

# HEXIMA LIMITED

# ASX ANNOUNCEMENT



02 November 2021

## 2021 ANNUAL GENERAL MEETING – CEO’S PRESENTATION

MELBOURNE, AUSTRALIA (2 December 2021): Hexima Limited (ASX:HXL) provides the attached CEO Presentation to be delivered at today’s Annual General Meeting commencing at 11.00am AEDT.

The AGM can be joined at <https://meetings.linkgroup.com/HXL21>

**This announcement is authorised for release to ASX by Michael Aldridge, Managing Director & CEO.**

### ***Enquiries:***

Dr Nicole van der Weerden  
Chief Operating Officer  
[n.vanderweerden@hexima.com.au](mailto:n.vanderweerden@hexima.com.au)

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### **ABOUT HEXIMA**

Hexima (ASX:HXL) is a clinical stage, anti-infectives focused biotechnology company engaged in the research and development of defensin peptides for applications as human therapeutics. Our lead product candidate, pezadeftide (HXP124) applied in a topical formulation, is a potential new prescription treatment for toenail fungal infections (or onychomycosis). Hexima is currently conducting an Australian phase IIb clinical trial testing pezadeftide for the treatment of onychomycosis. Hexima holds granted, long-life patents protecting pezadeftide in major markets globally. For additional information please visit [www.hexima.com.au](http://www.hexima.com.au). You can also find us on [Twitter](#) and [LinkedIn](#) or email us at [info@hexima.com.au](mailto:info@hexima.com.au).

