (101 MHz, CDCl₃): δ = 153.0, 150.5, 139.7, 135.7, 127.7, 32.1, 31.9, 30.9, 30.1, 29.9, 29.8, 29.8, 29.7, 29.7, 29.7, 29.6, 29.6, 29.6, 29.5, 29.5, 22.9, 14.3; HRMS (ESI⁺): m/z calcd for C₄₁H₇₆BrN₆ 662.5234 [M+H]⁺, found 662.5245.

2.6 General Procedure for Synthesis of Tris[µ-pyridinato-N¹,C²]tri-gold(I) complexes 1

The synthesis was performed following a modified procedure by Vaughan et al. To a stirred solution of 2-bromopyridine 3 in dry THF (40–60 µmol/mL) was added n-BuLi (0.3–1.6 M solution in n-hexane, 1.0 eq.) dropwise at −40 to −25 °C, and the resulting orange or red reaction mixture was stirred for 1 h. Chloro(triphenylarsine)gold(I) (1.0 eq.) was suspended in THF (50–100 µmol/mL) at −40 °C and added to the reaction mixture. The resulting suspension was allowed to warm to room temperature over a period of 1–3 h, and the mixture was concentrated in vacuo. The residue was filtered through a plug of alumina and either purified by recrystallization from toluene and benzene or column chromatography (alumina, cyclohexane/toluene or cyclohexane/CH₂Cl₂) to yield pyridinates 1 in yields of 6–37%. Yields are not optimized.
2.6.1 Tris[µ-(4-hexylpyridinato-N¹,C²)]tri-gold(I) (1b)

According to the general procedure, 2-bromo-4-hexylpyridine (3a, 78.0 mg, 322 µmol, 1.0 eq.) in dry THF (7 mL) was reacted with n-BuLi (1.7 M solution in n-hexane, 0.19 mL, 320 µmol, 1.0 eq.) and chloro(triphenylarsine)gold(I) (173 mg, 322 µmol, 1.0 eq.) in dry THF (3 mL). Two consecutive column chromatography steps (alumina, cyclohexane/CH₂Cl₂ and alumina, cyclohexane/benzene) yielded 1b (42.1 mg, 39.1 µmol, 36%) as a colorless wax. Rᵣ 0.17 (alumina, cyclohexane/CH₂Cl₂: 5/1); ¹H NMR (400 MHz, C₆D₆): δ = 8.13 (d, J = 5.9 Hz, 3H), 7.67 (d, J = 1.8 Hz, 3H), 6.26 (dd, J = 5.9 Hz, 1.8 Hz, 3H), 2.15 (t, J = 7.7 Hz, 6H), 1.38–1.17 (m, 6H), 1.25–1.10 (m, 18H), 0.87 (t, J = 7.2 Hz, 9H); ¹³C NMR (101 MHz, C₆D₆): δ = 184.6, 151.3, 150.5, 138.7, 121.2, 35.4, 32.0, 30.1, 29.3, 22.9, 14.3; MS (ESI+): m/z calcd for C₃₃H₄₈Au₃N₃ 1100.3 [M+Na]+, found 1100.4.

2.6.2 Tris[µ-(4,5-dihexanylpyridinato-N¹,C²)]tri-gold(I) (1c)

According to the general procedure, 2-bromo-4,5-dihexylpyridine (3b, 40 mg, 120 µmol, 1.0 eq.) in dry THF (4 mL) was reacted with n-BuLi (0.28 M solution in n-hexane, 0.44 mL, 120 µmol, 1.0 eq.) and chloro(triphenylarsine)gold(I) (66 mg, 120 µmol, 1.0 eq.) in dry THF (3 mL). The residue was purified by several column chromatography steps (alumina, cyclohexane/toluene) to yield 1c (3.0 mg, 6.8 µmol, 6%) as a colorless wax. Significant amounts of AsPh₃ and traces of other unknown impurities could not be completely removed. Rᵣ 0.35 (alumina, toluene); ¹H NMR

(500 MHz, C₆D₆): δ = 8.27 (s, 3H), 7.79 (s, 3H), 2.34 (t, J = 7.8 Hz, 6H), 2.28 (t, J = 7.8 Hz, 6H), 1.51–1.43 (m, 6H), 1.37–1.14 (m, 42H), 0.85–0.93 (m, 18H); ¹³C NMR (126 MHz, C₆D₆): δ = 181.1, 151.6, 148.9, 138.5, 133.4, 32.0, 32.0, 32.0, 30.8, 30.0, 29.9, 29.7, 23.0, 22.9, 14.4, 14.3; HRMS (ESI+): m/z calcd for C₅₁H₈₄Au₃N₃ 1352.5554 [M+Na]+, found 1352.5535.

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2.6.3  Tris[µ-(4-dodecylpyridinato-N¹,C²)]tri-gold(I) (1d)

According to the general procedure, 2-bromo-4-hexanylpyridine (3c, 178 mg, 545 µmol, 1.0 eq.) in dry THF (12 mL) was reacted with n-BuLi (1.6 M solution in n-hexane, 0.34 mL, 550 µmol, 1.0 eq.) and solid chloro(triphenylarsine)gold(I) (294 mg, 545 µmol, 1.0 eq.). Two consecutive column chromatography steps (alumina, cyclohexane/CH₂Cl₂ and Al₂O₃, cyclohexane/benzene) and several recrystallization steps from toluene yielded 1d (17.1 mg, 12.8 µmol, 7%) as white solid. Rf 0.2 (alumina, cyclohexane/CH₂Cl₂: 5/1); ¹H NMR (400 MHz, C₆D₆): δ = 8.12 (d, J = 5.8 Hz, 3H), 7.68 (d, J = 1.7 Hz, 3H), 6.27 (dd, J = 5.9 Hz, 1.8 Hz, 3H), 2.19 (t, J = 7.8 Hz, 6H), 1.44–1.15 (m, 60H), 0.92 (t, J = 7.2 Hz, 9H); ¹³C NMR (101 MHz, C₆D₆): δ = 184.4, 151.3, 150.6, 138.7, 121.3, 35.4, 32.4, 30.2, 30.2, 29.9, 29.9, 29.8, 29.6, 22.9, 14.3; MS (MALDI–TOF): m/z calcd for C₅₁H₈₄Au₃N₃ 1329.6 [M⁺], found 1329.6.

