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## **Project Website**

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# **SLAMF7 CAR-T CELLS** IN MULTIPLE MYELOMA



This project has received funding from the European Union's Horizon 2020 research and innovation programme under grant agreement No 754658

## Background and aims

Multiple myeloma is a rare blood cancer with around 40.000 new cases diagnosed in Europe each year. Although it is currently incurable, the treatment of multiple myeloma has improved substantially in the last years.

## **CAR-T** cells - a new form of immunotherapy

CAR-T cells, also known as 'Chimeric Antigen Receptor' T cells, have the potential to fundamentally change the treatment of multiple myeloma and could ignite a new era of chemotherapy-free cancer medicine.

CAR-T cells are part of a patient's immune system. The modification of these T cells with a CAR enables them to recognise a protein expressed on the surface of cancers cells and, through this, to find and destroy them.

CAR-T cells have been used for the treatment of other blood cancers, demonstrating impressive clinical responses and improved outcomes in patients. The CARAMBA project is now assessing this innovative approach in multiple myeloma.

# Multiple myeloma-specific

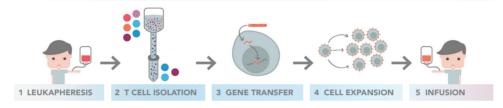
The CARAMBA project is running the first CAR-T cell clinical trial targeting the multiple myeloma-specific protein SLAMF7 which can be found on the surface of myeloma cells. T cells are therefore collected from the patient and equipped with CAR, redirecting the T cells to recognize the SLAMF7 protein and destroy the multiple myeloma cells.

#### Mission

The ultimate goal of the CARAMBA project is to promote a chemotherapy-free and more curative treatment approach for multiple myeloma and to establish SLAMF7 CAR-T cells as a broadly applicable therapeutic concept for multiple myeloma patients in Europe.

## SLAMF7 CAR-T clinical trial – 4 sites 30 patients in Europe

The clinical trial within CARAMBA is designed as a Phase I/II trial and will start in the second half of 2019. Phase I is a dose escalation study and will explore the most effective dose of the CAR-T cell product. For the Phase II part of the trial, 19 patients will be treated with the maximum tolerated dose of SLAMF7 CAR-T cells. The treatment is carried out as follows:



- 1 White blood cells of a patient are extracted by leukapheresis, a procedure in which white blood cells are separated from a sample of blood.
- 2 T cells are isolated from the white blood cells.
- **3** In the next step, the gene sequence for the SLAMF7 CAR is inserted into the DNA of the T cells. The CAR enables the T cells to recognize the SLAMF7 protein on myleoma cells.
- 4 The modified T cells are then expanded outside the patient's body.
- 5 CAR-T cells are infused back into the patients, where they can multiply when they encounter their target protein SLAMF7 and subsequently kill myeloma cells.



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