



# FTCD rabbit pAb

|                           |   |
|---------------------------|---|
| <b>Catalog No</b>         | BYab-08520  |
| <b>Isotype</b>            | IgG   |
| <b>Reactivity</b>         | Human; Mouse;Rat  |
| <b>Applications</b>       | WB  |
| <b>Gene Name</b>          | FTCD  |
| <b>Protein Name</b>       | FTCD  |
| <b>Immunogen</b>          | Synthesized peptide derived from human FTCD AA range: 157-207   |
| <b>Specificity</b>        | This antibody detects endogenous levels of FTCD at Human/Mouse/Rat  |
| <b>Formulation</b>        | Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.   |
| <b>Source</b>             | Polyclonal, Rabbit,IgG  |
| <b>Purification</b>       | The antibody was affinity-purified from rabbit serum by affinity-chromatography using specific immunogen.   |
| <b>Dilution</b>           | WB 1: 500-2000  |
| <b>Concentration</b>      | 1 mg/ml   |
| <b>Purity</b>             | ≥90%  |
| <b>Storage Stability</b>  | -20°C/1 year  |
| <b>Synonyms</b>           |   |
| <b>Observed Band</b>      |   |
| <b>Cell Pathway</b>       | Cytoplasm, cytosol . Golgi apparatus . Cytoplasm, cytoskeleton, microtubule organizing center, centrosome, centriole . More abundantly located around the mother centriole. .   |
| <b>Tissue Specificity</b> |   |
| <b>Function</b>           | catalytic activity:5-formimidoyltetrahydrofolate + L-glutamate = tetrahydrofolate + N-formimidoyl-L-glutamate.,catalytic activity:5-formimidoyltetrahydrofolate = 5,10-methenyltetrahydrofolate + NH(3).,catalytic activity:5-formyltetrahydrofolate + L-glutamate = tetrahydrofolate + N-formyl-L-glutamate.,cofactor:Pyridoxal phosphate.,disease:Defects in FTCD are the cause of glutamate formiminotransferase deficiency [MIM:229100]; also known as formiminoglutamicaciduria (FIGLU-uria). It is an autosomal recessive disorder. Features of a severe phenotype, include elevated levels of formiminoglutamate (FIGLU) in the urine in response to histidine administration, megaloblastic anemia, and mental retardation. Features of a mild phenotype include high urinary excretion of FIGLU in the absence of histidine administration, mild developmental |

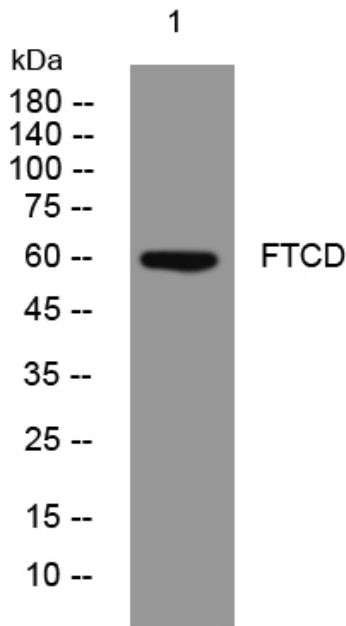
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delay, and no hematological abnormalities.,function:Folate-d

|                                  |  |
|----------------------------------|--|
| <b>Background</b>                | The protein encoded by this gene is a bifunctional enzyme that channels 1-carbon units from formiminoglutamate, a metabolite of the histidine degradation pathway, to the folate pool. Mutations in this gene are associated with glutamate formiminotransferase deficiency. Alternatively spliced transcript variants have been found for this gene.[provided by RefSeq, Dec 2009], |
| <b>matters needing attention</b> | Avoid repeated freezing and thawing!   |
| <b>Usage suggestions</b>         | This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.  |

## Products Images



Western blot analysis of lysates from HeLa cells, primary antibody was diluted at 1:1000, 4° over night

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