2.6.4  Tris[µ-(4,5-dioctadecanylpyridinato-N¹,C²)]tri-gold(I) (1e)

According to the general procedure, 2-bromo-4,5-dioctadecanylpyridine (3e, 135 mg, 204 µmol, 1.0 eq.) in dry THF (5 mL) was reacted with n-BuLi (0.30 M solution in n-hexane, 0.68 mL, 204 µmol, 1.0 eq.) and chloro(triphenylarsine)gold(I) (110 mg, 204 µmol, 1.0 eq.) in dry THF (4 mL). The residue was recrystallized twice from toluene (2.5 mL), decanted and washed with cold toluene (4 × 0.5 mL). The remaining solid was dissolved in hot toluene and filtered through a plug of alumina at 50 °C. The crude product was recrystallized from benzene, filtered through a plug of alumina and dried in vacuo to yield 1e (24.7 mg, 10.6 µmol, 16%) as a white solid. Mp 70 °C (decomposition at 128 °C); ¹H NMR (500 MHz, C₆D₆, 313 K): δ = 8.31 (s, 3H), 7.80 (s, 3H), 2.43 (t, J = 7.9 Hz, 6H), 2.37 (t, J = 8.0 Hz, 6H), 1.59–1.52 (m, 6H), 1.46–1.31 (m, 186H), 0.93 (t, J = 6.9 Hz, 18H); ¹³C NMR (126 MHz, C₆D₆, 313 K): δ = 181.2,
151.7, 148.8, 138.6, 133.4, 32.4, 32.1, 31.0, 30.3, 30.3, 30.2, 30.2, 30.1, 30.1, 30.0, 30.0, 30.0, 29.9, 29.9, 29.8, 23.1, 14.4; MS (MALDI–TOF): m/z calcd for C₆₉H₁₂₀Au₃N₃ 2338.69 [M]⁺, found 2338.76.
3 NMR titration experiments

Binding constants were determined through NMR titrations in C₆D₆ and CDCl₃ according to a procedure by Thordarson et al. [11] All titration experiments were carried out at constant concentration of the donor 1, therefore all solutions of acceptors 2 were prepared using a stock solution of the donor 1. Experiments were carried out as follows: A stock solution of donor 1 (ca. 4 mL) was prepared, and four NMR-tubes were filled with different amounts of this stock solution (0.5 to 0.15 mL). Three titration solutions containing different concentrations of the acceptors 2 were prepared from the initial donor stock solution. Each NMR-tube was then filled to a total volume of 0.5 mL. ¹H-NMR spectra were measured, and four to six small portions of acceptor solution were successively added. Each new sample was analyzed by ¹H NMR spectroscopy. All volumes were quantified by weighing using Hamilton syringes, and the weight differences were estimated and used for calculation of the concentrations. The measured data was fitted to a 1:2 binding model, using either the website supramolecular.org or a MatLab code provided by Thordarson et al. [11] All titrations between donor 1b and acceptors 2a–c were performed three times, and the data are listed in the following.
3.1 Titration of pyridinate 1b with gold pyrazolate 2a

3.1.1 Data of experiment 1

Figure S1. $^1$H NMR spectra for the titration of 1b with 2a (Experiment 1).
Table S 1. Measured data of the titration experiment between 1b and 2a, fitted to a non-linear 1:1 binding model (Experiment 1).

<table>
<thead>
<tr>
<th>$K_{11}$ [L mol$^{-1}$]</th>
<th>$\delta_H$</th>
<th>$\delta_{HG}$</th>
<th>RMS</th>
<th>Cov</th>
</tr>
</thead>
<tbody>
<tr>
<td>$1.50 \times 10^3$</td>
<td>proton b</td>
<td>8.134</td>
<td>7.842</td>
<td>$1.79 \times 10^{-2}$</td>
</tr>
<tr>
<td>$K_{11}$ error [%]</td>
<td>proton a</td>
<td>7.663</td>
<td>7.192</td>
<td>$6.52 \times 10^{-2}$</td>
</tr>
<tr>
<td>66.5</td>
<td>proton c</td>
<td>6.261</td>
<td>6.019</td>
<td>$1.10 \times 10^{-2}$</td>
</tr>
<tr>
<td>total</td>
<td></td>
<td></td>
<td></td>
<td>$3.95 \times 10^{-2}$</td>
</tr>
</tbody>
</table>

Table S 2. Measured data of the titration experiment between 1b and 2a, fitted to a non-linear 1:2 binding model (Experiment 1).

<table>
<thead>
<tr>
<th>$K_{11}$ [L mol$^{-1}$]</th>
<th>$K_{12}$ [L mol$^{-1}$]</th>
<th>$\delta_H$ [ppm]</th>
<th>$\delta_{HG}$ [ppm]</th>
<th>$\delta_{HG2}$ [ppm]</th>
<th>RMS</th>
<th>Cov</th>
</tr>
</thead>
<tbody>
<tr>
<td>$1.52 \times 10^4$</td>
<td>446</td>
<td>proton b</td>
<td>8.133</td>
<td>7.881</td>
<td>7.802</td>
<td>$2.06 \times 10^{-3}$</td>
</tr>
<tr>
<td>$K_{11}$ error [%]</td>
<td>$K_{12}$ error [%]</td>
<td>proton a</td>
<td>7.662</td>
<td>7.342</td>
<td>7.032</td>
<td>$3.15 \times 10^{-3}$</td>
</tr>
<tr>
<td>6.46</td>
<td>2.15</td>
<td>proton c</td>
<td>6.260</td>
<td>6.042</td>
<td>5.995</td>
<td>$1.68 \times 10^{-3}$</td>
</tr>
<tr>
<td>total</td>
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<td></td>
<td></td>
<td></td>
<td>$2.38 \times 10^{-3}$</td>
</tr>
</tbody>
</table>
Figure S 2. Binding isotherms of the titration experiment between 1b and 2a, fitted to a 1:1 binding model, and the corresponding residuals between fitted and measured values (Experiment 1).
Figure S 3. Binding isotherms of the titration experiment between 1b and 2a, fitted to a 1:2 binding model, and the corresponding residuals between fitted and measured values (Experiment 1).
3.1.2 Data of experiment 2

Figure S 4. $^1$H NMR spectra for the titration of 1b with 2a (Experiment 2).

Table S 3. Measured data of the titration experiment between 1b and 2a, fitted to a non-linear 1:2 binding model (Experiment 2).

<table>
<thead>
<tr>
<th>$K_1$ [L mol$^{-1}$]</th>
<th>$K_{12}$ [L mol$^{-1}$]</th>
<th>$\delta_H$ [ppm]</th>
<th>$\delta_{HG}$ [ppm]</th>
<th>$\delta_{HG2}$ [ppm]</th>
<th>RMS</th>
<th>Cov</th>
</tr>
</thead>
<tbody>
<tr>
<td>5.09 x 10$^4$</td>
<td>574</td>
<td>proton b</td>
<td>8.127</td>
<td>7.909</td>
<td>7.790</td>
<td>2.70 x 10$^{-3}$</td>
</tr>
<tr>
<td>$K_1$ error [%]</td>
<td>$K_{12}$ error [%]</td>
<td>proton a</td>
<td>7.661</td>
<td>7.039</td>
<td>7.048</td>
<td>4.25 x 10$^{-3}$</td>
</tr>
<tr>
<td>14.4</td>
<td>3.29</td>
<td>proton c</td>
<td>6.254</td>
<td>6.051</td>
<td>5.992</td>
<td>4.92 x 10$^{-3}$</td>
</tr>
<tr>
<td>total</td>
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<td></td>
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<td></td>
<td>4.06 x 10$^{-3}$</td>
<td>1.17 x 10$^{-3}$</td>
</tr>
</tbody>
</table>
Figure S 5. Binding isotherms of the titration experiment between 1b and 2a, fitted to a 1:2 binding model and the corresponding residuals between fitted and measured values (Experiment 2).
3.1.3 Data of experiment 3

Figure S 6. $^1$H NMR spectra for the titration of 1b with 2a (Experiment 3).

Table S 4. Measured data of the titration experiment between 1b and 2a, fitted to a non-linear 1:2 binding model (Experiment 3).

