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Meta-substituted benzamide oligomers that complex mono-, di- and tricarboxylates: folding-induced selectivity and chirality

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Fig. S1 Partial <sup>1</sup>H NMR spectra of T2 at different concentrations in DMSO- $d_6$  at 25 °C.



Fig. S2 Partial NOESY spectrum (400 MHz) of the mixture of T1 (10 mM) + 10 (10 mM) in DMSO- $d_6$  at 25 °C (mixing time = 0.3 s).



**Fig. S3** Partial NOESY spectrum (400 MHz) of the mixture of **T3** (10 mM) + **10** (10 mM) in DMSO- $d_6$  at 25 °C (mixing time = 0.3 s).



**Fig. S4** Benesi-Hildebrand (BH) plots using <sup>1</sup>H NMR titration data of (a) **T1-10**, (b) **T2-10**, (c) **T3-10**, (d) **T3-11**, (e) **T3-12**, (f) **T3-13**, (g) **T3-14** and (h) **T3-15**. All fit to 1:1 binding model.



Fig. S5 CD spectrum of L-16 (20.0 mM) in CHCl<sub>3</sub> at 25 °C.



**Fig. S6** Partial NOESY spectrum (400 MHz) of **T1** (10 mM) in DMSO- $d_6$  at 25 °C (mixing time = 0.3 s).



Fig. S7 Partial NOESY spectrum (400 MHz) of T2 (10 mM) in DMSO- $d_6$  at 25 °C (mixing time = 0.3 s).



Fig. S8 Partial NOESY spectrum (400 MHz) of T3 (10 mM) in DMSO- $d_6$  at 25 °C (mixing time = 0.3 s).



Fig. S9 Partial COSY spectrum (400 MHz) of T1 (10 mM) in DMSO-d<sub>6</sub> at 25 °C.



Fig. S10 Partial COSY spectrum (400 MHz) of the mixture of T1 (10 mM) + 10 (10 mM) in DMSO- $d_6$  at 25 °C.



Fig. S11 Partial COSY spectrum (400 MHz) of T2 (10 mM) in DMSO-d<sub>6</sub> at 25 °C.



Fig. S12 Partial COSY spectrum (400 MHz) of the mixture of T2 (10 mM) + 10 (10 mM) in DMSO- $d_6$  at 25 °C.



Fig. S13 Partial COSY spectrum (400 MHz) of T3 (10 mM) in DMSO-d<sub>6</sub> at 25 °C.



Fig. S14 Partial COSY spectrum (400 MHz) of the mixture of T3 (10 mM) + 10 (10 mM) in DMSO- $d_6$  at 25 °C.



Fig. S15 <sup>1</sup>H NMR and <sup>13</sup>C NMR spectra of compound **3** in CDCl<sub>3</sub>.



Fig. S16 <sup>1</sup>H NMR and <sup>13</sup>C NMR spectra of compound T1 in DMSO- $d_6$ .



Fig. S17 <sup>1</sup>H NMR and <sup>13</sup>C NMR spectra of compound 6 in DMSO- $d_6$ .



Fig. S18 <sup>1</sup>H NMR and <sup>13</sup>C NMR spectra of compound 7 in acetone- $d_6$ .



Fig. S19 <sup>1</sup>H NMR and <sup>13</sup>C NMR spectra of compound T2 in DMSO- $d_6$ .



Fig. S20 <sup>1</sup>H NMR and <sup>13</sup>C NMR spectra of compound 8 in CDCl<sub>3</sub>.

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Fig. S21 <sup>1</sup>H NMR and <sup>13</sup>C NMR spectra of compound 9 in DMSO- $d_6$ .



Fig. S22 <sup>1</sup>H NMR and <sup>13</sup>C NMR spectra of compound T3 in DMSO- $d_6$ .

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Fig. S23 <sup>1</sup>H NMR spectrum of compound 11 in DMSO- $d_6$ .



Fig. S24 <sup>1</sup>H NMR spectrum of compound 12 in DMSO- $d_6$ .



Fig. S25 <sup>1</sup>H NMR spectrum of compound 13 in DMSO- $d_6$ .

![](_page_18_Figure_3.jpeg)

Fig. S26 <sup>1</sup>H NMR spectra of compound 14 in DMSO- $d_6$ .

![](_page_19_Figure_1.jpeg)

Fig. S27 <sup>1</sup>H NMR spectrum of compound 15 in DMSO- $d_6$ .

![](_page_19_Figure_3.jpeg)

Fig. S28 <sup>1</sup>H NMR spectrum of compound L-16 in DMSO- $d_6$ .