

Lxbio Pharmaceuticals is an innovative biotechnology company and a global leader in bacteriophage technology and antibody discovery for oncology and ophthalmology.

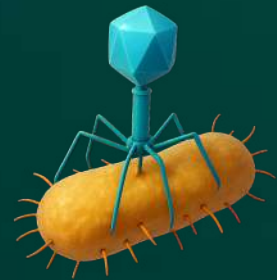
Leveraging its own CRO specialized in antibody and CAR-T cell discovery and production, Lxbio provides both external services and drives its internal R&D programs. Its therapeutic focus spans ophthalmology, oncology, and dermatology, supported by a robust and innovative pipeline.

Key programs include TP-102, the first bacteriophage-based therapy for diabetic foot infection (Phase 2b clinical trials), the first antibody-loaded nanoparticles in eyedrops and the first bacteriophage-based eyedrops for blepharitis in ophthalmology, and bacteriophage-corticosteroid combination for atopic dermatitis in dermatology.

Furthermore, Lxbio is advancing a proprietary phage delivery platform for oncology antibodies, reinforcing its leadership in next-generation biologics and precision medicine.



# Technology Platforms



## BacterioPhage for AMR Therapeutics

Bacteriophage therapeutics provide a targeted approach **against antimicrobial resistance**, precisely eliminating pathogens.

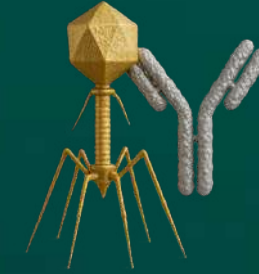
**Promising alternative** for infectiology, including diabetic foot, pulmonary, ocular, and other hard-to-treat infections.



## Anti-VEGF in Nanoparticles

Eydrops Anti-VEGF encapsulated in nanoparticles enables targeted, sustained drug delivery for **treating retinal diseases**.

Nanoparticles **enable precise ocular drug delivery**, enhancing therapeutic efficacy while minimizing side effects in eye disease treatment.



## Oncology Phage Antibody Delivery

Phage-Antibody Conjugate represents a **next-generation targeted oncology platform** with applications in both human and veterinary oncology.

Engineered M13 bacteriophages as a biocompatible and **customizable drug delivery vehicle**.



## Biologics R&D and Production Unit

Antibody and CAR-T cell R&D services span discovery to preclinical optimization, accelerating immunotherapy development.

Small-scale clinical-grade production supports translational research and early-phase trials.

# Innovative Pipeline

We are advancing first-in-class biologics, integrating bacteriophage therapies for infectious and inflammatory diseases, nanoparticle-encapsulated antibodies for improved ocular bioavailability, and novel phage delivery platforms to revolutionize targeted oncology antibody and other therapies.

Product	Indication	Product	Proof of concept	Preclinical development	Clinical trials Phase 1	Clinical trials Phase 2	Clinical trials Phase 3
Topical Bacteriophages cocktail Diabetes	Diabetic foot infection	<b>TP102</b>	COMPLETED				IN DEVELOPMENT
Anti-VEGF encapsulated in nanoparticles eyedrops	Macular degeneration & Diabetic retinopathy	<b>LXOP02</b>	COMPLETED				IN DEVELOPMENT
Eyedrops gel Bacteriophages	Blepharitis & ocular infection	<b>LXOP01</b>	COMPLETED	IN DEVELOPMENT			
Phage-based Delivery Systems	Several therapeutical targets	<b>LXON01</b>	COMPLETED				IN DEVELOPMENT



# TP-102

## The Potential to Transform the Care Of Diabetic Foot Infections

Precision against pathogens. Healing beyond antibiotics.



### **TP-102 is a Breakthrough Bacteriophage Therapy for Diabetic Foot Ulcers**

TP-102 is the first topical bacteriophage cocktail specifically developed to treat diabetic foot ulcers and chronic wound infections. Composed of five lytic bacteriophages, TP-102 offers a targeted, microbiome-sparing approach to eradicate multidrug-resistant pathogens, a growing concern in the management of chronic wounds.

Developed by a Portuguese biopharmaceutical company, TP-102 has demonstrated excellent safety and efficacy in clinical trials, promoting accelerated healing and bacterial clearance with no systemic adverse effects.

TP-102 has received FDA fast track and is expected to complete phase 2b by the end of November 2025.

This innovative platform has the potential to redefine the standard of care for diabetic foot infections, offering a next-generation solution where traditional antibiotics often fail.

R&D Stage

**Finishing Phase 2b**

**Human Clinical Trials**

# TP-102 The Product



TP-102 is being developed for topical treatment of patients with wound infections including chronic ulcers.