<table>
<thead>
<tr>
<th>$K_{11}$ [L mol$^{-1}$]</th>
<th>$K_{12}$ [L mol$^{-1}$]</th>
<th>$\delta_{11}$ [ppm]</th>
<th>$\delta_{HG}$ [ppm]</th>
<th>$\delta_{HG2}$ [ppm]</th>
<th>RMS</th>
<th>Cov</th>
</tr>
</thead>
<tbody>
<tr>
<td>2.41 $\times$ 10$^4$</td>
<td>526</td>
<td>proton b</td>
<td>8.132</td>
<td>7.896</td>
<td>7.800</td>
<td>1.44 $\times$ 10$^{-3}$</td>
</tr>
<tr>
<td>$K_{11}$ error [%]</td>
<td>$K_{12}$ error [%]</td>
<td>proton a</td>
<td>7.666</td>
<td>7.365</td>
<td>7.045</td>
<td>2.39 $\times$ 10$^{-3}$</td>
</tr>
<tr>
<td>4.03</td>
<td>1.69</td>
<td>proton c</td>
<td>6.260</td>
<td>6.051</td>
<td>5.993</td>
<td>1.74 $\times$ 10$^{-3}$</td>
</tr>
<tr>
<td>total</td>
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<td></td>
<td></td>
<td></td>
<td>1.90 $\times$ 10$^{-3}$</td>
<td>2.58 $\times$ 10$^{-4}$</td>
</tr>
</tbody>
</table>

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Figure S 7. Binding isotherms of the titration experiment between 1b and 2a, fitted to a 1:2 binding model, and the corresponding residuals between fitted and measured values (Experiment 3).
3.2 Titration of pyridinate 1b with copper pyrazolate 2b

3.2.1 Data of experiment 1

![Chemical structures of 1b and 2b](image)

Figure S 8. $^1$H NMR spectra for the titration of 1b with 2b (Experiment 1).

Table S 5. Measured data of the titration experiment between 1b and 2b, fitted to a non-linear 1:2 binding model (Experiment 1).

<table>
<thead>
<tr>
<th>$K_{11}$ [L mol$^{-1}$]</th>
<th>$K_{12}$ [L mol$^{-1}$]</th>
<th>$\delta_a$ [ppm]</th>
<th>$\delta_{HG}$ [ppm]</th>
<th>$\delta_{HG2}$ [ppm]</th>
<th>RMS</th>
<th>Cov</th>
</tr>
</thead>
<tbody>
<tr>
<td>1.00 × 10$^5$</td>
<td>225</td>
<td>proton b</td>
<td>8.134</td>
<td>7.704</td>
<td>7.540</td>
<td>5.66 × 10$^{-3}$</td>
</tr>
<tr>
<td>$K_{11}$ error [%]</td>
<td>$K_{12}$ error [%]</td>
<td>proton a</td>
<td>7.663</td>
<td>7.358</td>
<td>7.099</td>
<td>3.63 × 10$^{-3}$</td>
</tr>
<tr>
<td>28.23</td>
<td>3.78</td>
<td>proton c</td>
<td>6.261</td>
<td>6.055</td>
<td>6.010</td>
<td>2.94 × 10$^{-3}$</td>
</tr>
<tr>
<td>total</td>
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<td></td>
<td></td>
<td></td>
<td>4.24 × 10$^{-3}$</td>
<td>8.49 × 10$^{-4}$</td>
</tr>
</tbody>
</table>
Figure S 9. Binding isotherms of the titration experiment between 1b and 2b, fitted to a 1:2 binding model, and the corresponding residuals between fitted and measured values (Experiment 1).
3.2.2 Data of experiment 2

Figure S 10. $^1$H NMR spectra for the titration of 1b with 2b (Experiment 2).

Table S 6. Measured data of the titration experiment between 1b and 2b, fitted to a non-linear 1:2 binding model (Experiment 2).

<table>
<thead>
<tr>
<th>$K_{11}$ [L mol$^{-1}$]</th>
<th>$K_{12}$ [L mol$^{-1}$]</th>
<th>$\delta_{1b}$ [ppm]</th>
<th>$\delta_{1bG}$ [ppm]</th>
<th>$\delta_{1bG2}$ [ppm]</th>
<th>RMS</th>
<th>Cov</th>
</tr>
</thead>
<tbody>
<tr>
<td>1.65 $\times$ 10$^5$</td>
<td>341</td>
<td>proton b</td>
<td>8.127</td>
<td>7.727</td>
<td>7.541</td>
<td>3.38 $\times$ 10$^{-3}$</td>
</tr>
<tr>
<td>$K_{11}$ error [%]</td>
<td>$K_{12}$ error [%]</td>
<td>proton a</td>
<td>7.661</td>
<td>7.383</td>
<td>7.119</td>
<td>2.49 $\times$ 10$^{-3}$</td>
</tr>
<tr>
<td>17.8</td>
<td>3.45</td>
<td>proton c</td>
<td>6.254</td>
<td>6.059</td>
<td>5.999</td>
<td>4.90 $\times$ 10$^{-3}$</td>
</tr>
<tr>
<td>total</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>3.73 $\times$ 10$^{-3}$</td>
</tr>
</tbody>
</table>
Figure S11. Binding isotherms of the titration experiment between 1b and 2b, fitted to a 1:2 binding model, and the corresponding residuals between fitted and measured values (Experiment 2).
### 3.2.3 Data of experiment 3

![Figure S12. ¹H NMR spectra for the titration of 1b with 2b (Experiment 3).](image)

#### Table S7. Measured data of the titration experiment between 1b and 2b, fitted to a non-linear 1:2 binding model (Experiment 3).

<table>
<thead>
<tr>
<th></th>
<th></th>
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<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>2.05 × 10⁵</td>
<td>214</td>
<td>8.132</td>
<td>7.728</td>
<td>7.506</td>
<td>5.98 × 10⁻³</td>
<td>2.46 × 10⁻³</td>
</tr>
<tr>
<td>$K_{11}$ error [%]</td>
<td>$K_{12}$ error [%]</td>
<td>proton b</td>
<td>proton a</td>
<td>proton c</td>
<td></td>
<td></td>
</tr>
<tr>
<td>25.8</td>
<td>3.76</td>
<td>7.666</td>
<td>7.377</td>
<td>6.260</td>
<td>4.43 × 10⁻³</td>
<td>1.69 × 10⁻³</td>
</tr>
<tr>
<td></td>
<td></td>
<td>6.055</td>
<td>6.003</td>
<td>total</td>
<td>4.44 × 10⁻³</td>
<td>1.19 × 10⁻³</td>
</tr>
</tbody>
</table>
Figure S 13. Binding isotherms of the titration experiment between 1b and 2b, fitted to a 1:2 binding model, and the corresponding residuals between fitted and measured values (Experiment 3).
3.3 Titration of pyridinate 1b with silver pyrazolate 2c

3.3.1 Data of experiment 1

![Chemical structures of 1b and 2c]

Figure S 14. $^1$H NMR spectra for the titration of 1b with 2c (Experiment 1).

Table S 8. Measured data of the titration experiment between 1b and 2c, fitted to a non-linear 1:2 binding model (Experiment 1).

