

## Anti-ATP6V1E1 Antibody

Catalog #	Source	Reactivity	Applications
CQA2931	Rabbit	H, M, R	WB, IF/IC, IP
<b>Description</b>	Rabbit polyclonal antibody to ATP6V1E1		
<b>Immunogen</b>	Recombinant full length protein of human ATP6V1E1		
<b>Purification</b>	The antibody was purified by immunogen affinity chromatography.		
<b>Specificity</b>	Recognizes endogenous levels of ATP6V1E1 protein.		
<b>Clonality</b>	Polyclonal		
<b>Conjugation</b>			
<b>Form</b>	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, and 0.01% sodium azide.		
<b>Dilution</b>	WB (1/500 - 1/2000), IF/IC (1/50 - 1/200), IP (1/20 - 1/50)		
<b>Gene Symbol</b>	ATP6V1E1		
<b>Alternative Names</b>	ATP6E; ATP6E2; V-type proton ATPase subunit E 1; V-ATPase subunit E 1; V-ATPase 31 kDa subunit; p31; Vacuolar proton pump subunit E 1		
<b>Entrez Gene</b>	529 (Human); 11973 (Mouse); 297566 (Rat)		
<b>SwissProt</b>	P36543 (Human); P50518 (Mouse); Q6PCU2 (Rat)		
<b>Storage/Stability</b>	Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid freeze/thaw cycles.		

**Application key:** E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, CHIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference

**Species reactivity key:** H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb- Rabbit, S- Sheep, Z- Zebrafish

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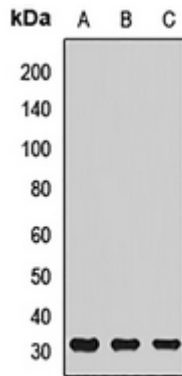
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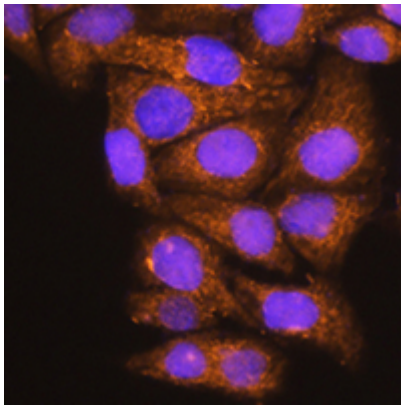
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## Product Data Sheet



Western blot analysis of ATP6V1E1 expression in LO2 (A), mouse liver (B), rat brain (C) whole cell lysates. (Predicted band size: 22; 23; 26 kD; Observed band size: 31 kD)



Immunofluorescent analysis of ATP6V1E1 staining in HeLa 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 AF594-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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