

Product Datasheet

ARID1A Antibody (JJ09-01)

NBP2-67886

Unit Size: 100 ul

Store at 4C short term. Aliquot and store at -20C long term. Avoid freeze-thaw cycles.

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NBP2-67886

ARID1A Antibody (JJ09-01)

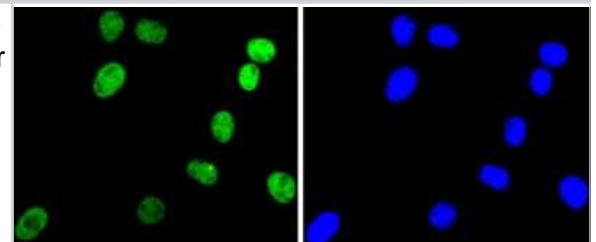
Product Information	
Unit Size	100 ul
Concentration	1 mg/ml
Storage	Store at 4C short term. Aliquot and store at -20C long term. Avoid freeze-thaw cycles.
Clonality	Monoclonal
Clone	JJ09-01
Preservative	0.05% Sodium Azide
Isotype	IgG
Purity	Protein A purified
Buffer	TBS (pH7.4), 0.05% BSA, 40% Glycerol

Product Description	
Host	Rabbit
Gene Symbol	ARID1A
Species	Human, Mouse, Rat
Immunogen	Recombinant protein within Human ARID1A aa 1,184-1,334 / 2,285. (SwissProt: O14497 Human; SwissProt: A2BH40 Mouse; Unigene: 61077 Rat)

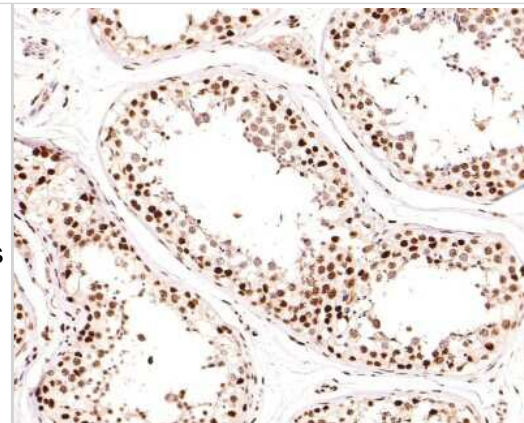
Product Application Details	
Applications	Immunocytochemistry/ Immunofluorescence, Immunohistochemistry, Immunohistochemistry-Paraffin
Recommended Dilutions	Immunohistochemistry, Immunocytochemistry/ Immunofluorescence 1:50-1:200, Immunohistochemistry-Paraffin 1:50-1:400

Images

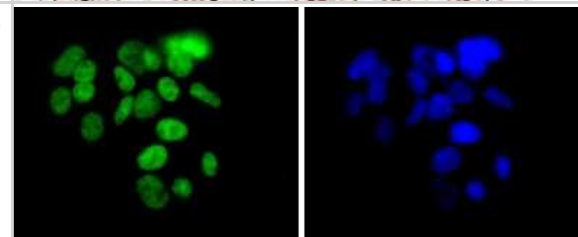
Immunocytochemistry/Immunofluorescence: ARID1A Antibody (JJ09-01) [NBP2-67886] - Staining ARID1A in SH-SY-5Y cells (green). The nuclear counter stain is DAPI (blue). Cells were fixed in paraformaldehyde, permeabilised with 0.25% Triton X100/PBS.



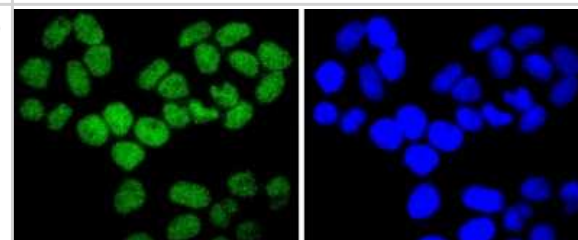
Immunohistochemistry-Paraffin: ARID1A Antibody (JJ09-01) [NBP2-67886] - Analysis of paraffin-embedded rat testis tissue using anti-ARID1A antibody. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0) for 20 minutes. The tissues were blocked in 1% BSA for 30 minutes at room temperature, washed with ddH₂O and PBS, and then probed with the primary antibody (1/100) for 30 minutes 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.



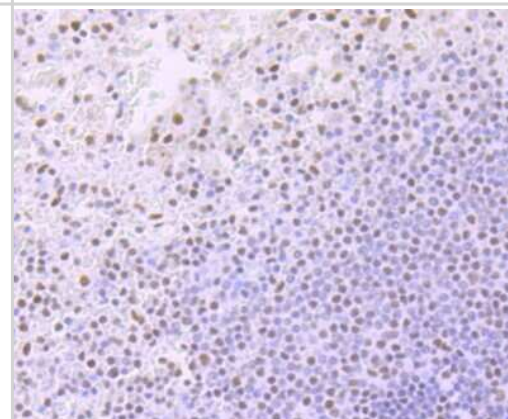
Immunocytochemistry/Immunofluorescence: ARID1A Antibody (JJ09-01) [NBP2-67886] - Staining ARID1A in 293 cells (green). The nuclear counter stain is DAPI (blue). Cells were fixed in paraformaldehyde, permeabilised with 0.25% Triton X100/PBS.