<table>
<thead>
<tr>
<th>$K_{11}$ [L mol$^{-1}$]</th>
<th>$K_{12}$ [L mol$^{-1}$]</th>
<th>$\delta_H$ [ppm]</th>
<th>$\delta_{HG}$ [ppm]</th>
<th>$\delta_{HG2}$ [ppm]</th>
<th>RMS</th>
<th>Cov</th>
</tr>
</thead>
<tbody>
<tr>
<td>2.91 $\times$ 10$^5$</td>
<td>1.42 $\times$ 10$^3$</td>
<td>7.662</td>
<td>7.357</td>
<td>7.289</td>
<td>2.26 $\times$ 10$^{-3}$</td>
<td>7.75 $\times$ 10$^{-4}$</td>
</tr>
<tr>
<td>$K_{11}$ error [%]</td>
<td>$K_{12}$ error [%]</td>
<td>proton a</td>
<td>7.662</td>
<td>7.357</td>
<td>7.289</td>
<td>2.26 $\times$ 10$^{-3}$</td>
</tr>
<tr>
<td>19.8</td>
<td>15.9</td>
<td>proton c</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
Figure S 15. Binding isotherms of the titration experiment between 1b and 2c, fitted to a 1:2 binding model, and the corresponding residuals between fitted and measured values (Experiment 1).
3.3.2 Data of experiment 2

Figure S 16. $^1$H NMR spectra for the titration of 1b with 2c (Experiment 2).

Table S 9. Measured data of the titration experiment between 1b and 2c, fitted to a non-linear 1:2 binding model (Experiment 2).

<table>
<thead>
<tr>
<th>$K_{11}$ [L/mol$^{-1}$]</th>
<th>$K_{12}$ [L/mol$^{-1}$]</th>
<th>$\delta_H$ [ppm]</th>
<th>$\delta_{HG}$ [ppm]</th>
<th>$\delta_{HG2}$ [ppm]</th>
<th>RMS</th>
<th>Cov</th>
</tr>
</thead>
<tbody>
<tr>
<td>2.90 $\times$ 10$^5$</td>
<td>5.12 $\times$ 10$^3$</td>
<td>8.126</td>
<td>7.441</td>
<td>7.321</td>
<td>1.35 $\times$ 10$^{-2}$</td>
<td>4.90 $\times$ 10$^{-2}$</td>
</tr>
<tr>
<td>$K_{11}$ error [%]</td>
<td>$K_{12}$ error [%]</td>
<td>proton b</td>
<td>7.661</td>
<td>7.317</td>
<td>7.324</td>
<td>1.23 $\times$ 10$^{-2}$</td>
</tr>
<tr>
<td>36.7</td>
<td>26.9</td>
<td>proton a</td>
<td>6.254</td>
<td>6.082</td>
<td>6.082</td>
<td>9.58 $\times$ 10$^{-3}$</td>
</tr>
<tr>
<td>proton e</td>
<td></td>
<td>proton c</td>
<td>2.172</td>
<td>2.245</td>
<td>2.329</td>
<td>5.34 $\times$ 10$^{-3}$</td>
</tr>
<tr>
<td>total</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>1.07 $\times$ 10$^{-2}$</td>
<td>1.21 $\times$ 10$^{-3}$</td>
</tr>
</tbody>
</table>
Figure S 17. Binding isotherms of the titration experiment between 1b and 2c, fitted to a 1:2 binding model, and the corresponding residuals between fitted and measured values (Experiment 2).
3.3.3 Data of experiment 3

Figure S 18. $^1$H NMR spectra for the titration of 1b with 2c (Experiment 3).

Table S 10. Measured data of the titration experiment between 1b and 2c, fitted to a non-linear 1:2 binding model (Experiment 3).

<table>
<thead>
<tr>
<th>$K_{11}$ [L mol$^{-1}$]</th>
<th>$K_{12}$ [L mol$^{-1}$]</th>
<th>$\delta_H$ [ppm]</th>
<th>$\delta_{HG}$ [ppm]</th>
<th>$\delta_{HG2}$ [ppm]</th>
<th>RMS</th>
<th>Cov</th>
</tr>
</thead>
<tbody>
<tr>
<td>$2.70 \times 10^5$</td>
<td>$2.72 \times 10^3$</td>
<td>proton b</td>
<td>8.122</td>
<td>7.423</td>
<td>7.305</td>
<td>$1.87 \times 10^{-2}$</td>
</tr>
<tr>
<td>$K_{11}$ error [%]</td>
<td>$K_{12}$ error [%]</td>
<td>proton a</td>
<td>7.662</td>
<td>7.323</td>
<td>7.327</td>
<td>$1.49 \times 10^{-2}$</td>
</tr>
<tr>
<td>43.8</td>
<td>35.8</td>
<td>proton c</td>
<td>6.260</td>
<td>6.047</td>
<td>6.093</td>
<td>$1.20 \times 10^{-2}$</td>
</tr>
<tr>
<td>total</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>$1.54 \times 10^{-2}$</td>
<td>$3.89 \times 10^{-3}$</td>
</tr>
</tbody>
</table>
Figure S 19. Binding isotherms of the titration experiment between 1b and 2c, fitted to a 1:2 binding model, and the corresponding residuals between fitted and measured values (Experiment 3).
3.4 Titration of pyridinate 1d with silver pyrazolate 2c

Figure S 20. $^1$H NMR spectra for the titration of 1d with 2c.
Table S11. Measured data of the titration experiment between 1d and 2c, fitted to a non-linear 1:2 binding model.

<table>
<thead>
<tr>
<th>$K_{11}$ [L mol$^{-1}$]</th>
<th>$K_{12}$ [L mol$^{-1}$]</th>
<th>$\delta_{44}$ [ppm]</th>
<th>$\delta_{HG}$ [ppm]</th>
<th>$\delta_{HG2}$ [ppm]</th>
<th>RMS</th>
<th>Cov</th>
</tr>
</thead>
<tbody>
<tr>
<td>5.81 × 10$^5$</td>
<td>9.43 × 10$^3$</td>
<td>proton b</td>
<td>8.136</td>
<td>7.473</td>
<td>7.331</td>
<td>1.52 × 10$^{-2}$</td>
</tr>
<tr>
<td>$K_{11}$ error [%]</td>
<td>$K_{12}$ error [%]</td>
<td>proton a</td>
<td>7.677</td>
<td>7.342</td>
<td>7.344</td>
<td>8.99 × 10$^{-3}$</td>
</tr>
<tr>
<td>32.3</td>
<td>32.7</td>
<td>proton c</td>
<td>6.282</td>
<td>6.062</td>
<td>6.108</td>
<td>6.80 × 10$^{-3}$</td>
</tr>
<tr>
<td>total</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>RMS</td>
<td>Cov</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>1.09 × 10$^{-2}$</td>
<td>1.91 × 10$^{-3}$</td>
</tr>
</tbody>
</table>
Figure S 21. Binding isotherms of the titration experiment between 1b and 2c, fitted to a 1:2 binding model, and the corresponding residuals between fitted and measured values.
4 Stability of pyridinate 1b

Scheme S 1. Decomposition of pyridinate 1b in solution under sunlight exposition, forming 2,2'-bipyridine 10 and colloidal gold.

Figure S 22. $^1$H NMR spectra of pyridinate 1b over a period of three days, showing successive decomposition to 2,2'-bipyridine 10 due to sunlight exposure.
5 NMR spectra

Figure S 23. $^1$H NMR spectrum of 5a in CDCl$_3$ (400 MHz).

