

Jun 26, 2024

Version 1

DOI

dx.doi.org/10.17504/protocols.io.n2bvjnd7ngk5/v1

Samantha Goetting¹

¹Johns Hopkins University



Samantha Goetting

Johns Hopkins University

Create & collaborate more with a free account

Edit and publish protocols, collaborate in communities, share insights through comments, and track progress with run records.

Create free account

OPEN ACCESS



DOI: https://dx.doi.org/10.17504/protocols.io.n2bvjnd7ngk5/v1

Protocol Citation: Samantha Goetting 2024. Immunofluorescence staining on larval and adult Drosophila gonads. **protocols.io** https://dx.doi.org/10.17504/protocols.io.n2bvjnd7ngk5/v1

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 26, 2024



Last Modified: June 26, 2024

Protocol Integer ID: 102463

Keywords: drosophila, immunofluorescence, staining protocol, larval, adult drosophila

Abstract

Immunofluorescence staining protocol for *Drosophila* gonads

Materials

1x PBS 0.3% PBTx (0.3% Triton-X in PBS) 1% PBTx (1% Triton-X in PBS) Paraformaldehyde or formaldehyde Normal serum (usually NGS) Primary antibodies Secondary antibodies DAPI

Troubleshooting

Before start

All steps are done with gentle rotation.



Day 1 2h 40m 1 Dissect tissue in 1x PBS. Transfer to a 1.5 mL tube containing 1x PBS. If it is a quick dissection (<20 mins), no ice needed. If more time is needed, keep samples on ice. 2 Remove PBS and add fixative. Fix in 4% paraformaldehyde in **0.3% PBTx** for 20 min RT 20m with gentle rotation. 500 uL fixative = 125 uL of 16% paraformaldehyde + 375 uL 0.3% PBTx 3 Aspirate the fixative and wash twice for 10 min in 1% PBTx. Not getting rid of fix will 20m affect your immunostaining. 4 Aspirate the supernatant and block/permeabilize for at least 2 hours in 1% PBTx + 5% 2h normal serum, or overnight at 4°C. 1 mL block solution = 50 uL NGS + 950 uL 1% PBTx 5 Primary antibodies are diluted in **0.3% PBTx** + 5% normal serum and incubate for 1 hour 12h at RT or overnight at 4°C. Overnight will give better staining. T Primary antibody mix in 0.3% PBTx + 15 uL NGS (300 uL total) Day 2 5h 20m 6 Remove the primary antibody mix and wash in **0.3% PBTx** three times for 20 min at RT. 1h 7 Wash in **0.3% PBTx** + 5% normal serum twice for 30 min at RT. 1h 1 mL wash solution = 50 uL NGS + 950 uL 0.3% PBTx 8 Secondary antibodies are diluted in 0.3% PBTx + 5% normal serum and incubated for ~2 2h hours at RT or overnight at 4°C. Keep tubes covered from light. 9 Aspirate the supernatant and add 500 uL DAPI to each tube. Incubate for 10 min at RT. 10m Keep tubes covered from light. 10 Aspirate the supernatant and wash in **0.3% PBTx** three times for 20 min at RT. 1h 11 Aspirate the supernatant and wash in **PBS** for 10 min at RT. 10m 12 Store in **PBS** at 4°C or proceed with mounting. Keep tubes covered from light.





Slaidina et al. (2020)