

New strategy for multiple myeloma immunotherapy

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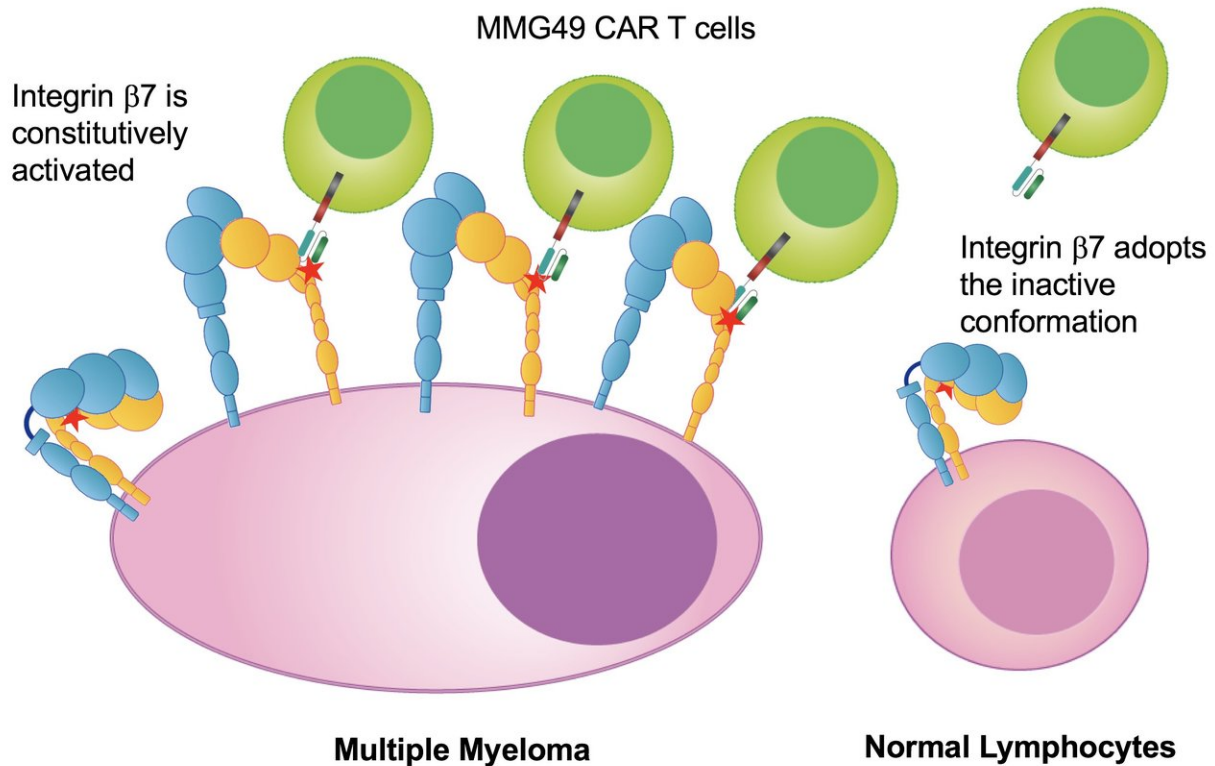


Fig.1.MMG49 CAR T cells target the activated conformation of integrin b7 expressed on MM cells. Credit: Osaka University

In recent decades monoclonal antibody-based treatment of cancer has been established as one of the most successful therapeutic strategies for both solid tumors and blood cancers. Monoclonal antibodies (mAb), as

the name implies, are antibodies that are made by clonal cells derived from a single parent cells and therefore share the identical amino acid sequences.

One of the leading technologies to emerge in mAb-based treatment is CAR-T, where CAR stands for "chimeric antigen receptor," and T represents T [cells](#), a type of [white blood cells](#) that have pivotal roles in immune defenses. CARs are produced by combining together the gene for an antibody that recognizes a tumor antigen with the gene for a receptor that resides on the surface of the T cells; insert this new gene into a T cell and it will be precisely targeted at the tumor.

Theoretically, new antigens – molecules capable of inducing an immune response to produce an antibody – that arise from cancer-specific mutations of cell-surface proteins are excellent targets. However, mAb therapy targeting such antigens is impractical because of these proteins' vast diversity within and between individual tumors, which renders identifying new cancer-specific target antigens difficult.