Figure S 24. $^{13}$C NMR spectrum of 5a in CDCl$_3$ (101 MHz).
Figure S 25. $^1$H NMR spectrum of 5b in CDCl$_3$ (500 MHz).

Figure S 26. $^{13}$C NMR spectrum of 5b in CDCl$_3$ (126 MHz).
Figure S 27. $^1$H NMR spectrum of 6b in CDCl$_3$ (400 MHz).

Figure S 28. $^{13}$C NMR spectrum of 6b in CDCl$_3$ (76 MHz).
Figure S 29. $^1$H NMR spectrum of 6c in CDCl$_3$ (400 MHz).

Figure S 30. $^{13}$C NMR spectrum of 6c in CDCl$_3$ (76 MHz).
Figure S 31. $^1$H NMR spectrum of 6d in CDCl$_3$ (400 MHz).

Figure S 32. $^{13}$C NMR spectrum of 6d in CDCl$_3$ (101 MHz).
Figure S 33. $^1$H NMR spectrum of 6f in CDCl$_3$ (400 MHz).

Figure S 34. $^{13}$C NMR spectrum of 6f in CDCl$_3$ (101 MHz).
Figure S 35. $^1$H NMR spectrum of 3b in CDCl$_3$ (400 MHz).

Figure S 36. $^{13}$C NMR spectrum of 3b in CDCl$_3$ (76 MHz).
Figure S 37. $^1$H NMR spectrum of 3c in CDCl₃ (300 MHz).

Figure S 38. $^{13}$C NMR spectrum of 3c in CDCl₃ (76 MHz).
Figure S 39. $^1$H NMR spectrum of 3d in CDCl$_3$ (400 MHz).

Figure S 40. $^{13}$C NMR spectrum of 3d in CDCl$_3$ (101 MHz).
Figure S 41. $^1$H NMR spectrum of 3f in CDCl$_3$ (400 MHz).

Figure S 42. $^{13}$C NMR spectrum of 3f in CDCl$_3$ (101 MHz).
Figure S 43. $^1$H NMR spectrum of 1b in CDCl$_3$ (400 MHz).

Figure S 44. $^{13}$C NMR spectrum of 1b in CDCl$_3$ (101 MHz).
Figure S 45. $^1$H NMR spectrum of 1c in CDCl$_3$ (500 MHz).

Figure S 46. $^{13}$C NMR spectrum of 1c in CDCl$_3$ (126 MHz).
Figure S 47. $^1$H NMR spectrum of 1d in CDCl$_3$ (400 MHz).

Figure S 48. $^{13}$C NMR spectrum of 1d in CDCl$_3$ (101 MHz).
Figure S 49. $^1$H NMR spectrum of 1e in CDCl$_3$ (500 MHz) at 313 K.

Figure S 50. $^{13}$C NMR spectrum of 1e in CDCl$_3$ (126 MHz) at 313 K.
6 UV/Vis spectroscopy

Solutions of donor pyridinate 1d, acceptor pyrazolates 2 and equal mixtures of both, donor and acceptor, were recorded by UV/Vis spectroscopy in n-hexanes, with concentrations between $1.0 \times 10^{-6}$ and $1.0 \times 10^{-4}$ mol/L. All spectra were measured at ten different concentrations, each was normalized to $\varepsilon_{\lambda}$, using the absorbance $E_{\lambda}$ and the concentration $c$. The solutions containing both the donor and the acceptor in equal concentrations were normalized to the concentration of pyridinate 1d.

$$\varepsilon_{\lambda} = \frac{E_{\lambda}}{c}$$

The analyzed samples were colorless, and no absorbance above $\lambda = 400$ nm was visible. A Charge-Transfer (CT) can be observed at $\lambda = 250$–350 nm.

![UV/Vis spectra](image)

Figure 5 51: UV/Vis spectra of pyridinate 1d (red), gold pyrazolate 2a (blue) and the 1:1 mixture of donor and acceptor (black).
Figure S 52 UV/Vis spectra of pyridinate 1d (red), copper pyrazolate 2b (blue) and the 1:1 mixture of donor and acceptor (black).

Figure S 53 UV/Vis spectra of pyridinate 1d (red), silver pyrazolate 2c (blue) and the 1:1 mixture of donor and acceptor (black).
7 Computational studies

7.1 General remarks

Our standard procedure for obtaining free energies $\Delta G$ is subdivided into three steps.$^{12}$ In step one, the electronic association energy $E_a$ is computed from equilibrium structures in gas phase. All molecules are optimized in gas phase employing the density functional PBEh-3c.$^{13}$ This composite method is designed to yield accurate molecular geometries and makes use of a modified double-zeta (DZ) basis set$^{14}$ (dubbed def2-mSVP) balanced with the D3-dispersion$^{15}$ correction, including the three body contributions (D3$^{ATM}$) and a geometrical counter-poise correction (gCP).$^{16}$ Single-point energies are calculated employing the hybrid functional PBE0$^{17,18}$_D3$^{ATM}$ combined with the quadruple-zeta basis set (QZ). At this QZ basis set level the basis set superposition errors are typically in the range of less than 2% of $\Delta E_a$ and can therefore be discarded.$^{19}$ To account for physically correct description of the long range correlation (mainly London dispersion) the pairwise additive D3 dispersion correction with Becke-Johnson damping$^{20,21}$ is used for all single-point calculations. The Axilrod-Teller-Muto type three-body$^{22,23}$ dispersion energy is always included and many-body dispersion contributions beyond the three body term are neglected. The electronic energy comprises of

$$E = E_{el} + E_{disp} + E_{disp}$$

where $E$ is the electronic energy of one species involved. The association energy $\Delta E_a$ in gas phase is calculated in the supramolecular approach

$$\Delta E_a = E_{product} - (E_{reactant1} + E_{reactant2})$$

where $E$ denotes the total electronic energy of the species. In step two, the thermo-statistical corrections from energy to free energy $G_{\text{RRHO}}^T$ are calculated for each molecule.$^{12}$ The thermo-statistical corrections are obtained in gas phase at a given temperature $T$ and normal pressure of 1 atm, including the zero-point vibrational energy. The computation of harmonic frequencies is carried out with the global hybrid PBEh-3c and for the larger systems with our newly developed tight binding method GFN-xTB.$^{24}$ The third step is the calculation of the solvation free energies $\delta G_{\text{solv}}^T(X)$ for each molecule at a given temperature $T$ and a solvent $X$. To get the solvation free energies COSMO-RS$^{25-27}$ is used in a black boxed manner and all results are implicitly converted to standard state conditions. The final free energy $\Delta G_a$ is the sum of the three contributions:

$$\Delta G_a = \Delta E_a + \Delta G_{\text{RRHO}}^T + \Delta \delta G_{\text{solv}}^T(X)$$
7.2 Technical details

All DFT calculations were performed using the TURBOMOLE 7.0.2 program package. The resolution-of-identity (RI) approximation for the Coulomb integrals was generally applied using matching default auxiliary basis sets. For the integration of the exchange-correlation contribution the numerical quadrature grids m3 in case of PBEh-3c and m4 in case of PBE0-D3^ATM and PW6B95-D3^ATM were employed. Single-point energies are calculated with PBE0-D3^ATM/(def2-QZ-f,g) and PW6B95-D3^ATM /def2-QZVP. The default convergence criteria for single-point energies as well as for geometry optimizations were used [10^7 E_h for energies and 10^5 E_h/Bohr for gradients]. The vibrational frequencies of PBEh-3c were scaled with a factor of 0.95. The conductor like screening model for real solvents (COSMO-RS) was used as implemented in COSMOtherm2016 employing the BP_TZVP_C30_1401-parametrization. For each solvation free energy value two single-point calculations with BP86/def-TZVP (one in the gas-phase and one in an ideal conductor) have to be performed on the optimized structures. These calculations are then used as input for the COSMOtherm program. The solvent for the calculations was benzene and the temperature was set to 298.15 K. All visualizations were done with USCF Chimera version 1.10.2.

