

• **One Control Set** of 10 culture plates
Innoculated with serially diluted, calibrated *E. Coli*
NOT IRRADIATED with UVC

• To compare 2 IDENTICAL Sets of PLATES where a $2 \times 2 \text{ cm}^2$ area is IRRADIATED with UVC
→ ensures **ACCURACY**

• REPEAT the experiment above TWICE → ensures **REPRODUCIBILITY**



Control + TWO IDENTICAL PLATES

No UVC #1 No UVC Irradiation	UVC1 #1 3 min Irradiation with 80 $\mu\text{W}/\text{cm}^2$	UVC2 #1
Control Set #1	IDENTICAL LactoB. Cultures, Exposed to the same DOSE	

0 **14.4 mJ/cm²**

No UVC #2 No UVC Irradiation	UVC1 #2 3 min Irradiation with 80 $\mu\text{W}/\text{cm}^2$	UVC2 #2
Control Set #2	IDENTICAL LactoB. Cultures, Exposed to the same DOSE	

0 **14.4 mJ/cm²**

STEP #1 DESIGN UVC LED IRRADIATION

1. GEOMETRY
→ DOME SHAPE
+ ALUMINUM SHIELDING FOR SAFETY



3. FOUR UVC LED's COVER 1/4 OF THE PLATE



4. WRAP DOME IN AL IN STERILE BSL II CLEAN ROOM WITH UVC GOGGLES, GLOVES, PPE



2. CULTURE PLATES INOCULATED WITH LACTO_B, 10 PLATES PER SET USING 10 DILUTIONS $1.0 - 10^{-9}$



5. PLACE CULTURE PLATE UNDER DOME

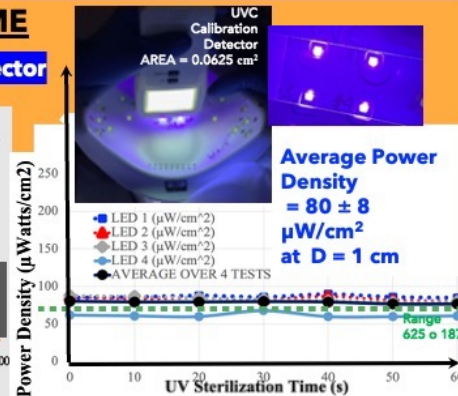
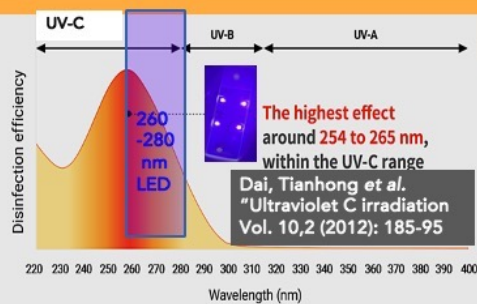


6. IRRADIATE FOR 3 MINUTES

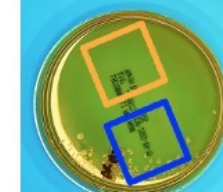
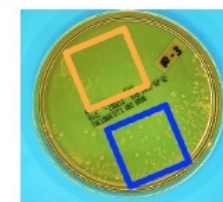
STEP #2 CALIBRATE UVC POWER

Four LEDs AFFIXED in A DOME

→ Measured Power Density via GaN Detector



#6 RESULTS OF ANALYSIS A. Two Examples of Results for Set #2, Dilutions = 10^{-3} and 10^{-4} after UVC irradiation of $2 \times 2 \text{ cm}^2$



Both culture plates (a) with 10^{-3} and (b) with 10^{-4} dilution

→ show that in the upper half

- where a $2 \times 2 \text{ cm}^2$ region was UVC irradiated,
- ALL bacteria were eradicated

In addition, the area AROUND the $2 \times 2 \text{ cm}^2$ show
→ UVC rays escaping from the $2 \times 2 \text{ cm}^2$ area eradicate LactoB. A. in an additional area

→ Compare CFU counts from UVC area with control set for 10^{-4}

