



8 May 2025

ASX Announcement

US non-deal roadshow presentation

MELBOURNE Australia, 6 May 2025: Australian antiviral drug development company, Island Pharmaceuticals Ltd (ASX: ILA; Island or the Company) is pleased to provide the attached copy of the Company's latest investor presentation, which will be used as part of an upcoming US non-deal roadshow.

Island advises that CEO and Managing Director, Dr David Foster will be undertaking a number of investor meetings in New York from Thursday, 8 May to Friday, 9 May with several US-based institutional investors, family offices and high net-worth individuals.

During meetings, Dr Foster will have the opportunity to provide US investors with a detailed overview of the clinical pathway for the Company's lead asset ISLA-101, prior to the pending release of unblinded results from the Phase 2a/b PROTECT clinical trial utilising ISLA-101 to combat dengue fever. High level results are anticipated later this month. Dr Foster will also provide an update to US investors around the Company's pipeline expansion opportunities, with particular focus on Galidesivir, a broad-acting antiviral molecule.

CEO and Managing Director, David Foster said: *"We are pleased to commence this roadshow, which has attracted strong interest from a diverse network of sophisticated US investors. We are confident that it will provide the Company with a fantastic opportunity to clearly articulate our value proposition at an exciting phase of our clinical development pathway. I look forward to meetings with a new investor network in New York ahead of the imminent release of our Phase 2a/b PROTECT trial results in the coming weeks."*

- Ends -

To subscribe to Island's monthly newsletter, [IslandWatch](#), and other forms of email communications, please visit [this page](#) of our website.



Approved for release to the ASX by:

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About Island Pharmaceuticals

Island (ASX: ILA) is a drug repurposing company, focused on areas of unmet need for antiviral therapeutics to address infectious diseases. Our lead asset is ISLA-101, a drug with a well-established safety profile, being repurposed for the prevention and treatment of dengue2 fever and other mosquito (or vector) borne diseases.

If ISLA-101 achieves FDA approval, and certain other criteria are met, Island may be eligible to obtain a "Priority Review Voucher" at the time of FDA approval. This means that as well as getting approval to manufacture and sell ISLA-101, the Priority Review Voucher (PRV) could permit Island to expedite the FDA approval process for a new drug or sell the PRV in a secondary market.

Island encourages all current investors to go paperless by registering their details with the Company's share registry, Automatic Registry Services, whose contact info is housed on the Shareholder Services page of the Company's website.

Visit www.islandpharmaceuticals.com for more on Island.



ISLAND

PHARMACEUTICALS

Antiviral therapeutics

COMBATting URGENT VIRAL DISEASE THREATS

US NON-DEAL ROADSHOW PRESENTATION

DR DAVID FOSTER, MANAGING DIRECTOR

May 2025

(ASX: ILA)

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Financial data All dollar values are in Australian dollars (\$) or A\$) unless otherwise stated. Any financial data in this presentation is unaudited. **Past performance** The operating and historical financial information given in this presentation is given for illustrative purposes only and should not be relied upon as (and is not) an indication of the Company's views on its future performance or condition. Actual results could differ materially from those referred to in this presentation. You should note that past performance of the Group is not and cannot be relied upon as an indicator of (and provides no guidance as to) future Group performance.

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This presentation contains certain "forward-looking statements". The words "expect", "anticipate", "estimate", "intend", "believe", "guidance", "propose", "goals", "targets", "aims", "outlook", "forecasts", "should", "could", "would", "may", "will", "predict", "plan" and other similar expressions are intended to identify forward-looking statements. Any indications of, and guidance on, future operating performance, earnings and financial position and performance are also forward-looking statements. Forward-looking statements in this presentation include statements regarding the Company's future growth options, strategies and new products. Forward-looking statements, opinions and estimates provided in this presentation are based on assumptions and contingencies which are subject to change without notice, as are statements about market and industry trends, which are based on interpretations of current market conditions.

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Nothing in this presentation will under any circumstances create an implication that there has been no change in the affairs of the Group since the date of this presentation.



