Anti-GST3 Antibody [JB40-79]

ET7107-71



| Product Type: | Recombinant Rabbit monoclonal IgG, primary antibodies |
|---|--|
| Species reactivity: | Human, Mouse, Rat |
| Applications: | WB, IHC-P, IF-Cell, IF-Tissue |
| Molecular Wt: | Predicted band size: 23 kDa |
| Clone number: | JB40-79 |
| Description: | Glutathione S-transferases (GSTs) function in the metabolic detoxification of various environmental carcinogens and lipid hydroperoxides. In response to oxidative stress, upregulation of the GST family member GSTP1 occurs, consistent with this function. Furthermore, the GSTP1 gene is subject to CpG island hypermethylation, a state that correlates with human prostatic carcinogenesis. GSTP1 gene hypermethylation can be detected in urine, ejaculate and plasma from men with prostate cancer, potentially making GSTP1 a useful biomarker for prostate cancer screening. |
| lmmunogen: | Recombinant protein within Human GST3aa 70-210 / 210. |
| Positive control: | A549 cell lysate, mouse liver tissue lysate, A431, A549, LOVO, rat epididymis tissue, human liver tissue, human placenta tissue, mouse fallopian tube tissue,human lung tissue,human lung squamous carcinoma tissue. |
| Subcellular location: | Cytoplasm, Mitochondrion, Nucleus. |
| Database links: | SwissProt: P09211 Human P19157 Mouse P04906 Rat |
| Recommended Dilutions: WB IF-Cell IF-Tissue IHC-P | 1:500-1:1,000 1:50-1:200 1:50-1:200 1:500 |
| Storage Buffer: | 1*TBS (pH7.4), 0.05% BSA, 40% Glycerol. Preservative: 0.05% Sodium Azide. |
| Storage Instruction: | Shipped at 4 $^\circ\!\!\mathbb{C}$. Store at +4 $^\circ\!\!\mathbb{C}$ short term (1-2 weeks). It is recommended to aliquot into single-use upon delivery. Store at -20 $^\circ\!\!\mathbb{C}$ long term. |
| Purity: | Protein A affinity purified. |

Hangzhou Huaan Biotechnology Co., Ltd.

Orders:0086-571-88062880

Technical:0086-571-89986345

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Applications:WB=Western blot IHC-P=Immunohistochemistry (paraffin) IF-Cell=Immunofluorescence (Cell) IF-Tissue=Immunofluorescence (Tissue) FC=Flow cytometry IP=Immunoprecipitation

Images

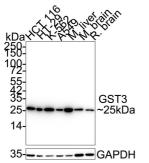


Fig1: Western blot analysis of GST3 on different lysates with Rabbit anti-GST3 antibody (ET7107-71) at 1/1,000 dilution.

Lane 1: HCT 116 cell lysate Lane 1: HT-29 cell lysate Lane 1: K-562 cell lysate Lane 1: A549 cell lysate Lane 2: Mouse liver tissue lysate Lane 2: Mouse brain tissue lysate Lane 2: Rat brain tissue lysate

Lysates/proteins at 20 µg/Lane1-4 and 40ug/Lane5-7.

Predicted band size: 23 kDa Observed band size: 25 kDa

Exposure time: 8 seconds; ECL: K1801;

4-20% SDS-PAGE gel.

Proteins were transferred to a PVDF membrane and blocked with 5% NFDM/TBST for 1 hour at room temperature. The primary antibody (ET7107-71) at 1/1,000 dilution was used in 5% NFDM/TBST at 4° C overnight. Goat Anti-Rabbit IgG - HRP Secondary Antibody (HA1001) at 1/50,000 dilution was used for 1 hour at room temperature.

Fig2: Western blot analysis of GST3 on different lysates with Rabbit anti-GST3 antibody (ET7107-71) at 1/1,000 dilution.

Lane 1: HCT 116-si NT cell lysate Lane 2: HCT 116-si GST3 cell lysate

Lysates/proteins at 10 µg/Lane.

Predicted band size: 23 kDa Observed band size: 23 kDa

Exposure time: 8 seconds; ECL: K1801;

4-20% SDS-PAGE gel.