## Targeted Antimicrobial Solution Against AMR Pathogens

Designed to combat multidrug-resistant bacteria. TP-102 phage cocktail offers precision killing where antibiotics fail, addressing a critical unmet need in the era of antimicrobial resistance.

## Topical, Non-Systemic Delivery with Rapid Local Action

Formulated for direct application to infected wounds, ensuring localized bacterial clearance, reduced systemic exposure, and a faster therapeutic response in the infected tissue microenvironment.

## Breakthrough Safety Profile

Comprised of naturally occurring, lytic-only bacteriophages with no transduction or lysogeny risk, and with no observed systemic toxicity or adverse events demonstrated in the completed REVERSE trial (TP-102).

## Demonstrated Efficacy in REVERSE Clinical Trial

Statistically significant reduction in bacterial load and wound size progression in DFI patients with resistant infections.

## First-in-Class Phage-Based Innovation for DFI

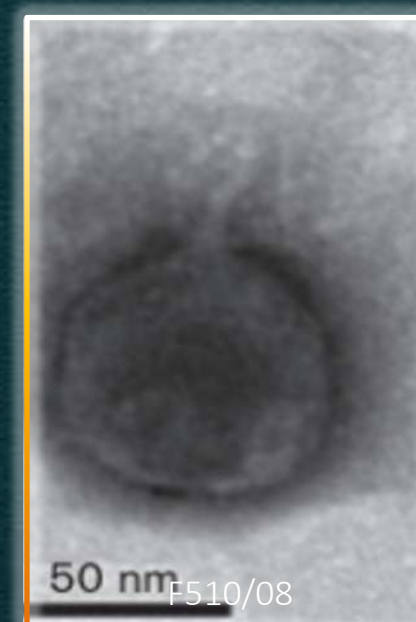
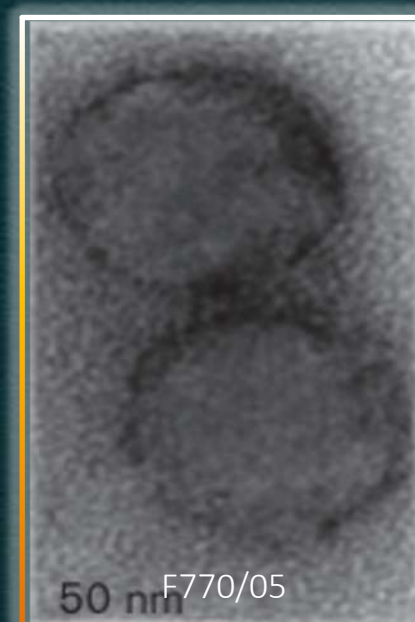
TP-102 is the first validated topical phage cocktail to complete a clinical trial for diabetic foot infections, creating strong IP positioning, early-mover advantage, and opportunities for regulatory fast-track and hospital formulary inclusion.

TP-102 is an innovative bacteriophage cocktail comprised of 5 lytic bacteriophages against:

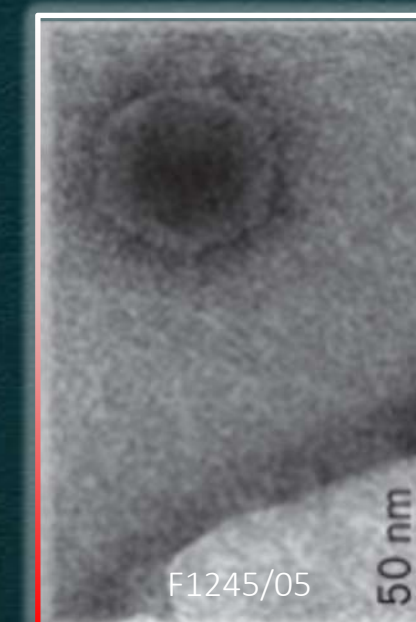
### *Staphylococcus aureus*



### *Pseudomonas aeruginosa*



### *Acinetobacter baumannii*





TP-102 is an innovative bacteriophage cocktail comprised of 5 lytic bacteriophages against:  
*Staphylococcus aureus* | *Pseudomonas aeruginosa* | *Acinetobacter baumannii*

Against AMR  
Pathogens

Topical with  
Rapid Local Action

First-in-Class  
Phage-Based



Fast Track Designation  
Granted to TP-102

PHASE I/IIa REVERSE study  
**COMPLETED**

NO TREATMENT RELATED ADVERSE EVENTS ASSOCIATED TO TP-102 REPORTED DURING THE TRIAL.  
TP-102 CAN BE EFFECTIVE IN REDUCTION OF DIABETIC FOOT ULCERS VOLUME.