**Island Pharmaceuticals
(ASX: ILA) is an antiviral
therapeutics company
targeting infectious diseases**



Dengue infects up
to 400m per year*



Major market
potential



Positive results in
aggressive models



Phase 2a/b PROTECT clinical
trial in dengue underway



Priority Review
Voucher potential



Pipeline expansion
pending with expedited
approval route defined



CORPORATE OVERVIEW



Snapshot

Share on issue ¹ :	210,259,700
Price per share ¹ :	\$0.18
Market capitalisation ¹ :	\$37.85m
Cash at bank (31 March 2025) ² :	\$4.82m
DoD grant funding to directly support the Phase 2a/b PROTECT clinical study	USD \$625k

Substantial shareholders

Dr William James Garner ³	15.79%
Jason Alan Carroll ⁴	14.73%
MWP Partners Limited ⁵	9.94%
Dr Daniel Tillett ⁶	6.72%

1. As at 5 May 2025

2. Does not take into consideration cash used since reporting date

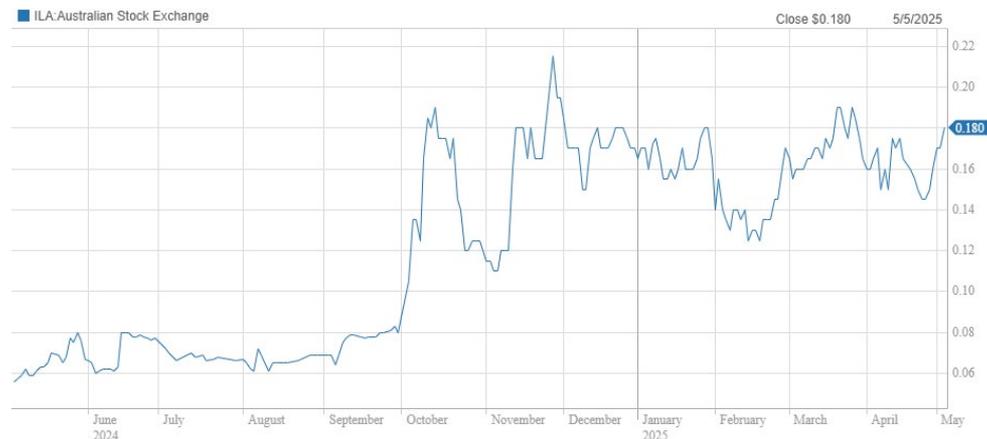
3 Per holding per Substantial interest notices lodged with ASX on 25 March 2025

4. Per holding per Substantial interest notice lodged with ASX on 25 March 2025

5 Per holding per Substantial interest notice lodged with ASX on 21 March 2025

6 Per holding per Substantial interest notices lodged with ASX on 26 March 2025

Price & volume (12 month)



Board of Directors

Phil Lynch, Executive Chairman

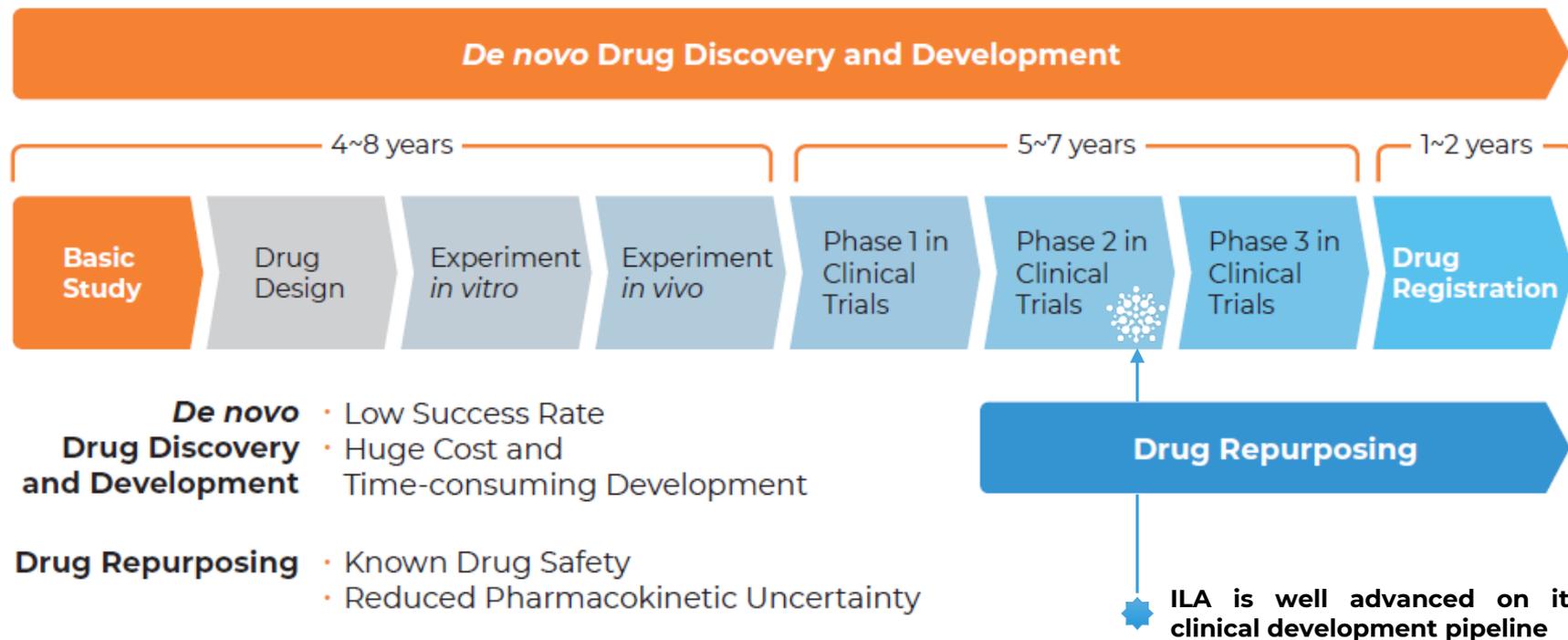
Dr David Foster, CEO and Managing Director

