

Recombinant human DLL4 protein (Fc Chimera Active) ab108557

7 References [画像数 6](#)

製品の詳細

製品名	Recombinant human DLL4 protein (Fc Chimera Active)
生理活性	Inhibits adipogenesis of 3T3L-1 cells and mesenchymal stem cells (MSCs). Induces the Notch target gene HES-1 when coated on a plate at 1 µg/ml.
精製度	>= 95 % SDS-PAGE. Purified using affinity chromatography.
エンドキシン・レベル	< 0.010 Eu/µg
発現系	HEK 293 cells
アクセッション番号	<u>Q9NR61</u>
タンパク質長	Protein fragment
Animal free	No
由来	Recombinant
生物種	Human
予測される分子量	80 kDa including tags
領域	1 to 529
タグ	Fc tag C-Terminus
配列の追加情報	Signal peptide and extracellular domain of human DLL4 (aa 1-529) are fused at the C-terminus to the Fc portion of human IgG1.

特性

Our **Abpromise guarantee** covers the use of **ab108557** in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

アプリケーション	Functional Studies SDS-PAGE
製品の状態	Lyophilized
備考	ab108557 interacts with Human Notch1 (as confirmed by Flow Cytometry).

前処理および保存

保存方法および安定性

Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.

Constituents: PBS, 0.5% Trehalose

ab108557 is 0.2 µm filtered.

This product is an active protein and may elicit a biological response in vivo, handle with caution.

再構成

Reconstitute with 100µl sterile water. PBS containing at least 0.1% BSA should be used for further dilutions. Working aliquots are stable for up to 3 months when stored at -20°C.

関連情報

機能

Plays a role in the Notch signaling pathway. Activates Notch-1 and Notch-4.

組織特異性

Expressed in vascular endothelium.

配列類似性

Contains 1 DSL domain.

Contains 8 EGF-like domains.

ドメイン

The Delta-Serrate-Lag2 (DSL) domain is required for binding to the Notch receptor.

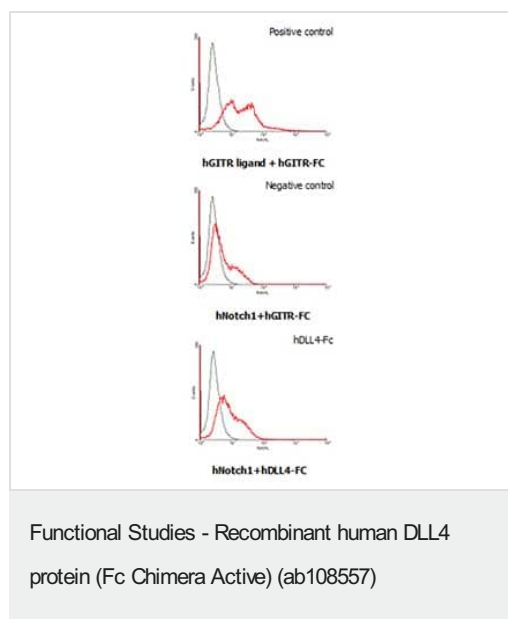
翻訳後修飾

Ubiquitinated by MIB (MIB1 or MIB2), leading to its endocytosis and subsequent degradation.

細胞内局在

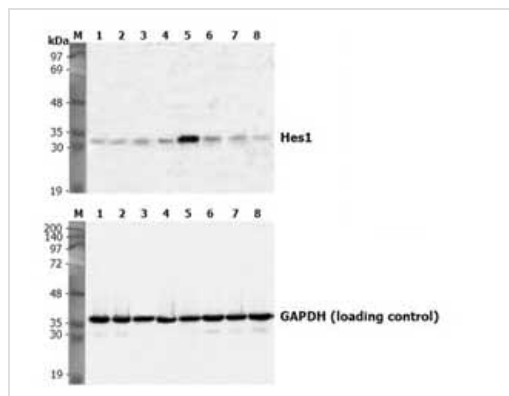
Membrane.

画像



Interaction of Human Notch1 with Human DLL4.

HEK293 cells transfected with a Human Notch1 or a Human GTR ligand expressing vector were incubated with 25 µg/ml of Human GTR-Fc or ab108557. Cells were stained with anti-Human IgG (Fc specific) FITC conjugate for DLL4-Fc binding.



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Induction of Hes-1 with the treatment of recombinant Human DLL4-Fc (ab108557).

