Supplementary Information

Microwave-enhanced antibacterial activity of polydopamine-silver hybrid

nanoparticles

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Fig S1. Impact of microwaves on PDA-Ag NPs. (a) UV-Vis spectra of PDA-Ag NPs after exposure to microwaves for 0 - 60 minutes. (b) Corresponding DLS size and zeta potential data of the samples are shown on the right. TEM images taken from sample at different intervals are shown in c-g.



Fig S2. Minimum bactericidal concentration images. Pictures of agar plates (1 of 3 biological

repeats) used to determine the MBC of PDA-Ag NPs for E.coli and B.subtilis.



Fig S3 Temperature measurements of test solutions during experiments. (a) Temperature measurements during experiments (0.1 mg/mL PDA-Ag + 10⁸ bacteria) to ensure temperature remains below thermal inactivation point. **(b)** Temperature measurements with 0.2 mg/mL PDA-Ag/PDA and PBS samples.



Fig S4. Bacterial cell viability and relative fluorescence of ROS indicators (showing bacterial ROS production). (a) Viability of *E.coli* and *B.subtilis* under microwave (MW) exposure, in presence of PDA-Ag NPs and MWs, and PDA-Ag NPs only. (b) Relative fluorescence of DCF (intracellular ROS indicator) for *E.coli* and *B.subtilis* in presence of microwaves (MW), PDA-Ag NPs and MW, and PDA-Ag NP only.



Fig S5. Live/dead assay images using SYTO 9 dye and propidium iodide staining. Fluorescent images showing *E.coli* and *B.subtilis*; control, microwave (MW) exposure, PDA-Ag and MW exposure or PDA-Ag only. SYTO 9 dye shows live cells in green, propidium iodide shows dead cells in red.



Fig S6. RNO assay used for the detection of singlet oxygen and hydroxyl. Mechanisms of (a) singlet

oxygen and **(b)** hydroxyl detection using RNO.



Fig S7. Amplex Red assay used for the detection of hydrogen peroxide and superoxide. Mechanism of hydrogen peroxide (H₂O₂) and superoxide sensing through the colorimetric detection of resorufin at 587 nm (with and without superoxide dismutase).