

Product datasheet

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ARG43066 anti-U2AF2 / U2AF65 antibody

Package: 50 μg Store at: -20°C

Summary

Product Description Rabbit Polyclonal antibody recognizes U2AF2 / U2AF65

Tested Reactivity Hu, Ms, Rat

Tested Application FACS, ICC/IF, IHC-P, WB

Host Rabbit

Clonality Polyclonal

Isotype IgG

Target Name U2AF2 / U2AF65

Species Human

Immunogen Recombinant protein corresponding to M238-H470 of Human U2AF2 / U2AF65.

Conjugation Un-conjugated

Alternate Names U2 auxiliary factor 65 kDa subunit; U2AF65; U2 snRNP auxiliary factor large subunit; Splicing factor

U2AF 65 kDa subunit; 65; hU2AF; hU2AF65

Application Instructions

Application table	Application	Dilution
	FACS	1:150 - 1:500
	ICC/IF	1:200 - 1:1000
	IHC-P	1:200 - 1:1000
	WB	1:500 - 1:2000
Application Note	IHC-P: Antigen Retrieval: Heat mediation was performed in EDTA buffer (pH 8.0). * The dilutions indicate recommended starting dilutions and the optimal dilutions or concentrations should be determined by the scientist.	

Properties

Form	Liquid	
Purification	Affinity purification with immunogen.	
Buffer	0.2% Na2HPO4, 0.9% NaCl, 0.05% Sodium azide and 4% Trehalose.	
Preservative	0.05% Sodium azide	
Stabilizer	4% Trehalose	
Concentration	0.5 mg/ml	
Storage instruction	For continuous use, store undiluted antibody at 2-8°C for up to a week. For long-term storage, aliquot and store at -20°C or below. Storage in frost free freezers is not recommended. Avoid repeated freeze/thaw cycles. Suggest spin the vial prior to opening. The antibody solution should be gently mixed	

For laboratory research only, not for drug, diagnostic or other use.

Bioinformation

Gene Symbol U2AF2

Gene Full Name U2 small nuclear RNA auxiliary factor 2

Background U2 auxiliary factor (U2AF), comprised of a large and a small subunit, is a non-snRNP protein required for

the binding of U2 snRNP to the pre-mRNA branch site. This gene encodes the U2AF large subunit which contains a sequence-specific RNA-binding region with 3 RNA recognition motifs and an Arg/Ser-rich domain necessary for splicing. The large subunit binds to the polypyrimidine tract of introns early during spliceosome assembly. Multiple transcript variants have been detected for this gene, but the full-

length natures of only two have been determined to date. [provided by RefSeq, Jul 2008]

Function Plays a role in pre-mRNA splicing and 3'-end processing (PubMed:17024186). By recruiting PRPF19 and

the PRP19C/Prp19 complex/NTC/Nineteen complex to the RNA polymerase II C-terminal domain (CTD), and thereby pre-mRNA, may couple transcription to splicing (PubMed:21536736). Induces cardiac troponin-T (TNNT2) pre-mRNA exon inclusion in muscle. Regulates the TNNT2 exon 5 inclusion through competition with MBNL1. Binds preferentially to a single-stranded structure within the polypyrimidine tract of TNNT2 intron 4 during spliceosome assembly. Required for the export of mRNA out of the nucleus, even if the mRNA is encoded by an intron-less gene. Represses the splicing of MAPT/Tau exon 10. Positively regulates pre-mRNA 3'-end processing by recruiting the CFIm complex to cleavage and

polyadenylation signals (PubMed:17024186). [UniProt]

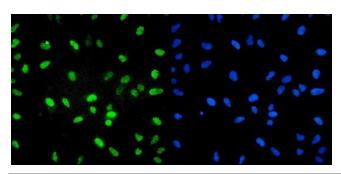
Calculated Mw 54 kDa

PTM Lysyl-hydroxylation at Lys-15 and Lys-276 affects the mRNA splicing activity of the protein, leading to

regulate some, but not all, alternative splicing events. [UniProt]

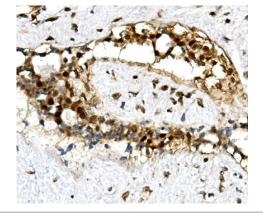
Cellular Localization Nucleus. [UniProt]

Images



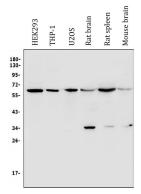
ARG43066 anti-U2AF2 / U2AF65 antibody ICC/IF image

Immunofluorescence: HeLa cells were blocked with 10% goat serum and then stained with ARG43066 anti-U2AF2 / U2AF65 antibody (green) at 2 μ g/ml dilution, overnight at 4°C. DAPI (blue) for nuclear staining.



