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A molecular view of  $TiOPc - C_{70}interface$  formation<sup>1</sup> YINYING WEI, University of Maryland, STEVEN ROBEY, National Institute of Standards and Technology, JANICE REUTT-ROBEY, University of Maryland — A key strategy for the improvement of organic electronic devices involves the optimization of chemical morphology for efficient charge separation. Fundamental studies of chemical morphology - electronic property relations, particularly along crucial domain boundaries, are needed to realize these goals. We present STM/STS studies of TiOPc:  $C_{70}$  films, prepared by vapor deposition on Ag (111). Sequential deposition of  $C_{70}$  onto the ordered honeycomb TiOPc phase generates extended domains of a co-crystalline  $C_{70(1)}$ TiOPc<sub>(2)</sub>monolayer phase, characterized as a hexagonal network (2.1 nm nearest-neighbor  $C_{70}$  packing). Subsequent deposition of  $C_{70}$  onto this network proceeds in a layer-by-layer growth process, generated molecularly abrupt interfaces. The transport gap of each interface is measured by STS, and correlated to the  $C_{70}$  packing density. The orderly structure evolution of the  $C_{70}$ -TiOPc is contrasted to that of the previously reported  $C_{60}$ -TiOPc and the implications for electronic transport are discussed.

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