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- Diagnostik & molekulare Diagnostik
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Lieferung & Zahlungsart

siehe unsere Liefer- und Versandbedingungen

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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Description

The PBMC Cytotoxicity Luciferase Assay Kit is a kit designed to determine the cytotoxicity profile of PBMCs (Peripheral Blood Mononuclear Cells) towards a target cell line expressing Firefly luciferase. It uses the luminescence signal from Firefly luciferase expressing target cells to measure the number of live target cells within a mixed cell population of PBMCs and target cells. The kit contains PBMCs, Thaw Medium and One-Step™ Luciferase Assay System.

Background

Lymphocyte-mediated cytotoxicity is a form of cellular immunity against intracellular pathogens, including viruses and certain bacteria and parasites. The most popular *in vitro* methods to monitor lymphocyte-mediated cytotoxicity on target cells are the cell-mediated cytotoxicity assays such as ADCC (antibody-dependent cellular cytotoxicity) and TDCC (T-cell dependent cellular cytotoxicity) in which immune effector cells and target cells are co-cultured. To analyze immune effector cell cytolytic activity in such heterogeneous cell population of effector and target cells, it is important to be able to discriminate between effector and target cell populations with distinct phenotypes. The use of luciferase allows for a clear separation between the effector and the surviving target cells. The instability of Firefly luciferase when released from dead target cells in cell culture gives it a half-life of approximately 2 hours, eliminating any residual luminescence signal generated from dead target cells. Cytotoxicity assays are crucial to understand the potency of CAR (chimeric antigen receptor) T and NK cells, and antibody-based immunotherapies.

Application(s)

- Luciferase-based analysis of live and dead target cells in cytotoxicity assays.
- Test the efficacy of multi-specific immune engager molecules.
- Assess the Fc effector function of candidate antibodies.

Supplied Materials

Catalog #	Name	Amount	Storage
79059	Normal Human Peripheral Blood Mononuclear Cells, Frozen	2 vials at 10 x 10^6 cells each	Liquid Nitrogen
60184	Thaw Medium 2	2 x 100 ml	4°C
60690-1	ONE-Step™ Luciferase Assay System	2 x 10 ml kit	-20°C

Materials Required but Not Supplied

- Target cell line of interest expressing Firefly luciferase.
- 96 well white, clear bottom plate
- T75 cell culture flask.
- Luminometer.

Storage Conditions

This assay kit will perform optimally for up to **6 months** from date of receipt when the materials are stored as directed.

Safety

This product is for research purposes only and not for human or therapeutic use. This product should be considered hazardous and is harmful by inhalation, in contact with skin, eyes, clothing, and if swallowed. If contact occurs, wash thoroughly.

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Assay Protocol

- This protocol is a general guideline only.
- This protocol is designed to perform the cytotoxicity assay in a 96-well plate. To perform the assay in different tissue culture formats, the cell number and reagent volume should be scaled appropriately.
- Each vial of human PBMCs (effector cells) is sufficient for 60 wells of a 96-well plate at an effector to target cell ratio (E:T) of 10:1 (1 x 10^5 effector cells:1 x 10^4 target cells). For a higher E:T cell ratio you may need to thaw both PBMC vials supplied.
- Any cell line that constitutively expresses the Firefly (*Photinus pyralis*) luciferase reporter and the desired antigen can be used as target cells in this assay. For example, Firefly Luciferase NALM6 Cell Line (BPS Bioscience #78494), expressing the firefly luciferase reporter under the control of a CMV promoter, can be used.
- The antibody dilution range should be optimized for your assay. A starting concentration of 50 nM is recommended as the highest value in the preparation of 5X antibody dilutions.
- We recommend the use of the following experimental controls:
    - Control 1: No antibody control. This control contains both PBMCs and target cells without antibody. This control is used to measure the maximum luminescence signal in the assay.
    - Control 2: PBMCs cells only. This control is used to determine the background luminescence signal.
    - Control 3: Antibody control. This control contains both effector and target cells in the presence of serial dilutions of a non-specific antibody (antibody of the same class and isotype as the specific antibody but unable to recognize the target).

One week prior to running the assay: Target Cell Thaw and Expansion

1. Thaw target cell line of interest.

2. Expand cells using the appropriate cell culture conditions for the cell line of interest.

3. Passage cells at least once to make sure they are healthy (2 x 10^6 cells are needed for the assay described below).

   Note: If using Firefly Luciferase NALM6 Cell line the cell culture and maintenance conditions can be found at Firefly Luciferase NALM6 Cell Line (bpsbioscience.com).

Day 1: PBMC Cell Preparation

1. Thaw one vial of PBMCs by swirling the vial of frozen cells for approximately 60 seconds in a 37°C water bath. As soon as the cells are thawed (it may be slightly faster or slower than 60 seconds), quickly transfer the entire content of the vial to a tube containing 10 ml of pre-warmed Thaw Medium 2.

