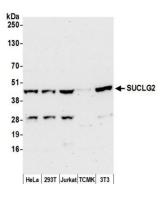
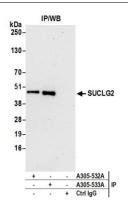
SUCLG2 Antibody

		-)				
Rabbit Polyclonal Antigen Affinity Purified			Protein ID	Q96199.2		
Catalog No. A305-533A		533A	GenelD	8801	DETLIVI	
Lot No.	A305-	533A-1				
APPLICATIONS		WB, IP				
SPECIES REACTIVITY		Human, Mouse				
PRESUMED REACTIVITY		Based on 100% sequence identity, this antibody is predicted to react with Bovine				
AMOUNT		100 µl				
CONCENTRATION		1000 µg/ml				
STORAGE/SHELF LIFE		2 - 8° C / 1 year from date of receipt				
PHYSICAL STATE		Liquid				
BUFFER		Tris-citrate/phosphate buffer, pH 7 to 8 containing 0.09% Sodium Azide				
ISOTYPE		lgG				
ORIGIN		USA				
PRODUCTION PROCEDURES		Antibody was affinity purified using an epitope specific to SUCLG2 immobilized on solid support.				
		The epitope recognized by A305–533A maps to a region between residue 382 to 432 of human Succinyl–CoA ligase [GDP–forming] subunit beta, mitochondrial using the numbering given in entry Q96199.2 (GeneID 8801).				
		Antibody concentration was determined by extinction coefficient: absorbance at 280 nm of 1.4 equals 1.0 mg of IgG.				
APPLICATIONS		Centrifuge tube to remove product from lid. Optimal working dilutions should be determined experimentally by the investigator. Prepare working dilution immediately before use.				
		Western Blot	1:1	,000 - 1:5,000		
		Immunoprecipit	ation 2 –	10 µg/mg lysate		
APPLICATION N	IOTES	Western blot of immunoprecipitates performed using Normal Pig Serum (Cat. No. S100–020), Goat anti-Rabbit Light Chain HRP Conjugate (Cat. No. A120–113P) and 4–20% SDS–PAGE (link to IP-western blot protocol in Additional Info section below).				
		Western blot of lysates performed using standard western blot reagents and 4-20% SDS-PAGE.				
ADDITIONAL IN	IFO	https://www.bethyl.com/product/A305–533A				
		Use the link above to view SDS, a current list of citations, and other product specific information. IP-western blot protocol: https://www.bethyl.com/content/protocol_IP_WB				
		IP-western blot p	protocol: http	os://www.bethyl.com/con	ent/protocol_IP_WB	
	_					

This document certifies that this product has met all of the quality control standards defined by Bethyl Laboratories, Inc. Eric McIntush, PhD | Chief Scientific Officer Date: June 21, 2019

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Detection of human and mouse SUCLG2 by western blot. Samples: Whole cell lysate (50 μ g) from HeLa, HEK293T, Jurkat, mouse TCMK-1, and mouse NIH 3T3 cells prepared using NETN lysis buffer. Antibody: Affinity purified rabbit anti-SUCLG2 antibody A305-533A (lot A305-533A-1) used for WB at 0.4 μ g/ml. Detection: Chemiluminescence with an exposure time of 30 seconds. Detection of human SUCLG2 by western blot of immunoprecipitates. *Samples:* Whole cell lysate (0.5 or 1.0 mg per IP reaction; 20% of IP loaded) from 293T cells prepared using NETN lysis buffer. *Antibodies:* Affinity purified rabbit anti-SUCLG2 antibody A305-533A (lot A305-533A-1) used for IP at 6 µg per reaction. SUCLG2 was also immunoprecipitated by rabbit anti-SUCLG2 antibody A305-532A. For blotting immunoprecipitated SUCLG2, A305-533A was used at 1 µg/ml. *Detection:* Chemiluminescence with an exposure time of 30 seconds.