Chris Ntoumenopoulos, Non-Executive Director

COMPANY OVERVIEW

- Lead asset, ISLA-101 repurposed for dengue fever and other mosquito (or vector) borne diseases
- ISLA-101 has been part of 40+ Phase I, II and III human clinical trials in indications including cancer and respiratory diseases, and deemed safe by regulators
- Pre-clinical work at Monash University demonstrated promise as an antiviral drug
- ISLA-101 has US government and military support
- Phase 2a/b PROTECT trial has prophylactic (preventative) and therapeutic (treatment) arms via two patient cohorts
- Promising Phase 2a (preventative) cohort data allowed for start of Phase 2b (treatment) cohort
- Phase 2a/b high level results expected in May 2025
- IND (Investigational New Drug) status with the FDA provides Island with a streamlined pathway to market
- Effort to expand asset portfolio is well advanced

BENEFITS OF DRUG REPURPOSING

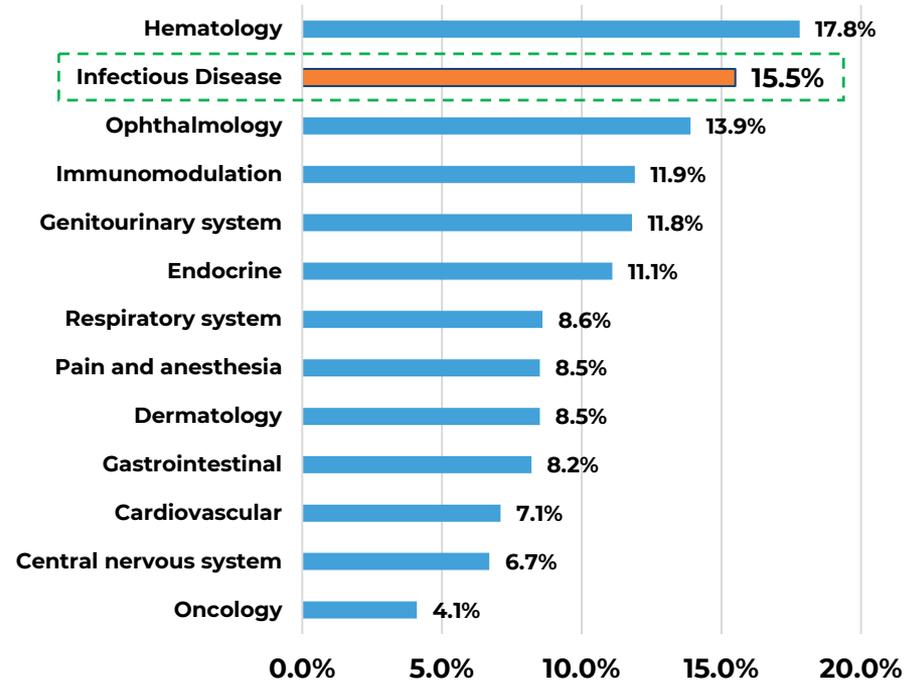


INCREASED LIKELIHOOD OF SUCCESS



- Treatments for infectious disease **have a statistically higher likelihood of overall success** in clinical trials
- Anti-infective treatments sit at the low end of the drug development cost curve across all therapeutic areas
- JAMA research shows that **anti-infective drugs were the least expensive to develop**¹
- Infectious disease treatments have the third-highest probability of phase II success (38.4%), behind hematology (48.1%) and metabolic (45%) treatments²
- Probability of a successful phase III transition (post phase II) for infectious disease treatments increases to 64%²

