



10 February 2022

ASX Announcement
**Updated investor presentation and
webinar invitation**

MELBOURNE Australia, 10 February 2022: Australian mid-clinical stage antiviral drug development company, Island Pharmaceuticals Ltd (ASX: ILA; "Island"; "the Company") is pleased to release a copy of an investor presentation that will be delivered to investors over the coming weeks.

In addition, Island announces its participation in the ShareCafe Small Cap "Hidden Gems" webinar, to be held tomorrow, Friday 11 February 2022 from 12:30pm AEDT / 9:30am AWST. Through the webinar, Chief Executive Officer Dr. David Foster will provide an update on Island's progress in moving lead drug candidate, ISLA-101 through to the clinic for dengue fever.

This webinar is able to be viewed live via Zoom and will provide viewers the opportunity to hear from, and engage with, a range of ASX-listed leading micro/mid cap companies. To access further details of the event and to register at no cost, please copy and paste the following link into your internet browser:

https://us02web.zoom.us/webinar/register/5416151767246/WN_rJUuE_AySLqmpK1pJMzpXg

A recorded copy of the webinar will be made available following the event.

Approved for release to the ASX by:

Dr Paul MacLeman
Executive Chairman
Island
Pharmaceuticals Ltd
info@islandpharmaceuticals.com

Investors and media, for further information, please contact:

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About Island Pharmaceuticals

Island (ASX: ILA) is a mid-clinical-stage drug repurposing company, focused on the topical area of antiviral therapeutics for infectious diseases. Our lead asset is ISLA-101, a drug with a well-established safety profile, being repurposed for the prevention and treatment of dengue fever and other mosquito (or vector) borne diseases. The Company is close to commencing a Phase 2a clinical trial in dengue-infected subjects.



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, Automic 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

SOLVING URGENT VIRAL DISEASE THREATS

(ASX: ILA)

Investor update – February 2022

DISCLAIMER

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Not an offer or financial product advice

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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.

Future performance

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.

Forward-looking statements, including projections, guidance on future operations, earnings and estimates (if any), are provided as a general guide only and should not be relied upon as an indication or guarantee of future performance. No representation is given that the assumptions upon which forward looking statements may be based are reasonable. This presentation contains statements that are subject to risk factors associated with the Group's industry. These forward-looking statements may be affected by a range of variables which could cause actual results or trends to differ materially, including but not limited to earnings, capital expenditure, cash flow and capital structure risks and general business risks.

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Subject to any continuing obligations under applicable law, the Company disclaims any obligation or undertaking to provide any updates or revisions to any forward-looking statements in this presentation to reflect any change in expectations in relation to any forward-looking statements or any change in events, conditions or circumstances on which any such statement is based.

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.

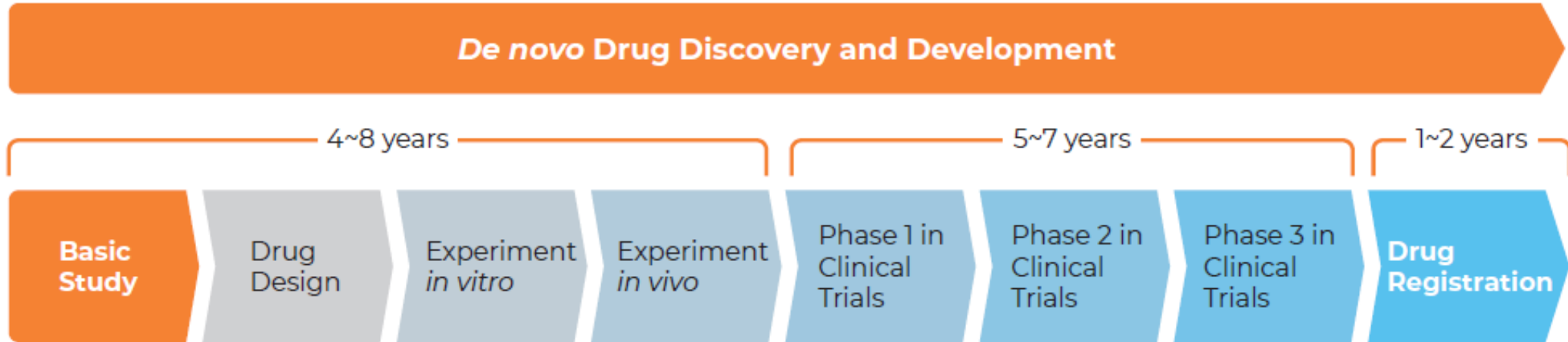
Island Pharmaceuticals (ASX: ILA) is a mid clinical-stage drug repurposing company, focused on the rapid development of antiviral therapeutics for infectious diseases

Island Pharmaceuticals lists on the ASX following oversubscribed A\$7.5m IPO in April 2021

Island's drug repurposing strategy enables rapid and efficient development of antiviral therapies

Initial focus is on mosquito borne diseases with a Phase II lead program in Dengue fever