PHASE IIb REVERSE 2 study  
**COMPLETED**

Clinical study CLOSED with 76 patient in 2 countries: USA and India

PHASE III  
REVERSE 3 study

estimated to start in  
**Q1 2026**

## TP-102 Clinical Case Compassionate Use

This report describes an application of phage therapy (TP-102) for diabetic foot ulcers in a compassionate use and offers data of efficacy and tolerability



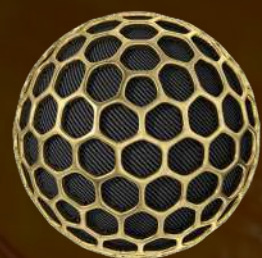
**START**  
Phage therapy



**2 Months**  
post phage therapy



**6 Months**  
post phage therapy  
**recovered**



# LXOP02

Anti-VEGF Lipid-based Nanoparticles in an Eyedrops Formulation

## From Needle to Drops

A Safer Future for Retinal Care



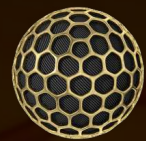
We are developing the first-ever **anti-VEGF eye drop formulation** designed specifically for the treatment of diabetic retinopathy. This innovative solution encapsulates the anti-VEGF agent within lipid-based nanoparticles, enabling effective topical delivery to the retina, a significant advancement over current invasive intravitreal injection therapies. By combining nanotechnology with a non-invasive administration route, this product aims to improve patient compliance, reduce treatment-associated risks, and expand access to care for millions affected by diabetic retinal diseases.

R&D Stage

**Finishing *In vitro***

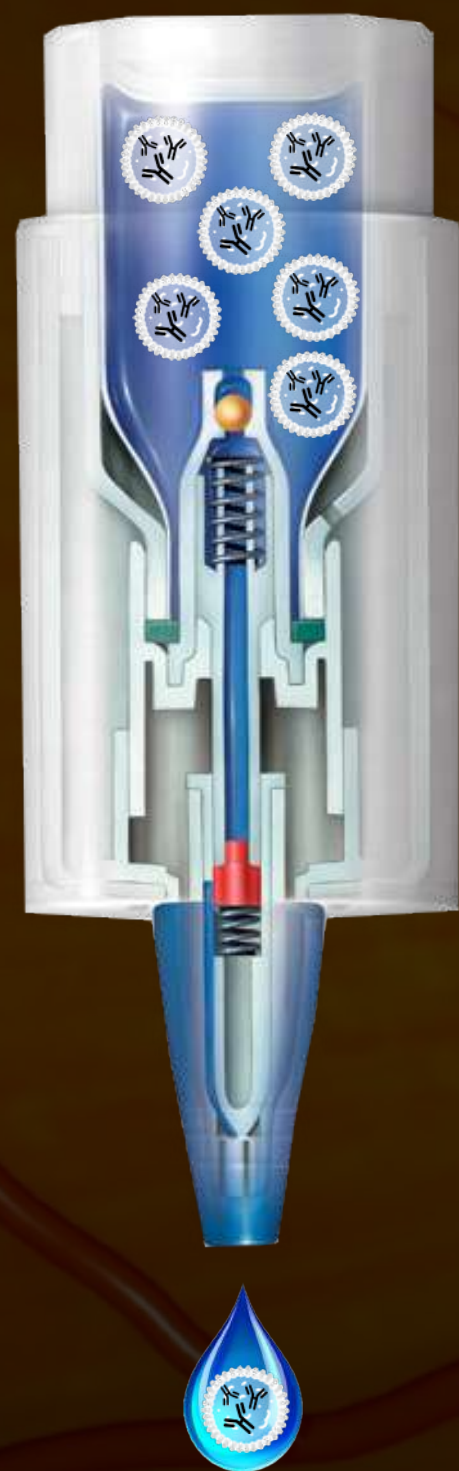
**Preclinical Trials**





# LXOP02 The Product

Anti-VEGF Lipid-based Nanoparticles in an Eyedrops Formulation



## Nano-encapsulation of Anti-VEGF Fragment

The formulation incorporates recombinant anti-VEGF antibody fragment within lipid-based nanoparticles (LNPs), enhancing molecular stability and protecting the biologic from enzymatic degradation upon topical administration

## Lipid-Based Nanoparticle Carrier System

The LNPs are composed of biocompatible lipids optimized for ocular permeability and mucoadhesion, promoting efficient transcorneal and transscleral penetration toward posterior segment targets

## Topical Ophthalmic Delivery Platform

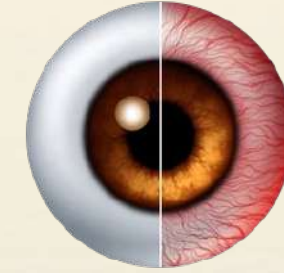
Designed as a sterile, aqueous eye drop, the formulation enables non-invasive administration of anti-VEGF therapy to the retina, aiming to reduce the need for frequent intravitreal injections in conditions such as nwAMD, DME, and RVO

