

Propidium Iodide Nucleic Acid Stain in Euplotes

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Abstract

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Protocol

Step 1.

Collect ~ 2 ml of Euplotes cells from a culture at 500 cells/ml, concentrate them by centrifugation, 7 min at 3000 rpm and remove the supernatant by pipetting.

Step 2.

Fix Euplotes cells with 1 volume of methanol for 10 min.

Step 3.

Remove as much as possible the solution and wash with 1X PBS (~1 ml) (washing step can be done by centrifugation).

Step 4.

Add Propidium Iodine 24 µg/ml and let it stain for 30 min.

Step 5.

Wash with 1X PBS (~1 ml) and remove as much as possible all the solution (washing step can be done by centrifugation).

Step 6.

Put 10 μl of the cells on a slide and observe them under the fluorescent microscope.