

Orders: orders@novusbio.com

Support: technical@novusbio.com

Web: www.novusbio.com

## NBP1-78965 Protocol

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

www.novusbio.com/NBP1-78965

## Serum protocol for TET1 Antibody (NBP1-78965)

TET1 Antibody: https://www.novusbio.com/products/tet1-antibody\_nbp1-78965 Immunohistochemistry-Paraffin Embedded Sections

## Antigen Unmasking:

Bring slides to a boil in 10 mM sodium citrate buffer (pH 6.0) then maintain at a sub-boiling temperature for 10 minutes. Cool slides on bench-top for 30 minutes.

## Staining:

- 1. Wash sections in deionized water three times for 5 minutes each.
- 2. Wash sections in wash buffer for 5 minutes.
- 3. Block each section with 100-400 ul blocking solution for 1 hour at room temperature.
- 4. Remove blocking solution and add 100-400 ul diluted primary antibody. Incubate overnight at 4 C.
- 5. Remove antibody solution and wash sections in wash buffer three times for 5 minutes each.
- 6. Add 100-400 ul biotinylated diluted secondary antibody. Incubate 30 minutes at room temperature.
- 7. Remove secondary antibody solution and wash sections three times with wash buffer for 5 minutes each.
- 8. Add 100-400 ul Streptavidin-HRP reagent to each section and incubate for 30 minutes at room temperature.
- 9. Wash sections three times in wash buffer for 5 minutes each.
- 10. Add 100-400 ul DAB substrate to each section and monitor staining closely.
- 11. As soon as the sections develop, immerse slides in deionized water.
- 12. Counterstain sections in hematoxylin.
- 13. Wash sections in deionized water two times for 5 minutes each.
- 14. Dehydrate sections.
- 15. Mount coverslips.

<sup>\*</sup>The above information is only intended as a guide. The researcher should determine what protocol best meets their needs. Please follow safe laboratory procedures.