

## Standard Laboratory Operating Procedure #200 Effectiveness of Personal Hygiene Products

Laboratory: Biotechnology SOP prepared by: R. Sanders and B. Wolfe Location: Lab Station Last Revision: 21 July 2013

**General:** The purpose of this protocol is to investigate the effectiveness of various types of hand cleaners that may have antimicrobial action.

Safety: Safety Glasses, Caution with the alcohol burners

## Materials:

E. Coli K12 Starter Plates (microbial culture)Small Beaker	
4-5 Various Hand Cleaners	Inoculating Loop
Filter Paper	Sterile Petri Dishes
Single Hole Punch	Sterile Forceps
70% IPA	Incubator
Alcohol Burner	

## Procedure:

- 1. Prepare nutrient agar plates per SOP #351, Agar Prep.
- 2. Prepare paper disc by sterilizing single-hole punch by dipping in 70% alcohol, then run through flame of alcohol burner, then punch 4-5 discs into appropriately labeled sterile petri dishes.
- 3. Add 1-2 mL of hand cleaners over sterile discs in appropriately labeled petri dishes and allow discs to soak 1-2 minutes in cleaners.
- 4. Using sterilized forceps, remove discs from cleaners and place into newly labeled sterile petri dishes to dry. Make sure petri dishes containing the soaked discs are placed next to a lit alcohol burner when drying to create an updraft limiting aerial contamination.
- 5. While discs are drying, flip 5 petri dishes over and divide the base of the plate into 4 quadrants drawing a cross with a marker. Label quadrants 1-4 along the edge of the plate.
- 6. Next, label along the edge of the base of petri dishes with the type of cleaner, technicians initials and date.
- 7. Flame an inoculating loop using an alcohol burner to sterilize. Select a colony from the E. Coli K12 starter plate to swab first test plate. Repeat this step until each test plate has been swabbed with E. Coli K12.
- 8. Using sterile forceps, place the dry treated paper disc in the middle of each quadrant of the appropriately labeled plates. Flame forceps each time to prevent cross-contamination.
- 9. Seal the edge of each petri dish with parafilm and place each petri dish upside down on shelf of incubator.
- 10. Incubate plates for 24 hours at 37 degrees C.
- 11. After 24 hours of incubation, remove plates from incubator and measure ring of inhibition of each trial.
- 12. Record measurements in data table of lab notebook.