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**Emergence of superconductivity, valence bond order and Mott insulators in Pd[(dmit)<sub>2</sub>] based organic salts<sup>1</sup>**

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The EtMe<sub>3</sub>P and EtMe<sub>3</sub>Sb nearly triangular organic salts are distinguished from most other Pd[(dmit)<sub>2</sub>] based salts, as they display valence bond and no long range order, respectively. Under pressure, a superconducting phase is revealed in EtMe<sub>3</sub>P near the boundary of valence bond order. We use slave-rotor theory with an enlarged unit cell to study competition between uniform and broken translational symmetry states, offering a theoretical framework capturing the superconducting, valence bond order, spin liquid, and metallic phases on an isotropic triangular lattice. Our finite temperature phase diagram manifests a remarkable resemblance to the phase diagram of the EtMe<sub>3</sub>P salt, where the re-entrant transitions of the type insulator-metal-insulator can be explained by an entropy difference between metal and the U(1) spin liquid. We find that the superconducting pairing symmetry is  $d \pm id$ , and predict different temperature dependences of the specific heat between the spin liquid and metal.

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