

## GITR Protein, Rat (HEK293, Fc)

<b>Cat. No.:</b>	HY-P75163
<b>Synonyms:</b>	Tumor necrosis factor receptor superfamily member 18; CD357; TNFRSF18; AITR; GITR
<b>Species:</b>	Rat
<b>Source:</b>	HEK293
<b>Accession:</b>	Q5M835 (E25-K121)
<b>Gene ID:</b>	/
<b>Molecular Weight:</b>	Approximately 44 kDa

### PROPERTIES

<b>AA Sequence</b>	<p>E E P S C G P G R V      R N G T G T N T R C      C S L C G P D K E D      C P K G R C I C V K</p> <p>P E Y H C E D P Q C      K T C K H Y P C Q P      G Q R V E S Q G N I      K F G F Q C V D C A</p> <p>M G T F S A G R E G      H C R L W T K</p>
<b>Biological Activity</b>	Measured by its binding ability in a functional ELISA. Immobilized Rat GITR at 2 µg/mL (100 µL/well) can bind Biotinylated Mouse GITR Ligand. The ED <sub>50</sub> for this effect is 82.41 ng/mL.
<b>Appearance</b>	Lyophilized powder.
<b>Formulation</b>	Lyophilized from a 0.2 µm filtered solution of PBS, pH 7.4.
<b>Endotoxin Level</b>	<1 EU/µg, determined by LAL method.
<b>Reconstitution</b>	It is not recommended to reconstitute to a concentration less than 100 µg/mL in ddH <sub>2</sub> O. For long term storage it is recommended to add a carrier protein (0.1% BSA, 5% HSA, 10% FBS or 5% Trehalose).
<b>Storage &amp; Stability</b>	Stored at -20°C for 2 years. After reconstitution, it is stable at 4°C for 1 week or -20°C for longer (with carrier protein). It is recommended to freeze aliquots at -20°C or -80°C for extended storage.
<b>Shipping</b>	Room temperature in continental US; may vary elsewhere.

### DESCRIPTION

<b>Background</b>	<p>GITR is expressed on regulatory T cells (Tregs) and some activated immune cells, including effector T lymphocytes, nature killer (NK) cells, and neutrophils<sup>[1]</sup>.</p> <p>The amino acid sequence of human GITR protein has low homology for mouse GITR protein.</p> <p>GITR does not have any enzymatic activity and signaling is propagated via recruiting TRAF-family members, specifically TRAF1, TRAF2 and TRAF5, to the GITR-signaling complex. The signaling is then mediated through NF-κB and MAPK pathways. GITR does not have any enzymatic activity and signaling is propagated via recruiting TRAF-family members,</p>
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specifically TRAF1, TRAF2 and TRAF5, to the GITR-signaling complex. The signaling is then mediated through NF- $\kappa$ B and MAPK pathways, protecting T cells from TCR activation-induced cell death<sup>[2]</sup>.

GITR (Glucocorticoid-induced TNFR-related protein, also known as TNFRSF18) is a type I transmembrane protein. GITR stimulates the proliferation of effector T-lymphocytes and partially reverses the immunosuppressive function of CD4+CD25+ Tregs<sup>[1]</sup>. GITR is activated by its ligand GITRL (TNFSF18). GITR induces NOS in murine macrophage in a time and dose-dependent manner<sup>[3]</sup>. GITR inhibits Multiple Myeloma (MM) cell proliferation in vitro and in vivo and induces apoptosis<sup>[4]</sup>.

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## REFERENCES

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- [1]. Tian J, et al. The Role of GITR/GITRL Interaction in Autoimmune Diseases. *Front Immunol*. 2020 Oct 9;11:588682.
- [2]. Krausz LT, et al. GITR-GITRL system, a novel player in shock and inflammation. *ScientificWorldJournal*. 2007 May 1;7:533-66.
- [3]. Shin HH, et al. Recombinant glucocorticoid induced tumor necrosis factor receptor (rGITR) induces NOS in murine macrophage. *FEBS Lett*. 2002 Mar 13;514(2-3):275-80.
- [4]. Liu Y, et al. Novel tumor suppressor function of glucocorticoid-induced TNF receptor GITR in multiple myeloma. *PLoS One*. 2013 Jun 13;8(6):e66982.
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**Caution: Product has not been fully validated for medical applications. For research use only.**

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