

# **Product datasheet for TA501890**

#### OriGene Technologies, Inc.

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## **GRHPR Mouse Monoclonal Antibody [Clone ID: OTI4B6]**

#### **Product data:**

**Product Type:** Primary Antibodies

Clone Name: OTI4B6

**Applications:** FC, IHC, WB

**Recommended Dilution:** WB: 1:2000, FLOW: 1:100, IHC: 1:50-1:150

Reactivity: Human, Mouse, Rat

Host: Mouse Isotype: IgG2a

Clonality: Monoclonal

Immunogen: Full length human recombinant protein of human GRHPR (NP\_036335) produced in HEK293T

cell

**Formulation:** PBS (pH 7.3) containing 1% BSA, 50% glycerol and 0.02% sodium azide.

**Concentration:** 0.84 mg/ml

**Purification:** Purified from mouse ascites fluids or tissue culture supernatant by affinity chromatography

(protein A/G)

Conjugation: Unconjugated

**Storage:** Store at -20°C as received.

**Stability:** Stable for 12 months from date of receipt.

**Predicted Protein Size:** 35.5 kDa

**Gene Name:** glyoxylate and hydroxypyruvate reductase

Database Link: NP 036335

Entrez Gene 680021 RatEntrez Gene 9380 Human

Q9UBQ7

**Background:** This gene encodes an enzyme with hydroxypyruvate reductase, glyoxylate reductase, and D-

glycerate dehydrogenase enzymatic activities. The enzyme has widespread tissue expression and has a role in metabolism. Type II hyperoxaluria is caused by mutations in this gene.

[provided by RefSeq]

Synonyms: GLXR; GLYD; PH2

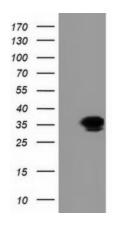


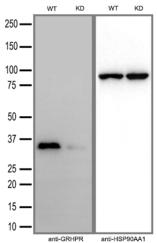


**Protein Families:** Druggable Genome

**Protein Pathways:** Glyoxylate and dicarboxylate metabolism, Metabolic pathways, Pyruvate metabolism

### **Product images:**

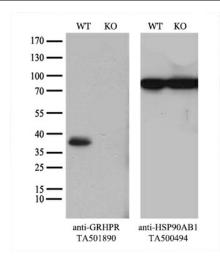




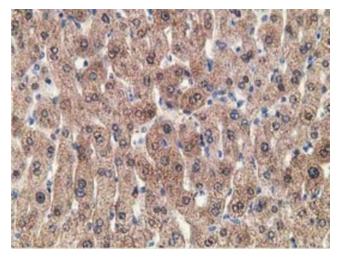
HEK293T cells were transfected with the pCMV6-ENTRY control (Left lane) or pCMV6-ENTRY GRHPR ([RC200963], Right lane) cDNA for 48 hrs and lysed. Equivalent amounts of cell lysates (5 ug per lane) were separated by SDS-PAGE and immunoblotted with anti-GRHPR. Positive lysates [LY415912] (100ug) and [LC415912] (20ug) can be purchased separately from OriGene.

Equivalent amounts of cell lysates (30 ug per lane) of wild-type HeLa cells (WT) and GRHPR-Knockdown HeLa cells (KD) were separated by SDS-PAGE and immunoblotted with anti-GRHPR monoclonal antibody TA501890 (1:5000). Then the blotted membrane was stripped and reprobed with anti-HSP90AA1 antibody as a loading control.

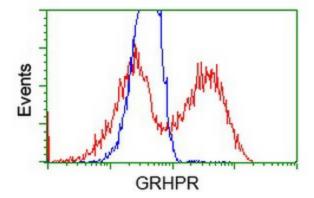




Equivalent amounts of cell lysates (10 ug per lane) of wild-type 293T cells (WT, Cat# LC810293T) and GRHPR-Knockout 293T cells (KO, Cat# [LC841943] ) were separated by SDS-PAGE and immunoblotted with anti-GRHPR monoclonal antibody TA501890 (1:200`). Then the blotted membrane was stripped and reprobed with anti-HSP90 antibody as a loading control.

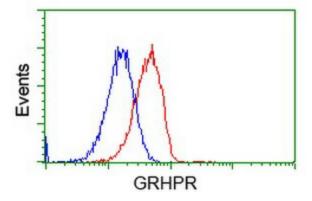


Immunohistochemical staining of paraffinembedded Human liver tissue within the normal limits using anti-GRHPR mouse monoclonal antibody at 1:150 (TA501890). Heat-induced epitope retrieval by EDTA solution buffer pH 8.0 at 120°C for 3 min.



HEK293T cells transfected with either [RC200963] overexpress plasmid (Red) or empty vector control plasmid (Blue) were immunostained by anti-GRHPR antibody (TA501890), and then analyzed by flow cytometry.





Flow cytometric Analysis of Hela cells, using anti-GRHPR antibody (TA501890), (Red), compared to a nonspecific negative control antibody (TA50011), (Blue).