Proteins were transferred to a PVDF membrane and blocked with 5% NFDM/TBST for 1 hour at room temperature. The primary antibody (ET7107-71) at 1/1,000 dilution was used in 5% NFDM/TBST at 4° C overnight. Goat Anti-Rabbit IgG - HRP Secondary Antibody (HA1001) at 1/50,000 dilution was used for 1 hour at room temperature.

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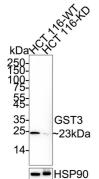
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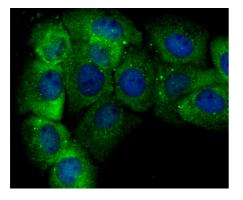


Fig3: ICC staining of GST3 in A431 cells (green). Formalin fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 10 minutes at room temperature and blocked with 1% Blocker BSA for 15 minutes at room temperature. Cells were probed with the primary antibody (ET7107-71, 1/50) for 1 hour at room temperature, washed with PBS. Alexa Fluor®488 Goat anti-Rabbit IgG was used as the secondary antibody at 1/1,000 dilution. The nuclear counter stain is DAPI (blue).

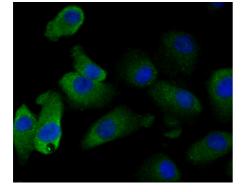


Fig4: ICC staining of GST3 in A549 cells (green). Formalin fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 10 minutes at room temperature and blocked with 1% Blocker BSA for 15 minutes at room temperature. Cells were probed with the primary antibody (ET7107-71, 1/50) for 1 hour at room temperature, washed with PBS. Alexa Fluor®488 Goat anti-Rabbit IgG was used as the secondary antibody at 1/1,000 dilution. The nuclear counter stain is DAPI (blue).

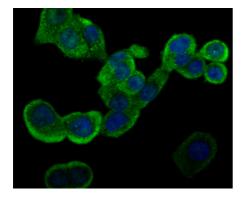


Fig5: ICC staining of GST3 in LOVO cells (green). Formalin fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 10 minutes at room temperature and blocked with 1% Blocker BSA for 15 minutes at room temperature. Cells were probed with the primary antibody (ET7107-71, 1/50) for 1 hour at room temperature, washed with PBS. Alexa Fluor®488 Goat anti-Rabbit IgG was used as the secondary antibody at 1/1,000 dilution. The nuclear counter stain is DAPI (blue).

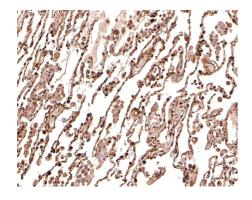


Fig6: Immunohistochemical analysis of paraffin-embedded human lung tissue with Rabbit anti-GST3 antibody (ET7107-71) at 1/500 dilution.

The section was pre-treated using heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0) for 20 minutes. The tissues were blocked in 1% BSA for 20 minutes at room temperature, washed with ddH₂O and PBS, and then probed with the primary antibody (ET7107-71) at 1/500 dilution for 1 hour at room temperature. The detection was performed using an HRP conjugated compact polymer system. DAB was used as the chromogen. Tissues were counterstained with hematoxylin and mounted with DPX.

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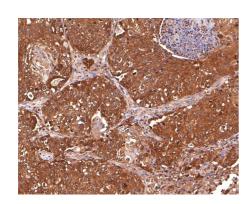


Fig7: Immunohistochemical analysis of paraffin-embedded human lung squamous carcinoma tissue with Rabbit anti-GST3 antibody (ET7107-71) at 1/500 dilution.

The section was pre-treated using heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0) for 20 minutes. The tissues were blocked in 1% BSA for 20 minutes at room temperature, washed with ddH₂O and PBS, and then probed with the primary antibody (ET7107-71) at 1/500 dilution for 1 hour at room temperature. The detection was performed using an HRP conjugated compact polymer system. DAB was used as the chromogen. Tissues were counterstained with hematoxylin and mounted with DPX.

Note: All products are "FOR RESEARCH USE ONLY AND ARE NOT INTENDED FOR DIAGNOSTIC OR THERAPEUTIC USE".

Background References

- 1. Sun K H et al. Glutathione-S-transferase P1 is a critical regulator of Cdk5 kinase activity. J Neurochem 118:902-914 (2011).
- 2. Kong K H et al. Tyrosine-7 in human class Pi glutathione S-transferase is important for lowering the pKa of the thiol group of glutathione in the enzyme-glutathione complex. Biochem Biophys Res Commun 184:194-197 (1992).

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