However, such challenges have driven researchers centered at Japan's Osaka University to think outside of the box; cancer-specific antigen formed by the modification of proteins during or after synthesis, such as glycosylation (attachment of sugar moieties to [protein](#)) or conformational changes, might have been missed in previous analyses. The team believed new antigen epitopes, which is the part of an antigen recognized by the immune cells, could be discovered by thoroughly searching for cancer-specific mAbs and characterizing the antigens they recognize.

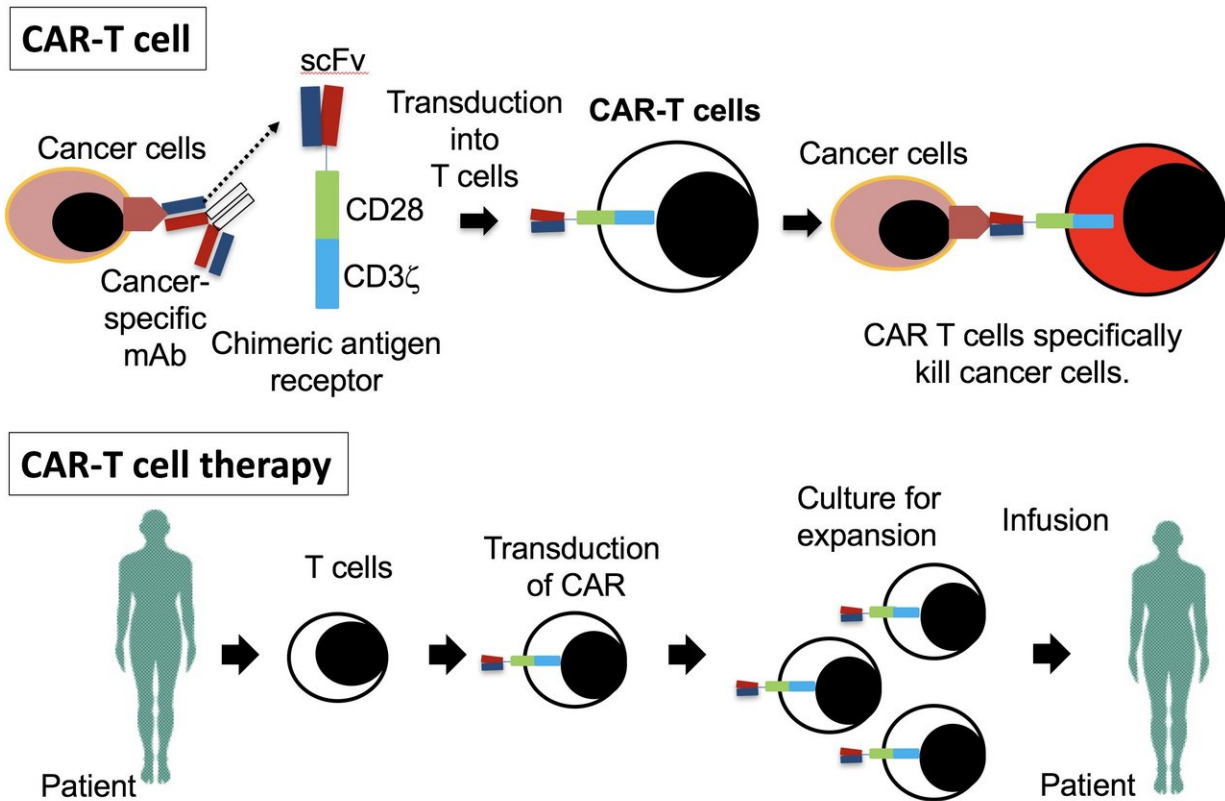


Fig.2. Outline of CAR T cell therapy. Credit: Osaka University

"We applied this strategy to identify novel therapeutic targets for multiple myeloma (MM), a cancer that forms in a type of white blood cell called a plasma cell," explains Naoki Hosen, lead author of the study, which was recently published in *Nature Medicine*. "Despite advances in MM treatment, relapse remains common. As such, there is an ongoing need for new therapeutic approaches, including mAb-based therapies."

