

# LXD231: A light activated treatment for reducing microbial bioburden in infected wounds

David Chisholm, Ryan Waite, Joy Paterson, Eva Dias, Alba Pujol, Cole Sims, Daniel Callaghan, Gary Sharples, Carrie Ambler

## 1. Introduction

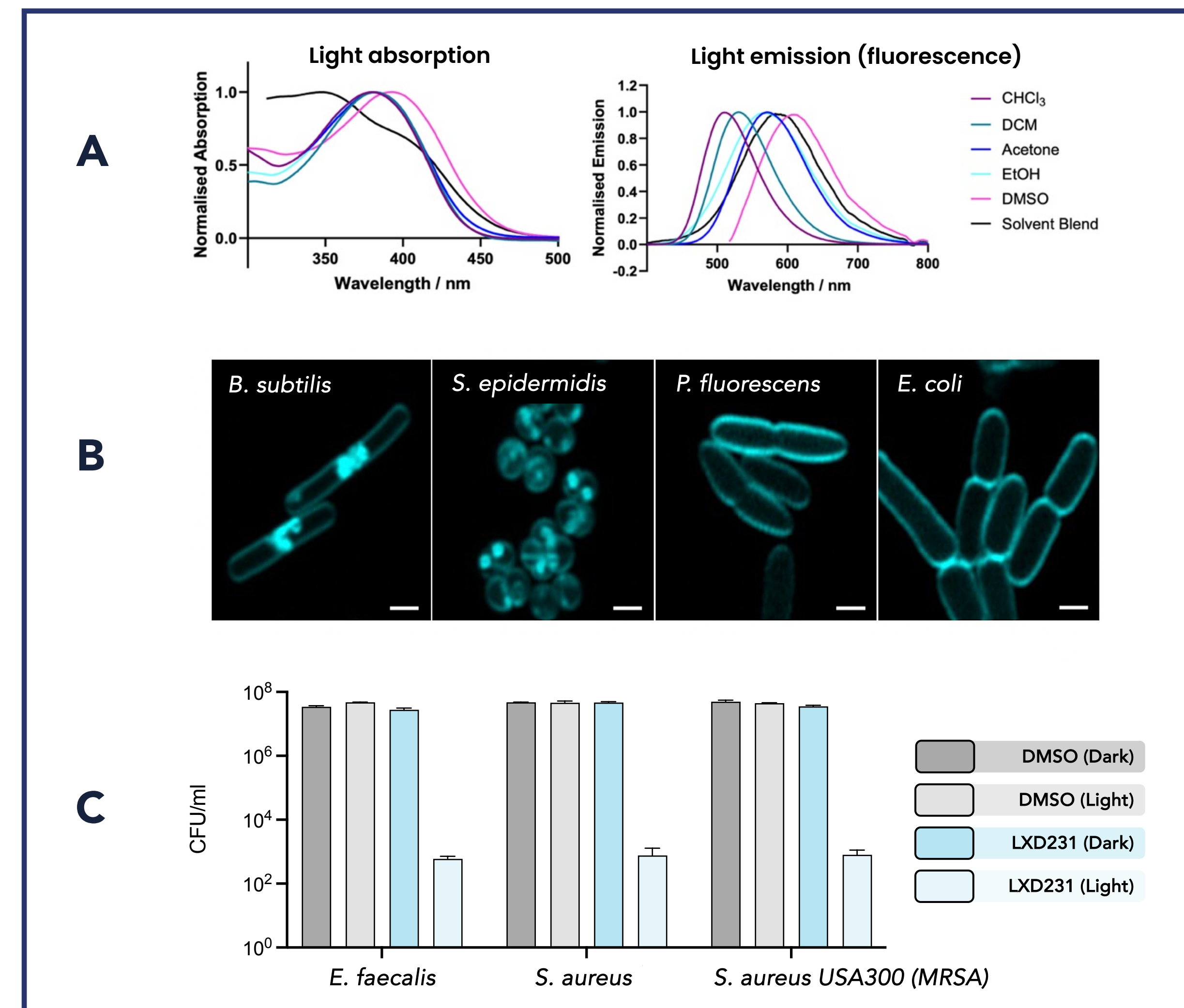
- **Infections that arise within wounds delay healing** and increase the likelihood of complications such as gangrene and amputation. Management of infected wounds is costly and time-consuming.
- Antibiotics, the tools used to fight entrenched infections arising in wounds, are becoming **increasingly ineffective due to antimicrobial resistance (AMR)**. It is crucial that we develop novel antimicrobial modalities that inhibit the establishment of resistant microbe colonies.
- LightOx are developing LXD231, a light-activated gel specifically designed to **eliminate antibiotic-resistant bacteria that arise in infected wounds**.

