

**Anti-GPC3 / Glypican-3 Reference Antibody (codrituzumab)  
Recombinant Antibody  
Catalog # APR10154****Specification**

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**Anti-GPC3 / Glypican-3 Reference Antibody (codrituzumab) - Product Information**

Application	FC, Kinetics, Animal Model
Primary Accession	<a href="#">P51654</a>
Reactivity	Human
Clonality	Monoclonal
Isotype	IgG1
Calculated MW	145.5 KDa

**Anti-GPC3 / Glypican-3 Reference Antibody (codrituzumab) - Additional Information****Target/Specificity**  
GPC3 / Glypican-3**Endotoxin**  
< 0.001EU/ µg, determined by LAL method.**Conjugation**  
Unconjugated**Expression system**  
CHO Cell**Format**  
Purified monoclonal antibody supplied in PBS, pH6.0, without preservative. This antibody is purified through a protein A column.**Anti-GPC3 / Glypican-3 Reference Antibody (codrituzumab) - Protein Information****Name** GPC3**Synonyms** OCI5**Function**  
Cell surface proteoglycan (PubMed: [14610063](http://www.uniprot.org/citations/14610063)). Negatively regulates the hedgehog signaling pathway when attached via the GPI- anchor to the cell surface by competing with the hedgehog receptor PTC1 for binding to hedgehog proteins (By similarity). Binding to the hedgehog protein SHH triggers internalization of the complex by endocytosis and its subsequent lysosomal degradation (By similarity). Positively regulates the canonical Wnt signaling pathway by binding to the Wnt receptor Frizzled and stimulating the binding of the Frizzled receptor to Wnt ligands (PubMed: [16227623](http://www.uniprot.org/citations/16227623), PubMed: [24496449](http://www.uniprot.org/citations/24496449)). Positively regulates the non-canonical Wnt signaling pathway (By similarity). Binds to CD81 which decreases

the availability of free CD81 for binding to the transcriptional repressor HHEX, resulting in nuclear translocation of HHEX and transcriptional repression (By similarity). Inhibits the dipeptidyl peptidase activity of DPP4 (PubMed:<a href="http://www.uniprot.org/citations/17549790" target="\_blank">17549790</a>). Plays a role in limb patterning and skeletal development by controlling the cellular response to BMP4 (By similarity). Modulates the effects of growth factors BMP2, BMP7 and FGF7 on renal branching morphogenesis (By similarity). Required for coronary vascular development (By similarity). Plays a role in regulating cell movements during gastrulation (By similarity).

#### Cellular Location

Cell membrane; Lipid-anchor, GPI-anchor {ECO:0000250|UniProtKB:P13265}; Extracellular side {ECO:0000250|UniProtKB:P13265}

#### Tissue Location

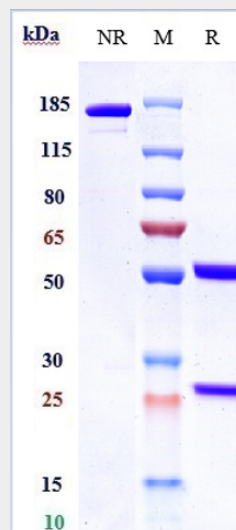
Detected in placenta (at protein level) (PubMed:32337544). Highly expressed in lung, liver and kidney

### Anti-GPC3 / Glypican-3 Reference Antibody (codrituzumab) - Protocols

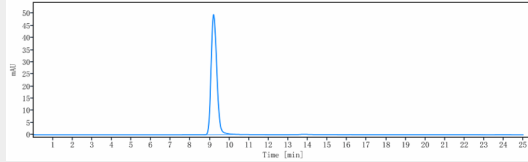
Provided below are standard protocols that you may find useful for product applications.

