## Radical polymers as interfacial layers in inverted hybrid perovskite solar cells

**Luyao Zheng**<sup>2</sup>, Iz40@zips.uakron.edu, Sanjoy Mukherjee<sup>3</sup>, Kai Wang<sup>2</sup>, Martha E. Hay<sup>3</sup>, Bryan W. Boudouris<sup>1</sup>, Xiong Gong<sup>2</sup>. (1) 2148 Forney Hall of Chem Eng, Purdue University, West Lafayette, Indiana, United States (2) Department of Polymer Engineering, University of Akron, Akron, Ohio, United States (3) Charles D. Davidson School of Chemical Engineering, Purdue University, West Lafayette, Indiana, United States

We report high performance hybrid perovskite solar cells (PSCs) through implementation of a radical polymer-based copolymer that contains a second moiety capable of undergoing crosslinking through simple exposure to ultraviolet (UV) light, poly (2,2,6,6-tetramethylpiperidinyloxy-4-yl methacrylate)-co-(4-benzoylphenyl methacrylate) (PTMA-BP). The PTMA-BP thin film engineered the surface of a poly(3,4-ethylenedioxythiophene):polystyrene sulfonate (PEDOT:PSS) hole extraction layer (HEL). Systematic investigations indicate that PTMA-BP can induce better band alignment between PEDOT:PSS and perovskite hybrids, reduce interfacial charge carrier recombination, and improve crystallization of perovskite hybrids that are cast on top of the HEL. As a result, the stable PSCs incorporated with PTMA-BP exhibit a 15% power conversion efficiency, which is more than a 15% enhancement compared to cells that lacked the PTMA-BP interfacial modifying layer.

