

# Instructions For Use TEAK-IFU

205 South 600 West Logan, Utah 84323, U.S.A. - Tel. (800) 729-8350 - Tel. (435) 755-9848 - Fax (435) 755-0015 - www.scytek.com Rev. 1, 4/30/2025

# **TEA Stain Kit**

### **Description and Principle**

TEA Stain Kit is intended for use in cytological applications. TEA stain solutions #1 and #2 are used to stain a smeared slide for microscopic examination during rapid on-site evaluation (ROSE), with the option for permanent staining.

# **Expected Results**

Nuclei: Blue Cytoplasm: Light pink

 Kit Contents
 Storage

 1. TEA Stain #1
 18-25°C

 2. TEA Stain #2
 18-25°C

### Suggested Controls (not provided)

Slide smear.

## **Uses/Limitations**

For In-Vitro Diagnostic use only.
Do not use if reagents become cloudy or precipitate
Do not use past expiration date.
Use caution when handling reagents.
Non-Sterile
Intended for cytology specimens.

### Storage

Store kit and all components at room temperature (18-25°C).

# **Safety and Precautions**

Please see current Safety Data Sheets (SDS) for this product and components GHS classification, pictograms, and full hazard/precautionary statements

### **Procedure for TEA Staining:**

- 1. Prepare a slide smear.
- 2. Ensure that the smear size is appropriate (covers 1/2 to 2/3 of the slide) to achieve optimal stain exposure.

### 3. Staining with TEA Solution:

- Either spray TEA Stain #1 onto the slide smear or dip it in a jar containing solution for at least 3 seconds. (Note: The duration of exposure to TEA Stain #1 determines the color of the cytoplasm. A shorter exposure is desirable to prevent RBCs from picking up the stain.)
- Add 2-3 drops of TEA Stain #2, allow it to sit for a minimum of 3 seconds. (Note: Longer exposure to TEA Stain #2 retains more blue color in the nuclei and background. A shorter exposure is desirable to prevent the background from retaining excess stain.)
- 4. Place a coverslip on top of the stained slide.
- 5. Gently apply pressure to the coverslip using paper towels or gauze, guiding any excess stain to the edges.
- 6. Slide is ready for microscopic examination.



### **Slide Aftercare Following ROSE:**

- 1. Place the specimen slide with coverslip in an alcohol container. This step aids in the de-staining process.
- 2. Send the container to the cytology lab for re-staining with PAP stain. The coverslip will fall off or peel off easily with gentle traction.
- Note: Re-staining with PAP stain or using the slide for other ancillary testing, such as IHC or molecular tests, can be performed because TEA stain solutions are alcohol-based and do not interfere with these studies. For optimum results, place the slide in an alcohol-based fixative solution not greater than 48 hours.
- Use you regular in-house de-staining procedure, if necessary or if the coverslip did not fall off, to process the slide for re-staining with PAP or ancillary testing.

### **Procedure for Permanent TEA Staining:**

- 1. Follow steps 1-3 of the TEA Staining Procedure.
- 2 Dip the slide 10 times in de-ionized water to rinse off the TEA stains.
- 3. Apply Xylene or a similar product to the top of the slide.
- 4. Gently apply pressure to the coverslip using paper towels or gauze.
- 5. The permanent slide is now ready for microscopic examination.

ScyTek Laboratories, Inc. 205 South 600 West Logan, UT 84321 U.S.A.

C€

IVD

Ec\_REP]
Emergo Europe
Westervoortsedijk 60
6827 AT Arnhem,The Netherlands