

PZT

Subjects: [Chemistry](#), [Organic](#)

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This designed bridging ligand may offer the benefits of polydentate ligands and minimisation of isomeric possibilities.

The tris(pyrazolyl)methane fragment is of interest, because the coordination chemistry of such systems has also been studied. It, too, is easy to prepare, by reaction of pyrazole with chloroform, and to derivatise, for example by reaction with formaldehyde, to give a molecule, 2,2,2-tris(1*H*-pyrazol-1-yl)ethanol, that can be readily joined to the other binding site, a terpyridine containing synthon.

terpyridine

pyrazolyl

ditopic bridging ligand

4'-(4-(2,2,2-tris(1*H*-pyrazol-1-ido)ethoxymethyl)phenyl)-2,2':6',2''-terpyridine (PZT)

Ramin Zibaseresht^{a,b*}

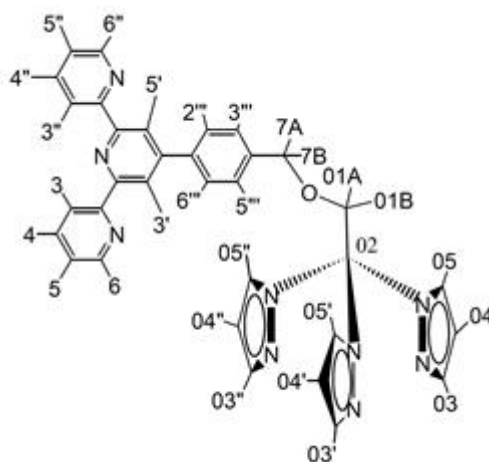
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The ligand synthesis is relatively straightforward. Syntheses of suitable precursor molecules for both ends of the ditopic ligand are available in the literature. 4'-(4-(bromomethyl)phenyl)-2,2':6',2''-terpyridine is readily prepared by radical bromination of the related alkane, and this was reacted with 2,2,2-tris(1*H*-pyrazol-1-yl)ethanol, in the presence of sodium hydride.

The solution of 2,2,2-tris(1*H*-pyrazol-1-yl)ethanol (tpe) (0.368 g, 1.505 mmol) and 4'-(*p*-bromomethylphenyl)-2,2':6',2''-terpyridine (btp) (0.605 g, 1.505 mmol) in dry THF (30 mL) was added dropwise to a suspension of NaH (0.15 g) in dry THF (20 mL) under argon over a period of 3 h. The mixture was stirred at reflux under argon for 24 h and then allowed to cool at room temperature. To this yellow solution was added enough water dropwise carefully to consume the excess NaH. The mixture was extracted from diethyl ether (4 × 50 mL), and the combined organic extracts were washed with 50 mL saturated NaHCO₃ solution, then with 50 mL saturated NaCl solution, and finally with 50 mL water. To the aqueous solution was added diethyl ether. The organic layer was extracted. After drying the combined extracts with anhydrous MgSO₄, filtered and the solvent was removed under reduced pressure to yield a pale yellow powder; mp 183-184°. Purification was achieved by recrystallisation of the crude material from CH₂Cl₂-hexanes (1:5) to afford (0.7190 g, 84%) as a pale yellow microcrystalline material. Crystals suitable for X-ray diffraction were obtained by layering a dichloromethane solution of the ligand with hexanes. ¹H NMR (500 MHz; solvent CDCl₃): δ 8.734 (2H, d, H₆, H₆''), 8.727 (2H, s, H₃', H₅'), 8.67 (2H, d, H₃, H₃''), 7.89 (2H, m, H₄, H₄''), 7.86 (2H, d, H₂'', H₆''), 7.68 (3H, d, H₀₃, H₀₃', H₀₃''), 7.46 (3H, d, H₀₅, H₀₅', H₀₅''), 7.36 (2H, m, H₅, H₅''), 7.30 (2H, d, H₃'', H₅''), 6.36 (3H, m, H₀₄, H₀₄', H₀₄''), 5.20 (2H, s, H₀₁), 4.61 (2H, s, H₇). ¹³C NMR (75 MHz; solvent CDCl₃): δ 156.12, 155.86, 149.81 (C₃, C₃''), 149.06, 141.35 (C₀₃, C₀₃', C₀₃''), 138.07, 138.04 (C₄, C₄''), 136.94, 130.90 (C₀₅, C₀₅', C₀₅''), 128.13 (C₃'', C₅''), 127.38 (C₂'', C₆''), 123.86 (C₅, C₅''), 121.38 (C₆, C₆''), 118.81 (C₃', C₅'), 106.55 (C₀₄, C₀₄', C₀₄''), 89.84 (C₀₂), 73.80 (C₇), 73.63 (C₀₁). ¹H NMR (500 MHz; solvent dms-*d*₆, see Figure above for labeling): δ 8.86 (2H, d, H₃, H₃''), 8.80 (2H, s, H₃', H₅'), 8.77 (2H, d, H₆, H₆''), 8.14 (2H, dd,

