

Product Sheet

H_BDCA2 Reporter DDX35™ Jurkat Cell Line

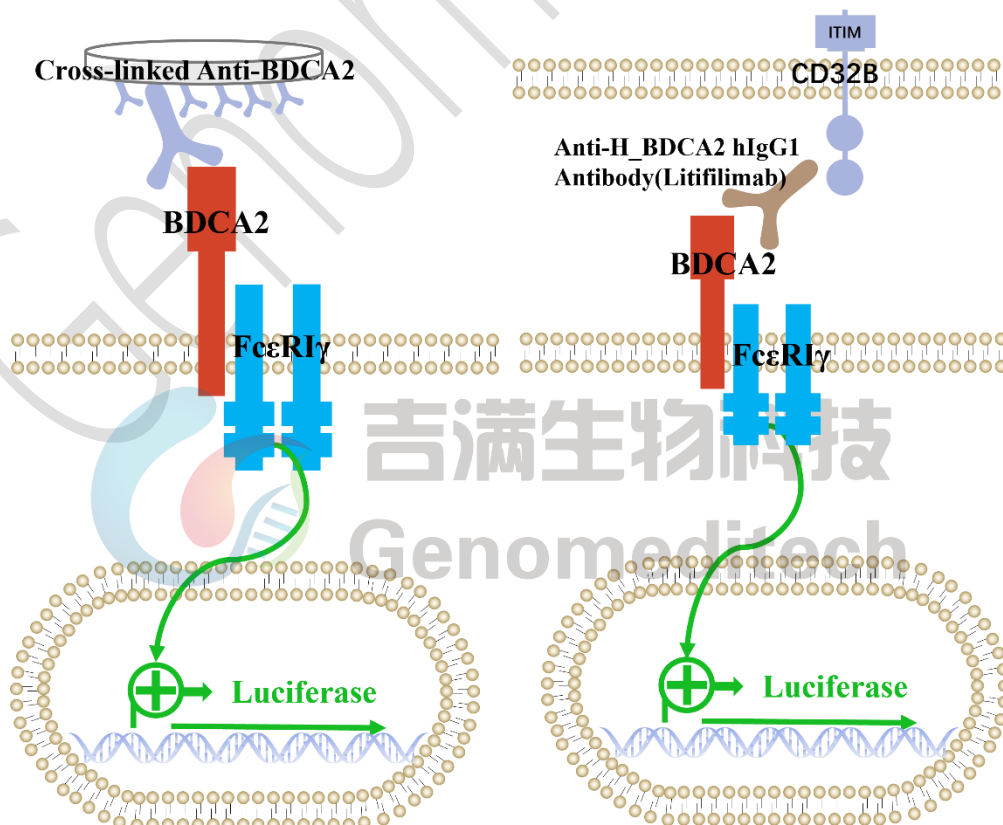
Catalog number: GM-C39599

Version 3.3.1.251212

Blood dendritic cell antigen 2 (BDCA2) is a C-type lectin expressed on plasmacytoid dendritic cells (pDCs) and is implicated in lupus pathogenesis. It consists of a single C-terminal extracellular carbohydrate recognition domain (CRD) of the class II C-type lectin family, a transmembrane region, and a short N-terminal cytoplasmic tail lacking a signaling motif. BDCA2 signals via the associated transmembrane adaptor FcεRIγ, triggering B cell receptor (BCR)-like signaling cascades. However, the ability of humanized anti-BDCA2 monoclonal antibodies to reduce disease activity in patients with cutaneous lupus remains poorly characterized.

H_BDCA2 Reporter DDX35™ Jurkat Cell Line is a clonal stable Jurkat cell line constitutively expressing human BDCA2 and Fc ε RIγ gene, along with signal-dependent expression of a luciferase reporter gene. When drug stimulation is applied, it activates downstream signaling pathways, leading to the expression of luciferase. The measurement of luciferase activity indicates the activation level of the signaling pathway and can thus be used to evaluate the in vitro effects of an antibody targeting BDCA2.

H_BDCA2 Reporter DDX35™ Jurkat Cell Line was obtained through extensive monoclonal screening and multiple rounds of monoclonal selection. It possesses high stability, high sensitivity, and high amplification properties, meeting the standards for customers' batch library construction and release experiments.