THE BENEFITS OF DRUG REPURPOSING



De novo Drug Discovery and Development

- Low Success Rate
- Huge Cost and Time-consuming Development

Drug Repurposing

- Known Drug Safety
- Reduced Pharmacokinetic Uncertainty

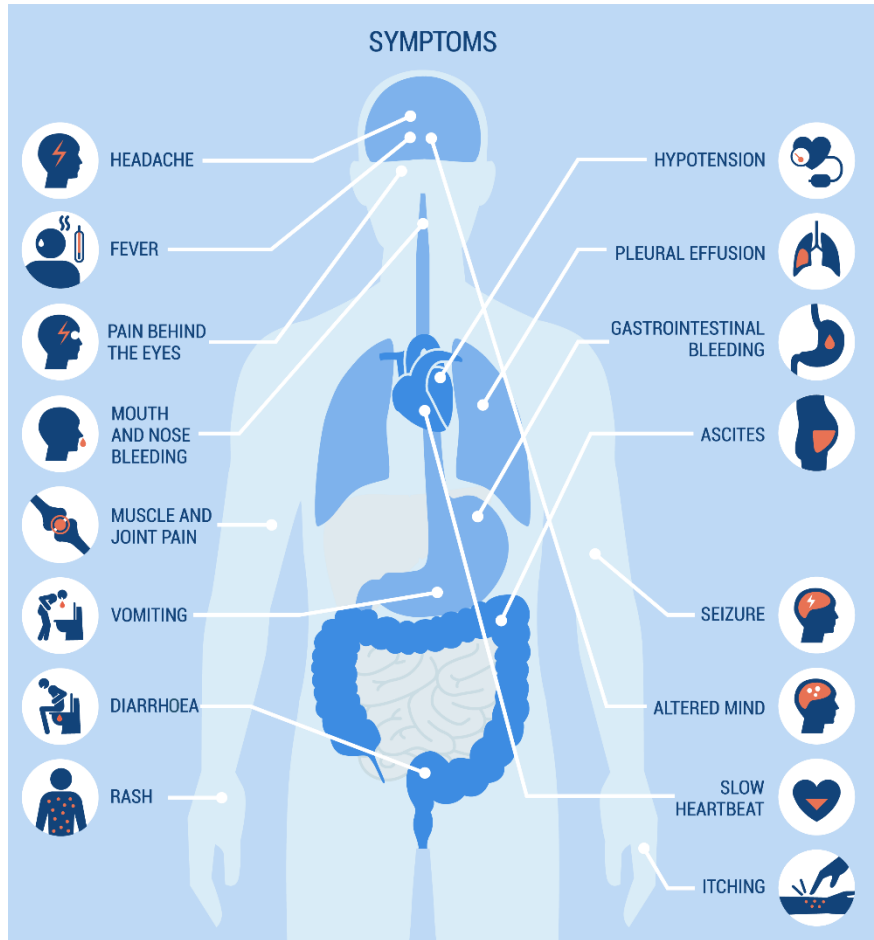


THE CHALLENGE



WHY DENGUE AS A FIRST TARGET FOR ISLA-101?

SIGNIFICANT UNMET NEED FOR DISEASE WITH INCREASING INCIDENCE



Significant unmet need (3.9 billion people at risk)

Increasing spread to US, EU and Australia

ISLA-101 has both therapeutic and prophylactic potential

Strong animal and human model results

First claim then springboard into other arboviruses

Priority Review Voucher eligibility

COMMUNITIES HIT BY TWO VIRUSES AT ONCE



Dec 1, 2021

Chikungunya, Zika, and Dengue virus incidence in Mexico may be higher than previously reported



The researchers found 2.4 times the rate of arbovirolosis as originally reported, including coinfections, suggesting underestimation of the incidence of the three viruses. However, future research is needed to provide up-to-date incidence estimates of each virus.



BRYTFMONLINE



Porto Alegre issues epidemic alert for dengue, Zika virus and chikungunya at the end of the year

Dec 30, 2021

Colombia issues dengue fever alert

by NEWS DESK

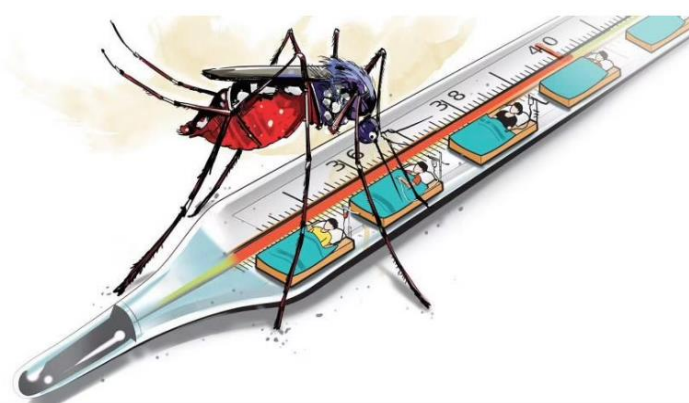
November 14, 2021



Rising dengue count a worry amid Covid

Nov 17, 2021

If the number of dengue cases continues to rise, complications and fatalities due to the disease are prone to occur, they opine.



