Kit Components (R37602)	Amount
Wash Buffer Fixative Solution Permeabilization Solution Blocking Solution	500 mL 20 mL 20 mL 12 x10 mL
Materials needed but not supplied	
6-well plate or 35-mm dish (protocol designed for ~ 10 cm² growth area)	

Primary & Secondary Antibodies

Dves



1. Remove media from cells



2. Add 1 mL of Fixative Solution



3. Incubate 15 min at 20-25°C



4. Remove Fixative Solution. Wash 3 times with 2 mL of Wash Buffer for 2-5 min



5. Add 1 mL of Permeabilization Solution



6. Incubate 15 min at 20-25°C



7. Remove Permeabilization Solution. Wash 3 times with 2 mL of Wash Buffer for 2-5 min



8. Add 2 mL of Blocking Solution



9. Incubate 60 min at 20-25°C



10. Proceed with desired cell staining and/or antibody labeling