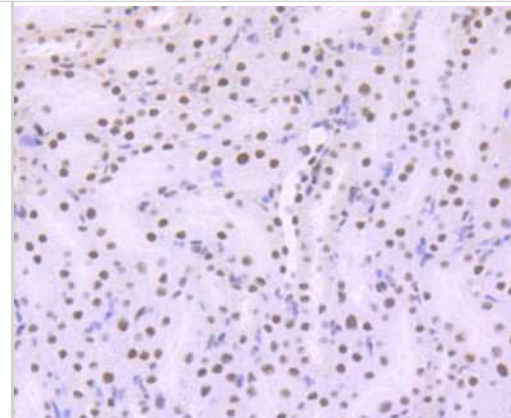
Immunocytochemistry/Immunofluorescence: ARID1A Antibody (JJ09-01) [NBP2-67886] - Staining ARID1A in Hela cells (green). The nuclear counter stain is DAPI (blue). Cells were fixed in paraformaldehyde, permeabilised with 0.25% Triton X100/PBS.



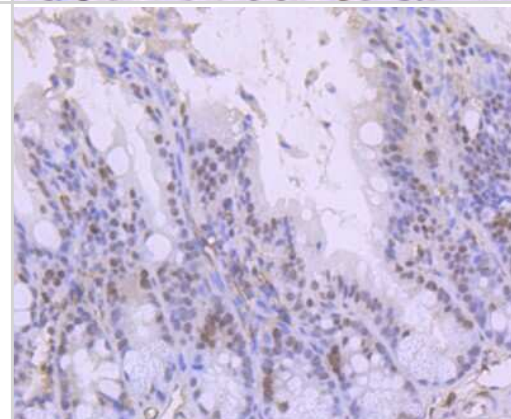
Immunohistochemistry-Paraffin: ARID1A Antibody (JJ09-01) [NBP2-67886] - Analysis of paraffin-embedded human spleen tissue using anti-ARID1A antibody. Counter stained with hematoxylin.



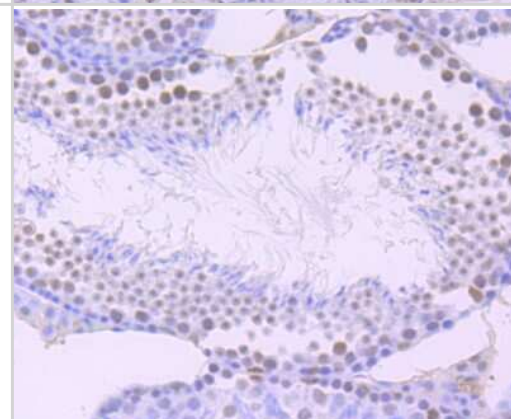
Immunohistochemistry-Paraffin: ARID1A Antibody (JJ09-01) [NBP2-67886] - Analysis of paraffin-embedded mouse brain tissue using anti-ARID1A antibody. Counter stained with hematoxylin.



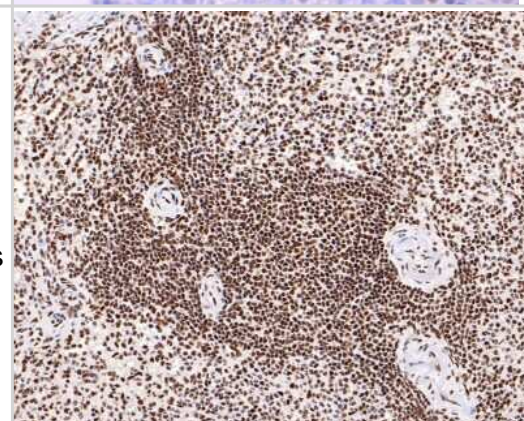
Immunohistochemistry-Paraffin: ARID1A Antibody (JJ09-01) [NBP2-67886] - Analysis of paraffin-embedded mouse colon tissue using anti-ARID1A antibody. Counter stained with hematoxylin.



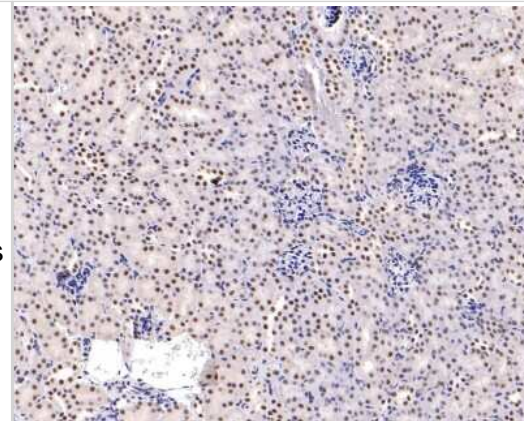
Immunohistochemistry-Paraffin: ARID1A Antibody (JJ09-01) [NBP2-67886] - Analysis of paraffin-embedded mouse testis tissue using anti-ARID1A antibody. Counter stained with hematoxylin.



Immunohistochemistry-Paraffin: ARID1A Antibody (JJ09-01) [NBP2-67886] - Analysis of paraffin-embedded human spleen tissue using anti-ARID1A antibody. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0) for 20 minutes. The tissues were blocked in 1% BSA for 30 minutes at room temperature, washed with ddH₂O and PBS, and then probed with the primary antibody (1/400) for 30 minutes 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.



Immunohistochemistry-Paraffin: ARID1A Antibody (JJ09-01) [NBP2-67886] - Analysis of paraffin-embedded mouse kidney tissue using anti-ARID1A antibody. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0) for 20 minutes. The tissues were blocked in 1% BSA for 30 minutes at room temperature, washed with ddH₂O and PBS, and then probed with the primary antibody (1/400) for 30 minutes 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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