The Sydney Morning Herald

World Asia Timor-Leste

East Timor's hospitals are fighting a deadly outbreak, but it's not COVID



By Chris Barrett
February 2, 2022 - 5:35pm

Save Share A A A



Nov 23, 2021

Upcoming mosquito season may be the worst yet. Here's how to prepare yourself

Australians are facing a brutal mosquito season bolstered by recent wet weather



ALJAZEERA

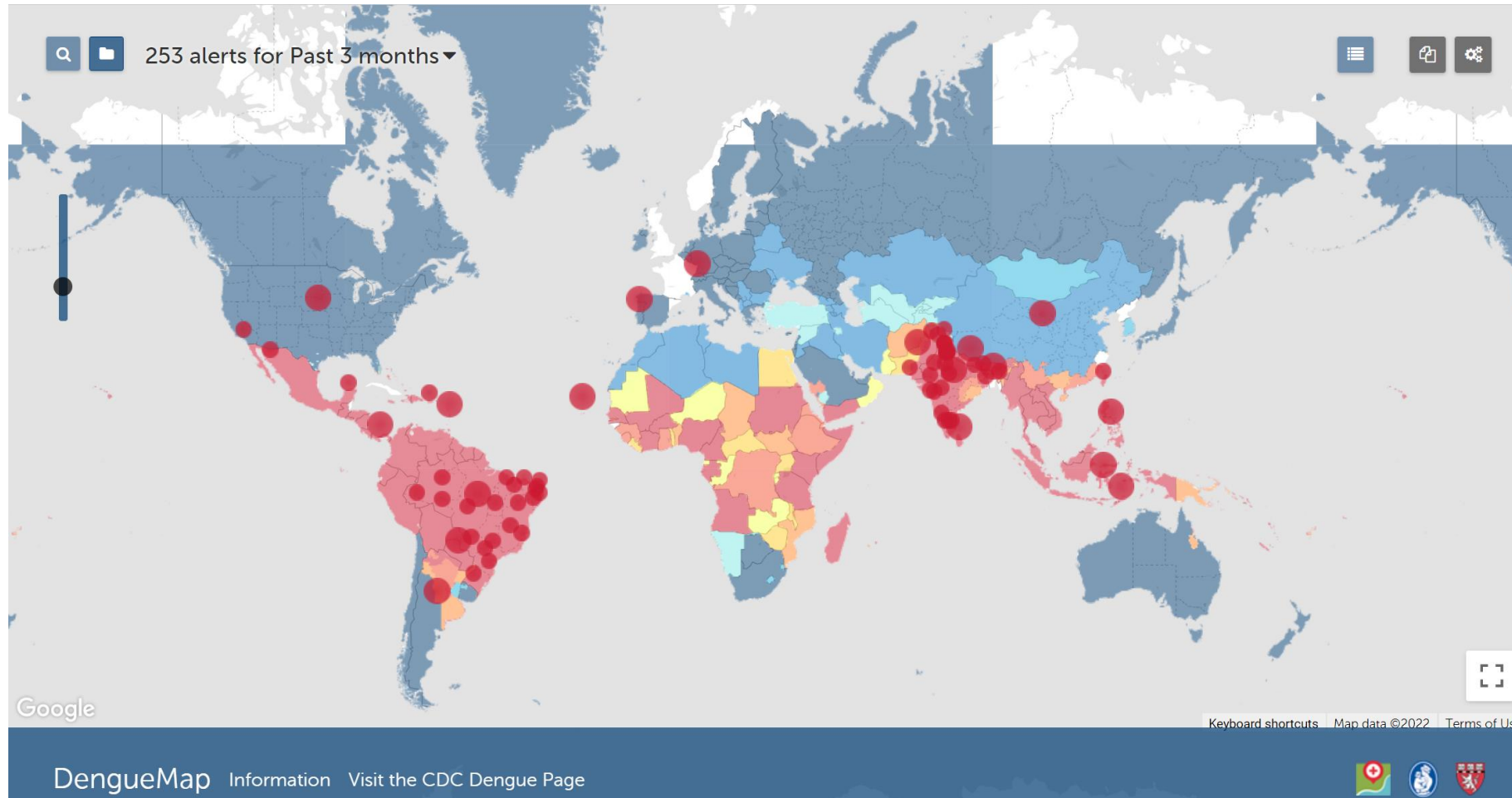
News | Health

India battles spike in dengue cases amid COVID pandemic

Nearly 1,170 dengue cases reported over the past week in New Delhi as the country faces yet another health crisis.



DENGUE IS A WIDESPREAD ISSUE



HealthMap Reports

Recent reports of local or imported dengue cases from official, newspaper, and other media sources.

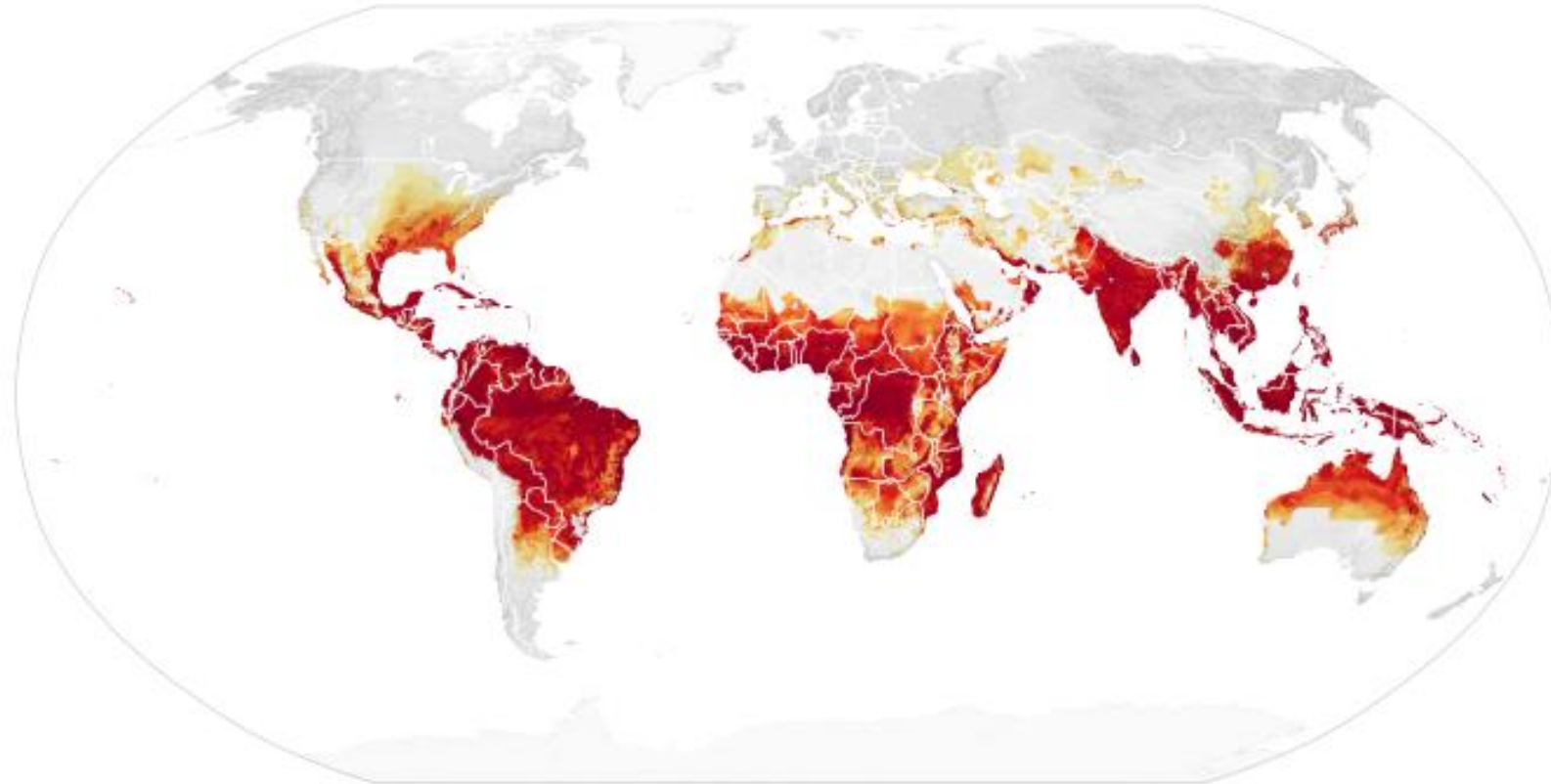
Source.

