

Supporting Information

Photodynamic Antimicrobial Textiles for Potential Combat and Clinical Applications

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1. Scanning Electron Microscopy (SEM) imagery of PVA-SbQ/RB-coated hemp

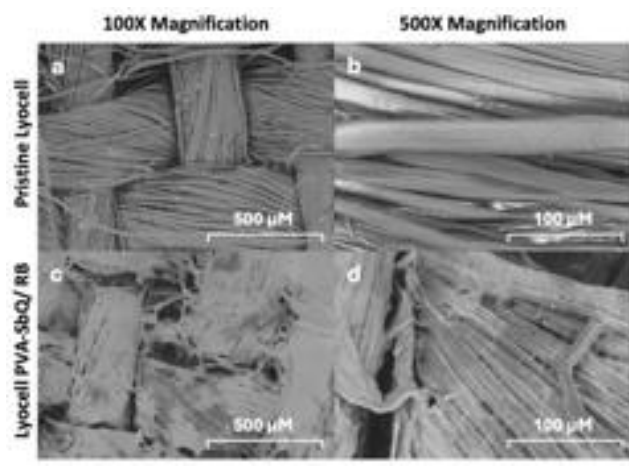


Fig S1 Representative scanning electron microscopy (SEM) images of hemp materials, comparing untreated (pristine; a, b) and PVA-SbQ/RB-coated samples (c, d). Images captured at 100× (left) and 500× (right) magnification illustrate surface morphology and effects of coating.

2. Energy Dispersive X-ray Spectroscopy imagery of PVA-SbQ/RB-coated hemp

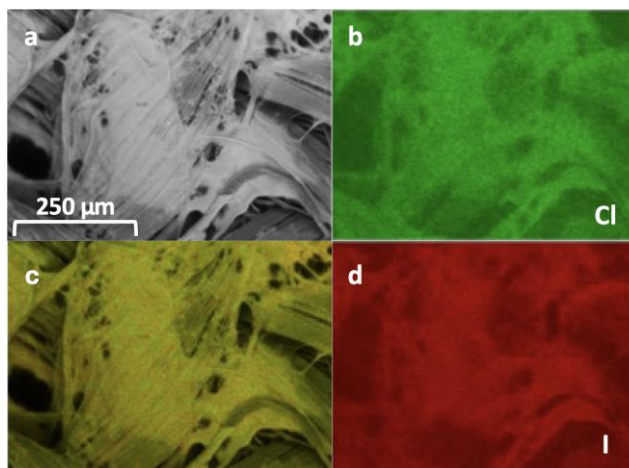


Fig S2 Scanning electron microscopy image (a) with energy dispersive spectroscopy elemental mapping of (b) Cl and (d) I for PVA-SbQ/RB hemp at 200× magnification. Quadrant (c) portrays the EDS maps overlaying the I and Cl signals.

3. Energy Dispersive X-ray Spectroscopy spectra of pristine, PVA-SbQ-coated, and PVA-SbQ/RB-coated lyocell/polyester and hemp

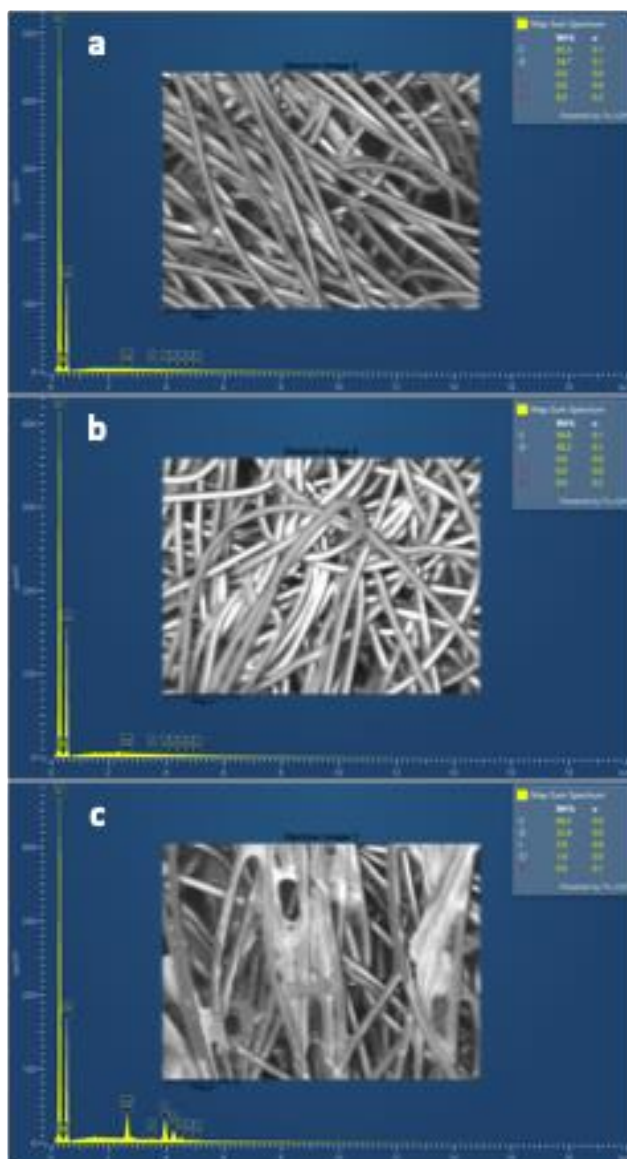


Fig S3 SEM micrographs with corresponding EDS spectra of (a) pristine lyocell/polyester, (b) PVA-SbQ-coated lyocell/polyester, and (c) PVA-SbQ/RB-coated lyocell/polyester.