Drug Development - Overall probability of success



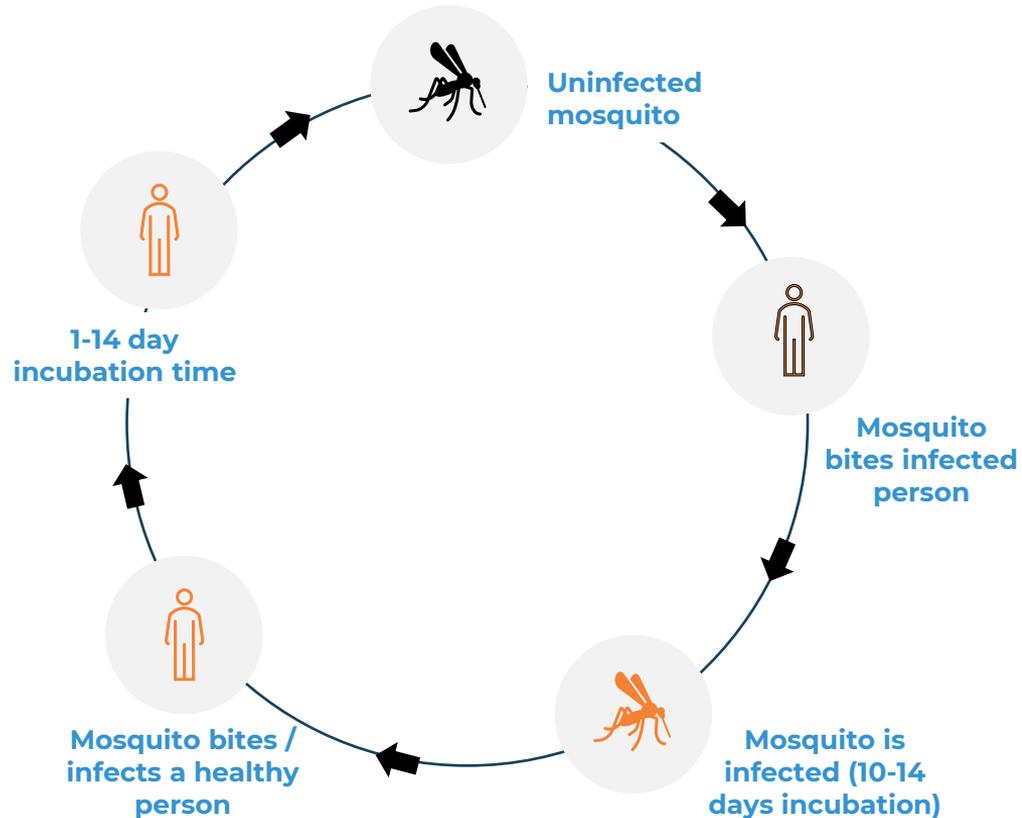
¹ JAMA Network: *Costs of Drug Development and Research and Development Intensity in the US, 2000-2018*

² Biotechnology Innovation Organisation: *Clinical development success rates and contributing factors 2011-2020*

DENGUE – INFECTION CAN LEAD TO LETHAL OUTCOMES



- Dengue is a viral infection transmitted to humans through the bite of infected mosquitoes
- It directly impacts white blood cell count and platelets - vital for body protective mechanisms
- Moderate to severe symptoms include high fever, muscle pain, shock, bleeding, vomiting and seizure amongst others
- There is no specific treatment for dengue
- Some vaccines have been shown to have preventative nature but are in limited supply
- ISLA-101 is scalable oral dosing solution which has demonstrated activity against dengue strains