● Country Level ● Local Level

■ Absent ■ Unlikely ■ Uncertain ■ Likely ■ Present

Dengue outbreaks occurred in many countries of the world in the Americas, Africa, the Middle East, Asia, and the Pacific Islands.

DENGUE IN 2050 – A GLOBAL DISEASE



Projected Environmental Suitability for Dengue in 2050



Prediction based on projections of future temperatures, rainfall, and mosquito populations (NASA Earth Observatory map by Lauren Dauphin based on data from Janey Messina, University of Oxford.)

LIMITED AVAILABLE SOLUTIONS

HIGHLY PREVALENT DISEASES WITH UNMET MEDICAL NEED



Worldwide prevalence

Dengue fever	West Nile	Zika Virus	Yellow fever	Japanese Encephalitis
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390 million	n/a	Up to 1.5 million	130,000	70,000
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Viral diseases are a leading cause of endemic and pandemic disease



Effective drug therapy

No	No	No	No	No
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Antimalarial drugs market is expected to reach US\$1B in 2026 providing guidance to potential market size



Vaccine

Limited	No	No	Limited	Limited
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Vaccine development potentially can exacerbate symptoms from infections by different strains

ILSA 101



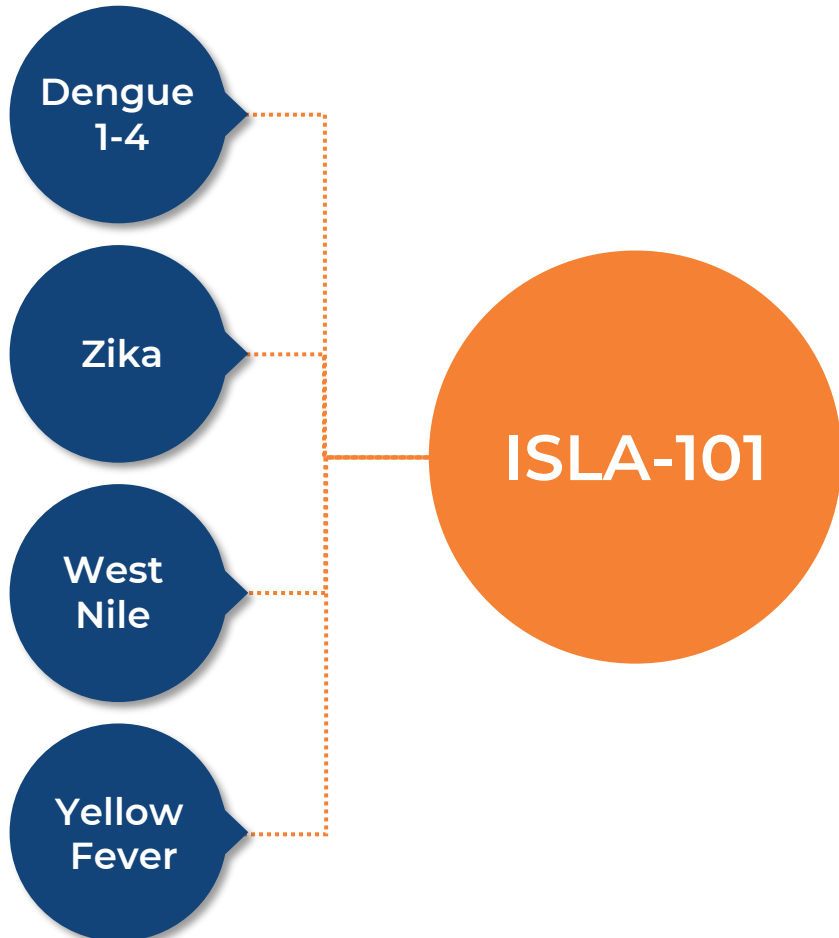
ISLAND

PHARMACEUTICALS

Antiviral therapeutics

ISLA-101 BROAD ACTIVITY EVIDENT

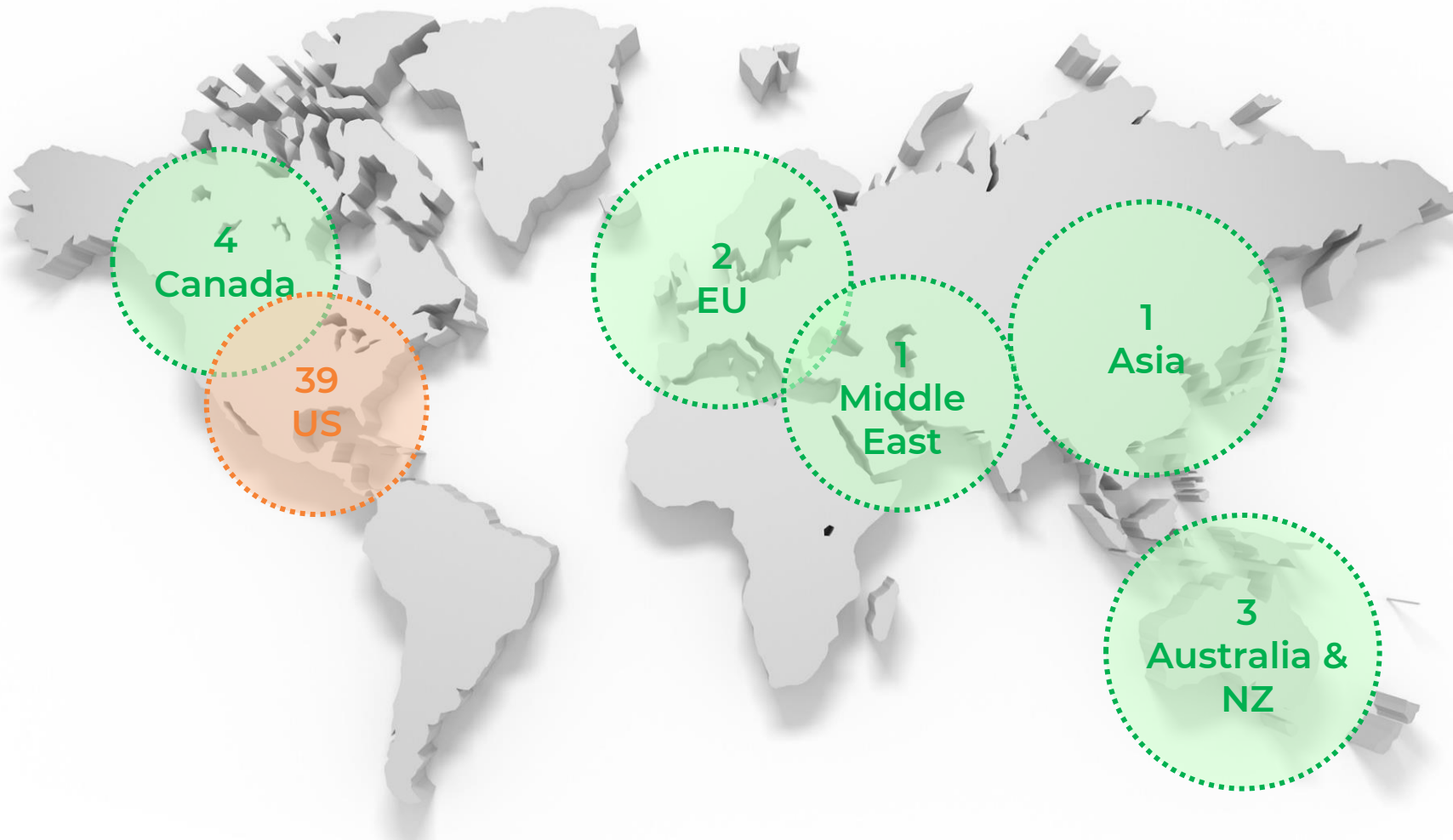
DEMONSTRATED ACTIVITY AGAINST FLAVIVIRUSES (A SUBGROUP OF ARBOVIRUSES) IN MULTIPLE MODELS OF INFECTION



- In *in-vitro* models using fresh human cells, ISLA-101 has demonstrated broad anti-viral activity
- In animal models, ISLA-101 is protective in dengue fever and Zika
- In extremely lethal animal models, ISLA-101 was shown to prevent death in 70% of subjects
- Increasing concentrations of ISLA-101 prevent death induced by an otherwise lethal dengue fever infection
- 45 HUMAN Clinical Studies of ISLA-101 completed in other indications

SAFETY PROFILE OF DRUG ESTABLISHED

45 HUMAN CLINICAL STUDIES OF ISLA-101 COMPLETED IN OTHER INDICATIONS



Verified as safe in humans by multiple regulators in other clinical indications

NEARING IND SUBMISSION

IND IS A CRITICAL REGULATORY MILESTONE, ENABLING PEACH STUDY COMMENCEMENT

Sponsor information (complete)

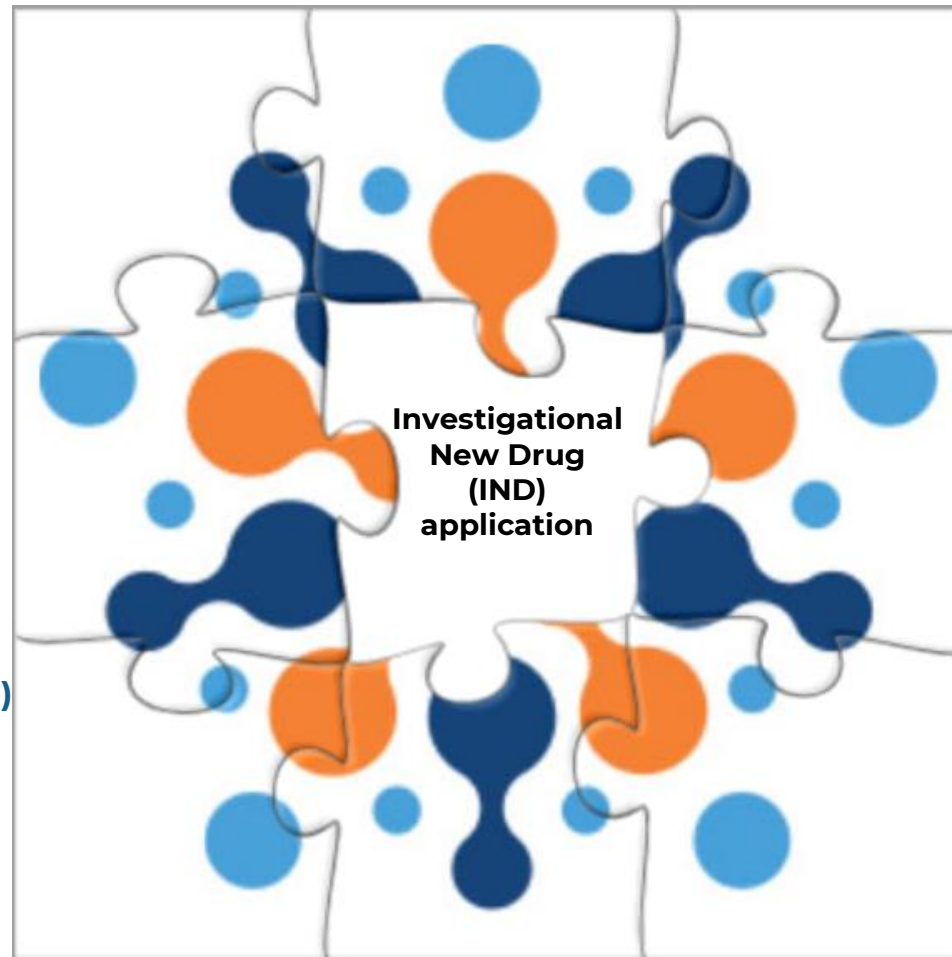
Summary of existing human data (complete)