   Note: Leaving the cells in the water bath at 37°C for too long will result in rapid loss of viability.

2. Spin down at 300 *x g* for 5 minutes, aspirate supernatant, and resuspend cell pellet in 10 ml of Thaw Medium 2 (1 x 10^6 cells/ml).

3. Plate cells in a T75 flask.

4. Incubate the flask overnight in a humidified 37°C incubator with 5% CO_2.

   Note: This step will enrich the lymphocyte population by depleting adherent cells.

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Day 2: Assay

For 96-well plate assays, each well will contain a final volume of 125 µl (25 µl of 5X antibody dilution, 50 µl PBMCs at desired E:T ratio and 50 µl of target cells).

1. Transfer 2 x 10^6 target cells to a clean 15 ml tube and centrifuge at 300 *x g* for 5 minutes.
2. Aspirate supernatant and resuspend target cells in 10 ml of Thaw Medium 2 (2 x 10^5 cells/ml).
3. Transfer cells to a solution reservoir.
4. Using a multichannel pipette, transfer 50 µl of target cell suspension (10,000 cells/well) to the Test Antibody, Control 1, and Control 3 wells.
5. Using a multichannel pipette, transfer 50 µl of Thaw Medium 2 to the Control 2 wells.
6. Keep the plate in a humidified 37°C incubator with 5% CO_2 while preparing PBMCs.
7. Collect PBMCs into a 15 ml tube and count cells.

   Note: Be careful to avoid detachment of the adherent cells by not shaking the T75 flask prior to or while transferring cells.
8. Centrifuge PBMCs at 300 *x g* for 5 minutes and aspirate the supernatant.
9. Dilute PBMCs in Thaw Medium 2 to 2 x 10^6 cells/ml.

   Note E:T ratio may need to be optimized in different experimental settings and cell density may need to be adjusted.
10. Add 50 µl of PBMC suspension to the Test Antibody, Control 1, Control 2, and Control 3 wells.
11. Keep the plate in a humidified 37°C incubator with 5% CO_2 while you are preparing antibody dilutions.
12. Prepare test antibody and antibody control dilutions at 5x the final concentrations to be tested, in Thaw Medium 2 (25 µl/well), starting at 50 nM.
13. Add 25 µl of antibody dilutions to the test antibody wells.
14. Add 25 µl of antibody control dilutions to the Control 3 wells.
15. Add 25 µl of Thaw Medium 2 to Control 1, and Control 2 wells.
16. Incubate the assay plate 24 hours in a humidified 37°C incubator with 5% CO_2.

    Note: The incubation time may need to be optimized for your assay.

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Example of Plate Schematic:

	1	2	3	4	5	6	7	8	9	10	11	12	
A	Dilu12	Dilu11	Dilu10	Dilu9	Dilu8	Dilu7	Dilu6	Dilu5	Dilu4	Dilu3	Dilu2	Dilu1	Test antibody
B	Dilu12	Dilu11	Dilu10	Dilu9	Dilu8	Dilu7	Dilu6	Dilu5	Dilu4	Dilu3	Dilu2	Dilu1	
C	Dilu12	Dilu11	Dilu10	Dilu9	Dilu8	Dilu7	Dilu6	Dilu5	Dilu4	Dilu3	Dilu2	Dilu1	Control 3
D	Dilu12	Dilu11	Dilu10	Dilu9	Dilu8	Dilu7	Dilu6	Dilu5	Dilu4	Dilu3	Dilu2	Dilu1	
E	Control 1	Control 2											
F	Control 1	Control 2											
G													
H													

Day 3: Luciferase Analysis

1. Thaw Luciferase Reagent Buffer (Component A) by placing the reagent in a Room Temperature (RT) water bath.

2. Equilibrate the buffer to RT and mix well before use.

3. Immediately before the experiment, prepare the Luciferase Assay Working Solution by diluting Luciferase Reagent Substrate (Component B) 100-fold with Luciferase Reagent Buffer (Component A), and mix well (you will need 125 µl/well).

   Note: Avoid exposure to excessive light. Only use enough of each component for the experiment, and store the remaining Component A and Component B separately at -20°C.

4. Remove the cells from the incubator and add 125 µl of Luciferase Assay Working Solution directly to the culture medium of each well.

