10320 Camino Santa Fe, Suite G
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Anti-DHFR Antibody Picoband™ (monoclonal, 3C8)

Catalog # ABO14841

Specification

Anti-DHFR Antibody Picoband™ (monoclonal, 3C8) - Product Information

Application	**WB, IHC**
Primary Accession	P00374
Host	**Mouse**
Isotype	**Mouse IgG2b**
Reactivity	**Rat, Human**
Clonality	**Monoclonal**
Format	**Lyophilized**

Description
Anti-DHFR Antibody Picoband™ (monoclonal, 3C8) . Tested in IHC, WB applications. This antibody reacts with Human, Rat.

Reconstitution
Add 0.2ml of distilled water will yield a concentration of 500 µg/ml.

Anti-DHFR Antibody Picoband™ (monoclonal, 3C8) - Additional Information

Gene ID 1719

Other Names
Dihydrofolate reductase, 1.5.1.3, DHFR

Calculated MW
22 kDa KDa

Application Details
Western blot, 0.1-0.5 µg/ml
 Immunohistochemistry (Paraffin-embedded Section), 0.5-1 µg/ml

Subcellular Localization
Mitochondrion. Cytoplasm

Tissue Specificity
Widely expressed in fetal and adult tissues, including throughout the fetal and adult brains and whole blood. Expression is higher in the adult brain than in the fetal brain.

Contents
Each vial contains 4mg Trehalose, 0.9mg NaCl, 0.2mg Na2HPO4, 0.05mg NaN3.

Immunogen
E.coli-derived human DHFR recombinant protein (Position: V2-D187). Human DHFRshares90%aminoacid (aa)sequenceidentitywith both mouseandrat DHFR.

Cross Reactivity
No cross-reactivity with other proteins.

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Storage	**Store at -20˚C for one year from date of receipt. After reconstitution, at 4˚C for one month. It can also be aliquotted and stored frozen at -20˚C for six months. Avoid repeated freeze-thaw cycles.**

Anti-DHFR Antibody Picoband™ (monoclonal, 3C8) - Protein Information

Name DHFR

Function
Key enzyme in folate metabolism. Contributes to the de novo mitochondrial thymidylate biosynthesis pathway. Catalyzes an essential reaction for de novo glycine and purine synthesis, and for DNA precursor synthesis. Binds its own mRNA and that of DHFR2.

Cellular Location
Mitochondrion {ECO:0000250|UniProtKB:P00375}. Cytoplasm {ECO:0000250|UniProtKB:P00375}

Tissue Location
Widely expressed in fetal and adult tissues, including throughout the fetal and adult brains and whole blood Expression is higher in the adult brain than in the fetal brain

Anti-DHFR Antibody Picoband™ (monoclonal, 3C8) - Protocols

Provided below are standard protocols that you may find useful for product applications.

- Western Blot
- Blocking Peptides
- Dot Blot
- Immunohistochemistry
- Immunofluorescence
- Immunoprecipitation
- Flow Cytomety
- Cell Culture