A Mouse preadipocyte cell line, 3T3L-1, was stimulated with 5 µg/ml of Human DLL4-Fc as in indicated time points and each cell lysate was prepared and subjected to western blot by using anti-Mouse Hes1 or GAPDH.

M: Marker.

Lane 1: hDLL4-Fc, 0 min.

Lane 2: hDLL4-Fc, 10 min.

Lane 3: hDLL4-Fc, 30 min.

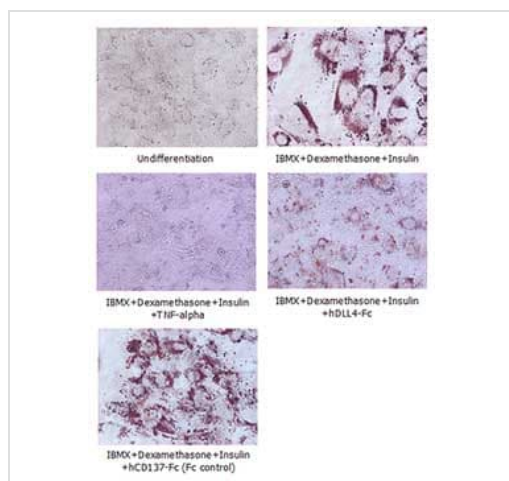
Lane 4: hDLL4-Fc, 1 hr.

Lane 5: hDLL4-Fc, 2 hr.

Lane 6: hDLL4-Fc, 4 hr.

Lane 7: hDLL4-Fc, 8 hr.

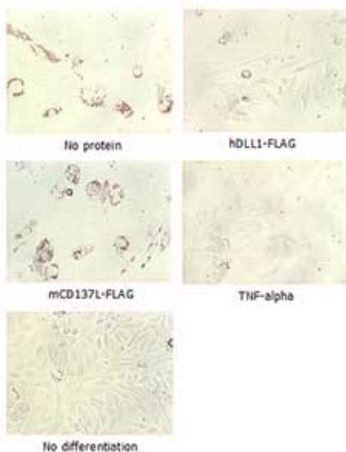
Lane 8: hDLL4-Fc, 24 hr.



Functional Studies - Recombinant human DLL4 protein (Fc Chimera Active) (ab108557)

Adipogenesis inhibition of 3T3L-1 cells.

3T3L-1 cells (mouse pre-adipocyte cells) were maintained in DMEM, supplemented with 10% fetal bovine serum and penicillin-streptomycin. For differentiation of 3T3L-1 cells, 3T3L-1 cells were cultured in adipogenic medium which was growth medium supplemented with 1 µM Dexamethasone, 0.5 mM IBMX, 10 µg/ml Insulin (day 0). Medium was changed every 2 days. Staining with Oil Red O was typically performed on day 7. Cells were washed twice with PBS, fixed with 3.7% formalin, and stained with 0.5% filtered Oil Red O in propylene glycol. For negative controls, mouse TNF-α (20 ng/ml) was added. Recombinant Human DLL4-Fc (ab108557) (5 µg/ml) dissolved in DPBS was added to the differentiation medium. These plates were then used to differentiate 3T3L-1 cells.



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Adipogenesis inhibition of MSCs.

MSCs (Mesenchymal stem cells) were maintained in DMEM, supplemented with 10% fetal bovine serum, penicillin streptomycin and glutamine. For differentiation of MSCs, MSCs were cultured in adipogenic medium which was growth medium supplemented with 1 μ M Dexamethasone, 0.5mM IBMX, 10 μ g/ml Insulin, 100 μ M Indomethacin (day 1). Medium was changed every 3 days. Staining with Oil Red O was typically performed on day 30. For negative controls, TNF- α (20 ng/ml) was added. To immobilize Notch ligands on the plastic surface of the culture plates, plates were incubated with a solution of ab108557 (5 μ g/ml) or mCD137-Fc (5 μ g/ml) in PBS for 2 hours at 37°C. Plates were then used to differentiate MSCs.



Functional Studies - Recombinant human DLL4 protein (Fc Chimera Active) (ab108557)

Adipogenesis inhibition of 3T3L-1 cells.



Functional Studies - Recombinant human DLL4 protein (Fc Chimera Active) (ab108557)

50 μ g of cell lysates derived from hDLL4-Fc (ab108557) or non-treated 3T3L-1 cells, which had been either differentiated or undifferentiated, and were subjected to Western blot by using a Mouse adiponectin antibody.

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