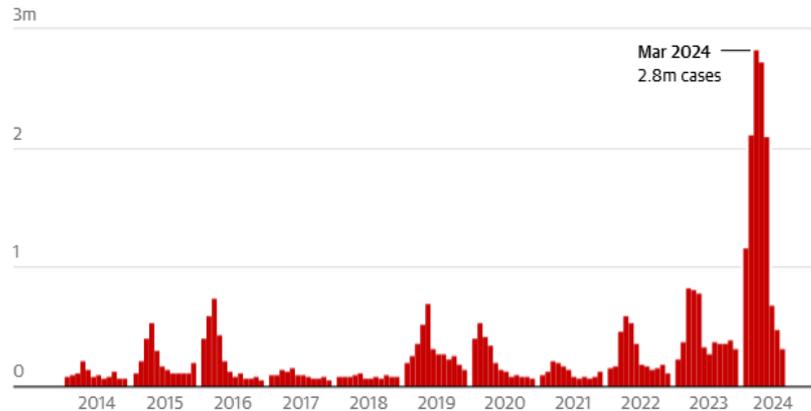


DENGUE - COMMON AND SPREADING

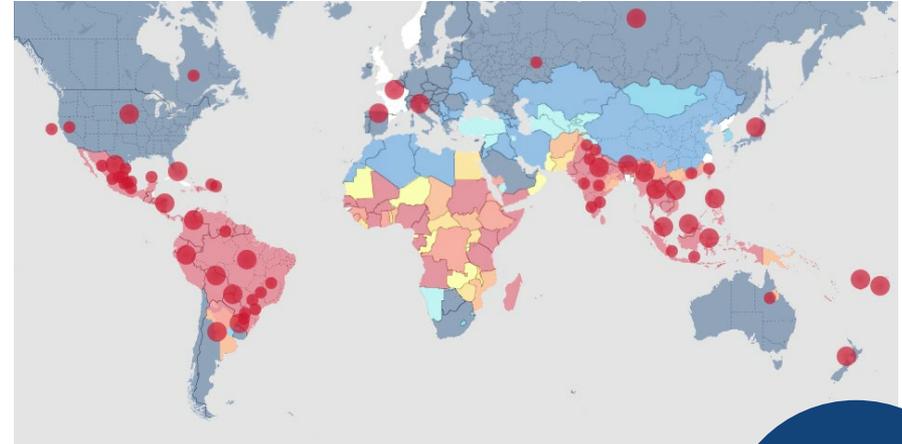


Global cases of dengue fever rose steeply in 2024

Monthly global cases, millions



Guardian graphic. Source: WHO. Note: case reporting requirements vary by country



Absent Unlikely Uncertain Likely Present
Country Level Local Level

HealthMap: recent reports of local or imported dengue cases (March 2025)

US\$8.9B
Estimated impact to the economy from dengue fever

“About half of the world's population is now at risk of dengue with an estimated 100 – 400 million infections occurring each year”

World Health Organisation, 30 May 2024

DENGUE BY 2050 IS MORE PREVALENT



Driven by:



Warmer temperatures

- Accelerating development
- Increases activity of female mosquitoes
- Reduces incubation time for mosquito to become infectious
- Allow mosquitoes to survive longer through winter



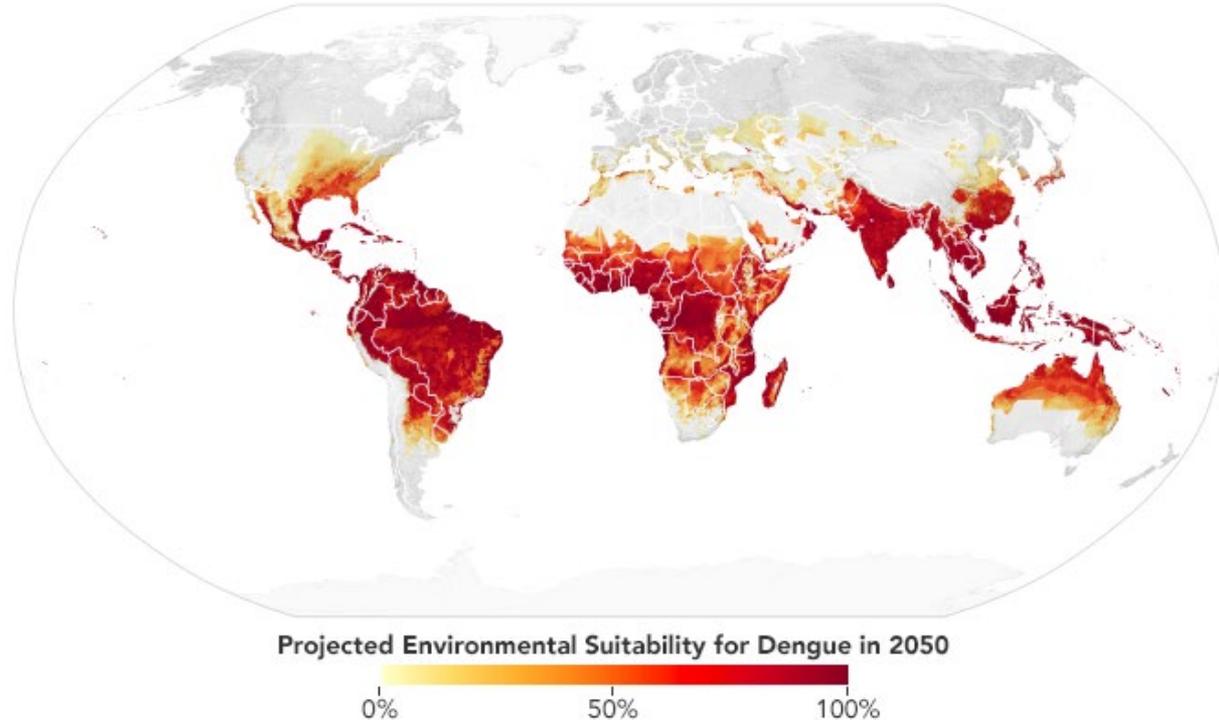
High humidity

- Improves mosquitoes' chance of survival



Extreme weather

- Disrupts water / sanitation
- Increased flooding can enhance breeding



NASA Earth Observatory map by Lauren Dauphin based on data from Janey Messina, University of Oxford - <https://earthobservatory.nasa.gov/features/disease-vector>

