

Jul 13, 2019

## Mouse Stellate Immunohistochemistry protocol

DOI

dx.doi.org/10.17504/protocols.io.2qggdtw

Pradeep Rajendran<sup>1</sup>

<sup>1</sup>[University of California, Los Angeles]



Scott John

## 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





DOI: https://dx.doi.org/10.17504/protocols.io.2qggdtw

Protocol Citation: Pradeep Rajendran 2019. Mouse Stellate Immunohistochemistry protocol. protocols.io https://dx.doi.org/10.17504/protocols.io.2qggdtw

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: May 09, 2019



Last Modified: July 13, 2019

Protocol Integer ID: 23016

Keywords: immunohistochemistry protocol, mouse

## Troubleshooting



- 1 See mouse steallte isolation protocol
- 2 Complete/intact stellate ganglia are fixed in 4% paraformaldehyde overnight at 4°C.
- 3 Fixed tissue is rinsed in phosphate buffered saline (PBS), and stored in PBS + 0.02% sodium azide.
- 4 The stellate ganglion whole mounts are permeabilized in 'block' solution. (10% normal donkey serum in 1.0% BSA + 0.4% Triton X-100 + PBS).
- 5 Ganglia are stained with antibodies directed to either tyrosine hydroxylase (TH) or protein gene product 9.5 (PGP 9.5). Secondary staining with streptavidin conjugated to ATTO-647N was used to visualize neurobiotin filling. Stained tissue was rinsed in PBS and mounted on glass slides.