## Sustained Release and Bioavailability Enhancement

The nanoparticle matrix allows for controlled release kinetics and prolonged ocular residence time, improving bioavailability and therapeutic exposure within intraocular tissues over extended period

## Preservative-Free Multidose Container System

The solution is filled in a preservative-free, multidose delivery container that utilizes a sterile barrier mechanism, minimizing ocular surface toxicity and enabling safe, long-term self-administration by patients



# LXOP01

The world's first bacteriophage-based eye drop.  
Target infection. Preserve vision. Restore comfort

This innovative formulation harnesses the natural antibacterial properties of lytic bacteriophages to selectively target and eliminate drug-resistant Staphylococcus and Pseudomonas species, a cause of Blepharitis and chronic eyelid inflammation.

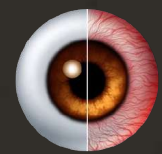
Unlike traditional antibiotics, phages do not disrupt the healthy ocular microbiome and do not contribute to antibiotic resistance. Developed using advanced stabilization and delivery technologies, the product ensures effective penetration of biofilms and sustained antimicrobial activity.



R&D Stage

**Finishing *prof of concept***

**Start Preclinical Trials**



# LXOP01 The Product

The world's first bacteriophage-based eye drop.

## Product Overview

*LX24OP01 is the first therapeutic product specifically developed for blepharitis using a bacteriophage cocktail. Formulated as a gel-based eye drop, ensuring enhanced ocular surface adherence and prolonged bioavailability.*

### Mechanism of Action

Comprises a proprietary cocktail of 4 lytic bacteriophages

Highly specific and potent lytic activity against *Staphylococcus aureus* and *Pseudomonas aeruginosa*

Selective bactericidal action minimizes disruption to the ocular microbiota

### Target Pathogens

***Staphylococcus aureus:***  
Predominant Gram-positive pathogen in blepharitis, often associated with inflammation and eyelid margin colonization

***Pseudomonas aeruginosa:***  
Opportunistic Gram-negative bacterium contributing to chronic cases and antibiotic-resistant infections

### Innovation and Advantages

***First-in-class phage gel*** eye drop developed for direct ocular application in blepharitis.

Phages replicate at the site of infection, amplifying their therapeutic effect

Non-antibiotic strategy reduces the risk of AMR development

Offers an alternative for patients with antibiotic-refractory blepharitis or those prone to recurrent infections

### Antibiotic Resistance Context

Rising resistance in *S. aureus* (e.g., MRSA) and *P. aeruginosa* limits the efficacy of conventional topical antibiotics

LX24OP01 circumvents this issue by using phages that target resistant bacterial strains without inducing cross-resistance

### Formulation Benefits

Gel-based vehicle ensures sustained release and prolonged contact time on the eyelid margin

Enhanced ocular retention vs traditional liquid drops, leading to improved patient compliance and efficacy

### Clinical Relevance

Designed for topical, non-invasive, and localized therapy of anterior and posterior blepharitis

Potential to become a paradigm shift in ocular infection management, especially in the era of antimicrobial resistance