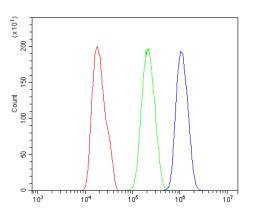
ARG43066 anti-U2AF2 / U2AF65 antibody IHC-P image

Immunohistochemistry: Paraffin-embedded Human mammary cancer tissue. Antigen Retrieval: Heat mediation was performed in EDTA buffer (pH 8.0). The tissue section was blocked with 10% goat serum. The tissue section was then stained with ARG43066 anti-U2AF2 / U2AF65 antibody at 1 μ g/ml dilution, overnight at 4°C.



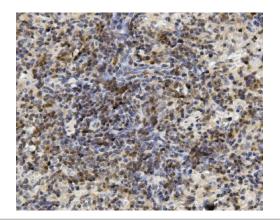
ARG43066 anti-U2AF2 / U2AF65 antibody WB image

Western blot: 50 μg of sample under reducing conditions. HEK293, THP-1, U2OS, Rat brain, Rat spleen and Mouse brain lysates stained with ARG43066 anti-U2AF2 / U2AF65 antibody at 0.5 $\mu g/ml$ dilution, overnight at 4°C.



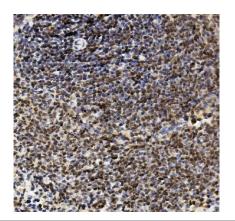
ARG43066 anti-U2AF2 / U2AF65 antibody FACS image

Flow Cytometry: PC-3 cells were blocked with 10% normal goat serum and then stained with ARG43066 anti-U2AF2 / U2AF65 antibody (blue) at 1 $\mu g/10^{\circ}6$ cells for 30 min at 20°C, followed by incubation with DyLight®488 labelled secondary antibody. Isotype control antibody (green) was rabbit IgG (1 $\mu g/10^{\circ}6$ cells) used under the same conditions. Unlabelled sample (red) was also used as a control.



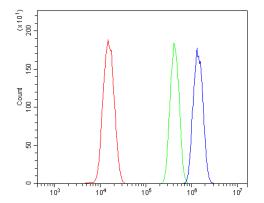
ARG43066 anti-U2AF2 / U2AF65 antibody IHC-P image

Immunohistochemistry: Paraffin-embedded Mouse spleen tissue. Antigen Retrieval: Heat mediation was performed in EDTA buffer (pH 8.0). The tissue section was blocked with 10% goat serum. The tissue section was then stained with ARG43066 anti-U2AF2 / U2AF65 antibody at 1 $\mu g/ml$ dilution, overnight at 4°C.



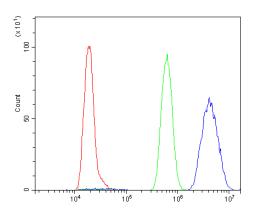
ARG43066 anti-U2AF2 / U2AF65 antibody IHC-P image

Immunohistochemistry: Paraffin-embedded Rat spleen tissue. Antigen Retrieval: Heat mediation was performed in EDTA buffer (pH 8.0). The tissue section was blocked with 10% goat serum. The tissue section was then stained with ARG43066 anti-U2AF2 / U2AF65 antibody at 1 μ g/ml dilution, overnight at 4°C.



ARG43066 anti-U2AF2 / U2AF65 antibody FACS image

Flow Cytometry: ANA-1 cells were blocked with 10% normal goat serum and then stained with ARG43066 anti-U2AF2 / U2AF65 antibody (blue) at 1 $\mu g/10^{\circ}6$ cells for 30 min at 20°C, followed by incubation with DyLight®488 labelled secondary antibody. Isotype control antibody (green) was rabbit IgG (1 $\mu g/10^{\circ}6$ cells) used under the same conditions. Unlabelled sample (red) was also used as a control.



ARG43066 anti-U2AF2 / U2AF65 antibody FACS image

Flow Cytometry: NRK cells were blocked with 10% normal goat serum and then stained with ARG43066 anti-U2AF2 / U2AF65 antibody (blue) at 1 $\mu g/10^6$ cells for 30 min at 20°C, followed by incubation with DyLight®488 labelled secondary antibody. Isotype control antibody (green) was rabbit IgG (1 $\mu g/10^6$ cells) used under the same conditions. Unlabelled sample (red) was also used as a control.