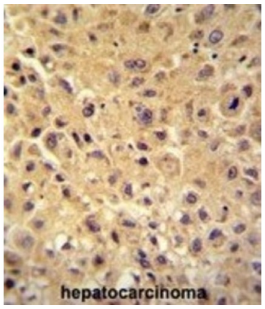
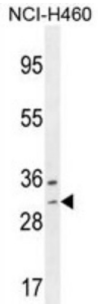
Williams-Beuren Syndrome Chromosomal Region 27 Protein (WBSCR27) Antibody

Catalogue No.:abx025778





WBSCR27 encodes a protein belonging to ubiE/COQ5 methyltransferase family. The gene is deleted in Williams syndrome, a multisystem developmental disorder caused by the deletion of contiguous genes at 7q11.22-q11.23.

Target:	Williams-Beuren Syndrome Chromosomal Region 27 Protein (WBSCR27)
Clonality:	Polyclonal
Reactivity:	Human
Tested Applications:	ELISA, WB, IHC
Host:	Rabbit
Recommended dilutions:	WB: 1/1000, IHC-P: 1/50 - 1/100. Not tested in IHC-F. Optimal dilutions/concentrations should be determined by the end user.
Conjugation:	Unconjugated
Immunogen:	KLH-conjugated synthetic peptide between 5-34 amino acids from the N-terminal region of human WBSCR27.
Isotype:	IgG
Form:	Liquid

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Purification:	Purified through a protein A column, followed by peptide affinity purification.
Storage:	Aliquot and store at -20°C. Avoid repeated freeze/thaw cycles.
UniProt Primary AC:	Q8N6F8 (UniProt, ExPASy)
Gene Symbol:	METTL27
KEGG:	hsa:155368
String:	9606.ENSP00000297873
Molecular Weight:	Calculated MW: 26.5 kDa
Buffer:	PBS containing 0.09% sodium azide.
Note:	THIS PRODUCT IS FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC, THERAPEUTIC OR COSMETIC PROCEDURES. NOT FOR HUMAN OR ANIMAL CONSUMPTION.

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