# Corneal Infection Therapy with Topical Bacteriophage

**Studies show evidence of effectiveness and safety of phages in *S. Aureus*-induced ocular infections.**

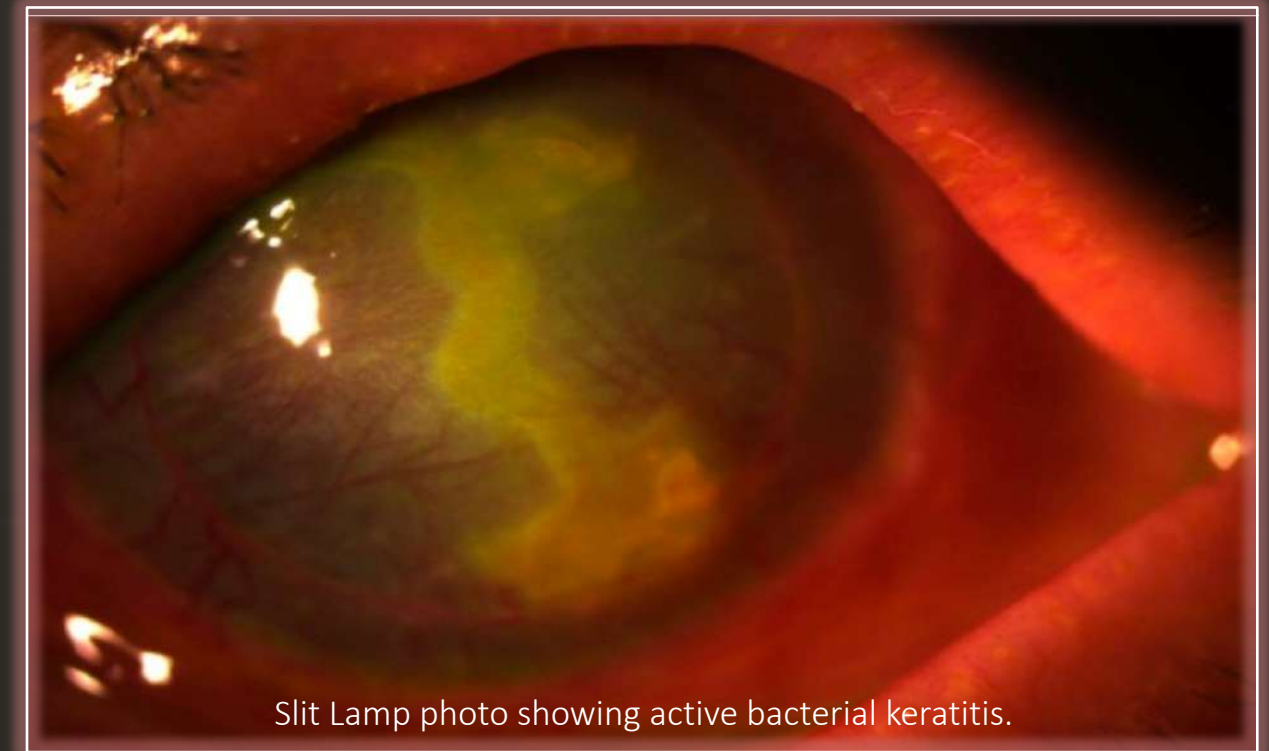
**Staphylococcus aureus is a primary pathogen in bacterial keratitis, a condition that can cause permanent visual impairment**

Report case of a 65-year-old woman  
Penetrating keratoplasty in the left eye for the post infectious corneal  
Started on broad-spectrum topical antibiotics has failed  
Microbial cultures were positive for VRSA.  
Persistent vancomycin-intermediate *S. aureus* (VISA) infection for 11 years

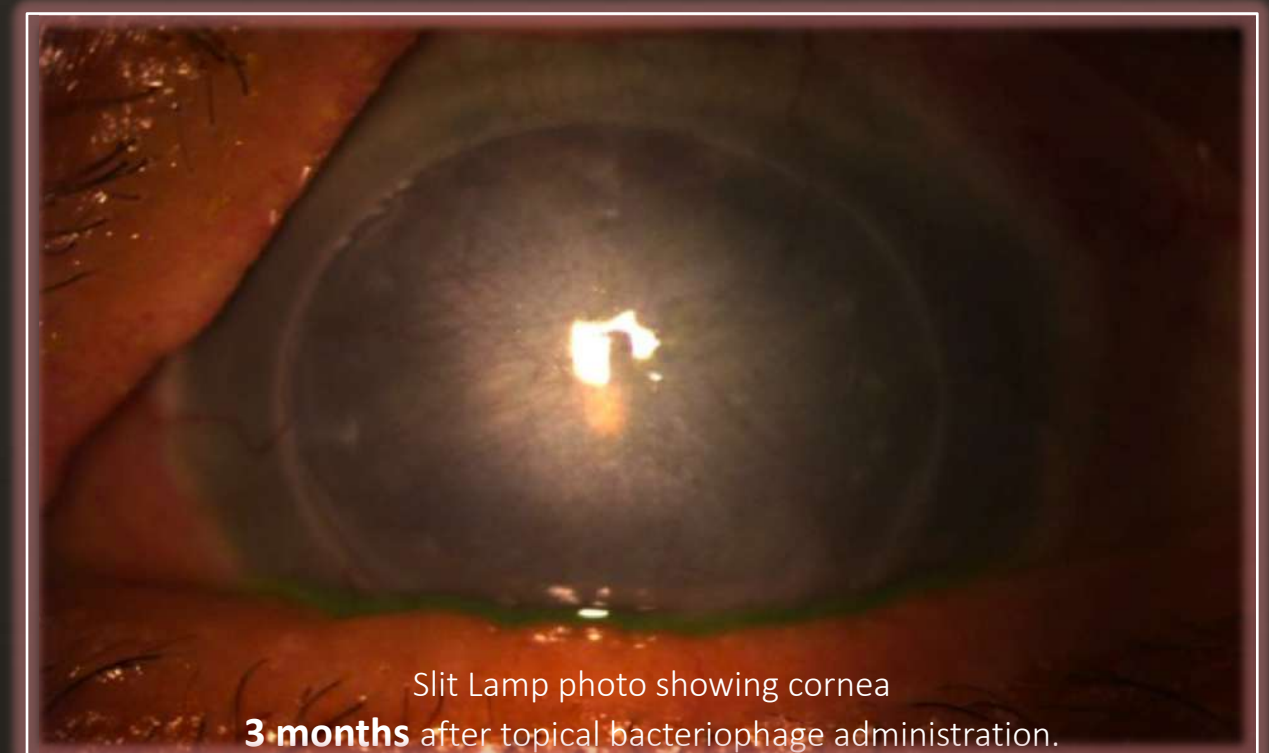
**Patient then decided to undergo Phage therapy  
Using the *S. aureus* bacteriophage (SATA-8505) for 4 weeks.**

After 3 months, normal ocular and nasal cultures were verified,  
indicating eradication of the infection.

