



Breakthrough Drug Delivery for Mucosal Surfaces Improving Patient Outcomes

Investment Opportunity

Building on over \$2M invested, we are **now raising \$1M** to reach key valuation inflection points – including First-in-Human clinical testing for OPT-101 and preclinical validation of our MNP platform across multiple indications & mucosal tissues

Platform Proof of Concept \$400K

Generate data across 5 drug classes: encapsulation, extended release, and PK

Fund two full-time scientists to define the API design space

Conduct preliminary toxicology for top candidates

CMC Acceleration \$450K

Parallelize CDMO work to support First-in-Human study of **OPT-101** (505(b)(2) glaucoma lead asset)

Complete formulation and scale-up for **OPT-201** (Dry Eye Disease in Dogs)

Scale-up manufacturing and qualify analytical methods

Strategic Partnering \$150K

Expand out-licensing outreach to pharma, animal health, and oncology partners

Develop investor materials, valuation model and data room

Position for a \$5-8M Series A in 2026

Outcomes of Investment

	Discovery	Preclinical	IND-Enabling	Clinical
Glaucoma OPT-101				Q3 2026
Dog Dry Eye OPT-201				Q4 2026
Partnered Indication OPT-102				Q4 2027
Additional Partnered Indication OPT-301				2028

Proven Platform

Demonstrated compatibility with multiple APIs and indications

Clinical Acceleration

Completed **OPT-101 FIH**; enables shorter clinical pathways for subsequent indications

Commercial Readiness

Enable short term revenue generation through partnerships and early stage out-licensing

Capital Efficiency

Achieve major inflection points with a minimal raise

Leadership Team



F. Lasowski, PhD
CEO

15 Years Experience leading translational biomaterials research into pharma innovation



R. Jacquemart, PhD, MBA
COO

25 Years Experience with biopharma product development & commercialization

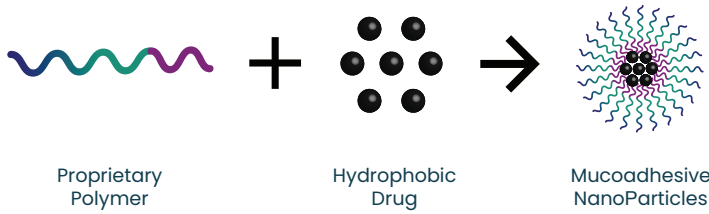


H. Sheardown, PhD
CSO

30 Years Experience with biomaterials & drug delivery research and development

OptimEyes' MNP platform delivers small-molecule drugs directly to mucosal surfaces via a mucoadhesive polymer system that enables sustained, localized drug release. This approach improves bioavailability, compliance and safety, and supports topical delivery for multiple indications. The technology is applicable to all mucosal surfaces, across ophthalmology, respiratory, urogenital, gastrointestinal, oncology and CNS pathologies.

Patented Polymer Technology



Platform Benefits

Compatible
with a wide range of small-molecule APIs

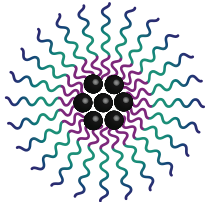
Scalable
manufacturing with modular CMC approach

De-Risked
regulatory pathway enabling speed & IP extensions

Low COGS
with simple supply chain

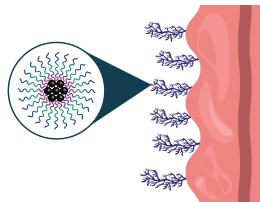
We transform proven molecules into next-generation therapies through localized, sustained delivery – unlocking opportunities across all mucosal surfaces

MNPs



Enable sustained release through encapsulation

Mucosal Binding



Polymeric MNPs adhere to mucosal surfaces

Localized Delivery

Delivered directly to the site of action to reduce off-target side effects

Reduced Dosing Frequency

Sustained release enabling fewer administrations, improved patient compliance and convenience

Greater Bioavailability at Target

Greater drug loading and prolonged surface interaction give more drug at the target tissue

Key Data & Milestones Achieved



Clear IP position based on novel polymer and formulation methods



Validated **encapsulation, extended release** and **pre-clinical studies** on 3 APIs



Regulatory plan confirmed in preliminary consultations



CDMO contracted: tech transfer for gmp manufacturing in preparation



Pre-clinical safety and glaucoma **efficacy** in canine



Glaucoma Clinical site and protocol for FiH study selected

Broad Applicability



Ocular



Nasal



Buccal



Urogenital



GI



Lung