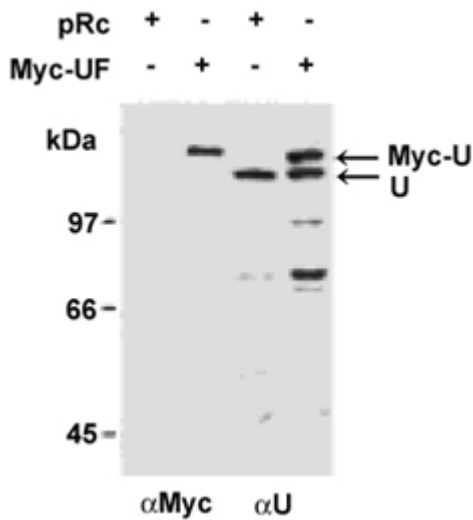


**anti-hnRNP-U / SAF-A antibody, rabbit serum**

<b>Product code</b>	70-415
<b>Size</b>	100 µl
<b>Storage</b>	Store 4°C for short term For long term storage store at -20°C. Aliquot to avoid repeated freezing and thawing.
<b>Concentration</b>	N/A
<b>Buffer</b>	0.05% sodium azide
<b>Purity</b>	Rabbit antiserum
<b>Immunogen</b>	Recombinant MBT-fused mouse hnRNP-U (aa 614-800)
<b>Isotype</b>	Rabbit IgG
<b>Reactivity</b>	Reacts with mouse and rat, and predicted to react with human from the amino acid sequence homology.
<b>Special notes</b>	N/A
<b>Application</b>	1. Western blotting (dilution: 1/3,000-1/1,000) 2. Immunocytochemistry (dilution: 1/1,000-1/500) 3. Immunoprecipitation
<b>Background</b>	<p><b>Heterogeneous nuclear ribonucleoprotein U (hnRNP-U</b>, also known as <b>scaffold attachment factor A, SAF-A</b>) is a nuclear matrix-associated protein that interacts with chromosomal DNA. <b>hnRNP-U</b> specifically binds to scaffold/matrix attachment region of DNA and could thus be involved in higher order chromatin structure. <b>hnRNP-U</b> is also a RNA binding protein and forms complexes with heterogeneous nuclear RNA (hnRNA) and plays an important role in pre-mRNA processing and transport.</p> <p><b>HnRNP-U</b> is reported to interact with necdin, a growth suppressor that is expressed in terminally differentiated neurons and skeletal muscle cells. It has been shown that <b>hnRNP-U</b> recruits necdin to the nuclear matrix where they form a stable complex. It is suggested that necdin suppresses cell proliferation through its interaction with <b>hnRNP-U</b> in the specific subnuclear structure (ref.2).</p>
<b>Data Link</b>	<a href="#">Q8VEK3</a> (mouse), <a href="#">Q00839</a> (human)
Please note: All products are FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES. NOT FOR MILITARY USE.	

Data Images: 70-415 anti-hnRNP-U / SAF-A antibody, rabbit serum

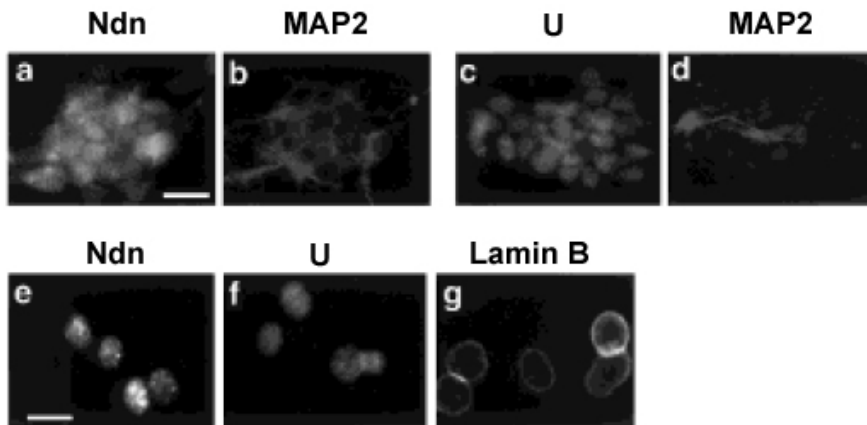


**Fig.1 Immunoblotting of hnRNP-U with this antibody (ref.2).**

Specificity of anti-hnRNP-U antibody, HUT.

Cell lysates were prepared from SAOS-2 cells transfected with pRc/CMV vectors (pRc) or pRc/CMV vectors expressing Myc-tagged hnRNP-U (Myc-UF). Exogenous Myc-tagged hnRNP-U (Myc-U) and endogenous hnRNP-U (U) proteins were detected by immunoblotting with anti-Myc antibody ( $\alpha$ Myc) or HUT ( $\alpha$ U).

This antibody recognized exogenous Myc-tagged hnRNP-U and endogenous ~120 kDa hnRNP-U proteins in SAOS-2 cells.



**Fig.2 Immunocytochemistry using this antibody, HUT (ref.2)**

Mouse P19 neurons were labeled with anti-neudin antibody (Ndn) (a) or with HUT for hnRNP-U (U) (c) in combination with anti-neuronal marker, MAP2, antibody for MAP2 (b, d). The nuclear matrix was prepared in situ and labeled for neudin (Ndn) (e), hnRNP-U (U) (f), and a nuclear matrix marker, lamin B (g).

Both necdin and hnRNP-U were localized to the nuclei of differentiated neurons, which express the neuronal marker MAP2 (**a-d**). Necdin was also distributed in the neuronal cytoplasm (**a**).

The immunocytochemical analysis of in situ extracted nuclear matrix revealed that both necdin and hnRNP-U were concentrated in intranuclear speckles throughout the nucleoplasm (**e, f**). Lamin B, a nuclear matrix marker, was localized to the nuclear lamina (**g**). These results suggest that both necdin and hnRNP-U are associated with the nuclear matrix of neurons.

**References:** This antibody was produced and used in ref.2.

1. Kiledjian M and Dreyfuss G (1992) "Primary structure and binding activity of the hnRNP U protein: binding RNA through RGG box." *EMBO J* 11: 2655-2664 PMID: [1628625](#)
2. Taniura H and Yoshikawa K (2002) "Necdin interacts with the ribonucleoprotein hnRNP U in the nuclear matrix." *J Cell Biochem* 84:545-555 PMID: [11813259](#)

**Related product:**

74-104 anti-APP (C-terminal) antibody, rabbit serum