Specifications

Quantity	5E6 Cells per vial, 1 mL
Product Format	1 vial of frozen cells
Shipping	Shipped on dry ice
Storage Conditions	Liquid nitrogen immediately upon receipt
Recovery Medium	RPMI 1640+10% FBS+1% P.S
Growth medium	RPMI 1640+10% FBS+1% P.S+3.5 µg/mL Blasticidin+400 µg/mL G418+0.75 µg/mL Puromycin
Note	None
Freezing Medium	90% FBS+10% DMSO
Growth properties	Suspension
Growth Conditions	37°C, 5% CO ₂
Mycoplasma Testing	The cell line has been screened to confirm the absence of Mycoplasma species.
Safety considerations	Biosafety Level 2
Note	It is recommended to expand the cell culture and store a minimum of 10 vials at an early passage for potential future use.

Materials

Reagent	Manufacturer/Catalogue No.
RPMI 1640	gibco/C11875500BT
Fetal Bovine Serum	ExCell/FSP500
Pen/Strep	Thermo/15140-122
Blasticidin	Genomeditech/ GM-040404
G418	Genomeditech/ GM-040402
Puromycin	Genomeditech/ GM-040401
H_FCGR2B(CD32B) CHO-K1 Cell Line	Genomeditech/ GM-C16925
Anti-H_BDCA2 hIgG1 Antibody(Litifilimab)	Genomeditech/ GM-31294AB
GMOne-Step 2.0 Luciferase Reporter Gene Assay Kit	Genomeditech/ GM-040513

Figures

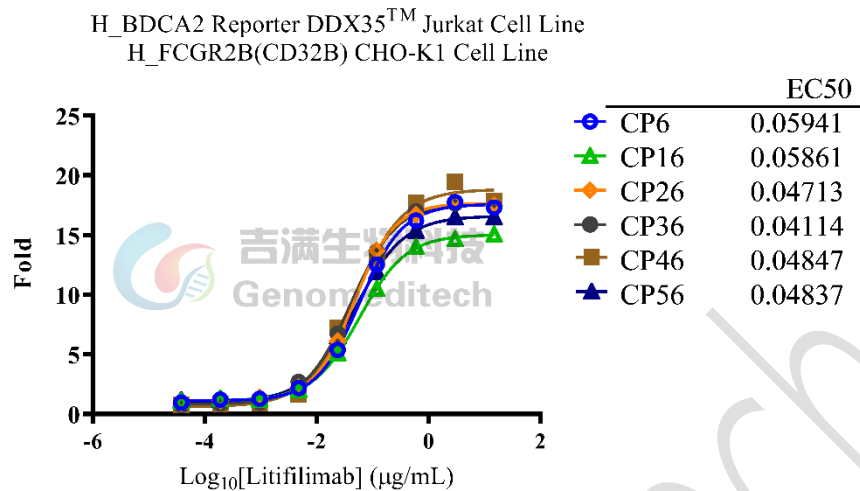


Figure 1 | The passage stability of response to Anti-H_BDCA2 hIgG1 Antibody(Litifilimab). Serial dilutions of the Anti-H_BDCA2 hIgG1 Antibody(Litifilimab) (Cat. [GM-31294AB](#)) and 1E5 cells/well of the passage 6, 16, 26, 36, 46 and 56 of the H_BDCA2 Reporter DDX35TM Jurkat Cell Line (Cat. GM-C39599) were added to 1E4 cells/well of H_FCGR2B(CD32B) CHO-K1 Cell Line (Cat. [GM-C16925](#)) for 16 hours. The firefly luciferase activity was measured using the Luciferase Reporter Assay Kit (Genomeditech). Data are shown by drug mass concentration.

