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ADDRESSING URGENT VIRAL DISEASE AND AND MEDICAL COUNTERMEASURES THREATS

ASSETS

Known small molecules with clinical history
Mid/late stage clinical or other abbreviated routes
Open USFDA INDs
Validated US Govt/military funding support

RAPID PATHS TO MARKET

Priority Review Voucher eligible

STRATEGIC CHECKLIST



- Clinical history
- ☐ Small molecule
- Proven preclinical anti-viral activity
- Eligible for Priority Review Vouchers
- National and military preparedness need
- Probable/confirmed non-dilutive funding to support: military, civilian, NGO
- ☐ Fit for our management and advisory team: mid/late stage drug development, extensive medical countermeasures experience and roles



STRATEGIC CONNECTIVITY BETWEEN ASSETS ISLA101 (FENRETINIDE) AND GALIDESIVIR



Both assets are:

- Phase II or later
- Have open Investigational New Drug applications with FDA
- Antivirals, therefore fast to develop and potential for early licensing if sought
- Supported by governments and military; needed for outbreaks and medical countermeasures
- Priority Review Voucher eligible (multiple options)
- Capable of multiple 'shots on goal': each has many potential disease targets



"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

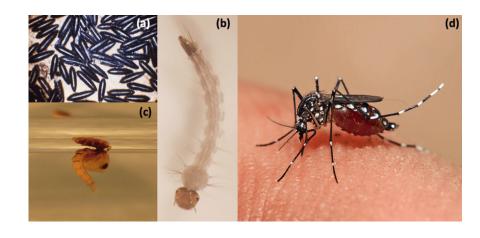


TARGETING KNOWN OR RE-PURPOSABLE DRUGS WITH REDUCED TIME TO MARKET



ISLA-101 – dengue, Zika, Yellow Fever, Chikungunya, other arboviruses

- 45+ human trials of ISLA-101 completed in other indications
- Island's recent SAD study reinforced safety and established the effective human EC50 dose
- Infectious disease, so trial endpoints are rapidly achieved
- Multiple PRV options
- CRADA with US military
- Entering Phase II now (Jul24)



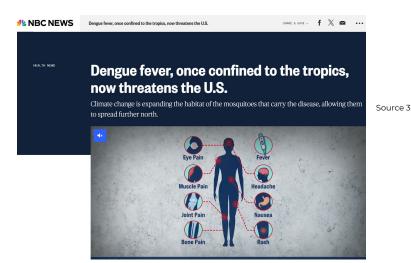
Single Ascending Dose study showed 2-3x bioavailability in fed subjects

Now exploring combined prophylactic / therapeutic Phase II study with FDA.

DENGUE IS CLOSE TO HOME









National



Environment

Not only is the virus causing people to fall extremely ill, in some instances, travel insurance costs can soar into the tens of thousands



World

Source 2



France warns of surge in imported dengue cases ahead of Olympics

Source 4

(Described as potential super-spreader event)

Article sources:

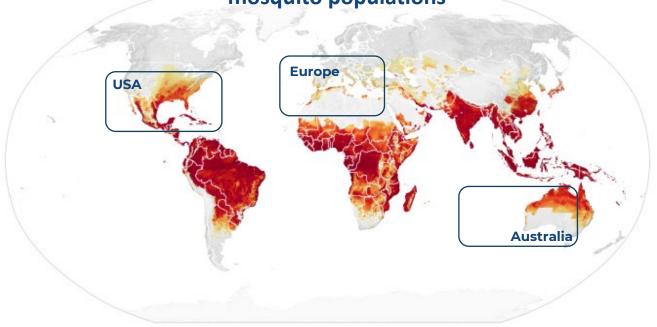
- https://www.news.com.au/travel/travel-updates/travel-stories/deadly-viral-infection-doubles-in-popular-tourists-spotsfor-aussies/news-story/fc32fa4fc37eeled4ca48a4deb44f5lc
- https://shorturl.at/NsvwY
- https://www.nbcnews.com/health/health-news/dengue-fever-climate-change-mosquitos-tropical-disease-rcna149366
- https://www.euronews.com/health/2024/04/24/france-warns-of-surge-in-imported-dengue-cases-ahead-of-olympics

ISLAND PHARMACEUTICALS **JULY 2024**

DENGUE IN 2050 – A GLOBAL DISEASE



WHO prediction based on projections of future temperatures, rainfall, and mosquito populations