A MULTI-BILLION-DOLLAR MARKET OPPORTUNITY



- There is currently no specific treatment for dengue – providing ILA with the opportunity to be first to market
- Takeda Pharmaceuticals project global sales (ex-USA) of its Qdenga dengue vaccine at US\$1.6bn to US\$2bn by 2030¹
- Qdenga is a prophylactic or preventative measure – ISLA-101 has the potential to become a treatment and/or prophylactic
- Quick establishment of the antimalarial drug market highlights potential for dengue drug development trends
- Antimalarial drug market was valued at US\$1.76Bn in 2024 with a potential to grow to US\$2.5Bn by 2030²

Dengue Fever treatment – Growth forecast (USD Bn)



Source: [Market Research Future](#) /Rahul Gotadki, May 2025)

¹Antimalarial Drugs Market by Drug Class, Drug Type, Route of Administration, Malaria Type, Distribution Channel, End User - Global Forecast 2025-20

²Fierce Biotech: [Takeda taps Biological E to ramp up Qdenga manufacturing capacity on quest to make 100M doses a year](#)

ISLA-101 – BROAD ACTIVITY EVIDENT

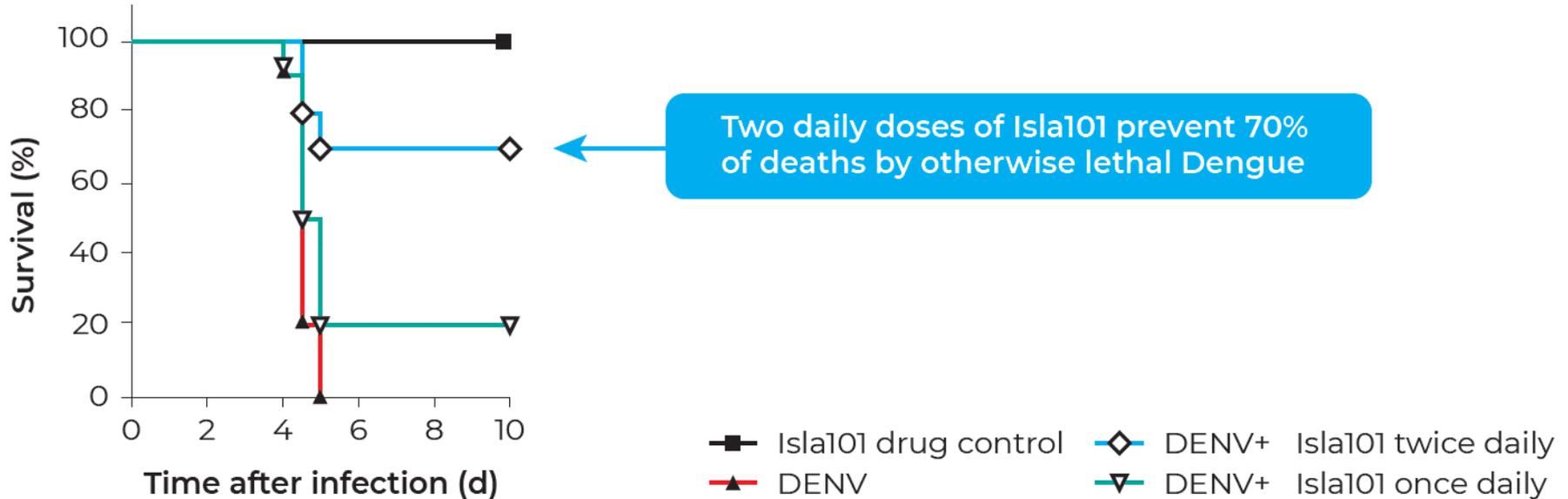


Demonstrated activity against flaviviruses (subgroup of arboviruses) in models of infection



- ISLA-101 has demonstrated broad anti-viral activity in *in-vitro* models
- Demonstrated potent anti dengue-1 activity in *in-vitro* models using fresh human cells
- Protective in dengue fever and Zika in animal models
- Shown to prevent death in 70% of subjects in extremely lethal animal models
- Increasing concentrations of ISLA-101 prevent death induced by an otherwise lethal dengue fever infection
- 48 human clinical studies completed in other indications
- ILA's Single Ascending Dose study and further modelling reinforced safety / tolerability and identified dosing for Phase 2 trial

PREVENTING ANIMAL DEATHS FROM LETHAL DENGUE AND PROTECTIVE AGAINST ZIKA



Survival curve showing protection from lethal dengue change by increasing dose of ISLA101 (mouse model).