The DLPNO-CCSD(T)[36,37] reference was calculated with the ORCA 4.0.0[38,39] implementation, i.e. employing the sparse maps infrastructure "TightPNO" setup (full LMP2 guess used, TCutPairs 1e-5 E_h, TCutPNO 1e-7, TCutDO 5e-3) and ORCA "TightSCF". The employed basis sets were: def2-TZVPP[14], def2-TZVPP/C[40] with corresponding ECPs for Ag and Au [Def2-ECP][41]; def2-QZVPP[14], def2-QZVPP/C[40] with corresponding ECPs for Ag and Au [Def2-ECP][41]. The ECP parameters for Au [Def2-ECP], Ag [Def2-ECP] have been obtained from TURBOMOLE 7.0.2. The CBS extrapolation has been performed with optimized exponents as proposed by Neese and Valeev. The electronic association energy ΔE_a includes the deformation energy as well (6.8 kcal mol⁻¹, mostly from the Ag complex (2c)), estimated error of the final ΔE_a= +/- 0.5 kcal mol⁻¹.

Table S 12. DLPNO-CCSD(T)/CBS(TZ/QZ) values for 1f•2c (all energies in kcal mol⁻¹).

<table>
<thead>
<tr>
<th>1f (1:1) with Accepter 2c</th>
<th>ΔE_int/CBS(TZ/QZ)</th>
<th>ΔE_deform/CBS(TZ/QZ)</th>
<th>ΔE_a (1f•2c) /CBS(TZ/QZ)</th>
</tr>
</thead>
<tbody>
<tr>
<td>2c (M = Ag)</td>
<td>-45.9</td>
<td>6.8</td>
<td>-39.15</td>
</tr>
</tbody>
</table>
Table S 13. D-A Free binding energies calculated with PW6B95-D3/QZ (all energies in kcal mol\(^{-1}\)).

<table>
<thead>
<tr>
<th>Molecule</th>
<th>$\Delta E$(PW6B95-D3/QZ)</th>
<th>$\Delta G_{\text{BH}}$(PBEh-3c)</th>
<th>$\Delta \delta G_{\text{sol}}$</th>
<th>$\Delta G_s$ (1f 2)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1f (1:1) with Acceptor 2</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>2a (M = Au)</td>
<td>−35.1</td>
<td>17.4</td>
<td>4.6</td>
<td>−13.2</td>
</tr>
<tr>
<td>2b (M = Cu)</td>
<td>−42.5</td>
<td>17.5</td>
<td>6.1</td>
<td>18.9</td>
</tr>
<tr>
<td>2c (M = Ag)</td>
<td>−43.7</td>
<td>17.2</td>
<td>6.0</td>
<td>−20.5</td>
</tr>
</tbody>
</table>

Table S 14: Dispersion-corrected interaction energies (all energies in kcal mol\(^{-1}\)).

<table>
<thead>
<tr>
<th>Molecule</th>
<th>$\Delta E$(D3(BJ)(^{\text{ATM}})) (PBE0)</th>
<th>$\Delta E$(DFT-D4 v 0.1)(PBE0) (^a)</th>
<th>$\Delta E$(PBE0/QZ-f-g)</th>
<th>$\Delta E$(PBE0-D3/QZ-f-g)</th>
<th>$\Delta E$(PBE0-D4v0.1)/QZ-f-g)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1f (1:1) with Acceptor 2</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>2a (M = Au)</td>
<td>−31.8</td>
<td>−35.5</td>
<td>−0.3</td>
<td>−32.1</td>
<td>−35.8</td>
</tr>
<tr>
<td>2b (M = Cu)</td>
<td>−31.2</td>
<td>−34.9</td>
<td>−6.8</td>
<td>−38.0</td>
<td>−41.7</td>
</tr>
<tr>
<td>2c (M = Ag)</td>
<td>−31.3</td>
<td>−34.6</td>
<td>−8.3</td>
<td>−39.5</td>
<td>−42.9</td>
</tr>
</tbody>
</table>

\(^a\) D4 v0.1 is a preliminary DFTD4 in-house development version that takes into account the influence of the electronic structure on the dynamic polarizability and thus the dispersion energy.[43]

Table S 15 Atomic C6(AA) coefficients of the metal centers calculated using D3 and D4.

<table>
<thead>
<tr>
<th>Molecule</th>
<th>C6(AA)/(au · bohr(^6)) D3(PBE0)[ATM, BJ]</th>
<th>C6(AA)/(au · bohr(^6)) D4(PBE0)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1f (M = Au)</td>
<td>317.2</td>
<td>306.3</td>
</tr>
<tr>
<td>2a (M = Au)</td>
<td>317.2</td>
<td>303.7</td>
</tr>
<tr>
<td>2b (M = Cu)</td>
<td>175.0</td>
<td>160.6</td>
</tr>
<tr>
<td>2c (M = Ag)</td>
<td>268.6</td>
<td>244.6</td>
</tr>
</tbody>
</table>
Table S 16. D-A Free binding energies calculated with PBE0-D3/QZ and thermostatistical contributions obtained with GFN-xTB (all energies in kcal mol$^{-1}$).

<table>
<thead>
<tr>
<th>1f (1:1) with Acceptor 2</th>
<th>$\Delta E$(PBE0-D3/QZ-f-g)</th>
<th>$\Delta G_{RRHO}$ (GFN-xTB)</th>
<th>$\Delta \delta G_{solv}$</th>
<th>$\Delta G_a$ (1f•2)</th>
</tr>
</thead>
<tbody>
<tr>
<td>2a (M = Au)</td>
<td>−32.1</td>
<td>22.3</td>
<td>4.6</td>
<td>−5.2</td>
</tr>
<tr>
<td>2b (M = Cu)</td>
<td>−38.0</td>
<td>Could not be obtained</td>
<td>6.1</td>
<td>–</td>
</tr>
<tr>
<td>2c (M = Ag)</td>
<td>−39.5</td>
<td>18.4</td>
<td>6.0</td>
<td>−15.2</td>
</tr>
</tbody>
</table>

Table S 17. A-D-A Free binding energies calculated with PBE0-D3/QZ and thermostatistical contributions obtained with GFN-xTB (all energies in kcal mol$^{-1}$).

