For Research Use Only
CoraLite®594-conjugated PHGDH
Monoclonal antibody
Catalog Number:CL594-67591

| Basic Information | Catalog Number: CL594-67591 | GenBank Accession Number: BC000303 | Purification Method: Protein G purification |
| :---: | :---: | :---: | :---: |
|  | $\begin{aligned} & \text { Size: } \\ & 1000 \mu \mathrm{~g} / \mathrm{ml} \end{aligned}$ | $\begin{aligned} & \text { GenelD (NCBI): } \\ & 26227 \end{aligned}$ | CloneNo.: 1E8B8 |
|  | Source: <br> Mouse | UNIPROT ID: $043175$ | Recommended Dilutions: <br> IF 1:50-1:500 |
|  | Isotype: $\operatorname{lgG1}$ | Full Name: phosphoglycerate dehydrogenase | Excitation/Emission maxima wavelengths: |
|  | Immunogen Catalog Number: AG6877 | Calculated MW: <br> 57 kDa | $588 \mathrm{~nm} / 604 \mathrm{~nm}$ |
|  |  | Observed MW: 57 kDa |  |

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Tested Applications:
IF/ICC
Species Specificity:
Human, Mouse, Rat

Applications

Background Information
PHGDH(D-3-phosphoglycerate dehydrogenase) is also named as 3-PGDH, PGDH3 and belongs to the D-isomer specific 2-hydroxyacid dehydrogenase family. It catalyzes the transition of 3-phosphoglycerate into 3phosphohydroxypyruvate, which is the first and rate-limiting step in the phosphorylated pathway of serine biosynthesis, using NAD+/NADH as a cofactor. 3-PGDH deficiency is a rare recessive inborn error in the biosynthesis of the amino acid L-serine characterized clinically by congenital microcephaly, psychomotor retardation, and intractable seizures(PMID:19235232 ).

Storage
Storage:
Store at $-20^{\circ} \mathrm{C}$. Avoid exposure to light.
Storage Buffer:
PBS with 50\% Glycerol, 0.05\% Proclin300, 0.5\% BSA, pH 7.3.
Aliquoting is unnecessary for $-20^{\circ} \mathrm{C}$ storage


Immunofluorescent analysis of ( $-20^{\circ} \mathrm{C}$ Ethanol)
fixed HeLa cells using CoraLite®594-conjugated PHGDH antibody (CL594-67591, Clone: 1E8B8) at dilution of 1:10.

