

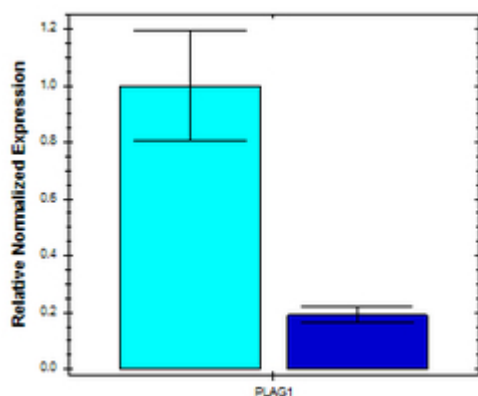
Product Data Sheet

Human PLAG1 Knockdown HEK293T Cell Lysate

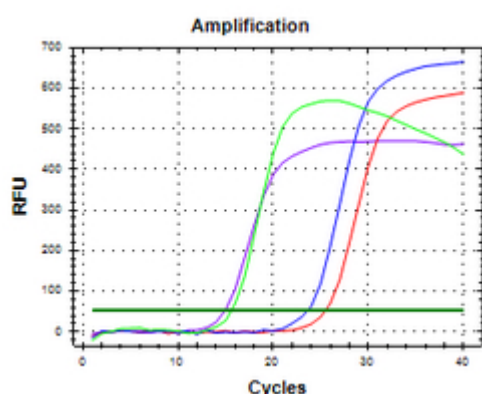
Catalog #	Source	Reactivity	Applications
CKD2040	HEK293T	Human	
Description	Human PLAG1 knockdown HEK293T cell lysate is engineered by our optimized transduction of the specific shRNA.		
Form	Liquid in 50 mM Tris-HCl, 2% SDS, 10% glycerol, 100 mM DTT, 0.01% Bromophenol blue		
Gene Symbol	PLAG1		
Alternative Names	Zinc finger protein PLAG1; Pleiomorphic adenoma gene 1 protein		
Entrez Gene	5324		
SwissProt	Q6DJT9		
KO/KD Validation	RT-qPCR		
Cell Line	HEK293T		
Components	Human PLAG1 knockdown HEK293T cell lysate x 1 vial; shControl HEK293T cell lysate x 1 vial		
Directions for Use	Thaw the vial on ice or at room temperature. Mix gently by pipetting or brief vortexing. Briefly centrifuge to collect the sample. Load the desired amount of lysate directly onto an SDS-PAGE gel. Perform SDS-PAGE and Western blot analysis using standard laboratory protocols. Note: This lysate has been pre-denatured in SDS sample buffer containing DTT and does not require additional reducing agent or boiling before electrophoresis.		
Storage/Stability	Shipped at 4°C. Upon delivery aliquot and store at -80°C for one year. Avoid freeze/thaw cycles.		

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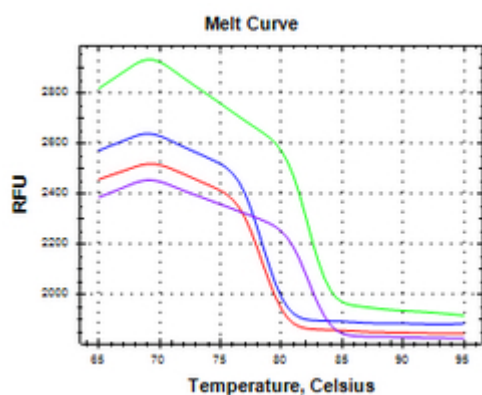
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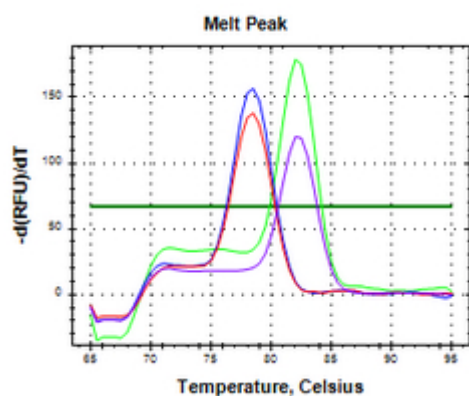
RT-qPCR analysis of PLAG1 knockdown efficiency. Compared with the 293T shRNA control (left), PLAG1 mRNA expression in the stable shRNA knockdown cell line (right) decreased, and the knockdown efficiency was 81%.



RT-qPCR amplification curves. The Ct values of the knockout sample significantly increased, confirming the effective knockdown of PLAG1 mRNA. [Red curve (knockdown sample), Blue curve (shRNA control), Purple and Green curve (reference gene GAPDH)]



RT-qPCR melt curve analysis demonstrated highly specific PCR amplification product without detectable nonspecific amplification or primer-dimer formation. [Red curve (knockdown sample), Blue curve (shRNA control), Purple and Green curve (reference gene GAPDH)]



This picture presents the raw fluorescence data (relative fluorescence units, RFU) used to generate the melt curves. [Red curve (knockdown sample), Blue curve (shRNA control), Purple and Green curve (reference gene GAPDH)]

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