PHASE 2A/B (PROTECT) STUDY OVERVIEW



Randomised, double blind, placebo-controlled dengue challenge study – prophylactic and treatment challenge:

- Study include a prophylactic (Phase 2a) and therapeutic (Phase 2b) arm
- Prophylactic Cohort- 2a: 4 subjects randomized 3:1
- Therapeutic Cohort: 2b: 10 subjects randomized 8:2
- Primary endpoint:
 - Assess effect of ISLA-101 on viremia after challenge with DENV-1-LVHC
- Secondary endpoints:
 - Characterise clinical, immunologic and virologic responses following ISLA-101 after challenge with DENV-1-LVHC
 - Assess effect of ISLA-101 on clinical signs and symptoms after challenge with DENV-1-LVHC
 - Assess safety of ISLA-101 in the challenge with DENV-1-LVHC
- **High level, unblinded results from both cohorts expected this month**



Trial being conducted at SUNY Upstate Medical University
Syracuse, New York.

PHASE 2A/B (PROTECT) DESIGN

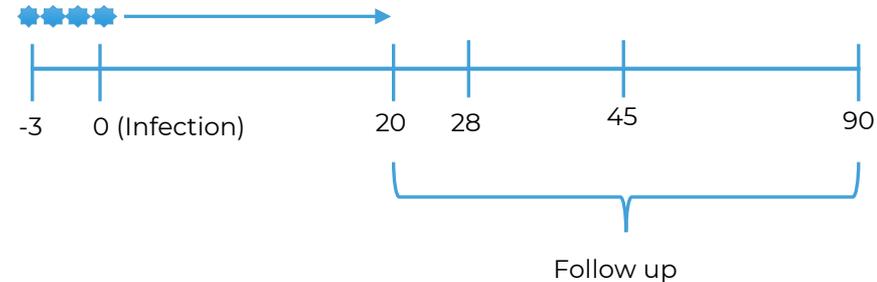


- Phase 1 (completed April 2024) achieved all study outcomes relating to safety and dosing, demonstrating benefit of Challenge study approach
- Phase 2a (prophylactic) subjects dosed in October 2024
- Safety Review Council review highlighted:
 - Administering ISLA-101 was safe
 - Study achieved appropriate ISLA-101 blood concentrations
 - Dosed subjects exhibited evidence of antiviral activity versus control
 - Unanimous decision to advance 2b cohort
- 2b (treatment) cohort administered ISLA-101 in February 2025
- Pharmacokinetic analysis of 2b cohort has shown target blood level concentration was achieved in all participants

High-level, unblinded results this month

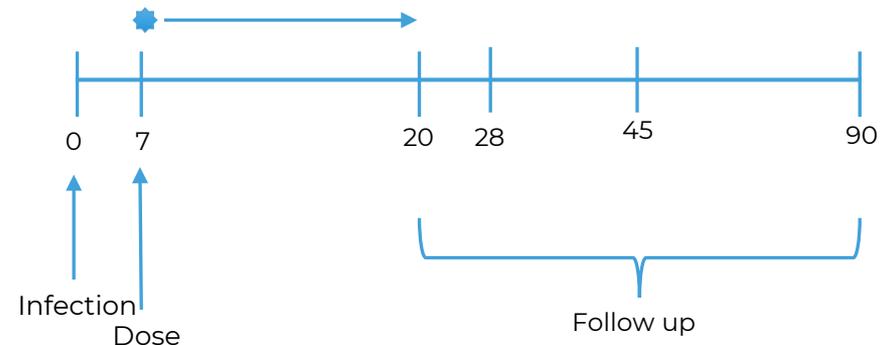
Phase 2A: Prophylactic (preventative) cohort