Projected Environmental Suitability for Dengue in 2050

0% 50% 100%

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

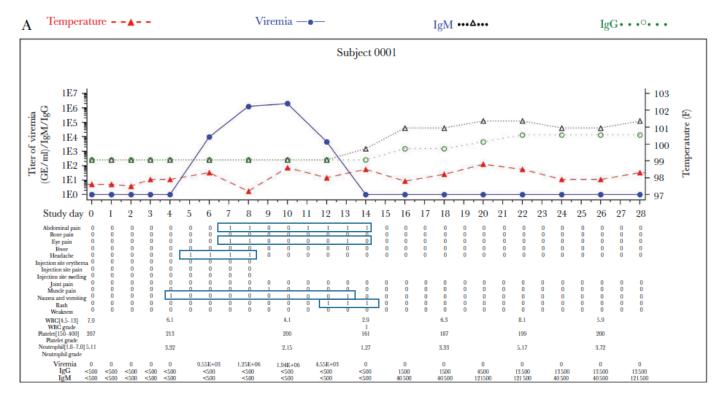
US ARMY CRADA PROVIDES CONTROL GROUP FOR PHASE 🎎 II STUDY (AND SUBSTANTIAL % OF PHASE II TRIAL COSTS)



P1 challenge study conducted by Walter Reed Army Institute of Research and SUNY is ILA's P2 study control data, and enables unprecedented ability to monitor dengue symptoms, including:

- Abdominal & eye pain
- Fever
- Headache
- Nausea and vomiting
- Rash

Island will use the same attenuated virus and approach in its current Phase 2 study.



Endy et al, J Inf Dis 2021

TARGETING KNOWN OR RE-PURPOSABLE DRUGS WITH REDUCED TIME TO MARKET



GALIDESIVIR – Ebola, Marburg, Zika & and most other RNA viruses

- 12 month, non-binding term sheet signed 3 July 2024 (AU)
- Phase I human safety data
- Proven protective efficacy in a number of lethal animal models
- If proceed to full option agreement, would be seeking to confirm continued access to FDA's **Animal Rule**
- Extensive US government funding to date



PRV eligible (multiple options)

COMMERCIAL OPPORTUNITIES



BOTH candidates have prophylactic and therapeutic potential







Military



National Outbreaks

Malaria is also a mosquito borne disease and therefore a proxy for other tropical diseases. Market for antimalarials is expected to reach US\$1B1 by 2026

1. https://www.alliedmarketresearch.com/anti-malarial-drua-market

Relationship with US Army in place (CRADA, ISLA101). Will continue discussions as programmes advance.

Proven interest in both assets.

Millions of patients in Latin
America & Asia offer potential
for sales in disease
suppression and treatment
during outbreaks of dengue,
Zika. Increasing rapidly
elsewhere.



Potential for countries to establish civilian and military drug stockpiles as happens with influenza.

Priority Review Vouchers

Last ten PRVs sold for an average of US~\$110M, with potential for ISLA-101 and galidesivir at the time of first FDA approval

FDA approval

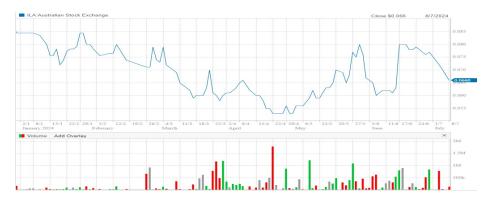
CORPORATE SNAPSHOT



Key data

Share price (AUD¹)	\$0.066
Market cap ¹	\$8.37m
Shares on issue ¹	126,767,093
Listed Options ²	32,506,360
Cash ³	\$1.6m

Recent ILA trading history



Board of Directors

Dr Paul MacLeman, Executive Chairman	
Dr David Foster, CEO and Managing Director	
Dr David Brookes, Non-Executive Director	
Mr Albert Hansen, Non-Executive Director	
Dr Anna Lavelle, Non-Executive Director	

Substantial shareholders

Shareholder	Ownership ⁴
Mr Jason Alan Carroll	24,100,000 (19.01%)
Dr William James Garner	22,056,105 (17.40%)
Albert Hansen / Kesa Partners	11,104,034 (8.68%)

Ownership breakdown

- Top 20¹: 72.3%
- Board and management¹: 13.3%

DoD grant funding

USD \$625k to directly support the Phase 2

^{1.} As at 8 July 2024 | 2. <u>ILAO Option terms:</u> Exercise price of \$0.06 expire 14 March 2025. | 3. As at 31 March 2024 – does not take into consideration cash burn since March 2024 or cash received from options exercised | 4. Shares held per ASX lodged substantial holder notices | Share price data sourced from asx.com.au

UPCOMING MILESTONES**



H1 FY 2024 (Jul - Dec 2023)

H2 FY 2024 (Jan - Jun 2024)

H1 FY 2025 (Jul – Dec 2024)

H2 FY 2025 (Jan - Jun 2025)

