

# **Quick Reference Guide**

# **Deposit and Surface Analysis Test Kit**

Product #: DSA-25 / DSA-100

NOTE: Please refer to Test Kit Instructions during first product use and for additional details including legal statements.



# Step 1 - UltraCheck™ 1 Calibration

Perform one UltraCheck 1 calibration per day or per each set of samples analyzed.

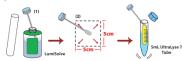


NOTE: If RLU<sub>ATP1</sub> ≤ 5,000 using a PhotonMaster or Lumitester C-110, rehydrate a new bottle of Luminase for maximum sensitivity.

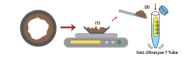
OR

# Step 2 - Sample Preparation → Select one of the following options:

**Option A - SURFACE SWAB** 

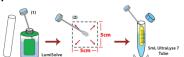


Option B: MEASURED DEPOSIT



**Option C: BIOFILM COLLECTOR** 





# Step 3 – Total ATP (tATP<sup>TM</sup>) Analysis $\rightarrow$ Then perform the following steps:

#### 3.1 - INCUBATION

Allow time for complete extraction.

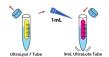






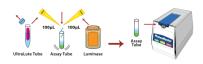
3.2 - DILUTION

Dilute out interferences.



#### 3.3 - ASSAY

Measure ATP concentration.



NOTE: If RLU<sub>tATP</sub> ≤ 10 using a PhotonMaster or Lumitester C-110, you are below the low- detection limit.

NOTE: If RLU<sub>tATP</sub> ≤ 50 using a PhotonMaster or Lumitester C-110, consider accounting for background (RLUbg). See Test Kit Instructions for guidance.

# **Calculations** $\rightarrow$ **Carry out calculations that correspond to the selected preparation method:**

## A - Surface Swab (Default $A_{sample} = 25 \text{cm}^2$ ):

$$ATP\left(pg\ ATP/cm^2\right) = \frac{RLU_{tATP}}{RLU_{ATP1}} \times \frac{50,000\left(pg\ ATP\right)}{A_{Sample}\left(cm^2\right)}$$

$$tATP\left(\frac{ME}{cm^2}\right) = tATP\left(\frac{pg\ ATP}{cm^2}\right) \times \frac{1\ ME}{0.001\ pg\ ATP} \\ tATP\left(\frac{ME}{g}\right) = tATP\left(\frac{pg\ ATP}{g}\right) \times \frac{1\ ME}{0.001\ pg\ ATP}$$

## B - Measured Deposit (Default m<sub>sample</sub> = 1g ):

$$tATP\left(pg\ ATP/cm^2\right) \ = \ \frac{RLU_{tATP}}{RLU_{ATP1}} \ \times \ \frac{50,000\left(pg\ ATP\right)}{A_{Sample}\left(cm^2\right)} \quad \text{OR} \qquad tATP\left(pg\ ATP/g\right) \ = \ \frac{RLU_{tATP}}{RLU_{ATP1}} \ \times \ \frac{50,000\left(pg\ ATP\right)}{m_{Sample}\left(g\right)} \quad \text{OR}$$

$$tATP\left(\frac{ME}{g}\right) = tATP\left(\frac{pg\ ATP}{g}\right) \times \frac{1\ ME}{0.001\ pg\ ATP}$$

### C - Biofilm Collector:

$$tATP\left(pg\ ATP\ /\ device\right) = rac{RLU_{IATP}}{RLU_{ATP1}} imes rac{50,000\left(pg\ ATP
ight)}{1\ device}$$

$$tATP\left(\frac{ME}{device}\right) = cATP\left(\frac{pg\ ATP}{device}\right) \times \frac{1\ ME}{0.001\ pg\ ATP}$$

NOTE: 1 ME (Microbial Equivalent) assumes 0.001 pg (1fg) ATP per cell

#### Interpretations Guidelines

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Application	Good Control (pg cATP/mL)	Preventative Action (pg cATP/mL)	Corrective Action (pg cATP/mL)
Surface, Deposits, Coupons*	< 10x	10x to 100x	> 100x

\*Guidelines are provided as a ratio of ATP on your surface/deposit/collector to bulk fluid ATP.

**NOTE: Interpretation Guidelines provided for general** guidance. For best results, establish your own baseline and control levels.