

## Product Datasheet

### Anti-RNF144A Antibody (orb214931)

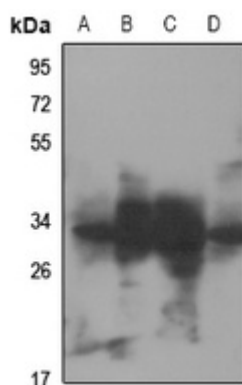
<b>Description</b>	Rabbit polyclonal antibody to RNF144A
<b>Species/Host</b>	Rabbit
<b>Reactivity</b>	Human, Mouse, Rat
<b>Conjugation</b>	Unconjugated
<b>Tested Applications</b>	IF, IH, WB
<b>Immunogen</b>	KLH-conjugated synthetic peptide encompassing a sequence within the center region of human RNF144A. The exact sequence is proprietary.
<b>Target</b>	RNF144A
<b>Preservatives</b>	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, and 0.01% sodium azide.
<b>Form/Appearance</b>	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, and 0.01% sodium azide.
<b>Storage</b>	Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.
<b>Note</b>	For research use only
<b>Clonality</b>	Polyclonal
<b>Source</b>	Rabbit
<b>Uniprot ID</b>	<b>P50876</b>
<b>Entrez</b>	<b>9781</b>
<b>Dilution Range</b>	WB: 1-500-1-1000, IHC-P: 1-100-1-200, IF/ICC: 1-100-1-500
<b>Expiration Date</b>	12 months from date of receipt.

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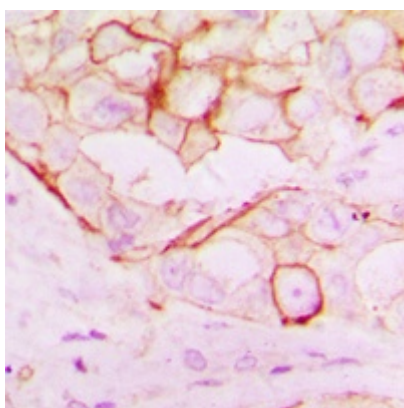
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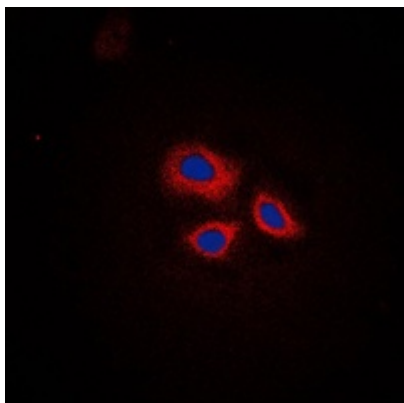
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Western blot analysis of RNF144A expression in mouse kidney (A), mouse testis (B), rat kidney (C), rat testis (D) whole cell lysates. (Predicted band size: 32 kD; Observed band size: 33 kD)



Immunohistochemical analysis of RNF144A staining in human breast cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of RNF144A staining in HepG2 cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a humidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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