

MycLight™ Rapid Fluorescence Bacterial Gram Stain Kit

 Catalog number: 22413
 Unit size: 100 Tests

Component	Storage	Amount
Component A: MycoLight™ Green	Freeze (< -15 °C), Minimize light exposure	1 vial (100 µL)
Component B: MycoLight™ Red	Freeze (< -15 °C), Minimize light exposure	1 vial (100 µL)

OVERVIEW

The MycoLight™ Rapid Fluorescence Bacterial Gram Stain Kit provides an easy and convenient way for determination of gram sign in live bacteria. Gram staining is a commonly used method in both clinical and research settings to taxonomically classify bacterial species into two large groups. Unfortunately, the traditional gram staining method is tedious and involves bacterial fixation which can be a significant drawback if the bacteria are to be characterized further. The MycoLight™ Rapid Fluorescence Bacterial Gram Stain Kit provides a one-step gram staining assay for live bacteria that overcomes the problems inherent in the traditional gram staining assays. The MycoLight™ Rapid Fluorescence Bacterial Gram Stain Kit utilizes two DNA dyes MycoLight™ Green and MycoLight™ Red with differential ability to stain gram positive and negative bacteria. MycoLight™ Green stains both gram positive and negative bacteria while MycoLight™ Red preferentially labels gram positive bacteria. The excitation/emission maxima for these two dyes are about 484/504 nm for MycoLight™ Green and 518/600 nm for MycoLight™ Red. Thus, when a mixture of gram positive and gram negative bacteria is stained with the dyes, gram positive bacteria will fluoresce red and gram negative bacteria will fluoresce green. The gram positive and negative staining can be monitored fluorimetrically with TRITC and FITC filter set respectively.

AT A GLANCE

Protocol Summary

1. Prepare bacterial samples
2. Prepare and add MycoLight™ dye working solution to bacteria samples
3. Incubate with MycoLight™ dye working solution at room temperature in dark for 15 minutes
4. Analyze sample by fluorescence microscope or fluorescence spectroscopy with FITC and TRITC filter sets

Important Thaw all the kit components at room temperature before use.

KEY PARAMETERS

Fluorescence microscope

Excitation	488/540 nm
Emission	530/620 nm
Recommended plate	Black wall/clear bottom
Instrument specification(s)	FITC/TRITC filters

PREPARATION OF WORKING SOLUTION

MycLight™ dye working solution

Mix equal volume of MycoLight™ Green (Component A) and MycoLight™ Red (Component B) in a tube and mix them well.

SAMPLE EXPERIMENTAL PROTOCOL

Preparation of Bacterial Samples

1. Prepare bacteria sample with concentration around 10^7 cells/ml. Grow bacteria into late log phase in appropriate medium. **Note:**

Measure the optical density of the bacterial culture at wavelength = 600 nm (OD600) to determine the cell number. For *E. coli* culture, OD600 = 1.0 equals 8×10^8 cells/ml.

2. Remove medium by centrifugation at 10,000 x g for 10 minutes and re-suspend the pellet in ddH₂O, adjust bacteria concentration to $\sim 10^7$ cells/ml.

Staining Protocol

1. Add 2 µL MycoLight™ dye working solution to 100 µL of the bacterial suspension.
2. Mix well and incubate in dark for 15 min at room temperature.
3. Monitor fluorescence of bacteria with a fluorescent microscope through FITC (Ex/Em = 488/530 nm) channel for gram-negative bacteria and TRITC (Ex/Em = 540/620 nm) channel for gram-positive bacteria. **Note:** The protocol only provides a guideline, should be optimized with different bacterial strains or other specific needs. **Note:** Relative ratio of gram positive and gram negative bacteria in a population can also be estimated with fluorescence spectroscopy with this kit. A sample analysis is included with the figures.

EXAMPLE DATA ANALYSIS AND FIGURES

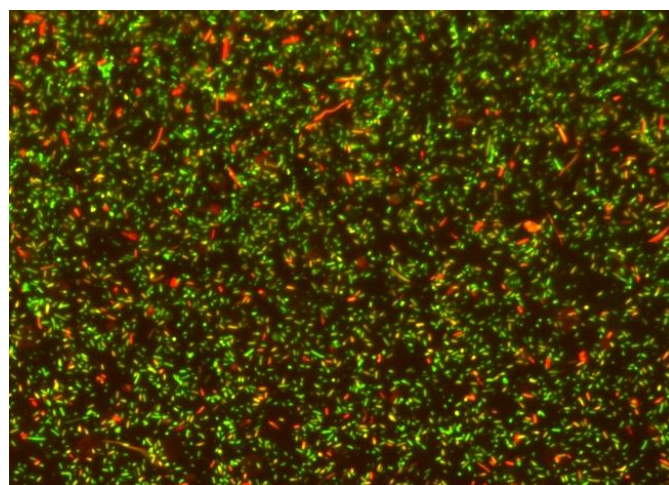


Figure 1. Mixture of *Escherichia coli* and *Bacillus subtilis* were stained with MycoLight™ Rapid Fluorescence Bacterial Gram Stain Kit. Red & Orange: Gram positive *Bacillus subtilis* cells; Green: gram-negative *Escherichia coli* cells.

DISCLAIMER

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