The team screened more than 10,000 anti-MM mAb clones and identified MMG49 as an MM-specific mAb specifically recognizing a subset of integrin $\beta 7$, a cell-surface receptors that facilitate cell-

extracellular matrix adhesion. MMG49 reacted to MM cells, but not other bone marrow cell types in MM patient samples. This prompted the researchers to design a CAR that incorporates a fragment derived from MMG49. The resulting MMG49 CAR T was found to have anti-MM effects without damaging normal blood cells.

"Our results also demonstrate that the active conformer of integrin $\beta 7$ can serve as an immunotherapeutic target against MM, even though the expression of the protein itself is not specific to MM," study coauthor Yukiko Matsunaga says. "Therefore it's highly plausible that there are other cancer immunotherapeutic targets that have yet to be identified in many cell-surface proteins that undergo conformational changes, even if the expression of the proteins themselves is not cancer-specific."

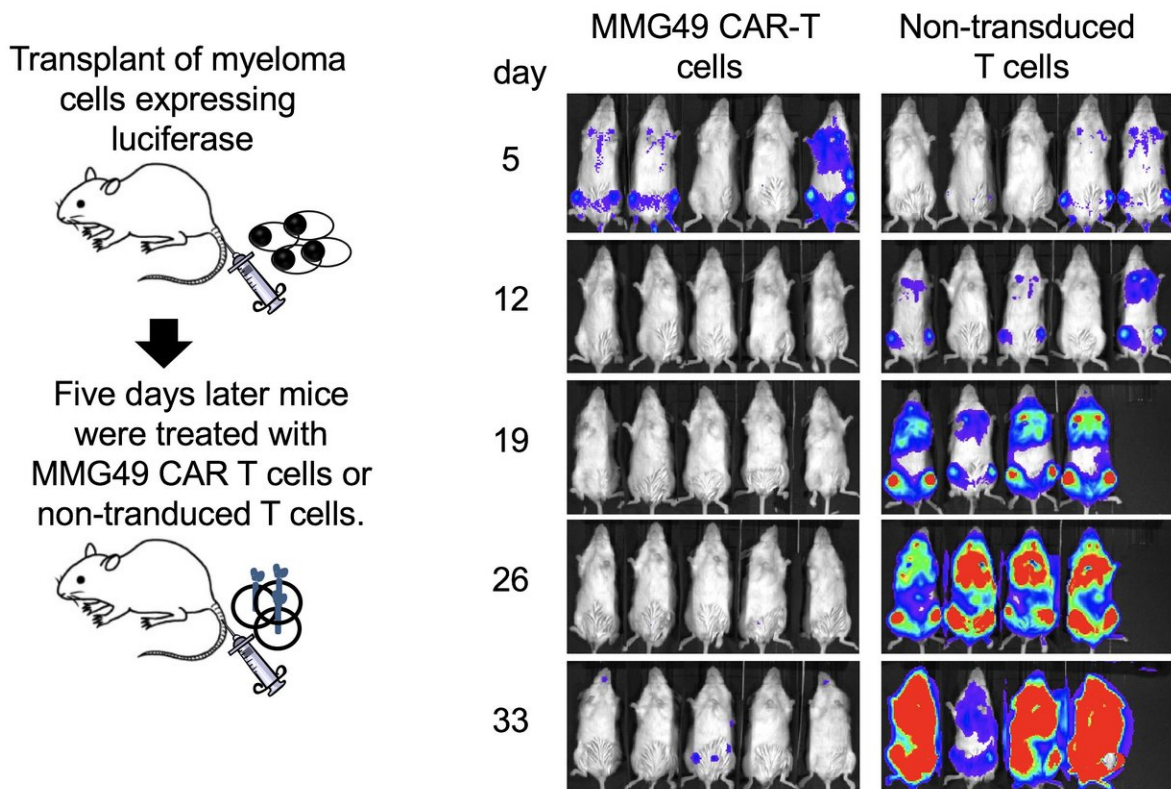


Fig.3. Anti-myeloma activity of MMG49 CAR T cells. Credit: Osaka University

More information: Naoki Hosen et al. The activated conformation of integrin $\beta 7$ is a novel multiple myeloma–specific target for CAR T cell therapy, *Nature Medicine* (2017). [DOI: 10.1038/nm.4431](https://doi.org/10.1038/nm.4431)

Provided by Osaka University

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