<table>
<thead>
<tr>
<th>1f • 2 with Acceptor to 2•1f•2</th>
<th>$\Delta E$(PBE0-D3/QZ-f-g)</th>
<th>$\Delta G_{RRHO}$ (GFN-xTB)</th>
<th>$\Delta \delta G_{solv}$</th>
<th>$\Delta G_a$ (2•1f•2)</th>
</tr>
</thead>
<tbody>
<tr>
<td>2a (M = Au)</td>
<td>−30.3</td>
<td>14.1</td>
<td>4.9</td>
<td>−11.4</td>
</tr>
<tr>
<td>2b (M = Cu)</td>
<td>−35.1</td>
<td>Could not be obtained</td>
<td>5.9</td>
<td>–</td>
</tr>
<tr>
<td>2c (M = Ag)</td>
<td>−36.4</td>
<td>11.0</td>
<td>4.9</td>
<td>−20.6</td>
</tr>
</tbody>
</table>

Figure S 54. Structures of stacked CTCs.
7.2.1 Qualitative evaluation of Charge-Transfer from the gold pyridinate complex 1f on a molecular extended graphite surface

To obtain a qualitative statement about the possible CT from 1f onto the HOPG, the complex 1f on a molecular extended graphite surface was optimized with PBE-D3/def2-SVP.\cite{note1} The CT was then calculated from the Natural Population Analysis, as well as from the total density and produced the same conclusion that only a charge of \(-0.00988\) is transferred onto the molecular graphene surface. In general CT interactions are ‘overestimated with GGAs’ therefore we qualitatively concluded that for the donor 1f CT is negligible small.

![Figure S 55: CTC 1f on molecular graphite surface.](image)

7.2.2 Association of 1f • 2a on a molecular extended graphite surface

The association of 1f with 2a on a molecular extended graphite surface in the gas phase was investigated. The structures were optimized with the global hybrid composite scheme PBEh-3c, and thermostatistical contributions were obtained with GFN-xTB.

For the complex on the graphite surface no charge transfer was found. A small dipole moment (0.13 debye) orthogonal to the graphite surface was found (for 1f on the surface), which indicates only a (very) small polarization.
Figure 56: Complex 1f • 2a on molecular graphite surface.

Listed below are the Cartesian coordinates of the optimized structures at the level of PBEh-3c.
1f-on molecular graphene surface:

<table>
<thead>
<tr>
<th>X</th>
<th>Y</th>
<th>Z</th>
<th>C</th>
<th>X</th>
<th>Y</th>
<th>Z</th>
</tr>
</thead>
<tbody>
<tr>
<td>11.3445368</td>
<td>-7.3819521</td>
<td>0.3530914</td>
<td>C</td>
<td>7.0845368</td>
<td>-9.8414621</td>
<td>0.3530914</td>
</tr>
<tr>
<td>12.0545368</td>
<td>-6.1521921</td>
<td>0.3530914</td>
<td>C</td>
<td>7.7945368</td>
<td>-8.6117021</td>
<td>0.3530914</td>
</tr>
<tr>
<td>11.3445368</td>
<td>-4.9224321</td>
<td>0.3530914</td>
<td>C</td>
<td>7.0845368</td>
<td>-7.3819521</td>
<td>0.3530914</td>
</tr>
<tr>
<td>12.0545368</td>
<td>-3.6926821</td>
<td>0.3530914</td>
<td>C</td>
<td>7.7945368</td>
<td>-6.1521921</td>
<td>0.3530914</td>
</tr>
<tr>
<td>11.3445368</td>
<td>-2.4629221</td>
<td>0.3530914</td>
<td>C</td>
<td>7.0845368</td>
<td>-4.9224321</td>
<td>0.3530914</td>
</tr>
<tr>
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8 STM measurements

Scanning tunnelling microscopy (STM) was performed under ambient conditions (room temperature) at the solution/solid interface, using 1-phenyloctane (PHO) as solvent and highly oriented pyrolytic graphite (HOPG) as substrate. The experimental setup consisted of an Agilent 5500 scanning probe microscope, that was placed on a Halcyonics actively isolated microscopy workstation. It was acoustically shielded with a home-built box. Scissor-cut Pt/Ir (80/20) tips were used and further modified after approach by applying short voltage pulses until the desired resolution was achieved. HOPG was obtained from Mikromasch and TipsNano in ZYB quality. All STM images (unless otherwise noted) were calibrated by subsequent immediate acquisition of an additional image at reduced bias voltage, therefore the atomic lattice of the HOPG surface was observed, which was used as a calibration grid. Data processing, also for image calibration, was performed using the SPIP 5 (Image Metrology) software package. (Supra-) molecular models were constructed using Spartan ’08 (Wavefunction, Inc.).

Three different sample preparation methods were used for performing the STM measurements presented here:

Method A (pure 1e): A solution of 1e in PHO was applied to HOPG at elevated temperature (80 °C), kept at this temperature for 20 s, and allowed to cool to rt before the STM measurements were performed with the tip immersed into the solution.

Method B (addition of 2c to a monolayer of 1e): 0.5 µL of a c1e = 3×10⁻⁶ M solution of 1e was dropped onto the HOPG substrate at elevated temperature (80 °C), kept at this temperature for 20 s, and allowed to cool to rt. Additional 0.5 µL of a 10⁻³ M or 10⁻⁴ M solution of 2c in PHO were added to the as-prepared monolayer, and the STM measurements were performed with the tip immersed into the solution. Consequently, the effective concentrations have to be corrected for the volume increase, and final concentrations are c₁e = 1.5×10⁻⁶ M and c₂c = 5×10⁻⁶ M (or 5×10⁻⁵ M).

Method C (co-adsorption of a mixture of 2c and 1e): 0.5 µL of a mixture of 1e and 2c in PHO was dropped onto HOPG substrate at elevated temperature (80 °C), kept at this temperature for 20 s, and allowed to cool to rt, and the STM measurements were performed with the tip immersed into the solution. Concentrations of 1e and 2c in the mixture are given.

All STM measurements were typically completed after 1 h after the aforementioned sample preparation procedures.

8.1 Concentration-dependent polymorphism

1e formed self-assembled monolayers at the interface of 1-phenyloctane (PHO) and highly oriented pyrolytic graphite (HOPG). At a (starting) concentration of 1×10⁻⁶ M to 3×10⁻⁶ M of
11e in the supernatant solution, a porous polymorph A is observed (overview STM images: Figure S 57a and b, high-resolution STM image: Figure S 57d and Figure 4 (Main Text)) to which a unit cell (containing two molecules) with lattice constants of \( a = b = (5.6\pm0.2) \) nm, \( \gamma(a,b) = (60\pm2)^{\circ} \) was indexed. All alkyl side chains of each molecule are aligned along the HOPG main axis directions, so that the unit cell vector \( a \) is oriented at an angle of \( \gamma(a,d_1) = (15\pm1)^{\circ} \) relative to the HOPG main axis direction \( d_1 \). Each of the \( C_{3v} \)-symmetric molecules (A) interacts with three adjacent molecules (B) by interdigitation of alkyl side chains in an ABAB packing scheme (cf. supramolecular and schematic models, Figure S 57e and f, respectively). A coordination number of \( CN = 3 \) (with respect to side chain interactions) results, and the overall packing can be viewed as a chiral honeycomb network.

