

# **Recombinant Human DLL1 Fc Chimera**

Catalog Number: 10184-DL

DESCRIPTION				
Source	Mouse myeloma cell line, NS0-derived human DLL1 protein			
	Human DLL1 (Ser22-Glu537) Accession # 000548	IEGRMDP	Human IgG <sub>1</sub> (Pro100-Lys330)	
	N-terminus		C-terminus	
N-terminal Sequence Analysis	Ser22			
Structure / Form	Disulfide-linked homodimer			
Predicted Molecular Mass	82 kDa			

SPECIFICATIONS		
SDS-PAGE	88-98 kDa, under reducing conditions	
Activity	Measured by the ability of the immobilized protein to enhance BMP-2 induced alkaline phosphatase activity in C3H10T1/2 mouse embryonic fibroblast cells. Nobta, M. <i>et al.</i> (2005) J. Biol. Chem. <b>280</b> :15842. The ED <sub>50</sub> for this effect is 0.1-0.5 μg/mL.	
Endotoxin Level	<0.10 EU per 1 $\mu$ g of the protein by the LAL method.	
Purity	>90%, by SDS-PAGE visualized with Silver Staining and quantitative densitometry by Coomassie® Blue Staining.	
Formulation	Lyophilized from a 0.2 µm filtered solution in HEPES and EDTA. See Certificate of Analysis for details.	

PREPARATION AND STORAGE			
Reconstitution	itution Reconstitute at 250 µg/mL in PBS.		
Shipping	The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below.		
Stability & Storage	Use a manual defrost freezer and avoid repeated freeze-thaw cycles.		
	12 months from date of receipt, -20 to -70 °C as supplied.		

- 1 month, 2 to 8 °C under sterile conditions after reconstitution.
- 3 months, -20 to -70  $^\circ\text{C}$  under sterile conditions after reconstitution.



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Recombinant Human DLL1 Fc Chimera (Catalog # 10184-DL) enhances Recombinant Human BMP-2 (Catalog # 355-BM) induced alkaline phosphatase activity in the C3H10T1/2 mouse embryonic fibroblast cell line. The ED<sub>50</sub> for this effect is 0.1-0.5 µg/mL.

#### SDS-PAGE



2 µg/lane of Recombinant Human DLL1 Fc Chimera was resolved with SDS-PAGE under reducing (R) and non-reducing (NR) conditions and visualized by Coomassie® Blue staining, showing bands at 88-98 kDa and 170-190 kDa, respectively.

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### BACKGROUND

Delta-like protein 1 (DLL1) is a 90-100 kDa type I transmembrane protein that belongs to the Delta/Serrate/Lag-2 (DSL) family of Notch ligands. Mature human DLL1 consists of a 528 amino acid (aa) extracellular domain (ECD) with one DSL domain and eight EGF-like repeats, a 23 aa transmembrane segment, and a 155 aa cytoplasmic domain (1). Within the ECD, human DLL1 shares 91% aa sequence identity with mouse and rat DLL1. It shares 26%, 37%, and 54% aa sequence identity with DLL2, 3, and 4, respectively. A 60 kDa ECD fragment released by ADAM9, 12, or 17 mediated proteolysis, promotes the proliferation of hematopoietic progenitor cells (2, 3). The residual membrane-bound portion of DLL1 can be cleaved by presenilin-dependent  $\gamma$ -secretase, enabling the cytoplasmic domain to migrate to the nucleus (4). DLL1 localizes to adherent junctions on neuronal processes through its association with the scaffolding protein MAGI1 (5). DLL1 is widely expressed, and it plays an important role in embryonic somite formation, cochear hair cell differentiation, plus B and T lymphocyte differentiation (6-11). The up-regulation of DLL1 in arterial endothelial cells following injury or angiogenic stimulation is central to postnatal arteriogenesis (12). DLL1 is also over-expressed in cervical carcinoma and glioma and contributes to tumor progression (1, 13). Soluble DLL-1 was shown to inhibit differentiation of hematopoietic precursor cells (14).

#### References:

- 1. Gray, G.E. et al. (1999) Am. J. Pathol. 154:785.
- 2. Dyczynska, E. et al. (2007) J. Biol. Chem. 282:436.
- 3. Karanu, F.N. et al. (2001) Blood 97:1960.
- 4. Ikeuchi, T. and S.S. Sisodia (2003) J. Biol. Chem. 278:7751.
- 5. Mizuhara, E. et al. (2005) J. Biol. Chem. 280:26499.
- 6. Takahashi, Y. et al. (2003) Development 130:4259.
- 7. Teppner, I. *et al*. (2007) BMC Dev. Biol. **7**:68.
- 8. Kiernan, A.E. *et al.* (2005) Development **132**:4353.
- 9. Schmitt, T.M. and J.C. Zuniga-Pflucker (2002) Immunity 17:749.
- 10. Hozumi, K. et al. (2004) Nat. Immunol. 5:638.
- 11. Santos, M.A. et al. (2007) Proc. Natl. Acad. Sci. 104:15454.
- 12. Limbourg, A. et al. (2007) Circ. Res. 100:363.
- 13. Purow, B.W. et al. (2005) Cancer Res. 65:2353.
- 14. Han. W. *et al*. (2000) Blood. **95(5)**:1616.

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