

## **Product Data Sheet**

### **Anti-CCDC44 Antibody**

Catalog #	Source	Reactivity	Applications
CQA6118	Rabbit	H, M, R	WB, IH
Description	ł	Rabbit polyclonal antibody t	o CCDC44
Immunogen	I	Recombinant fusion protein	of human CCDC44. The exact sequence is proprietary.
Purification	-	The antibody was purified by	immunogen affinity chromatography.
Specificity	I	Recognizes endogenous leve	ls of CCDC44 protein
Clonality	I	Polyclonal	
Conjugation			
Form	l	Liquid in 0.42% Potassium pl	nosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	ä	and 0.01% sodium azide.	
Dilution	١	WB (1/500 - 1/2000), IH (1/50	- 1/200)
Gene Symbol	-	TACO1	
Alternative Na	ames (	CCDC44; Translational activa	tor of cytochrome c oxidase 1; Coiled-coil
	(	domain-containing protein 4	4; Translational activator of mitochondrially-encoded
	(	cytochrome c oxidase I	
Entrez Gene	Ţ	51204 (Human); 70207 (Mot	ise); 360645 (Rat)
SwissProt	(	Q9BSH4 (Human); Q8K0Z7 (I	Mouse); B2RYT9 (Rat)
Storage/Stabil	lity S	Shipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid
	f	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

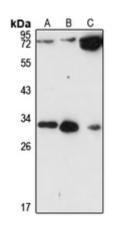
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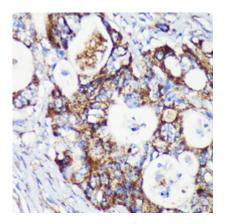


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Western blot analysis of CCDC44 expression in A375 (A), mouse heart (B), mouse kidney (C) whole cell lysates. (Predicted band size: 32 kD; Observed band size: 32 kD)



Immunohistochemical analysis of CCDC44 staining in human colon 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.

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