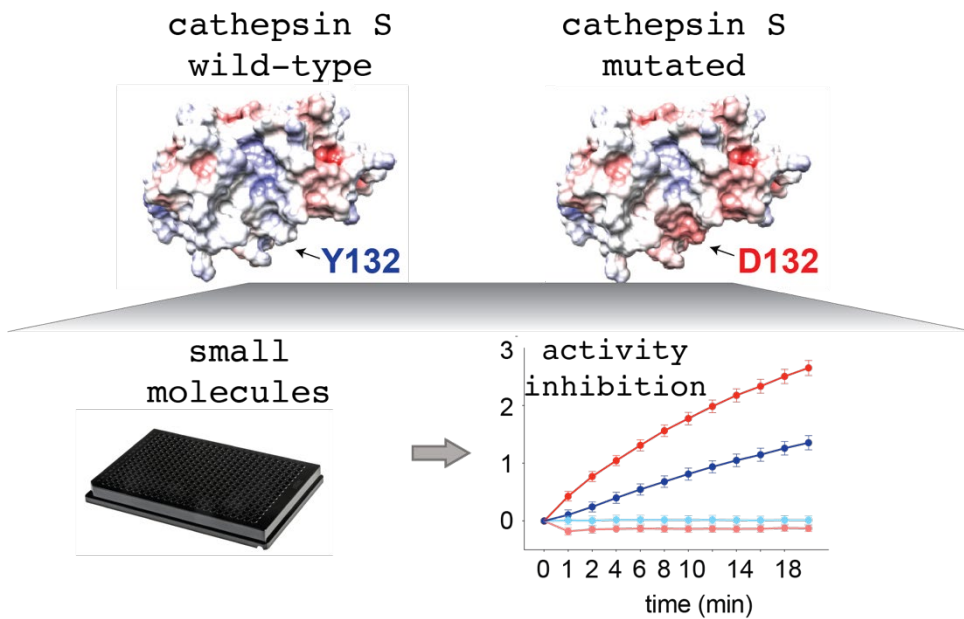


# Cathepsins inhibitors as anti-cancer therapies



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Cathepsin S, haematological, solid tumors, oncogenic mutation, Y132D

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## Description

Over the past years, cancer treatment has improved thanks to the introduction of targeted and immuno-therapies. However, tumor relapse and resistance to current therapies prompt to search of new therapeutic targets that could combine the benefit of both targeted and immuno-therapies. We identified the cysteine protease cathepsin S as an important therapeutic target in hematological malignancies and solid tumors.

In hematological malignancies, cathepsin S is over-activated by the oncogenic mutation Y132D and gene over-expression and loss of its activity induces activation of T-cells and anti-tumor immune responses. In addition, in solid tumors, inhibition of cathepsin S activity can limit the formation of metastasis. Therefore, we have developed a competitive image base assay to rapidly measure the activity of cathepsin S and with this assay we identified small molecules that can specifically inhibit the activity of cathepsin S.

## Advantages

- High level of specificity
- Rapid assessment of efficacy
- Low toxicity

## Applications

- Treatment of hematological cancer in early phase as single agent

