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Anti - CD57

Rabbit clonal antibody

CAT#

DB 216-1

CONCENTRATED READY TO USE (RTU)

DB 216-0.1 (100 μ l) DB 216-RTU-7 (7 ml) DB 216-0.2 (200 μ l) DB 216-RTU-15 (15 ml) DB 216-0.5 (500 μ l)

STORAGE AND APPLICATION

(1 ml)

CONCENTRATED READY TO USE (RTU)

Storage: +4°C, Do not freeze!

Application: IHC-P, dilution 1:100 - 1:200 Application: IHC-P, ready to use

PRODUCT INFORMATION

Clone: E20-l

Buffer: 20 mM Tris-HCl, pH 8.0 Stabilizer: 20 mg/ml BSA Preservative: 0.05% NaN₃

Specificity: Human, mouse, rat

Expiration: 24 months from the shipping date

Immunogen: Peptide derived from C-terminal sequence of human CD57. Antibody recognizes the epitope between

Lys316 - Val332.

Cellular localization: Golgi apparatus membrane

Positive control: brain tissue

Protein accession number: Q9P2W7 (h); Q9CW73 (m); O35789 (r)

IHC-P PROTOCOL - INSTRUCTION MANUAL

- 1. Deparaffinize the section in 3 changes of xylene, 10 minutes each.
- 2. Wash the section in 96%, 80% and 70% ethyl alcohol for 10 minutes each.
- 3. Rinse in distilled water, 2 x 5 minutes.
- 4. Block the endogenous peroxidase by incubating the tissue in 3% hydrogen peroxide (H_2O_2) for 10 minutes.
- 5. Wash in distilled water, 2 x 5 minutes.
- For antigen retrieval immerse the slide in citrate buffer, pH 6.0*, and incubate in water bath for 25-35 minutes at 96-98°C.
- Remove the staining to room temperature and let the slide to cool (in citrate buffer, pH 6.0) for 20 minutes.
- 8. Rinse in distilled water, 2 x 5 minutes.
- 9. Wash in PBS (buffer A), 2 x 5 minutes
- 10. CONCENTRATED:

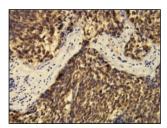
Incubate the section with primary antibody at the **dilution 1:100 - 1:200** for 1 hour in the closed wet chamber.

READY TO USE (RTU):

Incubate the section with primary antibody (ready to use) for 1 hour in a closed wet chamber.

- 11. Wash 3 x 5 minutes with buffer A.
- Apply the secondary antibody (the protocol depends on the supplier), and proceed to standard immunohistochemistry protocol (HRP - Peroxide - DAB).
 Micropolymer-HRP detection kit rabbit/mouse dual of DB Biotech is suggested (http://www.dbbiotech.com/products/detection-system.html).
- 13. Wash 3 x 5 minutes with buffer A.
- 14. Apply the chromogen (DAB), 1-3 minutes.
- 15. Wash in distilled water, 2 x 5 minutes.
- 16. Rinse in $CuSO_4.5H_2O$ solution /0,90g NaCl + 0,50g $CuSO_4.5\ H_2O$ in 100ml distilled water/
- 17. Wash in distilled water, 1 x 2 minutes.
- 18. Stain in hematoxylin for 5 minutes.
- 19. Wash in distilled water, 3 x 2 minutes.
- 20. Rinse in 37mM ammonium hydroxide solution.
- 21. Wash in distilled water, 1 x 2 minutes.
- 22. Mount the slide for observation.

* Citrate Buffer (10mM Citric Acid, 0.05% Tween-20, pH 6.0):



Poorly differentiated human neuroblastoma tissue stained with anti-CD57 (DB 216) monospecific clonal antibody. Formalin fixed, paraffin embedded human tissue (4 µm section) stained according to related DB Biotech datasheet.

PRECAUTIONS

- We strongly recommend to use DB Primary Antibody Diluent (catalog number DB D-125, or DB D-250), eventually alternative diluent (containing protease free BSA at the concentrations ≥ 1mg/ml) for dilution of concentrated antibodies, otherwise the warranty might be voided.
- 2. Centrifuge the vial before use.
- 3. Intended for professional In Vitro Diagnostic use in laboratories.
- Do not use after expiration date stamped on vial label.
- 5. Avoid contamination of the reagent.
- Any discrepancies in the recommended procedures stated in the working protocol may affect the final results.
- The reagent contains sodium azide (NaN₃) which is highly toxic in higher concentrations. The concentration in the reagent (0.05%) is not considered as hazardous.

Revision Date: 22.05.2017

- 8. Disposal of waste material must be conducted in accordance with local regulations.
- 9. Wear appropriate Personal Protective Equipment to avoid contact with eyes and skin.