

# Study reveals the structure of the second human cannabinoid receptor

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Cannabinoid receptors in humans respond to compounds found in cannabis. Credit: @tsarcyanide/MIPT

Studies indicate that CB2 is a promising target for immunotherapy, as well as treating inflammatory and neuropathic pain, and neurodegenerative diseases. It has also been shown that molecules blocking CB2 can reduce tumor growth.

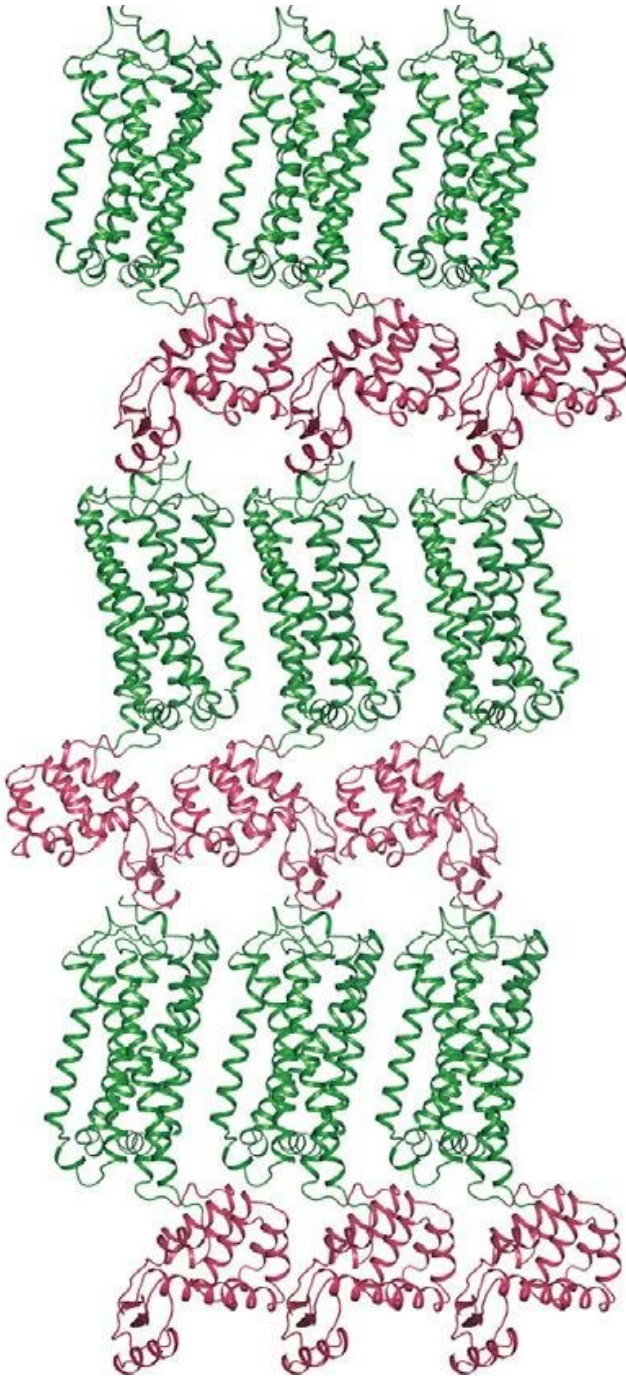
To effectively treat pathological conditions, drugs need to specifically target CB1 or CB2. However, the two receptors are very much alike. The [amino acid sequences](#) that encode them are 44 percent identical. So developing a selective medicine requires knowing the [structure](#) of both targets in great detail. Unlike CB1, the structure of CB2 has remained unknown up until now.

A Chinese research team joined forces with Russian and U.S. biologists to obtain the crystal structure of the human type 2 cannabinoid receptor. Their findings could lead to drugs against inflammatory, neurodegenerative, and other diseases. The authors of the paper, published in *Cell*, compare the newly discovered structure to that of the type 1 cannabinoid receptor, deeming the two receptors to be the yin and yang of the human endocannabinoid system.