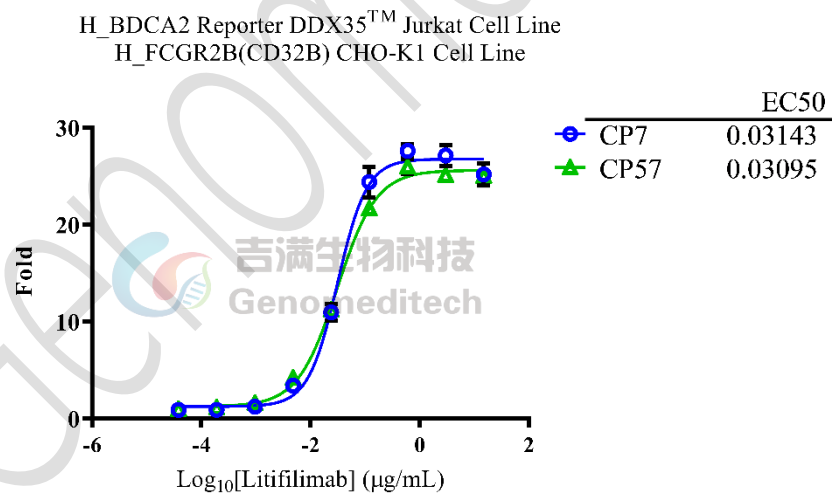


Figure 2 | The passage stability of response to Anti-H_BDCA2 hIgG1 Antibody(Litifilimab). Serial dilutions of the Anti-H_BDCA2 hIgG1 Antibody(Litifilimab)(Cat. [GM-31294AB](#)) and 1E5 cells/well of the passage 7 and 57 of H_BDCA2 Reporter DDX35TM Jurkat Cell Line (Cat. GM-C26024) were added to 1E4 cells/well of H_FCGR2B(CD32B) CHO-K1 Cell Line (Cat. [GM-C16925](#)) for 16 hours. The firefly luciferase activity was measured using the Luciferase Reporter Assay Kit (Genomeditech). Data are shown by drug mass concentration.

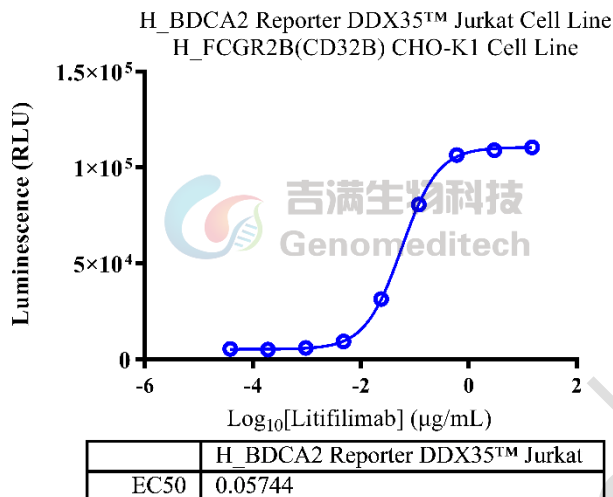


Figure 3 | Response to Anti-H_BDCA2 hIgG1 Antibody(Litifilimab). Serial dilutions of the Anti-H_BDCA2 hIgG1 Antibody(Litifilimab) (Cat. GM-C39599) and 1E5 cells/well of the H_BDCA2 Reporter DDX35™ Jurkat Cell Line (Cat. GM-C39599) were added to 1E4 cells/well of H_FCGR2B CHO-K1 Cell Line (Cat. M-C16925) for 16 hours. The firefly luciferase activity was measured using the Luciferase Reporter Assay Kit (Genomeditech). The maximum induction fold was approximately[22.4]. Data are shown by drug mass concentration.

