

A new fluorescent probe that can distinguish B cells from T cells

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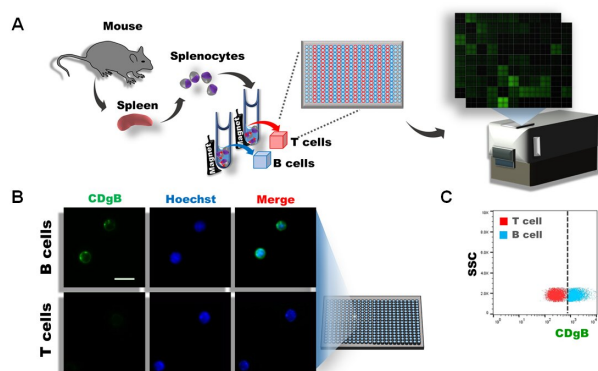


Figure 1. A) Splenocytes were isolated from mouse spleens, and T and B cells were separated using magnetic activated cell sorting (MACS). The cells were subsequently plated in 384 wells and over 10,000 different fluorescent molecules were screened using DOFLA. B) Fluorescence microscope image shows that CDgB stains B cells but not T cells. C) Flow cytometry graph of CDgB fluorescence vs SSC between T cell and B cell populations. Credit: Institute for Basic Science

Human blood contains several different components, including plasma, red blood cells (RBCs), white blood cells (WBCs), and platelets. Among these, WBCs are divided into numerous subcategories each with unique functions and characteristics, such as lymphocytes, monocytes, neutrophils, and others. Lymphocytes are further subdivided into T lymphocytes, B lymphocytes, and NK cells. Distinguishing and separating different types of these cells is highly important in carrying out studies in the field of immunology.

Discriminating different immune cell types is typically done by flow cytometry and fluorescence-activated cell sorting (FACS), which can identify distinct populations of [cells](#) according to their size, granularity, and fluorescence. While size and granularity alone cannot distinguish cells with similar physical parameters, different types of

immune cells display a distinct combination of immune receptors on the cell surfaces. For example, the T [lymphocytes](#) and B lymphocytes express CD3 and CD19, respectively. Therefore, fluorescently identifying immune cells have relied on staining the cells using multiple antibodies against different receptors. It has been long thought that it was impossible to distinguish different immune cell types without using these antibodies.

However, novel breakthrough research performed by the scientists at the Center for Self-assembly and Complexity within the Institute for Basic Science, South Korea, may have just changed this. The researchers employed a diversity oriented fluorescence library approach (DOFLA) to screen over 10,000 different fluorescent molecules using the B and T lymphocytes separated from mouse spleens. From this, they managed to discover a new fluorescent probe that can discriminate B lymphocytes over T lymphocytes without the cell receptor targeting antibodies.

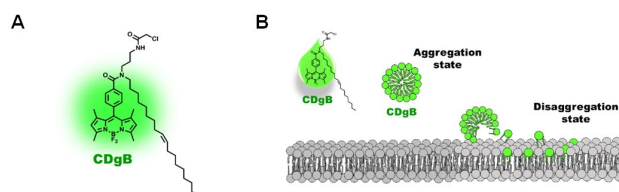


Figure 2. A) Structure of CDgB molecule. B) Free unbound CDgB forms aggregates in solution, which dissociates when they bind to lipid membranes. Credit: Institute for Basic Science

The researchers called the new probe CDgB, which stands for Compound of Designation green for B lymphocytes. CDgB is a lipophilic molecule that contains a fluorescent component attached to a hydrocarbon chain. As CDgB contains both a polar

fluorescent group and a hydrocarbon tail, it means that the free unbound CDgB dye molecules form aggregates similar to micelles in the solution and exhibit a low level of background fluorescence. When they attach to the cell surfaces, however, the aggregates dissociate and cause a spike in fluorescence signals. In addition, the lipophilic nature of the dye means that the dye does not bind to a protein target, and instead localizes to the lipid [membrane](#) structure directly. According to the researchers, this was "the first example to report such type of cell distinction mechanism."

The researchers further elucidated this mechanism by performing lipidomic analysis of B and T cell membranes. Phosphatidylcholine (PC) comprises the majority (> 60%) of the membrane phospholipids of both B and T lymphocytes. It was found that B lymphocytes in general had shorter PC's than that of the T lymphocytes. In addition, the membrane cholesterol content in the T lymphocytes was about twice higher than that of B lymphocytes. These factors give the B lymphocytes a more 'flexible' cell membrane, which was thought to be a crucial factor that explains why the CDgB molecules attach more readily to the cell membranes of the B lymphocytes over those of the T lymphocytes. Even among the B lymphocytes, it was found that the strength of the fluorescence was different based on the cell maturity. The B cell progenitors and immature B cells gave off much brighter fluorescence signals than mature B cells, which is most likely due to the higher membrane flexibility in the immature cells.

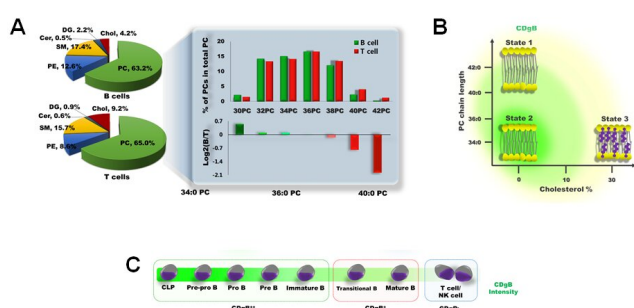


Figure 3. A) Difference in membrane lipid composition between B cells and T cells. B) CDgB selectively stains lipid membranes of B cells which possess shorter PC chain length and a lower percentage of cholesterol. C) CDgB shows higher fluorescence in immature B cells (CDgB++) than in mature B cells (CDgB+) or other lymphocytes (CDgB-). Credit: Institute for Basic Science

The researchers furthermore concluded that this new lipid oriented live cell distinction (LOLD) mechanism can supplement the existing cell distinction mechanism to improve our ability to distinguish specific cell types from complicated mixtures of different cells. This research was published in the *Journal of the American Chemical Society*.

More information: Lipid Oriented Live-Cell Distinction of B and T Lymphocytes, *Journal of American Chemical Society* (2021) doi.org/10.1021/jacs.1c00944

The CDgB is able to selectively target the cell membranes of B lymphocytes over T lymphocytes or NK cells. The researchers sought to optimize the selectivity of the CDgB by testing different derivatives of the molecules with various hydrocarbon chain lengths from 4 to 20 carbons. It was found that the CDgB derivatives with 14 to 18 carbons showed the highest selectivity towards the B lymphocytes, with C18 showing the best results. It became more difficult to distinguish the cells through fluorescence when the carbon length was increased beyond 20. The fact that carbon length matters in the selectivity hinted that the mechanism was dependent on the difference in the membrane structures between B and T lymphocytes.

Provided by Institute for Basic Science

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