## Drugs rely on knowing receptor structure

The two cannabinoid receptors, CB1 and CB2, belong to the so-called endocannabinoid system. This refers to a signaling system in the human body that regulates biological processes such as metabolism, pain sensation, neuronal activity, immune function, and others. It has been shown that the cannabinoid receptors can be targeted to alleviate certain pathological conditions, including chronic pain.

While the CB1 receptors are mostly found in the nervous system and are responsible for psychoactive effects, the CB2 receptors are predominantly present in the immune system.



Type 2 cannabinoid receptor crystal. Credit: Xiaoting Li *et al./Cell*

### X-raying CB2 crystal reveals its structure

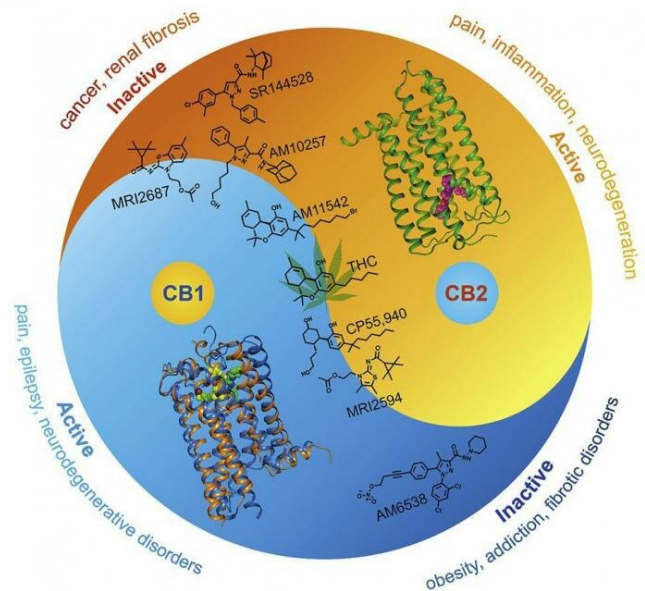
To identify the shape of an individual molecule, researchers make a crystal from many such molecules. When arranged in this highly ordered way, the molecules can be exposed to X-rays,

revealing their structure.

The team made a crystal (figure 1) from CB2 receptors bound to molecules blocking this receptor, which are potential [drug](#) candidates. That way, the X-ray analysis allowed the team to see both the structure of CB2 and how it connects to the blocking molecule, or antagonist.

However, receptors are unstable proteins by nature. To study them, they need to be modified using gene engineering. This involves introducing mutations that make the protein stable without changing its structure or function.

CompoMug is a software suite that predicts mutations potentially useful for stabilizing receptor [molecules](#). The mutations then need to be tested experimentally. This software was developed by two researchers from the Moscow Institute of Physics and Technology and the University of Southern California: visiting MIPT professor Vsevolod Katritch and Petr Popov from the Laboratory of Structural Biology of G Protein-Coupled Receptors.



Conditions that can be affected by activating or inhibiting type 1 (CB1) or type 2 (CB2) cannabinoid receptors independently. Credit: Xiaoting Li *et al./Cell*

In the study, the CB2 receptor had five mutations prompted by CompoMug.

### **Receptor structure discovery enables smarter drugs**

The researchers contrasted the structure of CB2 with that of CB1. They concluded that substances activating one of the receptors can actually weaken or inhibit the other, and vice versa. This opens up a possibility not just for drugs that target exclusively one receptor, but those that affect both receptors, but in different ways. Figure 2 shows the therapeutic implications.

"Every G protein-coupled [receptor structure](#) that is discovered has prospects for rational design of more efficient drugs," study co-author Petr Popov explains. "Now that the structures of both [cannabinoid receptors](#) are known, we can design selective compounds targeting only one of the receptors, as well as agents with a desired polypharmacological profile targeting both [receptors](#) at once."

**More information:** Xiaoting Li et al, Crystal Structure of the Human Cannabinoid Receptor CB2, *Cell* (2019). [DOI: 10.1016/j.cell.2018.12.011](https://doi.org/10.1016/j.cell.2018.12.011)

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