- ✓ ~45 previous clinical trials
- ✓ Human safety data

Trial monitoring and biostatistics plan (complete)

Preclinical work (complete)

- ✓ Animal pharmacology
- ✓ Toxicology
- ✓ pharmacokinetic data



Clinical trial protocol (almost complete)

Drug manufacturing (almost complete)

- ✓ Drug substance (API)
- Drug product

Investigator's brochure (almost complete)

- A major body of work – includes all instructions to enable the principal investigator to run the trial

Chemistry, manufacturing and control information (almost complete)

- ✓ Manufacturing method
- ✓ Manufacturing validation
- ✓ Drug substance manufacturing data for clinical batch
- Final clinical drug product for ISLA-101 trial manufactured

WORLD CLASS TEAM IN PLACE

TEAM ASSEMBLED WITH DOMAIN EXPERTISE IN CLINICAL TRIALS AND MANUFACTURING

Sponsor



Drug repurposing and dengue fever experts.

Key Island people overseeing the ISLA-101 PEACH study



Teresa Byrne, Vice President
Clinical Product Development.

Overseeing clinical development
of ISLA-101 in the upcoming
PEACH trial and other pipeline
programs.



Larry Norder, CMC Consultant

Overseeing manufacturing of
clinical drug substance, clinical
drug product and formulation
strategy. 25 years experience in
drug development, resulting in
several drug approvals.

Trial site and investigator



Clinical trial protocol, oversight and experience.
Dr Kristopher Paolino, MD has been appointed
Principal Investigator on the trial.

Data and CMC



Providing regulatory strategy and drafting
Investigational New Drug Application for the
use of ISLA-101 in PEACH study.

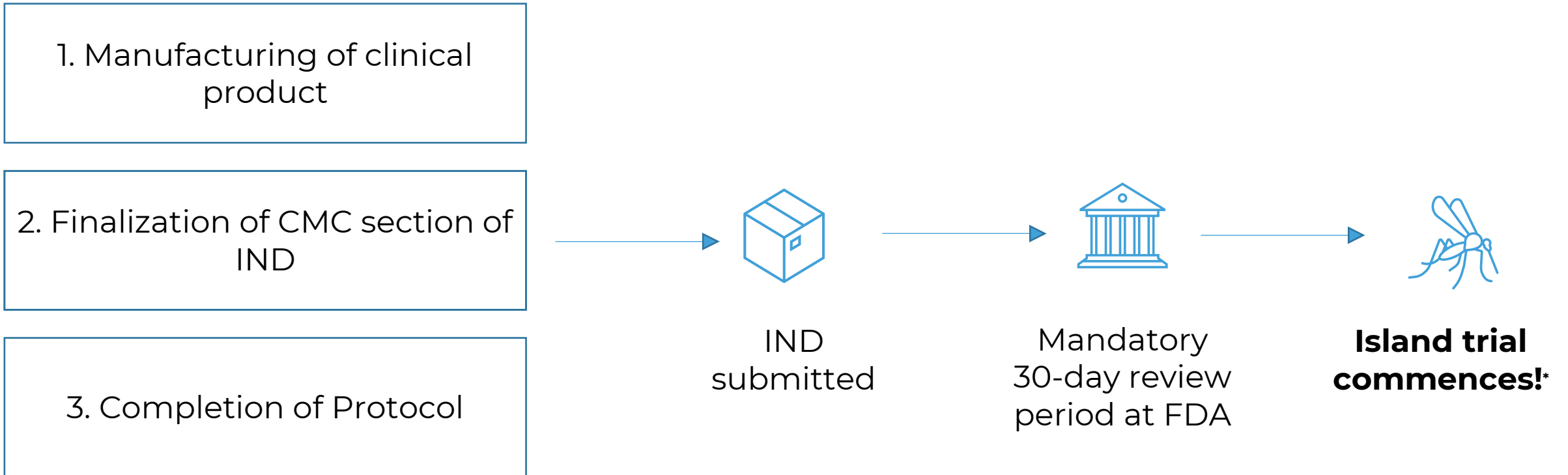
Clinical Research Organisation



Database creation and management. Clinical
trial support services. High quality CRO with
significant expertise in the dengue human
infection model (DHIM) that Island will be using
in the PEACH study.

