

# Multi-stimuli controlled release of a transmembrane chloride ion carrier from a sulfonium-linked procarrier

Sribash Das,<sup>a</sup> Oindrila Biswas,<sup>a</sup> Nasim Akhtar,<sup>a</sup> Anjali Patel<sup>b</sup> and Debasis Manna <sup>\*a</sup>

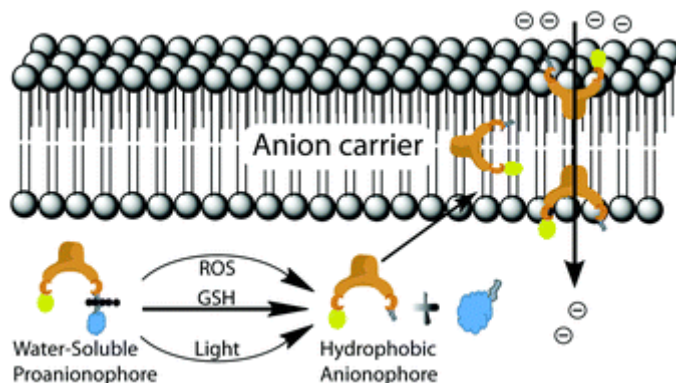
Department of Chemistry,<sup>a</sup> Department of Centre of Environment,<sup>b</sup> Indian Institute of Technology Guwahati, Assam 781039, India

E-mail: [dmanna@iitg.ac.in](mailto:dmanna@iitg.ac.in)

## Abstract

In recent times, anion transporters have received substantial consideration due to their ability to disrupt the ionic equilibrium across membrane bilayers. While numerous  $\text{Cl}^-$  ion transporters were developed for channelopathies, unfortunately, poor aqueous solubility precluded their bioapplicability. Herein, we demonstrate the development of a multi-stimuli activatable anion transport approach to induce regulated transport of  $\text{Cl}^-$  ions across membranes under specific conditions. The sulfonium-based procarrier was initially inactive, but the transmembrane transport of  $\text{Cl}^-$  ions was activated in the presence of stimuli such as glutathione (GSH), reactive oxygen species (ROS) and light. The release of the hydrophobic anionophore from the aqueous-soluble procarrier under specific conditions leads to the successful transport of  $\text{Cl}^-$  ions. Under physiological conditions, these anion carriers follow an antiport exchange mechanism to transport  $\text{Cl}^-$  ions across lipid bilayers. Such multi-stimuli activatable procarriers have great potential to combat various types of channelopathies, including cancer, cystic fibrosis, kidney stones, myotonia, and others.

Keywords:  $\text{Cl}^-$  ion transporters, Multi-stimuli (GSH, ROS and light), Channelopathies.



## Reference

1. Das, S., Biswas, O., Akhtar, N., Patel, A., & Manna, D. (2020). Multi-stimuli controlled release of a transmembrane chloride ion carrier from a sulfonium-linked procarrier. *Org. Biomol. Chem.*, 18(45), 9246-9252.