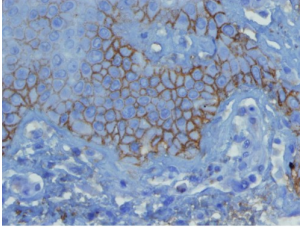


Complement Fragment C3d (C3d) Antibody

Catalogue No.: abx227113



IHC-P analysis of skin biopsy from the lesion of the early pemphigus vulgaris (without blister formation), showing strong positive intraepidermal intercellular immunostaining (4 μ m section).

Complement Fragment C3d (C3d) Antibody is a Rabbit Monoclonal antibody for the detection of C3d complement.

Target:	Complement Fragment C3d (C3d)
Clonality:	Monoclonal
Clone:	I808
Reactivity:	Human
Tested Applications:	IHC
Host:	Rabbit
Recommended dilutions:	IHC-P: 1/100 - 1/200. Optimal dilutions/concentrations should be determined by the end user.
Conjugation:	Unconjugated
Immunogen:	Synthetic peptide derived from the N-terminal region of human C3d complement fragment.
Isotype:	IgG
Form:	Liquid
Purification:	Purified from rabbit antiserum by proprietary techniques.
Storage:	Store at 2-8°C.
UniProt Primary AC:	P01024 (UniProt , ExPASy)
Buffer:	20 mM Tris-HCl, pH 8.0, containing 20 mg/ml BSA and 0.05% Na ₃ N.

Note: THIS PRODUCT IS FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC, THERAPEUTIC OR COSMETIC PROCEDURES. NOT FOR HUMAN OR ANIMAL CONSUMPTION.

Directions for use: Suggested IHC-P Protocol

1. Preparation of Tris-EDTA Buffer (10 mM Tris Base, 1 mM EDTA solution, 0.05% Tween-20, pH 9.0): Dissolve 1.21 g Tris and 0.37 g EDTA in 700 ml of distilled water and mix. Adjust pH to 9.0 with 1 M HCl and then add 0.5 ml of Tween-20 and mix thoroughly. Adjust the final volume to 1 L with distilled water. Store this solution at room temperature for up to 3 months or at 4 °C for long-term storage.
2. Preparation of Wash Buffer: Use 0.05 M Tris-HCl, pH 7.6, containing 0.2% Tween-20; or PBS containing 0.2% Tween-20.
3. Preparation of Copper Sulfate Solution: Dissolve 0.90 g NaCl and 0.50 g CuSO₄.5H₂O in 100 ml distilled water.
4. Deparaffinize the section in 3 changes of xylene, 10 minutes each.
5. Wash the section in 96%, 80% and 70% ethanol, 10 minutes each.
6. Rinse twice in distilled water, 5 minutes each.
7. Block endogenous peroxidase by incubating the tissue in 3% hydrogen peroxide (H₂O₂) for 10 minutes.
8. Rinse twice in distilled water, 5 minutes each.
9. Antigen retrieval: immerse the slide in Tris-EDTA buffer, pH 9.0, and incubate in a water bath for 20-25 minutes at 96-98 °C.
10. Remove the slide from the water bath and allow to stand at room temperature (in Tris-EDTA buffer, pH 9.0) for 20 minutes.
11. Rinse twice in distilled water, 5 minutes each.
12. Wash twice in Wash Buffer, 5 minutes each.
13. Incubate the section with primary antibody at 1/100 - 1/200 dilution for 1 hour in a closed wet chamber. It is recommended to use [abx291502](#) Primary Antibody Diluent or a diluent containing protease-free BSA (≥ 1 mg/ml) to dilute this antibody.
14. Rinse three times in distilled water, 5 minutes each.
15. Add the secondary antibody and proceed to standard immunohistochemistry protocol (HRP - Peroxide - DAB). It is recommended to use [abx291501](#) Rabbit and Mouse HRP/DAB Detection Kit.
16. Wash three times in Wash Buffer, 5 minutes each.
17. Apply the DAB chromagen for 1-3 minutes.
18. Wash twice in distilled water, 5 minutes each.
19. Rinse in Copper Sulfate Solution.
20. Wash in distilled water for 2 minutes.
21. Stain in hematoxylin for 5 minutes.
22. Wash three times in distilled water, 2 minutes each.
23. Rinse in 37 mM ammonium hydroxide solution for 1 minute.
24. Wash in distilled water for 2 minutes.
25. Mount the slide for observation.