# **INSTRUMENT NAME**: Comparison Microscope

II. <u>SUGGESTED USES</u>: Hair Analysis and Fiber Analysis.

### III. OPERATING PROCEDURES:

- A. START-UP AND ALIGNMENT
  - 1. Turn on the power switch for both microscope bases.
  - 2. Adjust to the desired light intensity with the rheostat control on each microscope base .
  - 3. Place a slide to be examined on the stage of both microscopes and adjust for "Koehler Illumination" (refer to operators manual for this procedure). illumination procedure is only performed as

This needed.

- B. COLLECTION AND STORAGE OF DATA
  - Observations of the material in question can be made with respect to but not limited to the following:
    - \* Physical Characteristics
- color
- crystal structure
- internal artifacts
- diameter
- cross-sectional shape
- surface texture
- texturizing
- \* Optical Properties
- extinction
- refractive indices
- birefringence
- sign of elongation
- pleochroism
- fluorescence (module A)
- fluorescence (module H3)
- \* Chemical Properties
- reactions to chemical tests
- solubility tests
- 2. Observations of the materials in question can be recorded by the following means:
  - \* Descriptive writing
  - \* Drawing or sketching
  - \* Polaroid or standard 35mm photography

# \* Digital image capture

## C. SHUTDOWN

- 1. Lower the light intensity for both microscope bases.
- 2. Turn off the power switch for both microscope bases.

# IV. SAFETY CONCERNS

- A. DO NOT SPEND LONG PERIODS OF TIME MAKING OBSERVATIONS AT HIGH LIGHT INTENSITIES.
- B. DO NOT OPERATE THE MICROSCOPE NEAR SINKS OR AREAS WHERE A POTENTIAL ELECTRICAL SHOCK CAN OCCUR.
- C. WHEN CHANGING BULBS, MAKE SURE THAT THE POWER CORD IS DISCONNECTED.
- D. DO NOT STARE DIRECTLY INTO THE FLUORESCENT LIGHT SOURCES.

### V. OTHER INFORMATION

NONE