## 2. LXD231

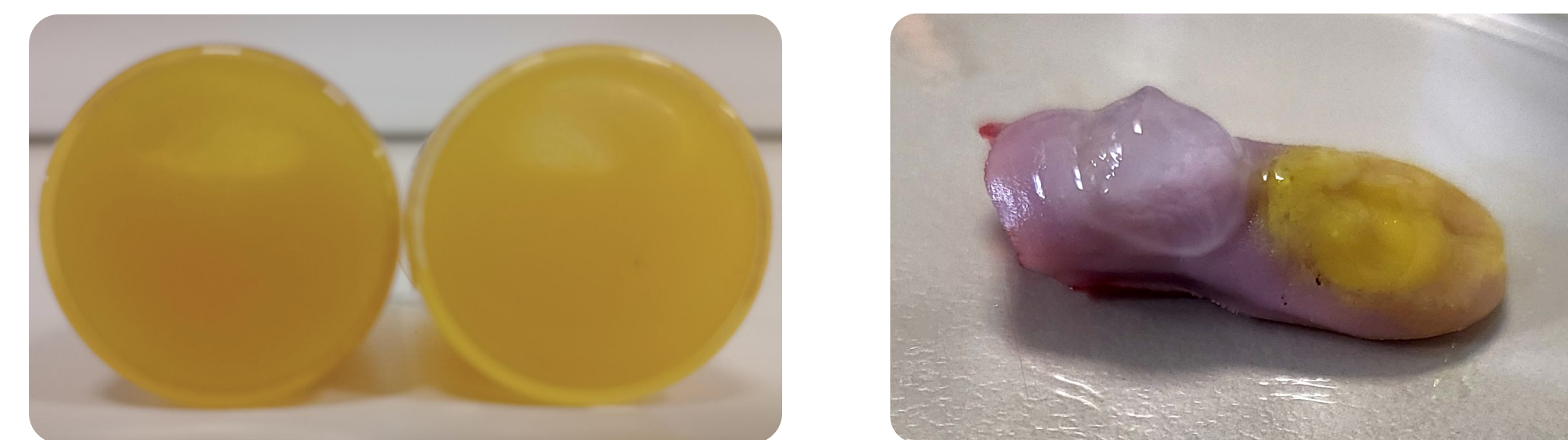
- **LXD231 is a small molecule drug that absorbs light of 380–450 nm** (violet to blue, Figure 1A) from simple LED light devices.
- The inherent fluorescence behaviour (Figure 1B) of LXD231 enables imaging of uptake into a range of bacteria, with incorporation into Gram-positive species and localisation onto the outer membrane surface of Gram-negative species.
- LXD231 generates radical oxygen species (ROS) when activated by light of 380–450 nm. ROS is destructive to bacteria and, furthermore, **bacteria are unable to develop resistance towards it**.
- Illumination of LXD231 via LED light devices activates the drug's antimicrobial activity (Figure 1C), **causing elimination of bacteria including antibiotic-resistant strains such as MRSA**.

## 3. Next steps

- We are in preclinical development, currently completing gel formulation of LXD231 (Figure 2) and exemplifying activity in *ex vivo* and *in vivo* models of an infected wound.
- LightOx are looking for feedback from practitioners in the wound care sector. We are targeting clinical trials in 2026–2028.



