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Immunocytochemistry

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Protocol status: Working

We use this protocol and it's working

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Aligning Science Across Parkinson's

Abstract

This is a protocol for immunocytochemistry for 24 and 96 well cell culture plates.

Materials

	A	B	C	D
	Vendor	Catalog#	Qty	Description
	RPI	A30075-100.0	1	Bovine Serum Albumin
	Electron Microscopy Services	15713	10 X 10ml	20% Paraformaldehyde



Preparation

- 1 Prepare 1x phosphate buffer saline (PBS)
- 2 Prepare fixation solution: 4% paraformaldehyde/4% sucrose in PBS

Note

can be stores at 4°C for up to 1 month

- 3 Prewarm the fixative solution to 37 degrees Celsius
- 4 Prepare blocking buffer: 3% Bovine Serum Albumin (BSA) in PBS
 - Seal the conical with parafilm to prevent evaporation and store at 4 degrees Celsius
- 5 Prepare 10% stock of Triton-X-100 in PBS
- 6 Prepare permeabilization buffer: 0.3% TX-100 in 3% BSA in PBS

Staining

- 7 Thoroughly aspirate media from each well without disturbing cells
- 8 Add pre-warmed fixative solution to the wells and incubate at room temperature for 15 minutes. (0.5 mL/well for 24-well plate, 100µL/well for 96-well plate).
- 9 Rinse 5X with PBS. This and subsequent washes can be done with the plate washer for 96-well plates.
- 10 Permeabilize neurons with a permeabilization buffer for 15 minutes at room temperature.



- 11 Rinse 3X with PBS.
- 12 Block neurons for a further 50 minutes with a blocking buffer at room temperature.
- 13 Dilute primary antibodies in blocking buffer.
- 14 Incubate cells with primary antibody for 2 hours at room temperature or overnight at 4°C.
- 15 Rinse 5X with PBS.
- 16 Dilute Alexa Fluor conjugated secondary antibodies corresponding to primary antibodies in blocking buffer in 1:500 dilution
- 17 Incubate cells in secondary antibodies for 1 hour at room temperature.

Note

Make sure that the secondary antibodies will not recognize each other or crossreact with the same primary antibodies!

- 18 Rinse 5X with PBS.
- 19 Mount coverslips onto glass slides with Prolong Gold mounting media. 96-well should be incubated with 100 μ L DAPI 1:10,000 in PBS.
- 20 Seal the 96-well plate with a black seal.
- 21 Visualize staining on a fluorescent microscope.