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During the last decades, polymers with highly extended p-electrons have attracted much attention because of their good conductivity and easy in band-gap/colour-tuning via structural control. PEDOT found several applications including antistatic coatings for photographic films, hole transport layers for light-emitting diodes, photovoltaic devices, organic thin film transistors, and sensors. Moreover, PEDOT various bioengineering applications such as biosensors, drug-releasing process and bioactuators due to its low structural defects, high aqueous stability and biocompatibility were reported. The nano-hybrid/conducting polymer composites have become an area of research for scientists during the past decade. Among the used nanomaterials to prepare the hybrid polymer composites, TiO₂ is one of the most investigated oxide semiconducting materials due to low cost, non-toxicity, and very good stability. TiO₂ finds a wide range of applications such as environmental cleaning, photocatalysts, photosplitting water, dye-sensitized solar cells, photochromic devices and gas sensing.

In this research, PEDOT-TiO₂ composite films were prepared by a simple mechanical mixture of TiO₂ and PEDOT:PSS under mechanical stirring, which was demonstrated to deliver an effectively combined network of both high electrical conductivity and superior electrocatalytic activity. The composite films have been explored as an alternative for the counter electrodes of dye-sensitized solar cells and scaffold of SAW sensor. It was manifested that plasma treated PEDOT-TiO₂ composite films displayed excellent performance comparable to Pt counter electrode in DSSCs due to the combined network endowing more favorable and efficient interfacial active sites. Consequently, plasma treated PEDOT-TiO₂ composite films exhibit a J_{sc} of 10.93 mA/cm², a V_{oc} of 0.74 V and a fill factor (FF) of 0.67 with an overall conversion efficiency of 5.84%. This result showed increasing of 15% electrocatalytic activity using plasma treated PEDOT-TiO₂ composite counter electrode.