With increasing concentration of 11e in the supernatant liquid phase, a more densely packed polymorph B was observed alongside with polymorph A (e.g., coexistence of both polymorphs A and B at \( c = 7\times10^{-6} \) M as seen in the overview image in Figure S 57c). The surface coverage of polymorph B gradually increases, and polymorph B is solely observed at a concentration as high as \( 1\times10^{-5} \) M (see high-resolution STM image in Figure S 57g). Two of the six alkyl side chains of each molecule of 11e point towards the solution phase, so that four alkyl side chains of each molecule are adsorbed on the HOPG surface and are aligned along two HOPG main axis directions so that hexagonal assemblies, formed from six molecules, are observed (schematic model in Figure S 57i), in which each two molecules interact via interdigitating alkyl side chains (with an ABAB packing scheme). These hexagonal assemblies pack densely to form a hexagonal pattern (Figure S 57h and i), to which a unit cell of \( a = b = (7.1\pm0.2) \) nm, \( \gamma(a,b) = (60\pm2)^{\circ} \) (containing six molecules) is indexed. The unit cell vector \( a \) is oriented at an angle of \( \gamma(a,d_1) = (19\pm2)^{\circ} \) relative to the HOPG main axis direction, \( d_1 \). As a result of the hierarchical packing of hexagonal units (each formed by six molecules), triples of bright spots, containing three neighboring backbones, are observed. The bright regions in the interior of some of the pores (indicated by arrow 1 in Figure S 57g) appeared blurred, which is attributed to additional, unspecifically adsorbed molecules. Even at this increased concentration, no evidence for the formation of multilayers, where molecules of 11e stack on top of each other, is observed in any of the images acquired.
Figure S 57. (a)–(c), (d), (g) STM images, (e), (h) supramolecular, and (f), (i) schematic models of self-assembled monolayers of 1e at the PHO/HOPG interface. (a)–(c) Overview STM images (100×100 nm², internal scanner calibration) of (a), (b) polymorph A (a: c = 1×10⁻⁶ M, b: c = 3×10⁻⁶ M, both: Vₛ = −0.8 V, lᵢ = 11 pA) and (c) polymorph A and B (c = 7×10⁻⁶ M, Vₛ = −0.8 V, lᵢ = 11 pA); (d) high-resolution STM image, (e) supramolecular model, and (f) schematic model of polymorph A (c: 20.0×20.0 nm², c = 1.5×10⁻⁶ M, Vₛ = −0.15 V, lᵢ = 24 pA), a = b = (5.6±0.2) nm, γ(a,b) = (60±2°), γ(a,d₁) = (15±1°); (g) high-resolution STM image, (h) supramolecular model, and (i) schematic model of polymorph B (g: 42.8×42.8 nm², c = 1×10⁻⁵ M, Vₛ = −0.8 V, lᵢ = 10 pA), a = b = (5.6±0.2) nm, γ(a,b) = (60±2°), γ(a,d₁) = (15±1°). The red and white (black) lines indicate unit cells and HOPG main axis directions, respectively.
8.2 Co-adsorption of 2c on 1e

We next investigated whether the concept of π-acid-base interactions could be applied to template the co-adsorption of 2c on a monolayer of 1e. The lower concentration range of $1\times10^{-6}$ M to $3\times10^{-6}$ M for 1e was chosen to make sure that all alkyl chains are aligned on the HOPG substrate.

Two different sample preparation methods, B and C, were used to investigate the co-adsorption of 2c on top of the backbones of 1e.

When 0.5 μL of a solution of 1e with $c_{1e} = 3\times10^{-6}$ M was dropped onto the HOPG substrate at elevated temperature (80 °C), kept at this temperature for 20 s, and allowed to cool to r.t., and 0.5 μL of a $10^{-3}$ M solution of 2c in PHO was added to the as-prepared monolayer (which was previously denoted as method B), the STM images shown in Figure S 58a–d were observed. Polymorph A of 1e is observed to cover the HOPG surface, while few additional bright features are observed which are marked by green circles. When a mixture of 1e ($c_{1e} = 1.5\times10^{-6}$ M) and 2c ($c_{2c} = 5\times10^{-4}$ M) was applied to the HOPG surface at r.t. (which was previously denoted as method C), the STM images shown in Figure S 58e–h were observed. The STM images again show the hexagonal packing of 1e in polymorph A, while parts of the HOPG surface remaining uncovered, and – again – additional bright features. In both cases, the bright features are clearly localized above the backbone regions of 1e (according to the lattice). We attribute these to the formation of supramolecular stacks, where 2c adsorbs on top of the backbone of 1e. Through its π-basicity, 1e acts as a secondary template for the adsorption of π-acidic 2c, dictated through the donor-acceptor interactions.
Figure S 58. STM images of self-assembled monolayers of 1e with coadsorbates of 2c (without and with markers), prepared by methods B and C. (a)–(d) Sample prepared by method B; 150x85 nm$^2$, $c_{1e} = 1.5 \times 10^{-6}$ M and $c_{2c} = 5 \times 10^{-4}$ M, $V_S = -0.8$ V, $I_t = 10$ pA; (e)–(h) sample prepared by method C; (e, f: 78x78 nm$^2$; g, h: 150x61 nm$^2$, $c_{1e} = 1.5 \times 10^{-6}$ M and $c_{2c} = 5 \times 10^{-4}$ M, $V_S = -0.9$ V, $I_t = 18$ pA. All image sizes are determined by internal scanner calibration.
In addition, we investigated whether 2c forms self-assembled monolayers under similar experimental conditions as used for the co-adsorption experiments, but without the template monolayer of 1e. Therefore, we prepared $10^{-3} - 10^{-6}$ M solutions of 2c in PHO, applied 0.5 µL of the solutions (at r.t.) to freshly cleaved HOPG surfaces (without further thermal annealing). Even under the highest concentrations ($10^{-3}$ M), no hints on any self-assembled monolayers or periodic patterns were observed by STM (Figure S 59).

![Typical STM image of the uncovered HOPG surface after applying a solution of 2c in PHO](image)

**Figure S 59.** Typical STM image of the uncovered HOPG surface after applying a solution of 2c in PHO ($221 \times 221$ nm$^2$ (internal scanner calibration), $c_{2c} = 10^{-3}$ M $V_s = -0.8$ V, $I_t = 20$ pA). No self-assembled monolayers are observed (natural step edges of the HOPG substrate are visible).

The results show that a template monolayer (such as 1e) is crucial for the adsorption of 2c (here).

### 8.3 Adsorption of 1e on Au(111)

We were interested to find out whether 1e forms self-assembled monolayers on Au(111). A substrate with several $10 \times 10$ nm$^2$ large terraces of Au(111) was obtained by flame-annealing commercially available glass slides covered with 250 nm gold with an 1 nm chromium adhesive layer (obtained from arrandee.com, Werther, Germany) with a butane flame according to literature procedures (see arrandee.com). After application of a $2.5 \times 10^{-6}$ M solution of 1e in PHO to the flame-annealed substrate at 80 °C, keeping at this temperature for 10 s and allowing the substrate to cool to r.t., we acquired the STM images shown in Figure S 60. Typical terraces of the Au(111) surface are seen, which are covered by medium bright features and bright features – the latter of which is attributed to the hexagonal packing of 1e similar to the porous polymorph A that was observed on HOPG.
Figure S 60. (a) STM image of 1e (c = 2.5×10⁻⁶ M) on a Au(111), 200×200 nm², internal scanner calibration, $V_S = -0.8$ V, $I_t = 5$ pA), (b) enlarged region (36×36 nm²) of (a).
9 References


[29] see also: Turbomole.com.


[38] ORCA – an ab initio, density functional and semiempirical program package, V. 4.0.0, F. Neese, MPI für Chemische Energiekonversion, Mülheim a. d. Ruhr (Germany), 2017.


