

Orders: orders@novusbio.com

Support: technical@novusbio.com

Web: www.novusbio.com

NBP1-79054 Protocol

 $\label{eq:protocols} \textbf{Protocols}, \textbf{Publications}, \textbf{Related Products}, \textbf{Reviews and more:}$

www.novusbio.com/NBP1-79054

Immunohistochemistry-Paraffin protocol for CD3 Antibody (NBP1-79054)

IHC-P Protocol (NBP1-79054): https://www.novusbio.com/products/cd3-antibody-n26-r nbp1-79054

- 1. Deparaffinize the section in 3 changes of xylene, 5 minutes each.
- 2. Wash the section in 96%, 80% and 70% benzyl alcohol for 5 minutes each.
- 3. Rinse in distilled water.
- 4. Block the endogenous peroxidase by incubating the tissue in 3% hydrogen peroxide (H2O2) for 10 minutes.
- 5. Wash in distilled water.
- 6. For antigen retrieval: immerse the slide in Tris-EDTA buffer, pH 9.0, 0.05% Tween- 20*, and incubate in microwave (850W) for 20 minutes. (Alternatively adjust to your own protocol, keeping the required pH)
- 7. Remove the staining to room temperature and let the slide to cool for 15 minutes.
- 8. Rinse in distilled water.
- 9. Wash in 0.05 M Tris-HCI, pH 7.6 buffer supplemented with 0.2% of Tween-20 (buffer A) for 5 minutes.
- 10. Incubate the section with primary antibody diluted in buffer A at the dilution 1:100 200 for 1 hour in the closed wet chamber.
- 11. Wash twice 5 minutes with buffer A.
- 12. Apply the secondary antibody (the protocol depends on the supplier), and proceed to standard immunohistochemistry protocol (HRP Peroxide DAB).
- 13. Wash twice 5 minutes with buffer A.
- 14. Apply the chromogen (DAB), 10 minutes.
- 15. Wash in water 10 minutes.
- 16. Stain in hematoxylin for 5 minutes.
- 17. Wash in water 10 minutes.
- 18. Dehydrate the section in 2 changes of 96% benzyl alcohol for 5 minutes each.
- 19. Wash the section in 2 changes of xylene for 2 minutes each.
- 20. Mount the slide for observation.