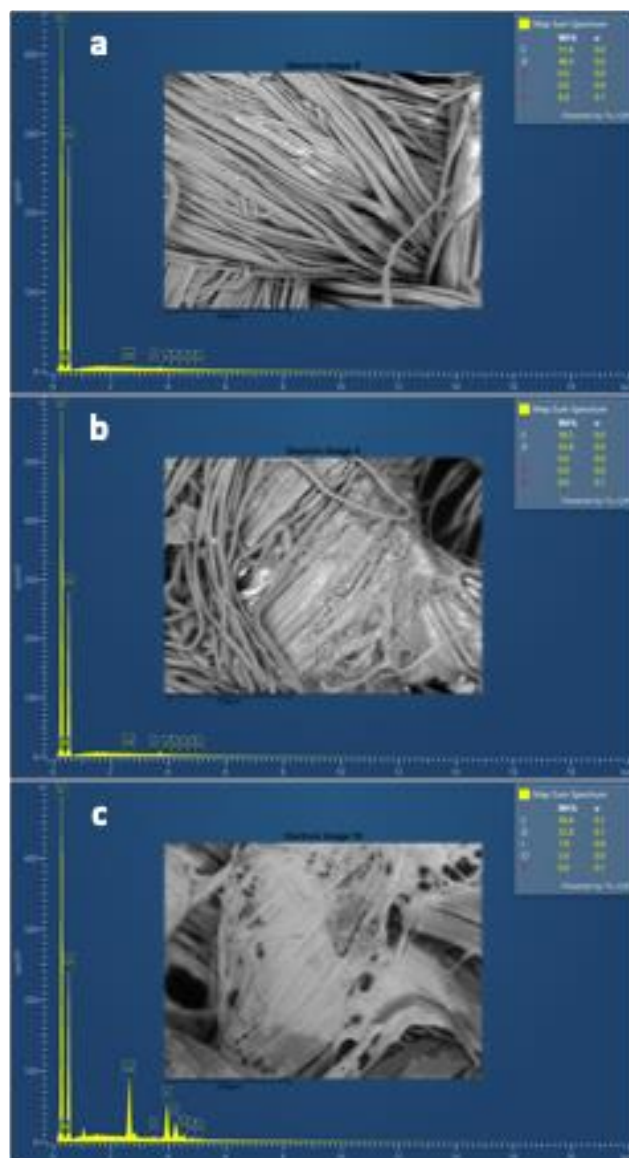


Fig S4 SEM micrographs with corresponding EDS spectra of (a) pristine hemp, (b) PVA-SbQ-coated hemp, and (c) PVA-SbQ/RB-coated hemp.

4. Time-of-Flight Secondary Ion Mass Spectrometry (TOF-SIMS) for PVA-SbQ/RB hemp.

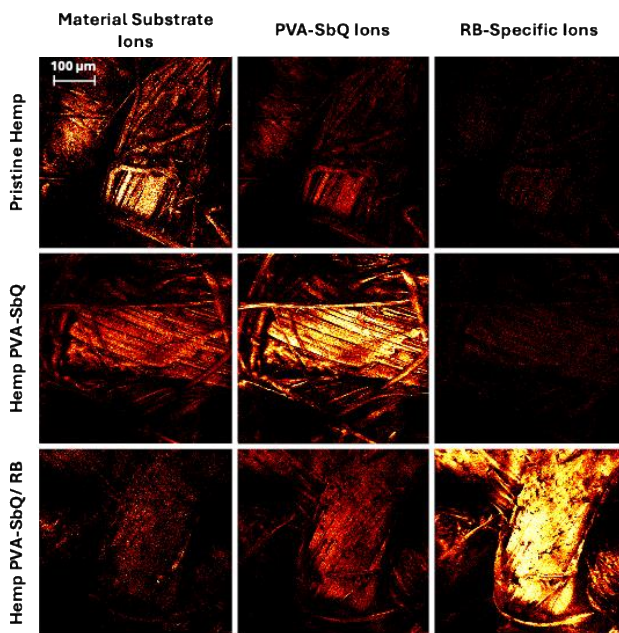


Fig S5 Heat maps generated via ToF-SIMS depicting the characteristic ions of pristine hemp (top row), PVA-SbQ-coated hemp (middle row), and PVA-SbQ/RB-coated hemp (bottom row).