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

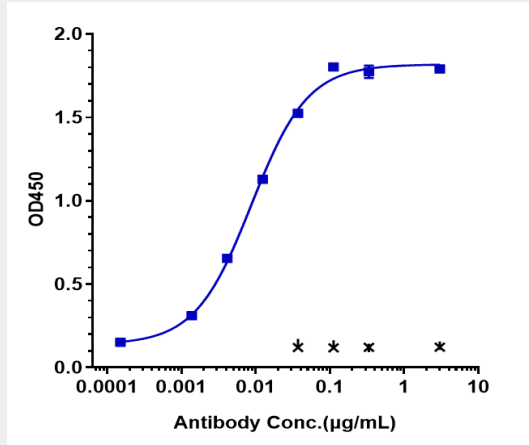
### Anti-GPC3 / Glypican-3 Reference Antibody (codrituzumab) - Images



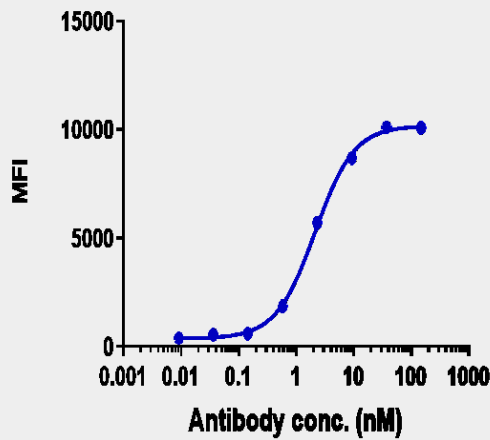
Anti-GPC3 / Glypican-3 Reference Antibody (codrituzumab) on SDS-PAGE under reducing (R) condition. The gel was stained with Coomassie Blue. The purity of the protein is greater than 95%



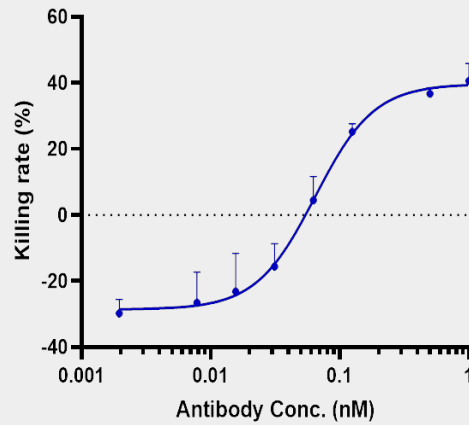
The purity of Anti-GPC3 / Glypican-3 Reference Antibody (codrituzumab) is more than 95%, determined by SEC-HPLC.



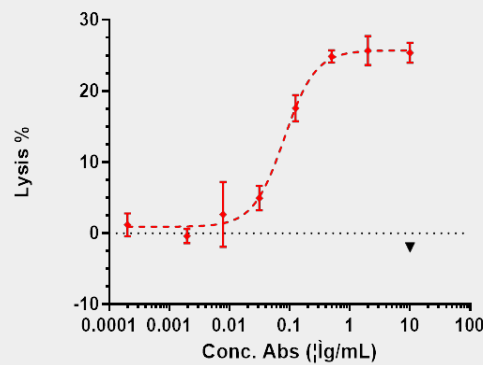
Immobilized human GPC3, His tag at 2 µg/mL can bind Anti-GPC3 / Glypican-3 Reference Antibody (codrituzumab)  $EC_{50}=0.008741 \mu\text{g/mL}$



Human GPC3 CHOS cells were stained with Anti-GPC3 / Glypican-3 Reference Antibody (codrituzumab) and negative control protein respectively, washed and then followed by PE and analyzed with FACS,  $EC_{210}=2.142 \text{ nM}$



The endocytosis ratio codrituzumab by HuH-7 increased with the increase of antibody concentration, and the Internalization Rate (%) reached 40% at antibody concentration of 1 nM.



Anti-GPC3 / Glypican-3 Reference Antibody (Codrituzumab)-induced ADCC activity was evaluated using human GPC3 CHOS Cell. The max Lysis rate was approximately 30%.