

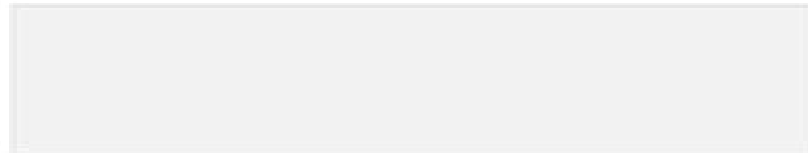
# Microbiology Lab Final

## Modules 2, 3, 5, & 7

### Module 2

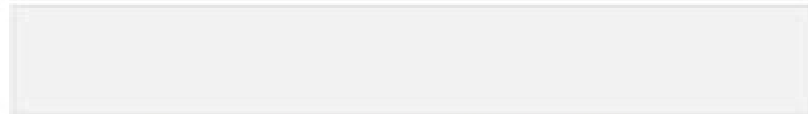
#### • Key Terminology

- Culture is a medium containing living microbes
  - Pure Culture: One species of Microorganisms
  - Mixed Culture: Two or more species of Microorganisms
  - Contaminated Culture: Contains one or more unknown organisms
- Streak Plate Inoculation
  - Purpose:
    - Obtain isolated colonies from a mixed culture -> Pure Culture
      - Colonies are accurately termed colony-forming units (CFUs)
  - Techniques include Quadrant and T-streak methods
    - Make sure to label the bottom of the plate with section, class, initials, and group table #
    - Draw T shape on T-streak plate
  - Typically performed on selective media



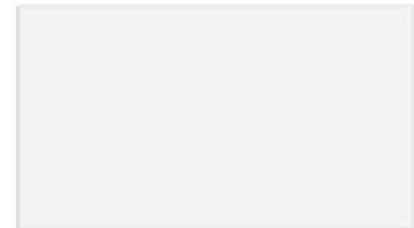
Make sure to incise the inoculating loop between streaks

Drag loop through the last streak to ensure some cells are carried over to the new streak.



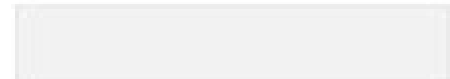
#### • Basic Characteristics of Agar Colony Growth

- Size
- Shape
- Margin
- Surface
- Texture
- Elevation
- Pigment/Optical Properties



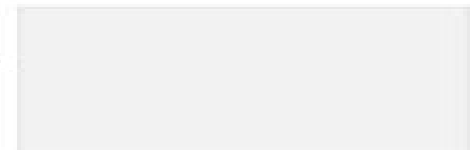
#### • Simple Stains

- A Stain is the solution of solvent and colored molecule (Chromogen that provides contrast)
  - Chromogen:
    - Chromophore gives color
    - Auxochrome gives charge



#### • Bacterial Smear Preparation

- Make sure you are wearing gloves and eye protection
- 1st: Place a small amount of distilled water on the slide (~5 drops) using the inoculating loop
- 2nd: Inoculate the loop and let cool for about 30 seconds, then grab a colony from the agar plate. Take the colony on the loop and swirl the colony in the water until no visible chunks are present
  - Spread the water to about the size of a dime
  - Inoculate loop once done
- 3rd: Allow the smear to air dry, if done correctly the



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**Lauren Gardner**



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