Anti-DHFR Antibody Picoband™ (monoclonal, 3C8) - Images

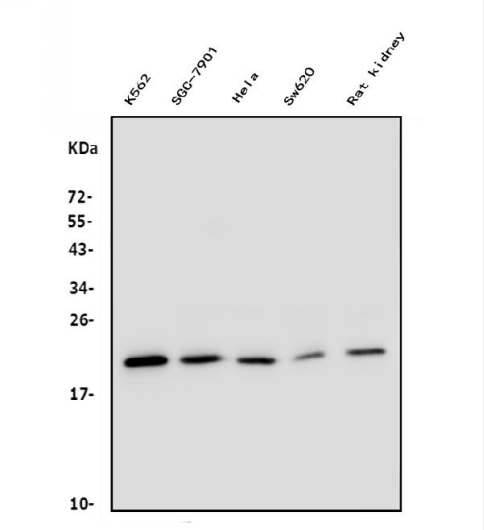

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Figure 1. Western blot analysis of DHFR using anti-DHFR antibody (M00813-1).
Electrophoresis was performed on a 5-20% SDS-PAGE gel at 70V (Stacking gel) / 90V (Resolving gel) for 2-3 hours. The sample well of each lane was loaded with 50ug of sample under reducing conditions.
Lane 1: human K562 whole cell lysates,
Lane 2: human SGC-7901 whole cell lysates,
Lane 3: human Hela whole cell lysates,
Lane 4: human SW620 whole cell lysates,
Lane 5: rat kidney tissue lysates.
After Electrophoresis, proteins were transferred to a Nitrocellulose membrane at 150mA for 50-90 minutes. Blocked the membrane with 5% Non-fat Milk/ TBS for 1.5 hour at RT. The membrane was incubated with mouse anti-DHFR antigen affinity purified monoclonal antibody (Catalog # M00813-1) at 0.5 µg/mL overnight at 4°C, then washed with TBS-0.1%Tween 3 times with 5 minutes each and probed with a goat anti-mouse IgG-HRP secondary antibody at a dilution of 1:10000 for 1.5 hour at RT. The signal is developed using an Enhanced Chemiluminescent detection (ECL) kit (Catalog # EK1001) with Tanon 5200 system. A specific band was detected for DHFR at approximately 22KD. The expected band size for DHFR is at 22KD.

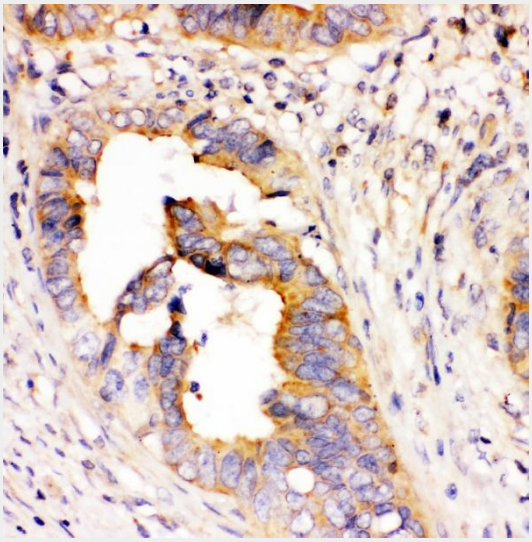

Figure 2. IHC analysis of DHFR using anti-DHFR antibody (M00813-1).
DHFR was detected in paraffin-embedded section of human rectal cancer tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 1 µg/ml mouse anti-DHFR Antibody (M00813-1) overnight at 4°C. Biotinylated goat anti-mouse IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using Strepavidin-Biotin-Complex (SABC) (Catalog # SA1021) with DAB as the chromogen.

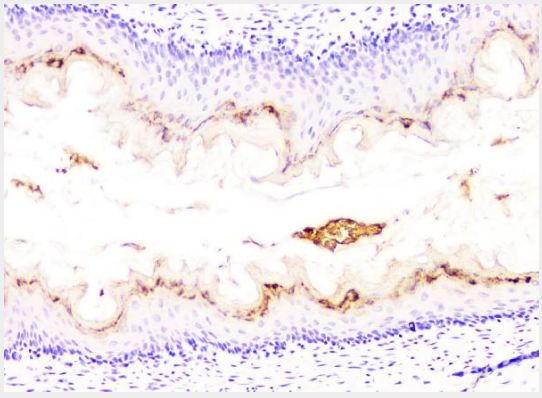

Figure 3. IHC analysis of DHFR using anti-DHFR antibody (M00813-1).
DHFR was detected in paraffin-embedded section of rat gaster tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 1 µg/ml mouse anti-DHFR Antibody (M00813-1) overnight at 4°C. Biotinylated goat anti-mouse IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using Strepavidin-Biotin-Complex (SABC) (Catalog # SA1021) with DAB as the chromogen.