H₄, H_{4''}), 7.99 (2H, d, H_{2'''}, H_{6'''}), 7.79 (3H, d, H₀₃, H_{03'}, H_{03''}), 7.63 (5H, m, H₀₅, H_{05'}, H_{05''}, H₅, H_{5'}), 7.48 (2H, d, H_{3'''}, H_{5'''}), 6.53 (3H, m, H₀₄, H_{04'}, H_{04''}), 5.17 (2H, s, H₀₁), 4.74 (2H, s, H₇). ¹³C NMR (75 MHz; solvent dms-*d*₆) δ 155.86, 155.07, 149.53 (C₃, C_{3''}), 149.33, 141.07 (C₀₃, C_{03'}, C_{03''}), 139.00, 137.71 (C₄, C_{4''}), 136.99, 131.23 (C₀₅, C_{05'}, C_{05''}), 128.56 (C_{3'''}, C_{5'''}), 127.07 (C_{2'''}, C_{6'''}), 124.77 (C₅, C_{5''}), 121.15 (C₆, C_{6''}), 118.07 (C_{3'}, C_{5'}), 106.65 (C₀₄, C_{04'}, C_{04''}), 89.42 (C₀₂), 72.80 (C₀₁), 72.46 (C₇). IR (KBr): /cm⁻¹ = 3140 m, 3055 m, 3017 m, 2939 m, 2885 w, 1983 w, 1929 w, 1790 w, 1728 w, 1697 w, 1582 m, 1520 m, 1466 m, 1389 s sh, 1312 s, 1258 m, 1196 m, 1126 s, 1096 s, 1042 w, 988 m, 949 w, 872 s sh, 895 m, 764 s, 656 m, 617 m, 525 m sh. ESI-MS: *m/z* 566.2931 ([M+H]⁺, 100%). Anal. Calc. for C₃₃H₂₇N₉O.0.5H₂O (574.65): C 68.98, H 4.91, N 21.94%; found: C 69.32, H 4.79, N 21.77%.

Crystals suitable for X-ray diffraction were obtained by layering a dichloromethane solution of the ligand with hexanes. The ligand was crystallised in orthorhombic *Pbca* space group ($R_1 = 0.06$). As shown in Figure 4.22, the three pyridyl rings are approximately coplanar, making an interannular angle with the central pyridine ring of 11.5°, with a *transoid* arrangement about each interannular C-C bond. This is in accord with the *transoid* configuration observed in the crystal structures of 2,2'-bipyridine, 2,2':6',2'':6'',2'''-quaterpyridine and 4'-phenyl-2,2':6',2''-terpyridine. The C-C and C-N distances within the phenyl and the pyridine rings are normal (average 1.307 and 1.351 Å, respectively). The interannular C-C bonds also closely resemble those seen in 2,2'-bipyridine, 2,2':6',2'':6'',2'''-quaterpyridine, and 4'-phenyl-2,2':6',2''-terpyridine with lengths of 1.498 Å.

The phenyl ring is not coplanar with the terpyridyl fragment, but is twisted about the interannular bond such that the plane of the phenyl ring makes an angle of 24.5° with the plane of the central pyridine ring. This represents a compromise between a coplanar arrangement in which π -conjugation and non-bonded H-H contacts between the *ortho* protons of the phenyl ring and H₂ H₄ are maximized. This angle is larger than that of 4'-phenyl-2,2':6',2''-terpyridine structure (10.9°). One of the pyrazole rings of the pyrazolyl site is approximately in the plane of C₇-O-C₀₁ linker with a slight twist of 10.6°. The angles between the remaining pyrazole rings and the plane of the etheric chain linker are 76.3° and 82.1°, respectively. The free nitrogen atoms of the pyrazole rings are all pointed out from each other in such a way that the interactions between the hydrogen atoms of the pyrazolyl rings and the free nitrogen atoms at the adjacent rings are small.

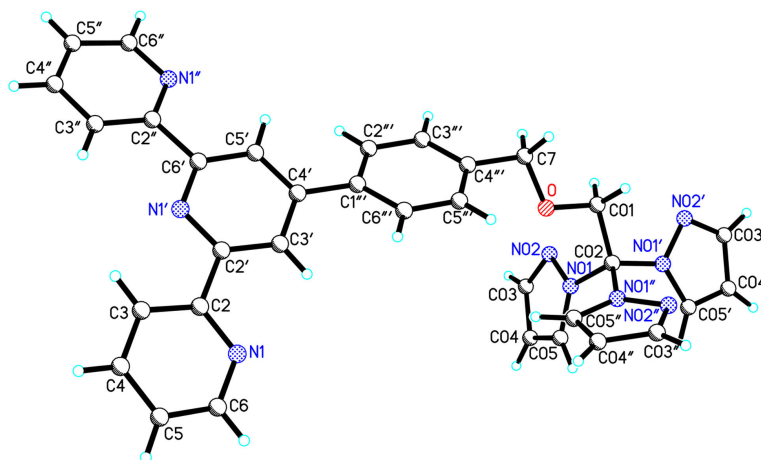


Figure 1. Molecular structure of ligand pzt. Selected bond lengths [Å] and angles [°]: N(1)-C(6) 1.340(4), N(1)-C(2) 1.358(4), N(1')-C(6') 1.348(4), N(1')-C(2') 1.361(4), N(1'')-C(2'') 1.338(4), N(1'')-C(6'') 1.358(4), N(01)-N(02) 1.369(4), N(01)-C(05) 1.392(4), N(01)-C(02) 1.490(4), N(02)-C(03) 1.349(5), N(01')-N(02') 1.354(4), N(01')-C(05') 1.370(4), N(01')-C(02) 1.447(4), N(01'')-C(05'') 1.369(4), N(01'')-N(02'') 1.379(4), N(01'')-C(02) 1.457(4), N(02'')-C(03'') 1.320(4), O-C(01) 1.414(4), O-C(7) 1.415(4), C(01)-O-C(7) 113.0(2), O-C(7)-C(4'') 107.6(3), O-C(01)-C(02) 105.5(3).

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