**This case suggests the therapeutic potential of phages  
in ocular disease, particularly in antibiotic-resistant  
cases.**



Slit Lamp photo showing active bacterial keratitis.



Slit Lamp photo showing cornea  
**3 months** after topical bacteriophage administration.



The LXON01, is a novel design Phage-powered modular therapeutic platform that represents a next-generation targeted oncology platform with applications in both human and veterinary oncology.

By combining tumour-specific antibody targeting with phage-based delivery, it achieves high tumour selectivity, enhanced intracellular drug accumulation, and superior therapeutic efficacy, while significantly reducing systemic toxicity.

The modular nature of this phage platform allows expansion to multiple tumour-associated antigens and diverse cytotoxic or immunomodulatory payloads, enabling a versatile pipeline for solid and hematologic malignancies.

R&D Stage

**Finishing *proof of concept***

**Start Preclinical Trials**



**LXbio**  
Pharmaceuticals



# LXON01

## Innovative Platform for Oncology Drug Delivery

### Phage-Based Delivery System

Utilizes engineered M13 bacteriophages as a biocompatible and customizable drug delivery vehicle. High payload capacity for cytotoxic agents or immune-modulatory molecules.

### Antibody-Phage Conjugation

Antibody conjugated to M13 phage surface proteins for targeted delivery. Stable and site-specific linkage ensures preservation of antibody affinity and phage integrity.

### Tumor Target Selectivity

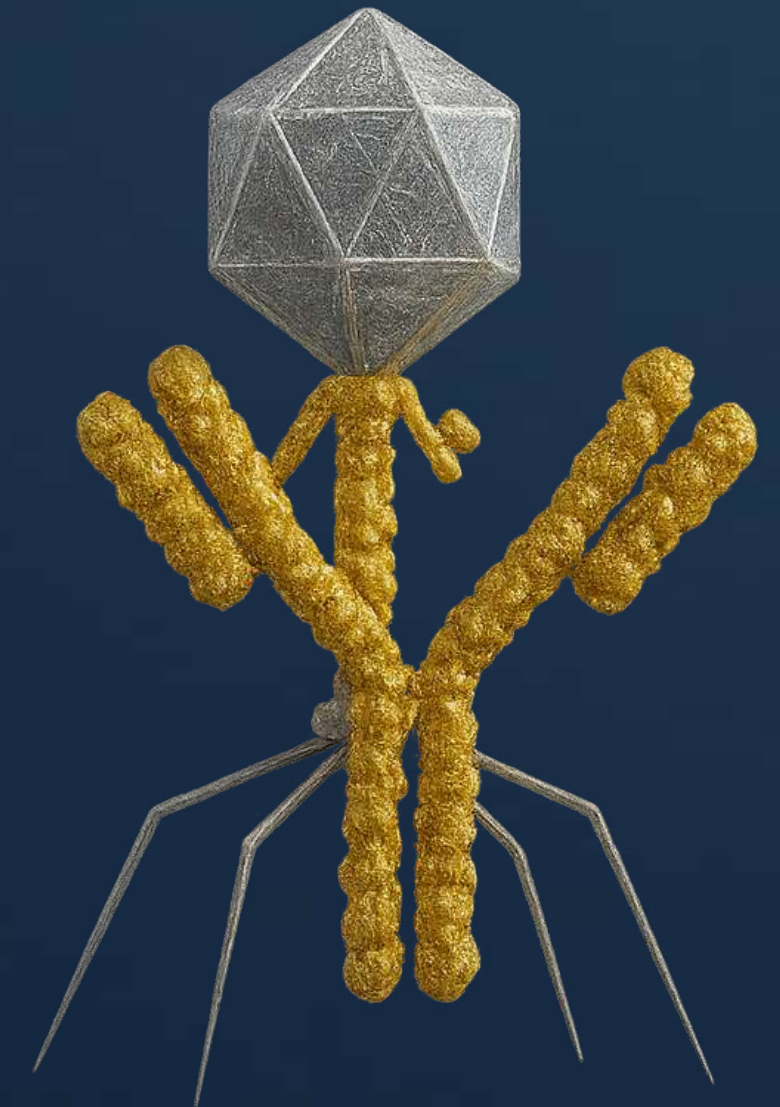
Antibody ensures highly selective binding to expressing tumour cells. M13 phage surface modifications further enhance tumour microenvironment tropism.

