

Bioorthogonal introduction of nitrite ions into cells for cancer therapy

November 25 2022







A team of researchers writing in the journal *Angewandte Chemie* has developed a bioorthogonal molecular system for the targeted introduction of nitrite ions into cells. Their system releases nitrite ions in cancer cells using a "click-to-release" strategy and these ions, along with other active ingredients, help to initiate cell death. The system could improve the synergistic effects of various cancer therapy drugs.

Cells rapidly convert nitrite ions into nitrogen monoxide (NO), which is involved in many cell processes. For example, it can enhance the effect of various <u>cancer</u> drugs by forming <u>reactive oxygen species</u>. However, the targeted introduction of nitrite to a specific location is complicated.

The research groups of Fude Feng at Nanjing University and Shu Wang at the Chinese Academy of Sciences, Beijing, China, have now developed a bioorthogonal system that selectively transports nitrite ions along with other active ingredients to the endoplasmic reticulum, where they are then released.

Bioorthogonal systems facilitate useful chemical reactions ("click reactions") in cells, without the risk of the reaction partners having adverse effects on the body on their journey to the target site. They have paved the way for an exciting array of novel disease treatment approaches. A testament to this is the fact that the 2022 Nobel prize in chemistry was awarded for the development of click chemistry and bioorthogonal chemistry.

To transport reaction partners to a target site without them participating in unwanted reactions, nitrite ions have to be bound to a carrier molecule as a nitro group. However, the conditions needed to release nitrite again when they reach their target are usually much harsher than those found in living cells. For this reason, the researchers designed two bioorthogonal precursors: one to transport the nitro group and other active ingredients, and another to carry out the click-to-release reaction



by reacting with the first precursor.

The first of the two precursors, ER-Non, performed a number of roles. Firstly, it is readily taken in by the endoplasmic reticulum. Not only do many important cell processes take place in this cell organelle, but it is also the site of action of a number of drugs. Secondly, alongside the nitro group, ER-Non transported the active substance novidamide, which triggers cellular stress responses at high doses and can thus cause cancer cells to initiate <u>cell death</u>.

The other molecular precursor, a dithiol, is activated by enzymes typical for cancer cells. In a click-to-release reaction, the activated molecule releases both the nitrite and the novidamide from ER-Non. The chemicals are not simply released; the reaction causes the new substance to fluoresce and, in so doing, to become a photosensitizer.

Under the action of light, it enhances the ability of the <u>nitrite</u> ion and the novidamide to generate reactive oxygen species and to thereby trigger cellular stress. This phenomenon of photosensitizing is used in photodynamic cancer therapy.

The researchers tested their bioorthogonal system on liver <u>cancer cells</u> and observed arrested growth of these <u>cells</u>. They also observed a notable increase in reactive oxygen species after adding both bioorthogonal components. Since none of the components alone would exert this effect, the team concluded that synergistic effects occur. This opens up new possibilities for more effective cancer therapies.

More information: Jian Sun et al, Dithiol-Activated Bioorthogonal Chemistry for Endoplasmic Reticulum-Targeted Synergistic Chemophototherapy, *Angewandte Chemie International Edition* (2022). DOI: 10.1002/anie.202213765



Provided by Wiley

Citation: Bioorthogonal introduction of nitrite ions into cells for cancer therapy (2022, November 25) retrieved 11 May 2024 from <u>https://phys.org/news/2022-11-bioorthogonal-introduction-nitrite-ions-cells.html</u>

This document is subject to copyright. Apart from any fair dealing for the purpose of private study or research, no part may be reproduced without the written permission. The content is provided for information purposes only.