

# DGCR2 Rabbit pAb

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Catalog # AP58944

## Product Information

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<b>Application</b>	WB
<b>Primary Accession</b>	<a href="#">P98153</a>
<b>Reactivity</b>	Human
<b>Predicted</b>	Mouse, Rat, Rabbit, Zebrafish, Orangutan
<b>Host</b>	Rabbit
<b>Clonality</b>	Polyclonal
<b>Calculated MW</b>	60811
<b>Physical State</b>	Liquid
<b>Immunogen</b>	KLH conjugated synthetic peptide derived from human DGCR2
<b>Epitope Specificity</b>	151-250/550
<b>Isotype</b>	IgG
<b>Purity</b>	affinity purified by Protein A
<b>Buffer</b>	0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol.
<b>SUBCELLULAR LOCATION</b>	Membrane; Single pass type I membrane protein.
<b>SIMILARITY</b>	Contains 1 C-type lectin domain. Contains 1 LDL-receptor class A domain. Contains 1 WW domain.
<b>Important Note</b>	This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.
<b>Background Descriptions</b>	The DGCR2 gene encodes a novel putative adhesion receptor protein, which may play a role in neural crest cells migration, a process which has been proposed to be altered in DiGeorge syndrome. Deletions of the 22q11.2 have been associated with a wide range of developmental defects (notably DiGeorge syndrome, velocardiofacial syndrome, conotruncal anomaly face syndrome and isolated conotruncal cardiac defects) classified under the acronym CATCH 22.

## Additional Information

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<b>Gene ID</b>	9993
<b>Other Names</b>	Integral membrane protein DGCR2/IDD, DGCR2, IDD, KIAA0163
<b>Target/Specificity</b>	Predominantly expressed in brain, heart, lung and fetal kidney. Low levels in liver and adult kidney.
<b>Dilution</b>	WB=1:500-2000
<b>Storage</b>	Store at -20 °C for one year. Avoid repeated freeze/thaw cycles. When reconstituted in sterile pH 7.4 0.01M PBS or diluent of antibody the antibody is stable for at least two weeks at 2-4 °C.

## Protein Information

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<b>Name</b>	DGCR2
<b>Synonyms</b>	IDD, KIAA0163
<b>Function</b>	Putative adhesion receptor, that could be involved in cell-cell or cell-matrix interactions required for normal cell differentiation and migration.
<b>Cellular Location</b>	Membrane; Single-pass type I membrane protein.
<b>Tissue Location</b>	Predominantly expressed in brain, heart, lung and fetal kidney. Low levels in liver and adult kidney

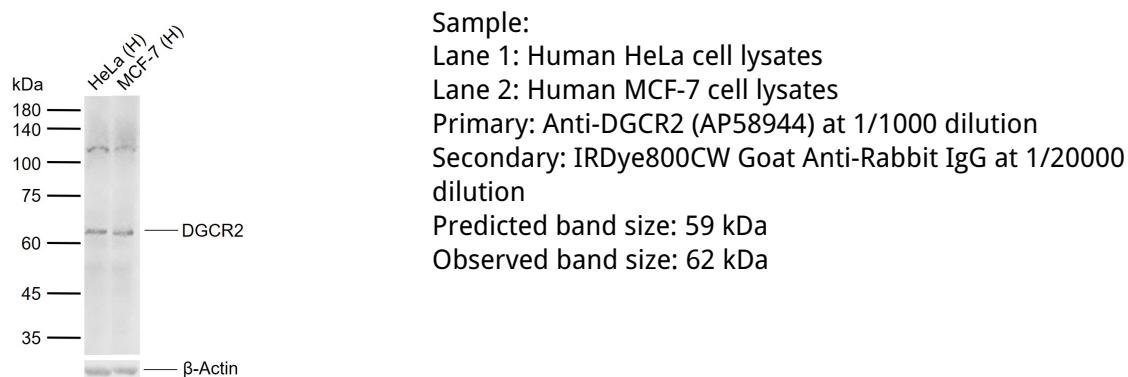
## Background

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The DGCR2 gene encodes a novel putative adhesion receptor protein, which may play a role in neural crest cells migration, a process which has been proposed to be altered in DiGeorge syndrome. Deletions of the 22q11.2 have been associated with a wide range of developmental defects (notably DiGeorge syndrome, velocardiofacial syndrome, conotruncal anomaly face syndrome and isolated conotruncal cardiac defects) classified under the acronym CATCH 22.

## Images

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Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.