

Anti-UBE2I antibody (1-100) [S9MR] (STJ11102069)

GENERAL INFORMATION

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

PRODUCT PROPERTIES

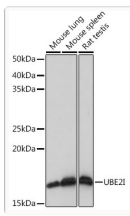
Clonality	Monoclonal
Concentration	Lot specific
Conjugation	Unconjugated
Purification	Affinity purification
Dilution Range	WB:1:500-1:1000 IHC-P:1:50-1:200 ELISA:Recommended starting concentration is 1 Mu g/mL. Please optimize the concentration based on your specific assay requirements.
Formulation	PBS with 0.02% Sodium Azide, 0.05% BSA, 50% Glycerol, pH 7.3.
Isotype	IgG
Molecular Weight	Protein Mw: 18kDa Observed Mw: 18kDa
Storage Instruction	Store at -20°C for up to 1 year from the date of receipt, and avoid repeat freeze-thaw cycles.

TARGET INFORMATION

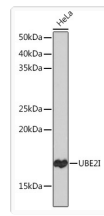
Gene ID	7329
Gene Symbol	UBE2I
UniProt ID	UBC9_HUMAN
Immunogen Region	1-100
Immunogen Sequence	MSGIALSRLAQRKAWRKDH PFGFVAVPTKNPDGTMNLMN WECAIPGKKGTPWEGGLFKL RMLFKDDYPSSPPKCKFEPP LFHPNVYPSGTVCLSILEED
Specificity	A synthetic peptide corresponding to a sequence within amino acids 1-100 of human UBE2I (P63279).

ADDITIONAL INFORMATION

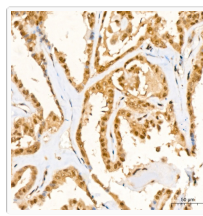
Note	STRICTLY FOR FURTHER SCIENTIFIC RESEARCH USE ONLY (RUO). MUST NOT TO BE USED IN DIAGNOSTIC OR THERAPEUTIC APPLICATIONS.
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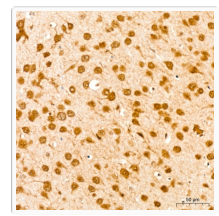
Western blot analysis of extracts of various cell lines, using UBE2I rabbit monoclonal antibody (STJ11102069) at 1:1000 dilution. Secondary antibody: HRP Goat Anti-rabbit IgG (H+L) (STJS000856) at 1:10000 dilution. Lysates/proteins: 25 μ g per lane. Blocking buffer: 3% non-fat dry milk in TBST. Detection: ECL Basic Kit. Exposure time: 1s.



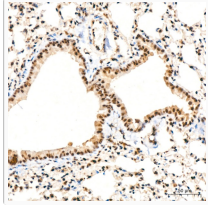
Western blot analysis of extracts of HeLa cells, using UBE2I rabbit monoclonal antibody (STJ11102069) at 1:1000 dilution. Secondary antibody: HRP Goat Anti-rabbit IgG (H+L) (STJS000856) at 1:10000 dilution. Lysates/proteins: 25 μ g per lane. Blocking buffer: 3% non-fat dry milk in TBST. Detection: ECL Basic Kit. Exposure time: 30s.



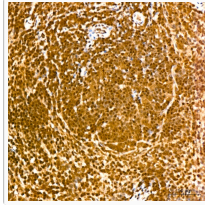
Immunohistochemistry analysis of UBE2I in paraffin-embedded human thyroid cancer tissue using UBE2I rabbit monoclonal antibody (STJ11102069) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to immunohistochemistry staining.



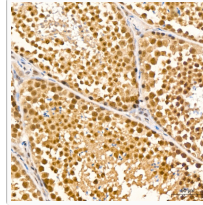
Immunohistochemistry analysis of UBE2I in paraffin-embedded mouse brain tissue using UBE2I rabbit monoclonal antibody (STJ11102069) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to immunohistochemistry staining.



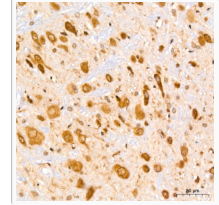
Immunohistochemistry analysis of UBE2I in paraffin-embedded mouse lung tissue using UBE2I rabbit monoclonal antibody (STJ11102069) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to immunohistochemistry staining.



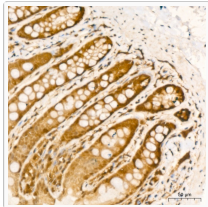
Immunohistochemistry analysis of UBE2I in paraffin-embedded mouse spleen tissue using UBE2I rabbit monoclonal antibody (STJ11102069) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to immunohistochemistry staining.



Immunohistochemistry analysis of UBE2I in paraffin-embedded mouse testis tissue using UBE2I rabbit monoclonal antibody (STJ11102069) 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 UBE2I in paraffin-embedded rat brain tissue using UBE2I rabbit monoclonal antibody (STJ11102069) 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 UBE2I in paraffin-embedded rat colon tissue using UBE2I rabbit monoclonal antibody (STJ11102069) 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.