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During the last few decades, the increase of infections by toxic pathogens/bio films leading to hospital-acquired infections (HAI) has motivated work in the area [1-2]. More advanced antibacterial films presenting uniform distribution, high adhesion to exible non-thermal resistant substrates, mechanical resistance and faster bacterial/bio film inactivation under light or in the dark are needed due to health concerns [1]. TiO_2 have been used under light >387 nm generating highly oxidative radicals as bactericide films for many years [2]. However, its restricted absorption of solar/visible light and slow bacterial inactivation kinetics has motivated workers to dope TiO_2 Cu or Ag to shift the absorption of the films to the visible region. This doping also precludes recombination of the photo-generated charges. Stable, adhesive uniform films of TiO_2 inactivated bacteria within 40 min [2]. But TiO_2/Cu (Cu 0.1%) films led to bacterial inactivation < 10 min under actinic light ($4\text{mW}/\text{cm}^2$) [3-4]. Next, the sputtered Cu for 5-10s (0.01% by weight/ppb levels) ZrO_2 layers on polyester [5] accelerated the kinetics by a factor of 3 with respect to films where the Cu was absent. The Cu intra-gap states seem to: a) accelerate the indirect transitions in the TiO_2

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