### Target Efficacy

Combines ADC cytotoxic potency with phage-mediated multivalency, increasing internalization and drug accumulation in tumour cells. Potential for immune activation via phage components, synergizing with tumour cell killing.

### Reduced Systemic Toxicity

Targeted delivery minimizes off-target effects and improves therapeutic index compared to conventional ADCs. Phage platform exhibits low immunogenicity and favourable safety profile in preclinical models.





# LXON01

## Innovative Platform for Oncology Drug Delivery

### Preliminary results in vivo of LX25300M01 Phage-ADC

#### Model & Study Design

Tumour induced in mouse models (HER2-positive).

Small batch of M13 Phage-Anti-HER2 ADC administered systemically.

#### Efficacy

Significant tumor **growth inhibition** observed.

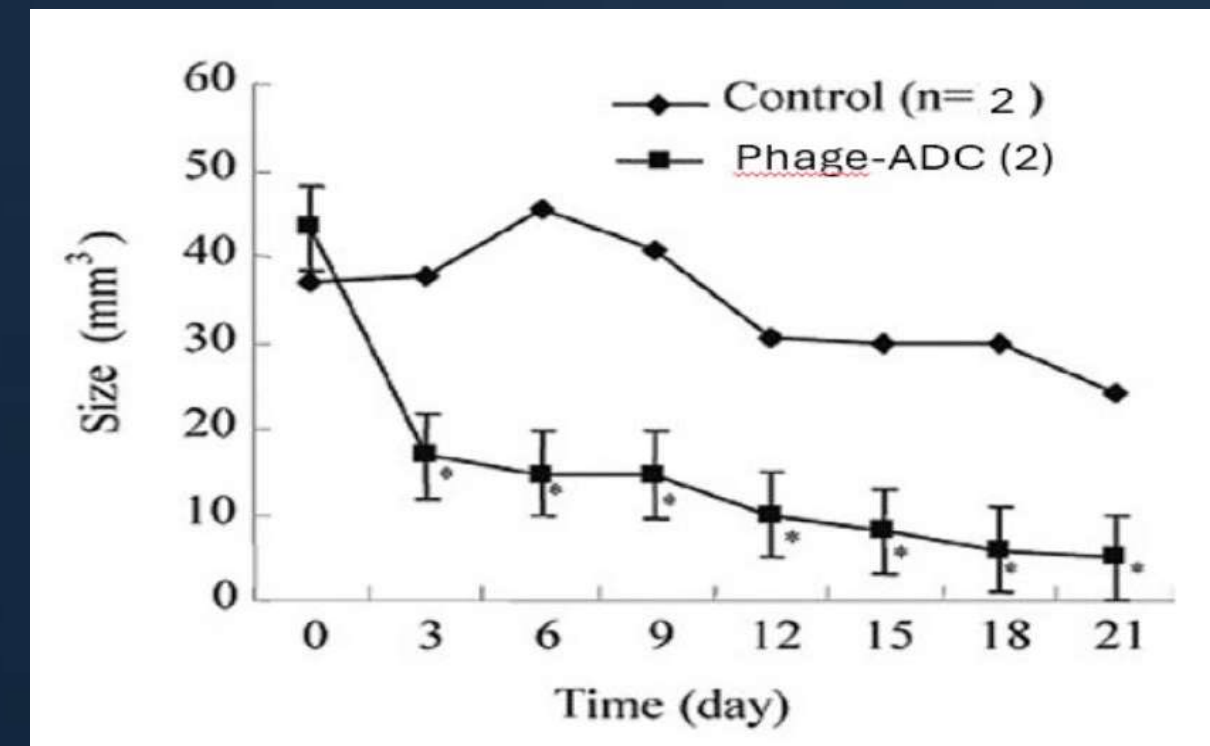
**Tumor volume reduction within 20 days** compared to control group.

Suggests **strong targeted anti-tumour activity** and proof of concept for therapeutic potential.

#### Safety

No major adverse effects or **systemic toxicity** observed in treated animals.

Stable weight and normal behavioural parameters indicate good tolerability.





## R&D UNIT

Antibody discovery and production,  
and CAR-T cell Technologies.

**YOUR IDEA,  
YOUR PROJECT,  
OUR MISSION**

### 1. End-to-End Antibody Engineering Platform

Comprehensive services covering the full antibody development cycle from target identification and antibody discovery (phage display and single B-cell) to humanization, affinity maturation, and Fc engineering, all under one roof.

