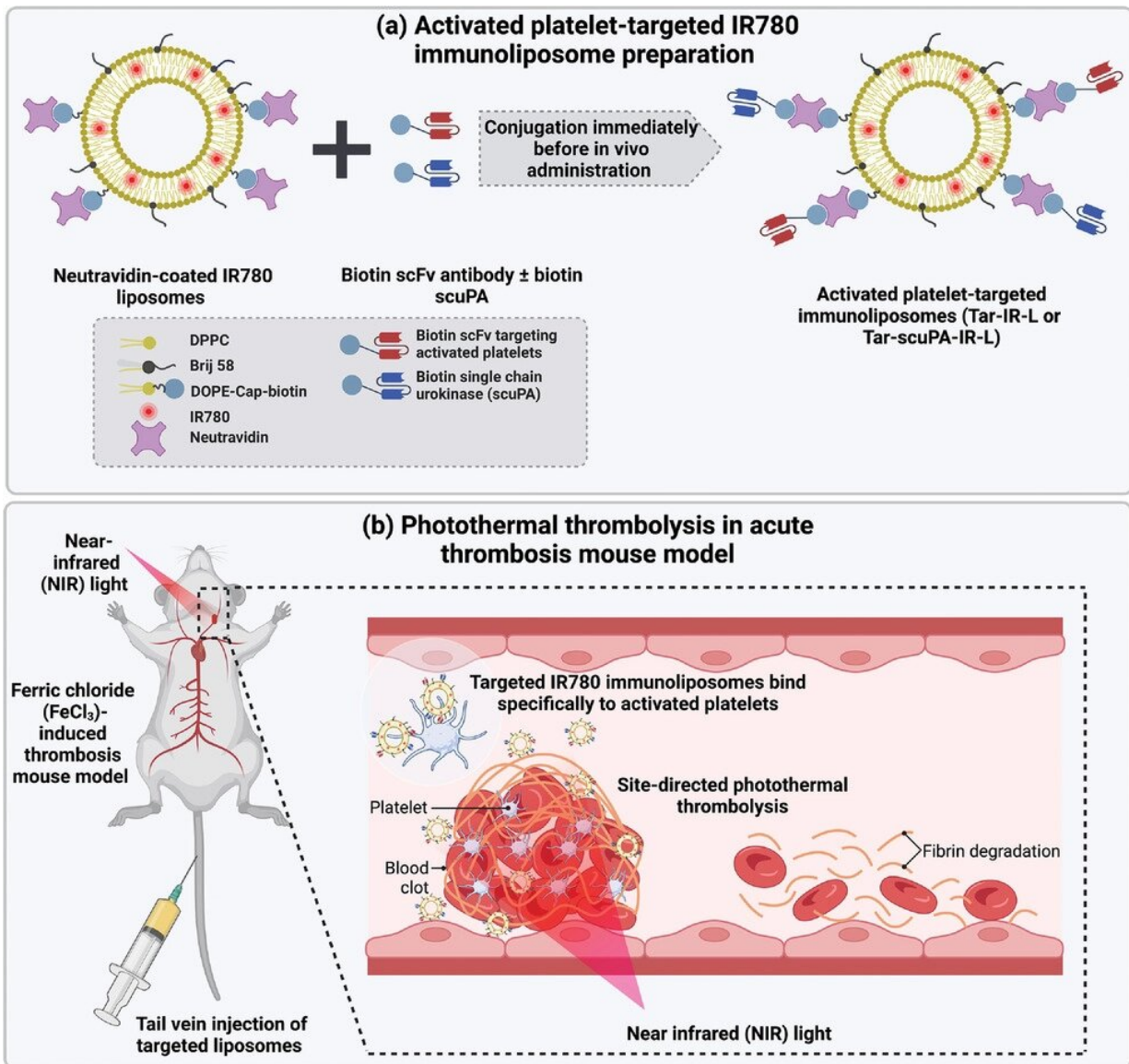


Targeted photothermal treatment for blood clots shows promise

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Schematic diagram of a) activated platelet-targeted IR780 immunoliposomes

(Tar-IR-L & Tar-scuPA-IR-L) preparation and b) their application in an acute thrombosis mouse model for photothermal thrombolysis. Credit: *Advanced Functional Materials* (2022). DOI: 10.1002/adfm.202209019

Nanoparticles loaded with dye that release heat upon near-infrared irradiation could provide the key to more targeted treatment for blood clots, according to new research by the Baker Institute and Swinburne University.

Acute blockage of vessels by [blood clots](#) are the underlying cause of heart attacks and strokes and a leading cause of death and disability globally, with current drug treatments associated with major side effects including potentially fatal bleeding.

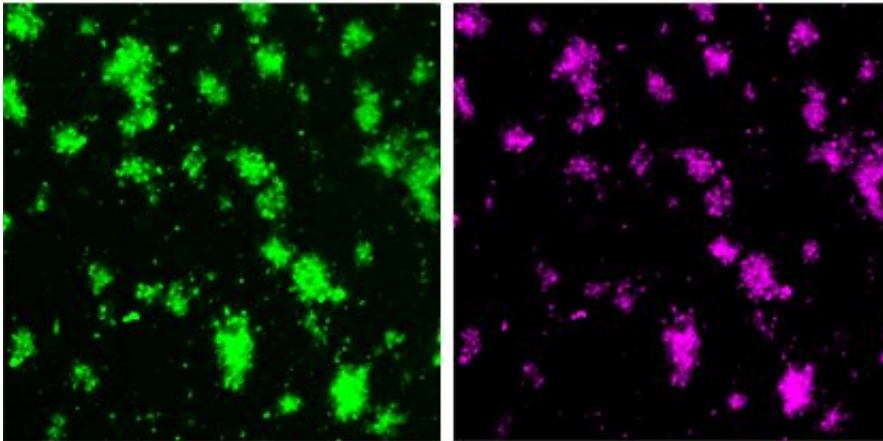
In this pre-clinical study, researchers trialed photothermal therapy, which converts [light energy](#) to [heat energy](#), to more safely and accurately target blood clots. This approach has already attracted attention as a new research frontier in [cancer treatment](#).

The mechanism for [drug delivery](#) was via clot-targeting liposomes, which are being investigated for their ability to deliver drug cargoes specifically to activated platelets, which cluster to form blood clots.

In their paper in *Advanced Functional Materials*, A/Prof Xiaowei Wang and colleagues also incorporated near-infrared dye within the liposome, providing the potential for faster and more potent thrombolysis under infrared-red irradiation.

Merging the bioengineering expertise of Swinburne University researchers together with the biotechnology capabilities of Baker Institute scientists, they were able to create nanoparticles that will home

themselves to blood clots. The targeting mechanism of drug cargoes meant that only a low dose is required, thereby their use will prevent potential bleeding complications.



The green image shows the platelet/clot material. The purple image shows particles that have bound to the clot. Credit: Baker Institute

They found this approach caused a significant reduction in the clot area compared to those areas treated with non-targeted liposomes, which may pave the way for a single dose treatment with fewer side effects.

They hope to build on these results to provide a viable new treatment approach to the current gold standard clot busting treatment involving plasminogen activators, which come with limitations such as poor clot penetration, the need for long infusions and subsequent risk of fatal systemic bleeding.

More information: Ahmed Refaat et al, Activated Platelet-Targeted IR780 Immunoliposomes for Photothermal Thrombolysis, *Advanced*

Functional Materials (2022). [DOI: 10.1002/adfm.202209019](https://doi.org/10.1002/adfm.202209019)

Provided by Baker Institute

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