

# Product datasheet for TA812319M

## UPP2 Mouse Monoclonal Antibody [Clone ID: OTI9G9]

### **Product data:**

Product Type:	Primary Antibodies
Clone Name:	OTI9G9
Applications:	WB
Recommended Dilution:	WB 1:2000
Reactivity:	Human, Mouse, Rat
Host:	Mouse
lsotype:	lgG1
Clonality:	Monoclonal
Immunogen:	Full length human recombinant protein of human UPP2 (NP_775491) produced in HEK293T cell.
Formulation:	PBS (pH 7.3) containing 1% BSA, 50% glycerol and 0.02% sodium azide.
Concentration:	1 mg/ml
Purification:	Purified from mouse ascites fluids or tissue culture supernatant by affinity chromatography (protein A/G)
Conjugation:	Unconjugated
Storage:	Store at -20°C as received.
Stability:	Stable for 12 months from date of receipt.
Predicted Protein Size:	35.3 kDa
Gene Name:	uridine phosphorylase 2
Database Link:	<u>NP_775491</u> <u>Entrez Gene 76654 MouseEntrez Gene 295620 RatEntrez Gene 151531 Human</u> <u>O95045</u>
Background:	Catalyzes the reversible phosphorylytic cleavage of uridine and deoxyuridine to uracil and ribose- or deoxyribose-1-phosphate. The produced molecules are then utilized as carbon and energy sources or in the rescue of pyrimidine bases for nucleotide synthesis. Shows substrate specificity and accept uridine, deoxyuridine, and thymidine as well as the two pyrimidine nucleoside analogs 5-fluorouridine and 5-fluoro-2(')-deoxyuridine as substrates. [UniProtKB/Swiss-Prot Function]



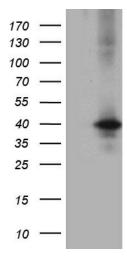
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	UPP2 Mouse Monoclonal Antibody [Clone ID: OTI9G9] – TA812319M
Synonyms:	UDRPASE2; UP2; UPASE2
Protein Pathway	s: Drug metabolism - other enzymes, Metabolic pathways, Pyrimidine metabolism

### **Product images:**



HEK293T cells were transfected with the pCMV6-ENTRY control (Cat# [PS100001], Left lane) or pCMV6-ENTRY UPP2 (Cat# [RC207260], Right lane) cDNA for 48 hrs and lysed. Equivalent amounts of cell lysates (5 ug per lane) were separated by SDS-PAGE and immunoblotted with anti-UPP2(Cat# [TA812319]). Positive lysates [LY406610] (100ug) and [LC406610] (20ug) can be purchased separately from OriGene.

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