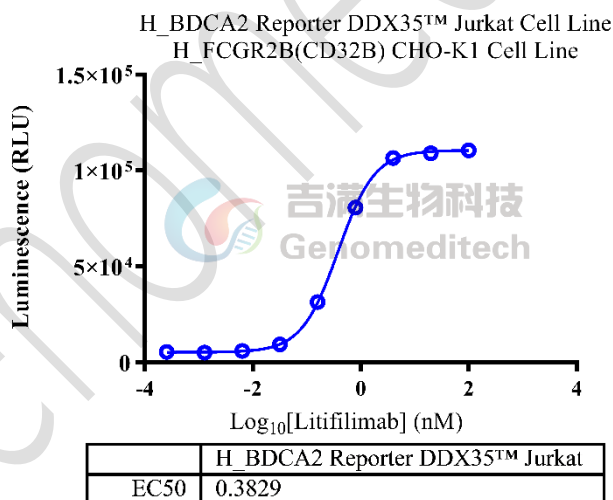


Figure 4 | Response to Anti-H_BDCA2 hIgG1 Antibody(Litifilimab). Serial dilutions of the Anti-H_BDCA2 hIgG1 Antibody(Litifilimab) (Cat. GM-C39599) and 1E5 cells/well of the H_BDCA2 Reporter DDX35™ Jurkat Cell Line (Cat. GM-C39599) were added to 1E4 cells/well of H_FCGR2B CHO-K1 Cell Line (Cat. M-C16925) for 16 hours. The firefly luciferase activity was measured using the Luciferase Reporter Assay Kit (Genomeditech). The maximum induction fold was approximately[22.4].data is shown by drug molar concentration.

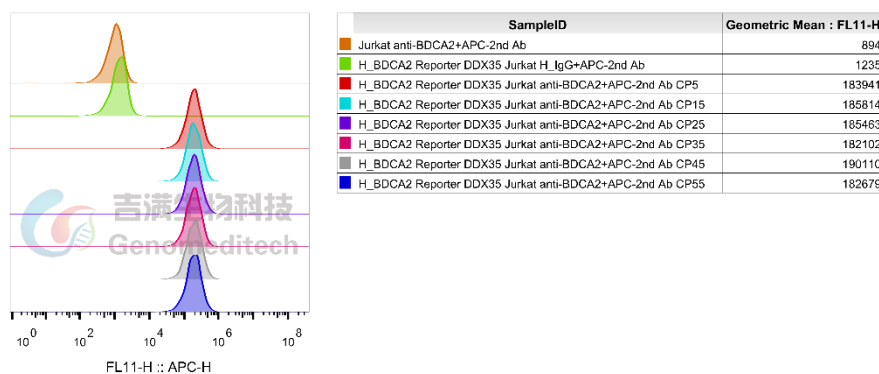


Figure 5 | The passage stability of the H_BDCA2 Reporter DDX35™ Jurkat Cell Line (Cat. GM-C39599) was determined by flow cytometry using AAnti-H_BDCA2 hIgG1 Antibody(Litifilimab) (Cat. [GM-31294AB](#)).

Cell Recovery

Recovery Medium: RPMI 1640+10% FBS+1% P.S

To insure the highest level of viability, thaw the vial and initiate the culture as soon as possible upon receipt. If upon arrival, continued storage of the frozen culture is necessary, it should be stored in liquid nitrogen vapor phase and not at -70°C. Storage at -70°C will result in loss of viability.

- Thaw the vial by gentle agitation in a 37°C water bath. To reduce the possibility of contamination, keep the O-ring and cap out of the water. Thawing should be rapid (approximately 2 - 3 minutes).
- Remove the vial from the water bath as soon as the contents are thawed, and decontaminate by dipping in or spraying with 70% ethanol. All of the operations from this point on should be carried out under strict aseptic conditions.
- Transfer the vial contents to a centrifuge tube containing 5.0 mL complete culture medium. And spin at approximately 176 x g for 5 minutes. Discard supernatant.
- Resuspend cell pellet with the recommended complete medium. And dispense the suspension into 1 - 2 T-25 culture flasks.
- Incubate the culture at 37°C in a suitable incubator. A 5% CO₂ in air atmosphere is recommended if using the medium described on this product sheet.

Cell Freezing

Freezing Medium: 90% FBS+10% DMSO

- Centrifuge at 176 x g for 3 minutes to collect cells.
- Resuspend the cells in pre-cooled freezing medium and adjust the cell density to 5E6 cells/mL.
- Aliquot 1 mL into each vial.
- Place the vial in a controlled-rate freezing container and store at -80°C for at least 1 day, then transfer to liquid nitrogen as soon as possible.

Cell passage

Growth medium: RPMI 1640+10% FBS+1% P.S+3.5 µg/mL Blasticidin+400 µg/mL G418+0.75 µg/mL Puromycin

Approximately 48-72 hours after the initial thawing, the cells can be passaged for the first time. After this initial passage, the culture medium can be adjusted to growth medium supplemented with antibiotics. If cells are not passaged within 48 hours, it is recommended to add some fresh recovery medium and place the flask horizontally.