Administer ISLA-101 daily



Phase 2B: Therapeutic (treatment) cohort

Administer ISLA-101 daily

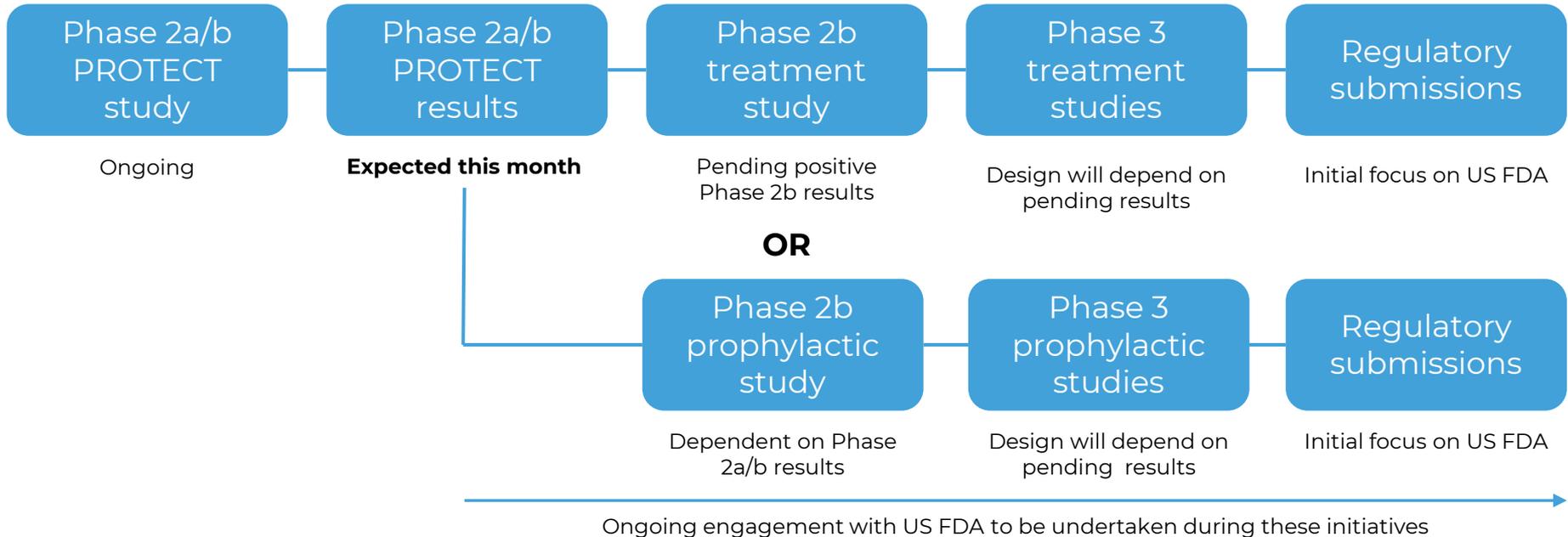


CLINICAL TRIAL AND REGULATORY PATHWAY



A defined clinical and regulatory route based on Phase 2a/b study results

- Two likely pathways depending on Phase 2a/b results
- Discussions advancing with multiple potential strategic partners for additional phase 2 and 3 clinical trials



GALIDESIVIR WIP - SIGNIFICANT PIPELINE OPPORTUNITY



Galidesivir – potential to tackle Ebola, Marburg, Zika & other RNA viruses

- 12-month binding LOI signed in Sept 2024
- Small molecule, re-purposable with reduced timeframe to market
- Substantial Phase 1 human safety data
- Demonstrated efficacy in multiple lethal animal models – may provide access to FDA’s Animal Rule
- Extensive US government funding to date
- PRV eligible across numerous potential indications
- Multiple commercial opportunities in travel, military, national safety and government stockpiling

POTENTIAL REGULATORY FAST TRACK



Existing Galidesivir data package and FDA Animal Rule unlock quicker approval path

- Data package includes successful non-human primate study in Marburg and two phase 1 safety studies
- FDA's Animal Rule allows for approval of drugs for in indications, based on animal efficacy data when human trials are unethical or infeasible, provided safety is shown in humans and the disease is well modelled in animals
- ILA may have the potential to undertake one successful animal study, which would allow for submission of a New Drug Application (NDA) with the FDA – this approval would unlock a Priority Review Voucher, valued in excess of US\$150m
- PRV's are an incentive granted by the FDA allowing the holder to expedite the review of a future drug application, typically awarded for developing treatments for neglected tropical diseases or medical countermeasures



NEAR TERM MILESTONES



A number of value catalysts pending in the coming months

Milestone	Timeframe
Advancement of Galidesivir opportunity	May 2025
High level, unblinded results from Phase 2a/b PROTECT study	May 2025
Meeting with US FDA to discuss Phase 2a/b PROTECT study results	Q3 CY2025
Completion of Phase 2/3 clinical trial pipeline planning	Q3 CY2025
Commencement of Phase 2/3 clinical trials based on Phase 2a/b PROTECT study results	Q4 CY2025
Ongoing engagement with potential partners for ISLA-101 clinical trial pathway	Ongoing
Assessment of additional pipeline opportunities to broaden asset portfolio	Ongoing

Dates are indicative only, based on current estimates and subject to change

Island Pharmaceuticals (ASX: ILA)



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