PATHWAY / TIMING TO CLINICAL TRIALS

MULTIPLE WORKSTREAMS WILL TAKE US TO FDA REVIEW OF OUR IND



PHASE II DENGUE (PEACH) TRIAL STUDY IN DETAIL



Antiviral therapeutics

“PEACH” STUDY- A PHASE 2A, RANDOMIZED, DOUBLE BLIND, PLACEBO-CONTROLLED STUDY FOR THE **P**ROPHYLACTIC **E**XAMINATION OF AN **A**NTIVIRAL IN A DENGUE **C**HALLENGE MODEL

Phase II trial protocol

Up to 4 cohorts/4 arms

Inclusion

- Healthy subjects
- Age 18-45
- Willing to use contraception for the duration of the study
- Informed consent

Exclusion

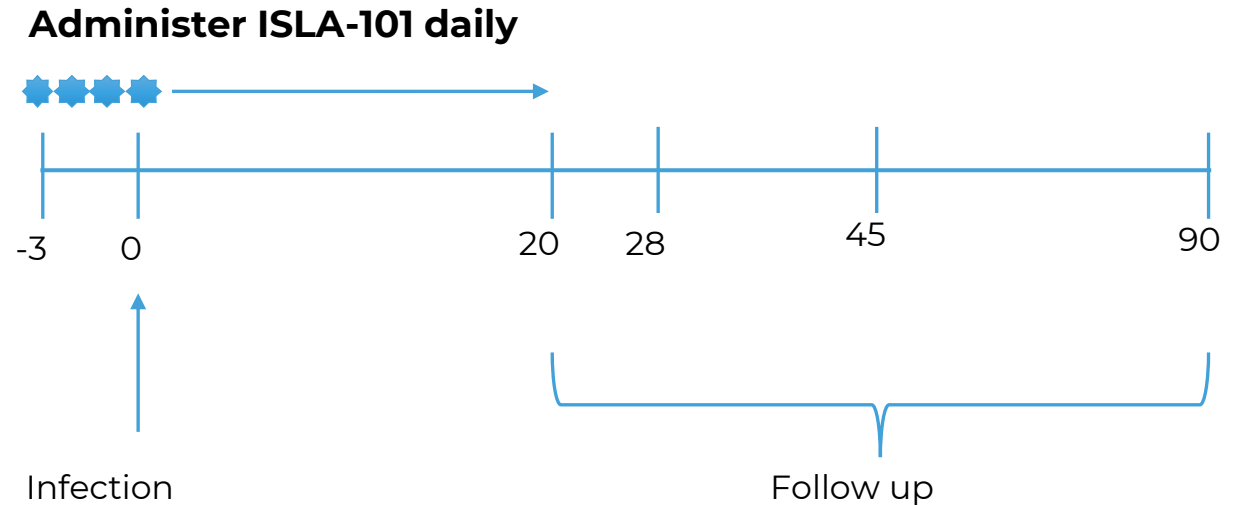
- Female: pregnant or lactating
- Prior infection with HIV, HCV, Flaviviruses
- Current, or a history of, auto-immune disease

Primary endpoint

- Assess the prophylactic effect of ISLA 101 on fever, clinical symptoms, laboratory abnormalities and viremia after challenge with DENV-1-LVHC

Secondary endpoints

- Characterise the clinical, immunologic and virologic responses following ISLA 101 after challenge with DENV-1-LVHC
- Assess the safety of ISLA 101 in the challenge with DENV-1-LVHC



The study will be run at SUNY Upstate Medical University Syracuse, New York

THE COMMERCIAL OPPORTUNITY



COMMERCIAL OPPORTUNITY



Prophylactic for travelers



Military



National outbreaks



Government Stockpiles



Priority Review Voucher



Tropical area travellers opportunity:

- Comparable to malaria market – expected to reach US\$1B in 2026*
- Increasing numbers of countries due to global warming

Military opportunity:

- Isla is partnering with army (CRADA in place) for Phase 2a clinical trial in Dengue Fever
- We will pursue a contract with the military as we get closer to approval

Endemic area opportunity:

- Many millions of patients in Central and South America
- Potential for sales for disease suppression and treatment during outbreaks
- Potential for endemic countries to establish and maintain drug stockpiles as happens with influenza

PRIORITY REVIEW VOUCHER ELIGIBILITY

- ISLA-101 is eligible for Neglected Tropical Disease designation for the treatment of dengue fever
- This designation means ISLA-101 has the opportunity to be awarded a Priority Review Voucher (PRV) from the FDA if first approved for dengue fever or Zika
- A PRV grants the holder an accelerated six month review of a drug application by the FDA
- As PRVs are transferable, they are highly valuable to drug development companies with numerous precedents for sales to biotech and pharma companies

