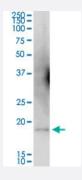


MaxPab@

WRB purified MaxPab mouse polyclonal antibody (B02P)

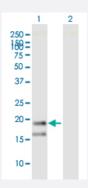
Catalog # H00007485-B02P Size 50 ug

Applications



Western Blot (Tissue lysate)

WRB MaxPab polyclonal antibody. Western Blot analysis of WRB expression in rat brain.



Western Blot (Transfected lysate)

Western Blot analysis of WRB expression in transfected 293T cell line (<u>H00007485-T03</u>) by WRB MaxPab polyclonal antibody.

Lane 1: WRB transfected lysate(19.80 KDa).

Lane 2: Non-transfected lysate.

Specification	
Product Description	Mouse polyclonal antibody raised against a full-length human WRB protein.
lmmunogen	WRB (NP_004618.2, 1 a.a. ~ 174 a.a) full-length human protein.
Sequence	MSSAAADHWAWLLVLSFVFGCNVLRILLPSFSSFMSRVLQKDAEQESQMRAEIQDMKQELSTVN MMDEFARYARLERKINKMTDKLKTHVKARTAQLAKIKWVISVAFYVLQAALMISLIWKYYSVPVAVV PSKWITPLDRLVAFPTRVAGGVGITCWILVCNKVVAIVLHPFS
Host	Mouse
Reactivity	Human, Rat



Product Information

Quality Control Testing	Antibody reactive against mammalian transfected lysate.
Storage Buffer	In 1x PBS, pH 7.4
Storage Instruction	Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

Applications

Western Blot (Tissue lysate)

WRB MaxPab polyclonal antibody. Western Blot analysis of WRB expression in rat brain.

Protocol Download

Western Blot (Transfected lysate)

 $We stern \ Blot \ analysis \ of \ WRB \ expression \ in \ transfected \ 293T \ cell \ line \ (\underline{H00007485-T03}) \ by \ WRB \ MaxPab \ polyclonal \ antibody.$

Lane 1: WRB transfected lysate(19.80 KDa).

Lane 2: Non-transfected lysate.

Protocol Download

Gene Info — WRB	
Entrez GeneID	<u>7485</u>
GeneBank Accession#	NM_004627.2
Protein Accession#	<u>NP_004618.2</u>
Gene Name	WRB
Gene Alias	CHD5
Gene Description	tryptophan rich basic protein
Omim ID	602915
Gene Ontology	<u>Hyperlink</u>
Gene Summary	This gene encodes a basic nuclear protein of unknown function. The gene is widely expressed in adult and fetal tissues. Since the region proposed to contain the gene(s) for congenital heart dise ase (CHD) in Down syndrome (DS) patients has been restricted to 21q22.2-22.3, this gene, which maps to 21q22.3, has a potential role in the pathogenesis of Down syndrome congenital heart disease. Alternatively spliced transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq



Product Information

Other Designations

congenital heart disease 5 protein