

**GPR84 Antibody (Extracellular Domain)**  
**Rabbit Polyclonal Antibody**  
**Catalog # ALS10096****Specification**

---

**GPR84 Antibody (Extracellular Domain) - Product Information**

Application	IHC-P
Primary Accession	<a href="#">O9NQS5</a>
Reactivity	Human, Monkey, Horse
Host	Rabbit
Clonality	Polyclonal
Calculated MW	44kDa KDa
Dilution	IHC-P~~N/A

**GPR84 Antibody (Extracellular Domain) - Additional Information****Gene ID** 53831**Other Names**

G-protein coupled receptor 84, Inflammation-related G-protein coupled receptor EX33, GPR84, EX33

**Target/Specificity**

Human GPR84. BLAST analysis of the peptide immunogen showed no homology with other human proteins.

**Reconstitution & Storage**

Long term: -70°C; Short term: +4°C

**Precautions**

GPR84 Antibody (Extracellular Domain) is for research use only and not for use in diagnostic or therapeutic procedures.

**GPR84 Antibody (Extracellular Domain) - Protein Information****Name** GPR84**Synonyms** EX33**Function**

G protein-coupled receptor that responds endogenously to dietary fatty acids or nutrient, specifically medium-chain free fatty acid (FFA) with carbon chain lengths of C9 to C14. Capric acid (C10:0), undecanoic acid (C11:0) and lauric acid (C12:0) are the most potent agonists (PubMed:&lt;a href="http://www.uniprot.org/citations/16966319" target="\_blank"&gt;16966319&lt;/a&gt;). In immune cells, functions as a pro- inflammatory receptor via 6-OAU and promotes the expression of pro-inflammatory mediators such as TNFalpha, IL-6 and IL-12B as well as stimulating chemotactic responses through activation of signaling mediators AKT, ERK and NF-kappa-B (By similarity). In addition, triggers increased bacterial adhesion and phagocytosis in macrophages and regulates

pro-inflammatory function via enhancing NLRP3 inflammasome activation (By similarity). Also plays an important role in inflammation by modulating neutrophil functions (By similarity). Mechanistically, promotes neutrophil chemotaxis, reactive oxygen species (ROS) production and degranulation via LYN-AKT/ERK pathway (By similarity). To regulate ROS, communicates with the two formyl peptide receptors FPR2 and FPR1 to control the NADPH oxidase activity in neutrophils (PubMed:<a href="http://www.uniprot.org/citations/33789297" target="\_blank">33789297</a>).

#### Cellular Location

Cell membrane; Multi-pass membrane protein

#### Tissue Location

Expressed predominantly in hematopoietic tissues. High levels detected in the bone marrow and lower levels in the peripheral leukocytes and lung. Also expressed in brain, heart, muscle, colon, thymus, spleen, kidney, liver, placenta and intestine. Within the leukocyte population expression is higher in neutrophils and eosinophils relative to T- or B-lymphocytes

#### Volume

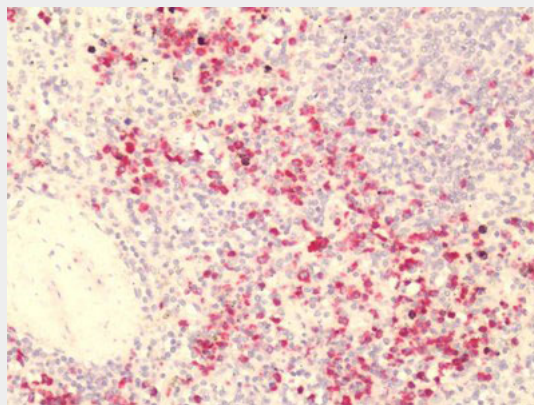
50 µl

### GPR84 Antibody (Extracellular Domain) - 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](#)

### GPR84 Antibody (Extracellular Domain) - Images



Human, Spleen: Formalin-Fixed Paraffin-Embedded (FFPE)

### GPR84 Antibody (Extracellular Domain) - Background

Receptor for medium-chain free fatty acid (FFA) with carbon chain lengths of C9 to C14. Capric acid (C10:0), undecanoic acid (C11:0) and lauric acid (C12:0) are the most potent agonists. Not activated by short-chain and long-chain saturated and unsaturated FFAs. Activation by medium-chain free fatty acid is coupled to a pertussis toxin sensitive G(i/o) protein pathway. May

have important roles in processes from fatty acid metabolism to regulation of the immune system.

#### **GPR84 Antibody (Extracellular Domain) - References**

Yousefi S.,et al.J. Leukoc. Biol. 69:1045-1052(2001).

Wittenberger T.,et al.J. Mol. Biol. 307:799-813(2001).

Kaighin V.A.,et al.Submitted (OCT-2008) to the EMBL/GenBank/DDBJ databases.

Takeda S.,et al.FEBS Lett. 520:97-101(2002).

Mural R.J.,et al.Submitted (JUL-2005) to the EMBL/GenBank/DDBJ databases.