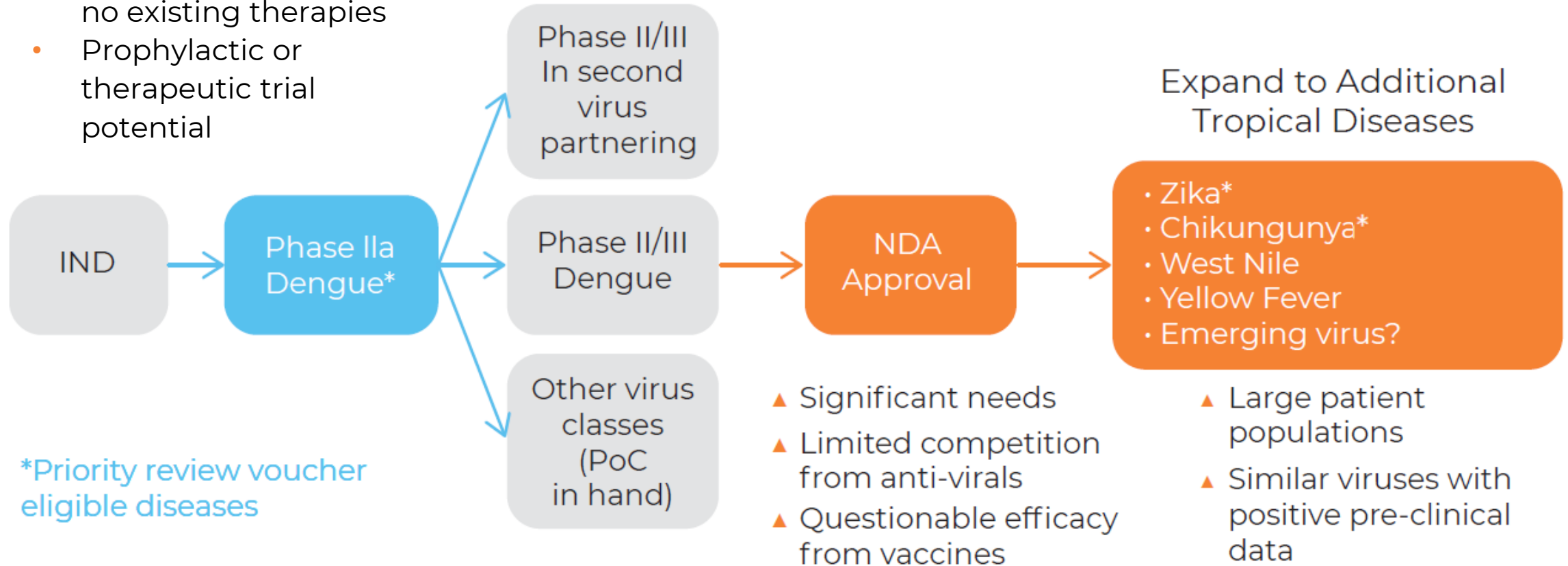
The last 10 PRV acquisitions

Date	Acquired by	Value
Q3 2019	Astra Zeneca	US\$95m
Q4 2019	Confidential	US\$95m
Q1 2020	Vifor Pharma	US\$111m
Q3 2020	Merck	US\$100m
Q4 2020	Abbvie	US\$95m
Q4 2020	United Therapeutics	US\$105m
Q1 2021	Alexion	US\$100m
Q3 2021	Undisclosed	US\$105m
Q3 2021	Undisclosed	US\$105m
Q4 2021	Undisclosed	US\$110m
Average		US\$102.1m

PLATFORM STRATEGY: MULTIPLE SHOTS ON GOAL

TARGETS

- Tropical diseases with no existing therapies
- Prophylactic or therapeutic trial potential



DRUG DEVELOPMENT PIPELINE

Program	Indication	Stage of Development				
		Preclinical	Phase I	Phase 2	Phase 3	FDA Review
ISLA-101	Dengue (PEACH)	Completed		To be initiated		
	Other mosquito (or vector) borne diseases	Completed				
Monash Collaboration	TBD					
Griffith Collaboration	TBD					

KEY COLLABORATIONS & ALLIANCES

SUPPORTING FUTURE PIPELINE DEVELOPMENT



Research Collaboration Agreement to screen thousands of known molecules against host targets building upon the Fenretinide (ISLA-101) discovery sourced from these laboratories that the Company has licensed for use against Flaviviruses



Research and development collaboration with Griffith University to screen for active anti-viral molecules in a rational repurposing strategy. The small molecule libraries for Drug Discovery (GRIDD) Compounds Australia facility, using highly sensitive assays



Research collaboration agreement signed with Australia's largest drug library containing millions of molecules that can be searched for drug re-purposing and pipeline development



Cooperative Research and Development Agreement (CRADA) with the US Army in preparation for its Phase II clinical study for ISLA-101



Supply agreement with Catalent for manufacture of Fenretinide softgels for dengue fever trial participants



Right to reference National Cancer Institute IND for Isla-101

BOARD, MANAGEMENT & SCIENTIFIC BOARD

ISLAND IS LED BY A HIGHLY CAPABLE, EXPERIENCED MANAGEMENT TEAM, BOARD OF DIRECTORS AND SCIENTIFIC ADVISORY BOARD WITH EXTENSIVE EXPERTISE IN DRUG REPURPOSING AND DEVELOPMENT, INFECTIOUS DISEASES AND EXECUTING SUCCESSFUL COMMERCIAL TRANSACTIONS.

MANAGEMENT TEAM & BOARD OF DIRECTORS



Dr. Paul MacLeman
Executive Chair



Dr. David Foster
CEO & Executive
Director



Dr. Anna Lavelle
Non-Executive
Director



Mr. Al Hansen
Non-Executive
Director



Dr. David Brookes
Non-Executive Director

SCIENTIFIC ADVISORY BOARD



Assoc. Prof. Leigh Farrell



Prof. Stephen Thomas MD



Dr. Simon Tucker

UPCOMING MILESTONES

H1 FY 2022

- ✓ Sign SUNY CTA
- ✓ Announce Principal Investigator
- ✓ Engaged CRO
- ✓ Drug substance (API) manufactured
- ✓ Advance research collaboration

H2 FY 2022

- Drug product manufactured
- File IND
- Open IND
- Screening subjects for PEACH* trial
- First subject in PEACH trial
- Advance through PEACH cohorts

H1 FY 2023

- Advance through PEACH cohorts
- Trial read out
- Meeting with FDA
- Identify lead molecules from research collaborations

* PEACH: Phase 2a, randomized, double blind, placebo-controlled study for the Prophylactic Examination of an Antiviral in a Dengue Challenge Model.

KEY STRENGTHS



Initially targeting dengue diseases with unmet need

Targeting diseases, starting with dengue fever, with a significant unmet medical need and growing economic burden



Drug repurposing strategy

Repurposing can save tens of millions of dollars and up to a decade of development time usually required to commercialise a new drug



Phase II ready asset

Lead compound, ISLA-101, has been in 45 clinical trials demonstrating an excellent safety profile in thousands of patients



Commercial upside

Potential 'platform in a pill' to treat tropical diseases. Approval of ISLA-101 by the US FDA could see company claim a Priority Review Voucher



Pipeline expansion strategy

Research collaboration agreements in place with Monash University and Griffith University to expand anti-viral pipeline beyond arbovirus indications



Highly experienced team

Experienced Board, Management Team & Scientific Advisory Board with extensive expertise in drug repurposing, infectious diseases and commercial transactions



ISLAND
PHARMACEUTICALS
Antiviral therapeutics

David Foster
Chief Executive Officer & Executive Director
david@islapharma.com

Authorized for release by Paul MacLeman, Executive Chairman

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Island Pharmaceuticals
Limited

ACN 641 183 842