# MANAGING DIRECTOR & CHIEF EXECUTIVE OFFICER PRESENTATION

MR MICHAEL ALDRIDGE



# MAJOR ACHIEVEMENTS FY2021

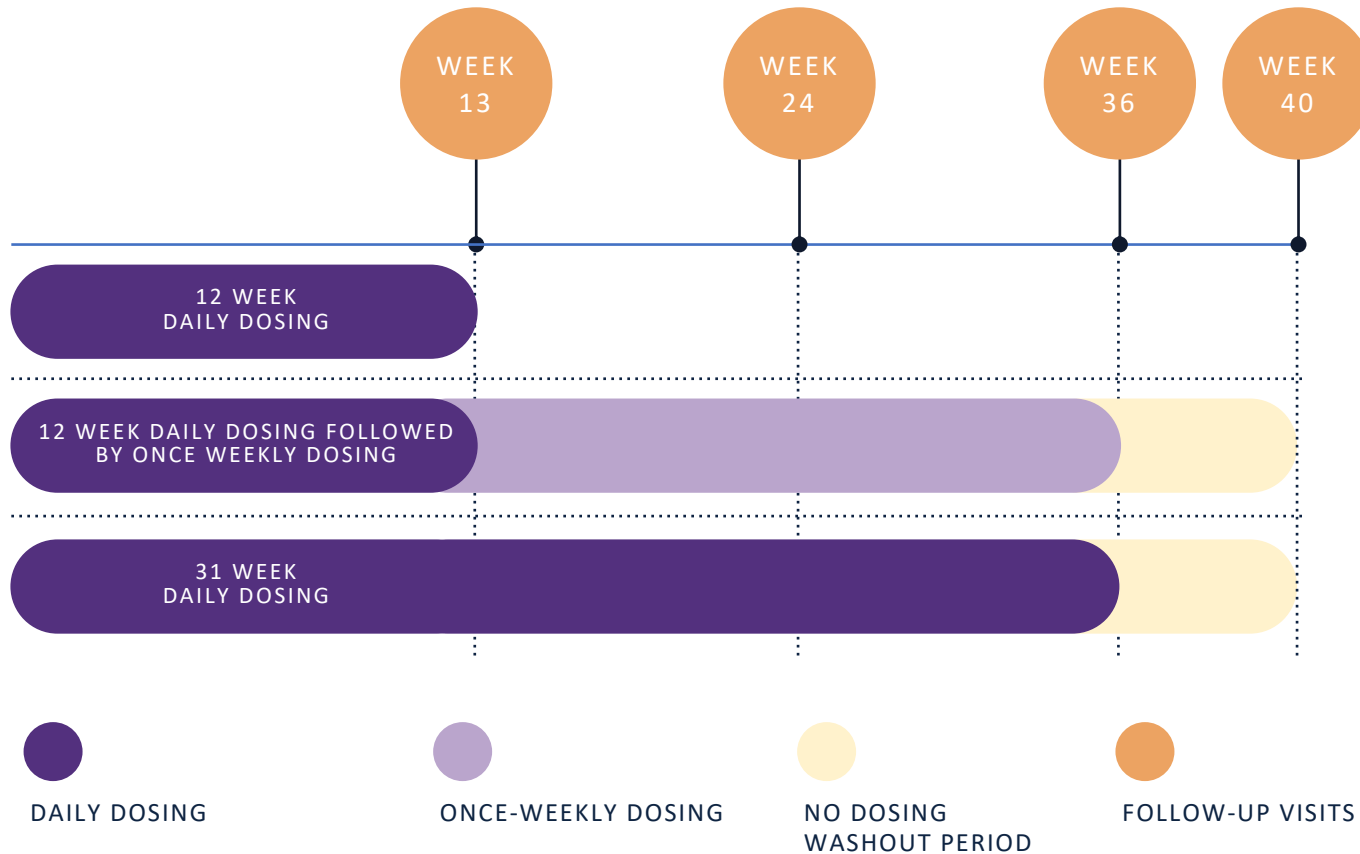
## Milestones & Achievements

- Public Offering and listing on ASX
  - \$3 million offering at A\$0.20 per share
- Phase IIb clinical trial
  - Initiation and completion of enrolment
- Intellectual property protection
- Scientific Advisory Board
  - Australian, US and Japan KOLs



# AUSTRALIAN PHASE IIB CLINICAL TRIAL

HXP124-ONY-002



NOTE: DAILY DOSING PERIODS INCLUDE 1-WEEK WASHOUTS EVERY 6 WEEKS

Enrolment completed July 2021

- Multi-center, randomised, double blind, vehicle-controlled study
- Primary endpoint safety and tolerability, secondary endpoints Mycological Cure and Clinical Efficacy
- Three active (2% pezadeftide) versus vehicle arms to test optimal dosing strategy
- Safety & efficacy assessed at 13, 24, 36 and 40 weeks, data expected Q2 2022



# Strong patent position

ADDITIONAL PROTECTION VIA  
FORMULATION PATENTS AND MARKET  
EXCLUSIVITY FOR BIOLOGICS

## Clearly defined growth strategy

- Develop independently in US and EU (ICH) markets
- License and collaborative development in Japan and potentially China - presently in preliminary discussions with multiple parties

Granted patents  
(exp 2035) in major  
markets covering the  
use of pezadeftide in  
the treatment of  
onychomycosis



Granted and  
pending patents  
covering stabilising  
formulation for  
pezadeftide



12-year US market  
exclusivity on FDA  
approval likely available  
as a biologic drug



# LOOKING FORWARD

A PIVOTAL YEAR IN GROWTH AND DEVELOPMENT

## Expected Milestones

- Ongoing US focused Phase III preparations
  - Completed \$11 million financing
- Phase IIb clinical trial – data Q2 2022
- Corporate partnership(s)
- Capital raising to fund phase III
- Expansion of development pipeline
- Initiate Phase III clinical trials - late CY2022



# Pezadeftide: a potential solution for a large and poorly served market



## POORLY SERVED MARKET

Affects 14% of the population  
Strong consumer preference for topical products  
Clear unmet medical need



## NEW AND UNIQUE

Novel molecule with unique mode of action  
Strong patent protection and long patent life



## SAFE

No systemic effects  
No local redness or irritation



## CONVENIENT

Easy to apply  
Short treatment duration  
Rapid clearing of infected nail



## EFFECTIVE

Efficiently penetrates the nail  
Rapidly kills fungus  
Best-in-class mycological cure