5. Wrap the plate with foil and gently rock it for ≥15 minutes at RT.

6. Measure firefly luminescence using a luminometer.

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Example Results

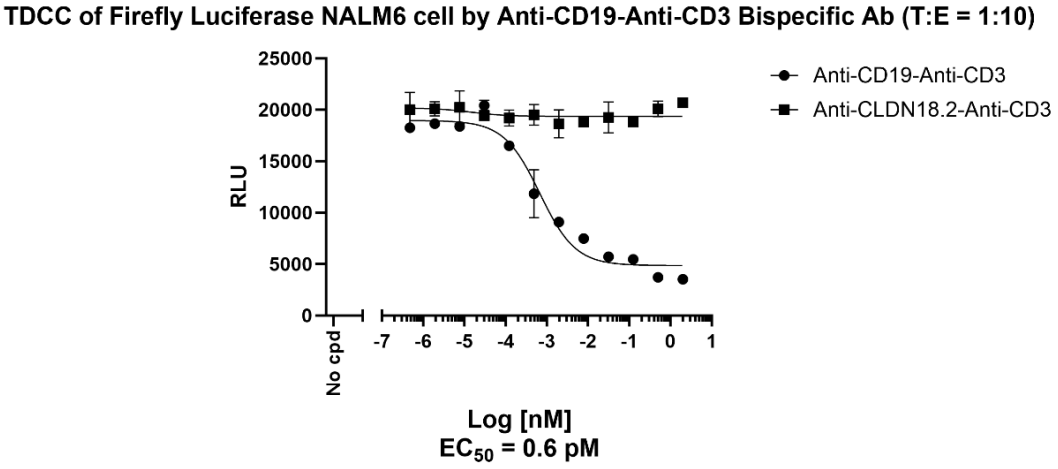

Figure 1. T cell-dependent cellular cytotoxicity (TDCC) of Firefly Luciferase NALM6 Cell Line when triggered by the Anti-CD19-Anti-CD3 Bispecific Molecule.
PBMCs and Firefly Luciferase NALM6 cells were combined at a 10:1 ratio in a 96-well white, clear bottom plate. The cells were incubated with a dilution series of Anti-CD19-Anti-CD3 Bispecific Molecule (BPS Bioscience #100441) or the antibody control, CLDN18.2-Anti-CD3 Bispecific Antibody (BPS Bioscience #101541), in a humidified 37°C incubator with 5% CO_2 for 24 hours. After incubation, luciferase activity was measured with One-Step™ Luciferase reagent. The raw luminescence data were fitted to a sigmoidal three-parameter curve using GraphPad Prism® software.

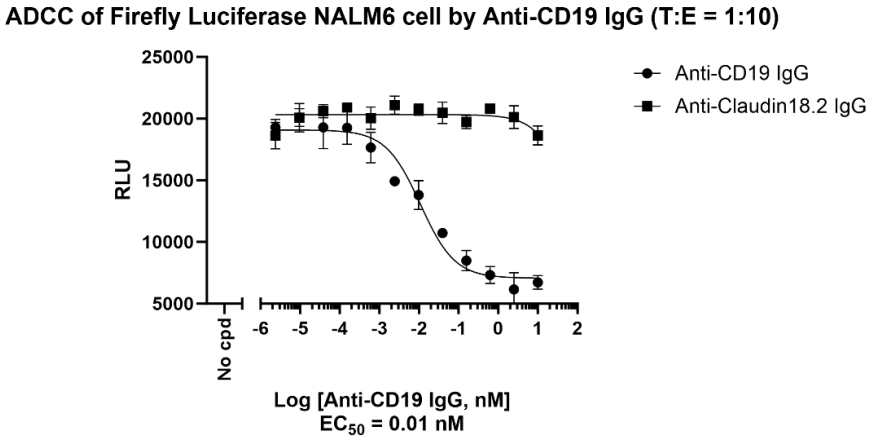

Figure 2. Antibody-dependent cellular cytotoxicity (ADCC) of Firefly Luciferase NALM6 Cell Line triggered by Anti-CD19 IgG.
PBMCs and Firefly Luciferase NALM6 cells were combined at a 10:1 ratio in a 96-well white, clear bottom plate. The cells were incubated with a dilution series of Anti-CD19 IgG (BPS Bioscience #100981) or the control Anti-Claudin-18 Isoform 2 IgG Antibody (BPS Bioscience #101564). After incubation for 24 hours in a humidified 37°C incubator with 5% CO_2, luciferase activity was measured with One-Step™ Luciferase reagent. The raw luminescence data were fitted to a sigmoidal three-parameter curve using GraphPad Prism® software.

Data shown is representative. For lot-specific information, please contact BPS Bioscience, Inc. at support@bpsbioscience.com.

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Troubleshooting Guide

Visit bpsbioscience.com/assay-kits-faq for detailed troubleshooting instructions. For all further questions, please email support@bpsbioscience.com

Related Products

Products	*Catalog #*	*Size*
PBMC Cytotoxicity Bioassay Kit (CFSE, 7-ADD)	82173	1 kit
PBMC Cytotoxicity Luciferase Assay Kit (NALM6)	82174	1 kit
Anti-CD4 Antibody, PE-Labeled	102010	25 µg/100 µg
Anti-CD8 Antibody, PE-Labeled	102011	25 µg/100 µg

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