5. Thermogravimetric Analysis (TGA) of hemp material.

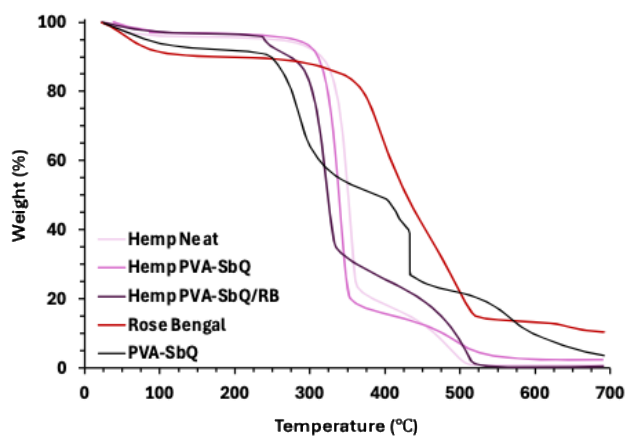


Fig S6 Thermogravimetric analysis (TGA) of pure Rose Bengal, neat PVA-SbQ, PVA-SbQ-coated, and PVA-SbQ/RB-coated hemp.

6. Determination of singlet oxygen production of hemp materials.

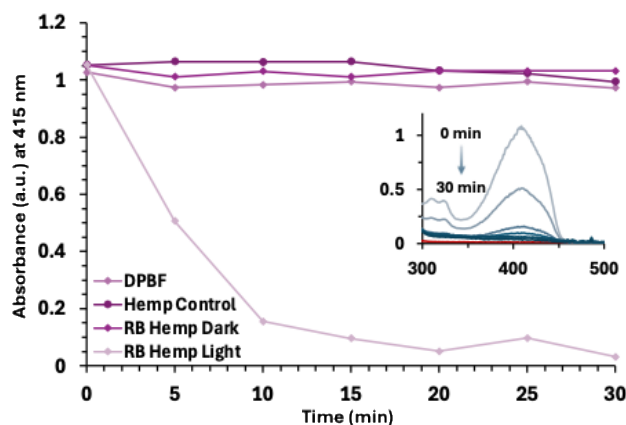


Fig S7 Photooxidation studies of DPBF by PVA-SbQ/RB-coated hemp. Each panel depicts the 415 nm absorbance of DPBF (0.05 mM) undergoing photooxidation upon illumination with visible light (532 nm, 68 ± 2 mW/cm²) in the absence of material (DPBF only), in the presence of pristine material (control), and in the presence of the PVA-SbQ/RB-coated material, as well as the latter in the absence of illumination (dark control). The insets depict time-dependent spectroscopic monitoring of DPBF photodegradation in the presence of PVA-SbQ/RB-coated materials illuminated with the aforementioned laser light source.

7. Antimicrobial Photodynamic Inactivation (aPDI) studies of hemp material.

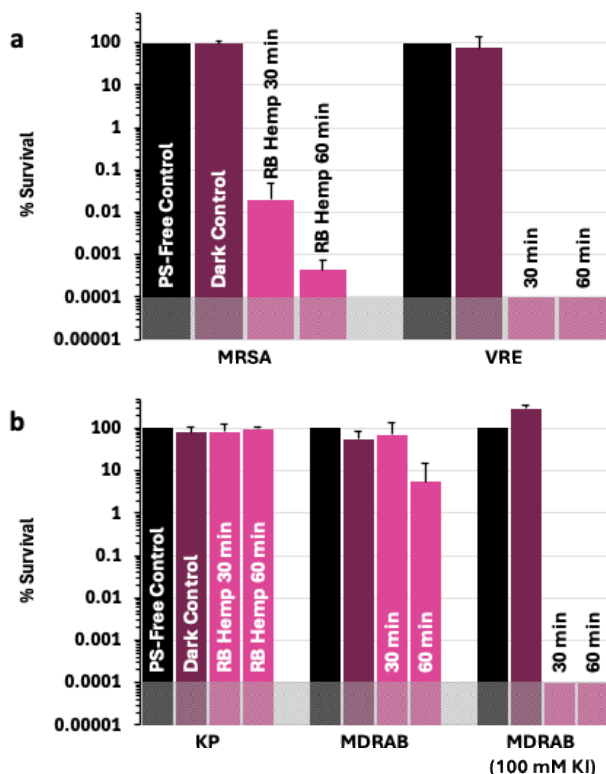


Fig S8 Antibacterial photodynamic inactivation (aPDI) studies against (a) Gram-positive MRSA and VRE, and (b) Gram-negative KP and MDRAB. The PVA-SbQ/RB-coated hemp dark control (dark magenta) and illumination studies (magenta) are shown as bacterial survival percentages when compared to the material-free (bacteria only) dark control (black). Where indicated, 100 mM potassium iodide (KI) was employed as a potentiator. All light studies were conducted under uniform illumination conditions (400–700 nm, 42 ± 5 mW/cm²) for 30 or 60 minutes. The limit of detection was 0.0001% survival, indicated by the shaded gray region.