### 2. Next-Generation Cell Line Technologies

Use of *state-of-the-art* expression-based platforms for high-yield, stable production of monoclonal, bispecific, and fragment antibodies optimized for R&D and preclinical.

### 3. Integrated CAR-T Discovery & Functional Screening

Advanced CAR design and validation workflows including screening, signalling domain optimization, and *in vitro* cytotoxicity assays, enabling rapid generation of functionally validated CAR-T constructs tailored for specific indications.

### 4. Flexible R&D-Grade Production Capabilities

Agile production for small antibody and CAR-T batches, enabling rapid prototyping and customized reagent generation to support biotech, academic, and translational R&D.

### 5. Scientific Expertise + Partnership Mindset

A team of multidisciplinary scientists with deep immunology, oncology, and cell therapy expertise, offering collaborative project design, data-driven decision support, and seamless integration with client pipelines from startup biotechs to large pharma.

## ✓ Efficient and Scalable Solutions

Our advanced platforms, including B cell technology, foster faster results without compromising quality.

## ✓ Proven Track Record

With a history of successful projects and strong client relationships, we bring credibility and trust to your ecosystem.

## ✓ Cost Effectiveness

Centralizing services can simplify administrative processes, such as legal agreements, and material transfers.

## ✓ Tailored Excellence Across Projects

We provide tailor-made solutions that align with the unique goals of each company.

## ✓ Synergistic Ecosystem Building

We can develop a deeper understanding of your research focus and provide tailored solutions for multiple companies.



### ANTIBODY DISCOVERY IN 6 WEEKS

Single-cell technology for precise identification and isolation of high-affinity antibodies directly from individual B-cells.

Our cutting-edge system enables ranking of thousands of antibody hits based on relative affinity, bypassing the need for traditional re-expression and characterization workstreams.

Production Stable cell lines

Discovery and Engineering Characterization

- Decrease timelines
- Unlock unprecedented specificity, speed, and therapeutic potential
- Simultaneous multiple target screening
- Native selections

Our workflow integrates single-cell isolation, high-throughput screening, and downstream functional assays. Patient-derived B cells are first enriched and then sorted based on their antigen-specific binding. Each single B cell's immunoglobulin repertoire is then profiled, allowing for efficient cloning and expression of candidate antibodies. This platform maximizes the probability of uncovering unique epitopes and neutralizing activities by capturing the natural diversity of the patient's antibody response. The process can be replicated for immunized animals, including mice, rabbits and camels, which yields the need for Humanization.

#### Engineering

Targeted modifications to the lead therapeutic molecule generate variants that are rapidly rated by high-throughput systems. The TPP-focused projects include, but are not limited to, **Antibody Humanization and Affinity Maturation.**

Mouse (0%Human) Chimeric (65%Human) Humanized (>80%Human) Fully Human (100%Human)

high potential for immunogenicity low

#### Stable CHO Cell Line Development in 4 WEEKS

Our single-cell platform delivers a faster, more precise, and scalable approach to antibody production, ideal for high-impact therapeutic applications. Our Single-cell precision identifies and ranks top antibody-producing cells, ensuring only the highest-yield and most effective antibodies advance.

LXbio

# Conclusion

1

## Leadership in Bacteriophage Therapeutics for AMR

Lxbio is advancing bacteriophage-based treatments for diabetic foot infection, blepharitis, offering targeted, microbiome-sparing solutions to combat antimicrobial resistance (AMR).

These programs address high-burden indications with limited effective therapies and are positioned for first-in-class differentiation.

2

## First-Ever Phage-Based Ophthalmic Eye Drop

Lxbio is developing the world's first bacteriophage eye drop, a disruptive, non-invasive formulation for ocular infections.

This innovation opens a novel therapeutic modality in ophthalmology, addressing unmet needs in microbial keratitis and chronic blepharitis with precision-targeted, preservative-free therapy.

3

## Breakthrough Anti-VEGF Nanoparticle Eye Drop

The company is the first to encapsulate anti-VEGF fragments in nanoparticles for topical ocular delivery.

This eye drop formulation is designed for diabetic retinopathy and AMD, replacing invasive intravitreal injections and improving compliance, access, and safety.

4

## Next-Generation Oncology Platform: Phage-Ab Conjugates

Lxbio's Phage-Antibody Conjugate technology is a cutting-edge platform for tumor-targeted biologics delivery.

With broad applicability in both human and veterinary oncology, it combines the specificity of phages with the potency of antibodies, enabling modular, scalable oncology solutions.

5

## Integrated CRO-Biotech Model with Biologics Expertise

Lxbio operates a fully integrated CRO and biotech platform offering discovery, engineering, and preclinical production of antibodies and CAR-T cells.

This dual capability accelerates internal pipeline development and generates revenue via contract research partnerships, de-risking the investment opportunity.

## CONTACT US

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