

Jun 25, 2024



# IF Labeling Protocol

DOI

#### dx.doi.org/10.17504/protocols.io.4r3l2q6rql1y/v1

Scott Vermilyea<sup>1</sup>

<sup>1</sup>University of Minnesota, Team Lee

ASAP Collaborative Rese...

Team Lee



#### Jane Balster

ASAP - Team Lee





DOI: dx.doi.org/10.17504/protocols.io.4r3l2q6rql1y/v1

Protocol Citation: Scott Vermilyea 2024. IF Labeling Protocol. protocols.io <a href="https://dx.doi.org/10.17504/protocols.io.4r3l2q6rql1y/v1">https://dx.doi.org/10.17504/protocols.io.4r3l2q6rql1y/v1</a>

License: This is an open access protocol distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited

Protocol status: Working We use this protocol and it's

working

Created: June 07, 2024

Last Modified: June 25, 2024

Protocol Integer ID: 102391

Keywords: Protein Labelling, Tetrafluorophenyl, Conjugation

**Funders Acknowledgement:** 

**ASAP** 

Grant ID: 000592



### **Abstract**

This protocol details the Labelling of Immunofluorescent Protein.

## Materials

XX Alexa Fluor™ 488 Microscale Protein Labeling Kit Thermo Fisher Catalog #A30006



### Immunofluorescent Protein Labeling Protocol

17m 15s

- 1 Alex Fluor 488 Microscale Protein Labelling Kit (Invitrogen, A30006).
- 1.1 According to manufacturer protocol, AF 488 reactive dye has Tetrafluorophenyl (TFP) ester moiety that is more stable in solution than the commonly used succinimidyl (NHS) ester. TFP ester react efficiently with primary amines of protein to form a stable dye-protein conjugate and is independent of pH between 4 and 10.
- 1.2 Single kit can label a total of  $\angle$  200 µg of  $\alpha$ S preformed fibrils.
- 1.3 Dilute Hu-WT-αS fibrils (PFFs) to a concentration of Δ 1 undetermined in PBS followed by sonication (1 Sec ON/1 Sec OFF) for 60 00:02:00 with 20% amplitude.

2m

1.4 Mix diluted PFFs with component B (NaHCO3), then added freshly prepared reactive dye (component A) and incubated for 00:15:00 at 8 Room temperature.

15m



- 1.5 Calculate amount of reactive dye using dye/protein molar ratio of 9.
- 1.6 Remove excess dye by resin gel (component E) purification.
- 1.7 Place conjugated reaction mixture of PFFs and reactive dye on the center of the resin bed surface, centrifuge at \$\mathbb{\mathbb{C}}\$ 16000 x g, 00:00:15 .

15s



1.8 Collect the filtrate as labeled PFFs.