

Anti-ABAT antibody (1-100) [S7MR] (STJ11102437)

GENERAL INFORMATION

Product Type	Primary antibodies
Applications	WB/IHC-P/ELISA
Host / Source	Rabbit
Reactivity	Human/Mouse/Rat

PRODUCT PROPERTIES

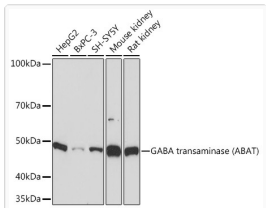
Clonality	Monoclonal
Concentration	Lot specific
Conjugation	Unconjugated
Purification	Affinity purification
Dilution Range	WB:1:500-1:2000 IHC-P:1:50-1:200 ELISA:Recommended starting concentration is 1 µg/mL. Please optimize the concentration based on your specific assay requirements.
Formulation	PBS with 0.02% Sodium Azide, 0.05% BSA, 50% Glycerol, pH 7.3.
Isotype	IgG
Molecular Weight	Protein Mw: 56kDa Observed Mw:
Storage Instruction	Store at -20°C for up to 1 year from the date of receipt, and avoid repeat freeze-thaw cycles.

TARGET INFORMATION

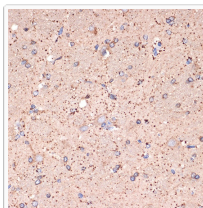
Gene ID	18
Gene Symbol	ABAT
UniProt ID	GABT_HUMAN
Immunogen Region	1-100
Immunogen Sequence	MASMLLAQRLACSFQHSYRL LVPGSRHISQAAAKVDVEFD YDGPLMKTEVPGPRSQELMK QLNIIQNAEAVHFFCNYEES RGNLYLDVDGNRMLDLYSQI
Specificity	A synthetic peptide corresponding to a sequence within amino acids 1-100 of human GABA transaminase (ABAT) (P80404).

ADDITIONAL INFORMATION

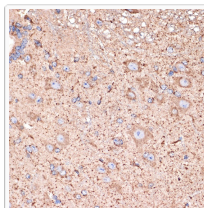
Note	STRICTLY FOR FURTHER SCIENTIFIC RESEARCH USE ONLY (RUO). MUST NOT TO BE USED IN DIAGNOSTIC OR THERAPEUTIC APPLICATIONS.
------	---



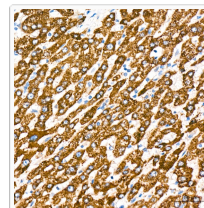
Western blot analysis of various lysates using GABA transaminase (GABA transaminase (ABAT)) Rabbit mAb (STJ11102437) at 1:1000 dilution. Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (STJS000856) at 1:10000 dilution. Lysates/proteins: 25 Mu g per lane. Blocking buffer: 3% nonfat dry milk in TBST. Detection: ECL Basic Kit Exposure time: 10s.



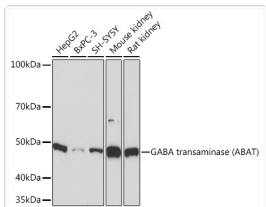
Immunohistochemistry analysis of paraffin-embedded Rat brain using GABA transaminase (GABA transaminase (ABAT)) Rabbit mAb (STJ11102437) at dilution of 1:100 (40x lens). Microwave antigen retrieval performed with 0. 01M Tris/EDTA Buffer (pH 9. 0) prior to immunohistochemistry staining.



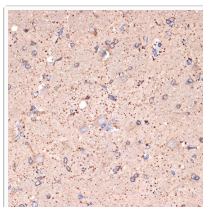
Immunohistochemistry analysis of paraffin-embedded Mouse spinal cord using GABA transaminase (GABA transaminase (ABAT)) Rabbit mAb (STJ11102437) at dilution of 1:100 (40x lens). Microwave antigen retrieval performed with 0. 01M Tris/EDTA Buffer (pH 9. 0) prior to immunohistochemistry staining.



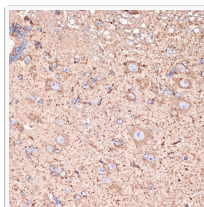
Immunohistochemistry analysis of paraffin-embedded Human liver tissue using GABA transaminase (ABAT) Rabbit mAb (STJ11102437) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0. 01 M citrate buffer (pH 6. 0) prior to immunohistochemistry staining.



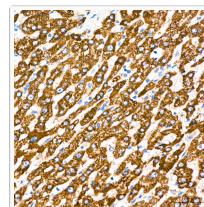
Western blot analysis of various lysates using GABA transaminase (GABA transaminase (ABAT)) Rabbit mAb (STJ11102437) at 1:1000 dilution. Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) (STJS000856) at 1:10000 dilution. Lysates/proteins: 25 Mu g per lane. Blocking buffer: 3% nonfat dry milk in TBST. Detection: ECL Basic Kit Exposure time: 10s.



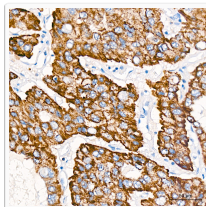
Immunohistochemistry analysis of paraffin-embedded Rat brain using GABA transaminase (GABA transaminase (ABAT)) Rabbit mAb (STJ11102437) at dilution of 1:100 (40x lens). Perform microwave antigen retrieval with 10 mM Tris/EDTA buffer pH 9. 0 before commencing with immunohistochemistry staining protocol.



Immunohistochemistry analysis of paraffin-embedded Mouse spinal cord using GABA transaminase (GABA transaminase (ABAT)) Rabbit mAb (STJ11102437) at dilution of 1:100 (40x lens). Perform microwave antigen retrieval with 10 mM Tris/EDTA buffer pH 9. 0 before commencing with immunohistochemistry staining protocol.



Immunohistochemistry analysis of paraffin-embedded Human liver tissue using GABA transaminase (ABAT) Rabbit mAb (STJ11102437) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0. 01 M citrate buffer (pH 6. 0) prior to immunohistochemistry staining.



Immunohistochemistry analysis of paraffin-embedded Human liver cancer tissue using GABA transaminase (ABAT) Rabbit mAb (STJ11102437) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0. 01 M citrate buffer (pH 6. 0) prior to immunohistochemistry staining.