- When the cell density reaches 1.5 - 2E6 cells/mL, subculture the cells. Do not allow the cell density to exceed 2E6 cells/mL.
- It is recommended to use T-25 flasks for subculturing.
- These cells are suspension cells, and it is recommended to use the "half-medium change" method to maintain optimal cell conditions during passaging.
- During passaging, you can directly add fresh growth medium to the culture flask, gently pipette to resuspend the cells, and then transfer the cell suspension to a new T-25 flask for continued culture.

Subcultivation Ratio: Maintain cultures at a cell concentraion between 3E5 and 1E6 viable cells/mL.

Medium Renewal: Every 2 to 3 days

Notes

- These cells are sensitive to density, so please ensure that the cell density is maintained within an appropriate range during culture and subculturing.
- During the first passage, pay attention to the nutrient supply; if not subculturing, make sure to add fresh recovery medium every other day as needed.

Related Products

CD40:CD40L	
H_CD40(TNFRSF5) Reporter 293 Cell Line	H_CD40(TNFRSF5) Reporter Jurkat Cell Line
Cynomolgus_CD40 CHO-K1 Cell Line	Cynomolgus_CD40L CHO-K1 Cell Line
H_CD40(TNFRSF5) CHO-K1 Cell Line	H_CD40(TNFRSF5) HEK-293 Cell Line
H_CD40L CHO-K1 Cell Line	H_CD40L HEK-293 Cell Line
Rabbit_CD40L NIH-3T3 Cell Line	
Anti-CD40L hIgG1 Reference Antibody (Frebio)	Anti-H_CD40 hIgG1 Antibody(APX005M)
Anti-H_CD40 hIgG1 Antibody(ravagalimab)	Anti-H_CD40L hIgG1 Antibody(dapirolizumab)
Anti-H_CD40L hIgG1 Antibody(frexalimab)	
Biotinylated Human CD40 Protein; His-Avi Tag	Cynomolgus CD40 Protein; His Tag
Human CD40 Protein; His Tag	Human CD40L Protein; His Tag
IFN-α	
IFNα Reporter HEK-293 Cell Line	IFNα Reporter MDCK Cell Line
IFNα Reporter THP1 Cell Line	
BCMA:BAFFR:TACI	

H_BAFFR Jurkat Blockade Reporter Cell Line	H_BAFFR Reporter Cell Line
H_BCMA Reporter Cell Line	H_TACI Reporter Cell Line
Cynomolgus_BCMA CHO-K1 Cell Line	Cynomolgus_BCMA HEK-293 Cell Line
H_BCMA CHO-K1 Cell Line	H_BCMA HEK-293 Cell Line
Anti-BAFF hIgG1 Antibody(belimumab)	Anti-BAFFR hIgG1 Antibody(ianalumab)
Anti-BCMA hIgG1 Antibody(Belantamab)	Anti-BCMA hIgG1 Antibody(SEA-BCMA)
Anti-BCMA hIgG4 Antibody(BCMB69)	
Biotinylated Human BAFF Protein; His-Avi Tag	Biotinylated Human BCMA Protein; His-Avi Tag
Cynomolgus BAFF Protein; His Tag	Cynomolgus BCMA Protein; hFc Tag
Cynomolgus BCMA Protein; His Tag	Human APRIL Protein; hFc Tag
Human BAFF Protein; His Tag	Human BCMA Protein; hFc Tag
Human BCMA Protein; His Tag	Mouse BAFF Protein; His Tag
BDCA2(CLEC4C)	
H_BDCA2 Reporter Jurkat Cell Line	Cynomolgus_BDCA2 CHO-K1 Cell Line
Cynomolgus_BDCA2 Jurkat Cell Line	H_BDCA2 CHO-K1 Cell Line
H_BDCA2 HEK-293 Cell Line	H_BDCA2 Jurkat Cell Line
Anti-H_BDCA2 hIgG1 Antibody(Litilimab)	
Cynomolgus BDCA2 Protein; His Tag	Human BDCA2 Protein; His Tag

License Agreement:

By purchasing and using this cell line product, the user voluntarily agrees to accept and abide by the following policies:

- This cell line product is restricted to research use only and shall not be used for any commercial purposes.
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