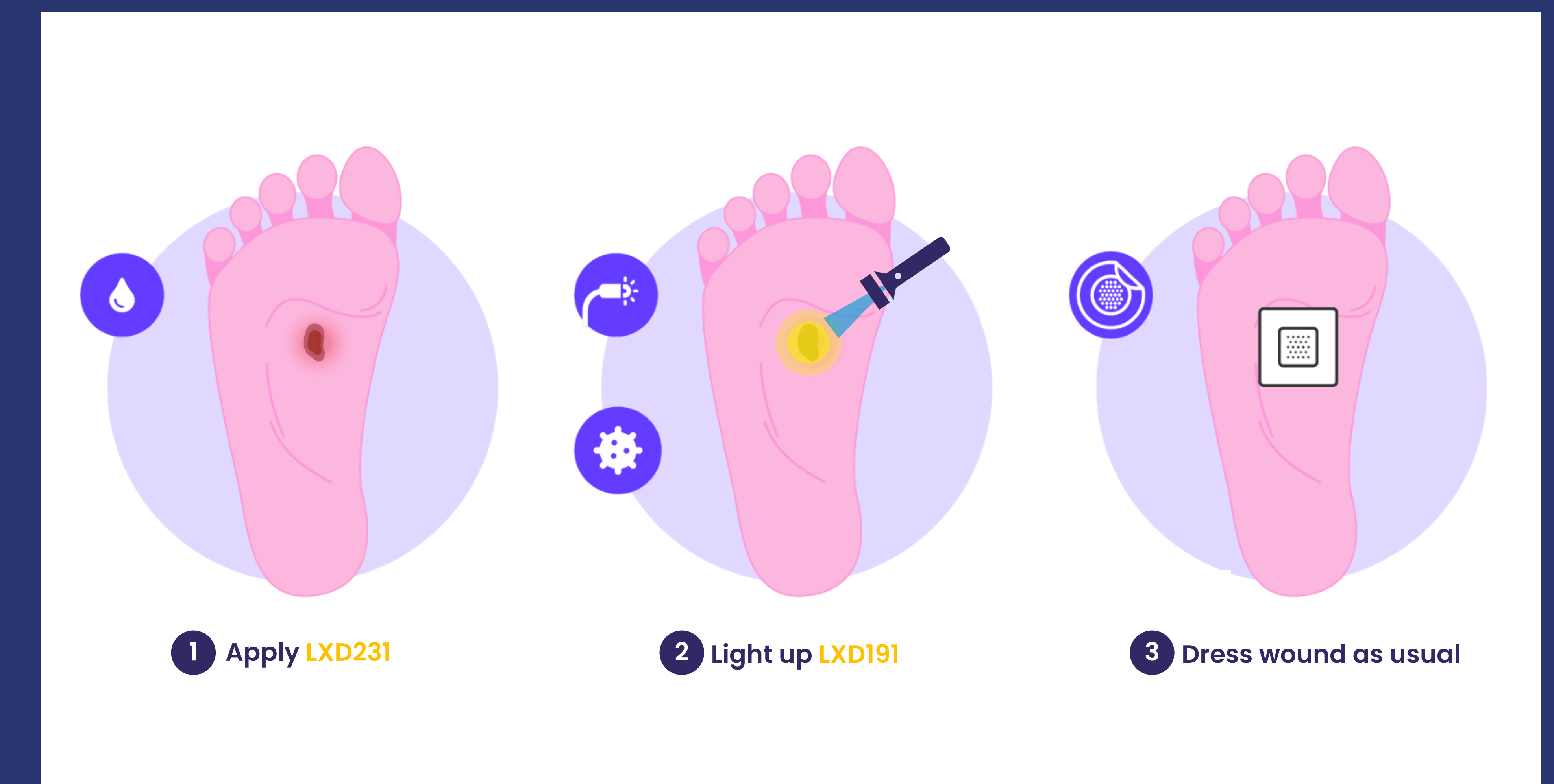
**Figure 1**  
(A) LXD231 photophysical behaviour.  
(B) Imaging localisation of LXD231 in a range of bacteria using confocal microscopy and LXD231's inherent fluorescence characteristics.  
(C) Light-activated antimicrobial activity of LXD231 (2  $\mu$ M) against a range of bacteria including antibiotic resistant MRSA.



**Figure 2**  
Development of gel formulations of LXD231 (left), enabling application to the wound surface (right).

# LIGHTOX<sup>®</sup>

## Eliminate antibiotic-resistant bacteria in infected wounds using a light activated gel treatment



Scan the QR code to learn more

[lightox.co.uk](http://lightox.co.uk)



**David Chisholm, PhD**  
Head of Technology

[david.chisholm@lightox.co.uk](mailto:david.chisholm@lightox.co.uk)