abcam

Product datasheet

Human PTGR1 knockout HeLa cell lysate ab258152

3 Images

Overview		
Product name	Human PTGR1 knockout HeLa cell lysate	
Product overview		
	Knockout cell lysate achieved by CRISPR/Cas9.	
Parental Cell Line	HeLa	
Organism	Human	
Mutation description	Knockout achieved by using CRISPR/Cas9, 2 bp deletion in exon2 and Insertion of the selection cassette in exon2.	
Passage number	<20	
Knockout validation	Sanger Sequencing, Western Blot (WB)	
Reconstitution notes	To use as WB control, resuspend the lyophilizate in 50 μ L of LDS* Sample Buffer to have a final concentration of 2 mg/ml. For reducing conditions, we recommend a final concentration of 0.1 M DTT.	
	*Usage of SDS sample buffer is not recommended with these lyophilized lysates.	
Notes	Lysate preparation: Our lysates are made using RIPA buffer to which we add a protease inhibitor cocktail and phosphatase inhibitor cocktail (ratio: 300:100:10). <i>This means that the protein of interest is denatured.</i> If you require a native form of the protein please use the live cell version - found <u>here</u> . Please refer to our lysis protocol for further details on how our lysates are prepared.	
	User storage instructions: Lyophilizate may be stored at 4°C. After reconstitution, store at - 20°C for short-term storage or -80°C for long-term storage.	
	Access thousands of knockout cell lysates, generated from commonly used cancer cell lines. See here for more information on knockout cell lysates.	
	Abcam has not and does not intend to apply for the REACH Authorisation of customers' uses of products that contain European Authorisation list (Annex XIV) substances. It is the responsibility of our customers to check the necessity of application of REACH Authorisation, and any other relevant authorisations, for their intended uses.	
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Tested applications	Suitable for: WB	

Properties

Storage instructions	Store at -80°C. Please refer to protocols.		
Components		1 kit	
ab261154 - Human PTGR1 knockout HeLa cell lysate		1 x 100µg	
ab255929 - Human wild-type HeLa cell lysate		1 x 100µg	
Cell type	epithelial		
Disease	Adenocarcinoma		
Gender	Female		
STR Analysis	Amelogenin X D5S818: 11, 12 D13S317: 12, 13.3 D7S820: 8, 12 D16S539: 9, 10 vWA: 16, 18 TH01: 7 TPOX: 8,12 CSF1PO: 9, 10		
Target			
Function	Functions as 15-oxo-prostaglandin 13-reductase and acts on 15-oxo-PGE1, 15-oxo-PGE2 and 15-oxo-PGE2-alpha. Has no activity towards PGE1, PGE2 and PGE2-alpha (By similarity). Catalyzes the conversion of leukotriene B4 into its biologically less active metabolite, 12-oxo-leukotriene B4. This is an initial and key step of metabolic inactivation of leukotriene B4.		
Tissue specificity	High expression in the kidney, liver, and intestine but not in leukocytes.		
Sequence similarities	Belongs to the NADP-dependent oxidoreductase L4BD family.		
Cellular localization	Cytoplasm.		

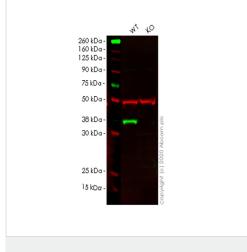
Applications

The Abpromise guarantee Our <u>Abpromise guarantee</u> covers the use of ab258152 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
WB		Use at an assay dependent concentration. Predicted molecular weight: 35 kDa.

Images



Western blot - Human PTGR1 knockout HeLa cell lysate (ab258152)

 Mut
 TCTAGGAGCTTCAGGATGGTTCGTACTAA--CATGGACCCTGAAGAAGCACTTTGTTGGC

 WIT
 TCTAGGAGCTTCAGGATGGTTCGTACTAAGACATGGACCCTGAAGAAGCACTTTGTTGGC

 Sanger Sequencing - Human PTGR1 knockout HeLa

cell lysate (ab258152)

Sanger Sequencing - Human PTGR1 knockout HeLa cell lysate (ab258152)

Lane 1:Wild-type HeLa cell lysate (20 ug) Lane 2:PTGR1 knockout HeLa cell lysate (20 ug)

ab181131 was shown to specifically react with PTGR1 in wild-type HeLa cells. Loss of signal was observed when knockout cell line **ab265169** (knockout cell lysate ab258152) was used. Wild-type and PTGR1 knockout samples were subjected to SDS-PAGE. **ab181131** and Anti-alpha Tubulin antibody [DM1A] - Loading Control (**ab7291**) were incubated overnight at 4°C at 1 in 1000 dilution and 1 in 20000 dilution respectively. Blots were developed with Goat anti-Rabbit IgG H&L (IRDye[®] 800CW) preadsorbed (**ab216773**) and Goat anti-Mouse IgG H&L (IRDye[®] 680RD) preadsorbed (**ab216776**) secondary antibodies at 1 in 20000 dilution for 1 hour at room temperature before imaging.

Allele-1: 2 bp deletion in exon2

Allele-2: Insertion of the selection cassette in exon2

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