Screening Libraries

Product Data Sheet

LILRB1/CD85j/ILT2 Protein, Human (HEK293, Fc)

Cat. No.: HY-P70182

Synonyms: rHuLIR-1/LILRB1, Fc; Leukocyte Immunoglobulin-Like Receptor Subfamily B Member 1; LIR-1;

> Leukocyte Immunoglobulin-Like Receptor 1; CD85 Antigen-Like Family Member J; Immunoglobulin-Like Transcript 2; ILT-2; Monocyte/Macrophage Immunoglobulin-Like

Receptor 7; MIR-7; CD85j; LILRB1; ILT2; LIR1; MIR7

Species: Human Source: HEK293

Accession: D9IDM8 (G24-H458)

Gene ID: 10859

Molecular Weight: 100-120 kDa

PROPERTIES

AA Sequence				
	G H L P K P T L W A E P G S V I T Q G S	PVTLRCQGGQ	ETQEYRLYRE	
	KKTAPWITRI PQELVKKGQF	PIPSITWEHA	GRYRCYYGSD	
	TAGRSESSDP LELVVTGAYI	KPTLSAQPSP	VVNSGGNVTL	
	QCDSQVAFDG FILCKEGEDE	HPQCLNSQPH	ARGSSRAIFS	
	V G P V S P S R R W W Y R C Y A Y D S N	SPYEWSLPSD	LLELLVLGVS	
	KKPSLSVQPG PIVAPEETLT	LQCGSDAGYN	RFVLYKDGER	
	D F L Q L A G A Q P Q A G L S Q A N F T	LGPVSRSYGG	QYRCYGAHNL	
	S S E W S A P S D P L D I L I A G Q F Y	DRVSLSVQPG	PTVASGENVT	
	LLCQSQGWMQ TFLLTKEGAA	DDPWRLRSTY	QSQKYQAEFP	
	M G P V T S A H A G T Y R C Y G S Q S S	KPYLLTHPSD	PLELVVSGPS	
	G G P S S P T T G P T S T S G P E D Q P	LTPTGSDPQS	GLGRH	
Appearance	Lyophilized powder.			
Formulation	Lyophilized from a 0.2 μm filtered solution of PBS, pH 7.4.			
Endotoxin Level	<1 EU/μg, determined by LAL method.			
Reconsititution	It is not recommended to reconstitute to a concentration less than 100 $\mu g/mL$ in ddH $_2$ O. For long term storage it is			
	recommended to add a carrier protein (0.1% BSA, 5% HSA, 10% FBS or 5% Trehalose).			
Storage & Stability	•	. 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.			
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Shipping	Room temperature in continental US; may vary elsewhere.			

DESCRIPTION

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Background

LILRB1 binds MHC class I and also contain immunoreceptor tyrosine-based inhibitory motifs involved in the intracellular transduction of inhibitory signaling, which establishes them as strong candidates for MHC class I-mediated suppression of phagocytosis $^{[1]}$.

LILRB1 and PD1 shows nonoverlapping expression patterns across CD8+ TEM and TEMRA subsets, and blocking both pathways synergistically enhanced CD8+ T cell function. LILRB1 is highly expressed by the CD8+ TEMRA subset, which is the most potent population for BiTE molecule–induced toxicity. LILRB1-expressing CD8+ T cells infiltrate solid tumors. LILRB1 blockade increases CD8+ T cell cytolytic activity in vitro^[3].

Caution: Product has not been fully validated for medical applications. For research use only.

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