- Key US and Australian patents granted for ISLA-
- Obtain HREC (i.e. IRB) approval
- Screen, enrol and dose volunteers in SAD study
- Successful progression through P1 cohorts

- Dose final Single
 Ascending Dose
 study subject
- Single Ascending

 Dose study read out
- FDA interaction on Phase 2 study protocol
- Pipeline expansion efforts
- Complete in silico modelling of multiple dosing regimen

- Obtain ethics approval for Phase 2 trial
- Initiate Phase 2 trial
- First patient dosed in Phase 2 trial
- Advances through Phase 2 study cohorts
- Ongoing DD on Galidesivir program
- Ongoing discussions with potential partners

- Data readout from Phase 2 study
- End of Phase 2 meeting anticipated with FDA
- Plans announced for next steps in clinical program
- Progress Ebola program
- Ongoing discussions with potential partners

^{**} Dates are indicative only, based on best estimates at the time of writing; subject to change.



SCIENTIFIC ADVISORY BOARD





Dr Leigh Farrell

Leigh has over 30 years' experience in the biotechnology and pharmaceutical industry and is Head of Health Security Systems Australia, a Division of DMTC Ltd, is a non-executive director of Pro Medicus Ltd, Ena Respiratory Pty Ltd and Axelia Oncology ty Ltd, and is a member of the Walter and Eliza Hall Institute of Medical Research Board Commercialisation Committee and a member of the Independent Advisory Council of Medicines Australia.

Leigh's past appointments include: Senior Vice
President, Commercial at Certara USA, Inc where he
was responsible for Asia Pacific Commercial and
global government engagement for the
preparedness, planning and response to major
health emergencies; Chairman & COO of d3
Medicine, LLC; Vice President of Business
Development at Biota Pharmaceuticals Ltd,
Research Manager Johnson & Johnson Research
and CEO of Gene Shears Pty Ltd. Leigh holds a PhD
in Biochemistry from Monash University.



Prof Stephen Thomas MD

US WRAIR – SUNY Upstate
Professor Stephen Thomas, MD has an international
leadership role as Lead Principal Investigator for
Pfizer/BioNTech global Phase III COVID-19 vaccine
trial now being deployed globally.

Prof. Thomas is a world-renowned virologist and vaccinologist and has authored numerous papers and articles on dengue fever, Zika and many other infectious diseases.

Chief, Division Of Infectious Diseases, New York
Upstate Medical University; Professor of Medicine,
Professor of Microbiology & Immunology, and
Infectious Diseases physician-scientist from the
State University of New York (SUNY), Upstate
Medical University; Chief, Division of Infectious
Diseases and Director, Institute for Global Health
and Translational Science (IGHTS.)

He had twenty years in the U.S. Army Medical Corps serving at the Walter Reed Army Institute of Research (WRAIR.)



Dr Amy Patick

BMS – Mayo Clinic
Amy Patick is a scientific consultant with deep expertise in antiviral drug discovery, development and viral resistance with broad know how in emerging virus epidemics and translational medicine.

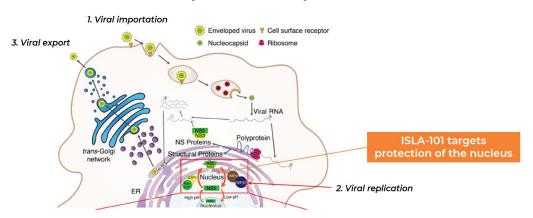
Previously, Dr. Patick has served as Vice President, Research at Adamas Pharmaceuticals, Vice President, Biological Sciences at Genelabs Technologies, Head of the Antiviral Biology Therapeutic Area at Pfizer, Inc. and Research Scientist at Bristol-Myers Squibb Company. Dr. Patick has also served as President for the International Society of Antiviral Research.

Dr. Patick was a postdoctoral fellow in immunology at the Mayo Clinic/Foundation in Rochester, MN and received her PhD in Medical Microbiology from the University of Wisconsin, Madison.

ISLA-101 PREVENTS VIRAL REPLICATION



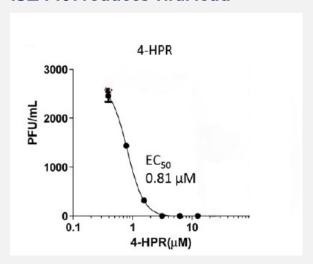
Mechanism of action (how it works)



ISLA-101 inhibits propagation of flaviviruses

- To replicate, the virus needs to hijack the nucleus of the host cell
- · Studies demonstrated ISLA-101 prevents this, therefore preventing virus replication
- Same mechanism of action for a therapeutic or prophylactic either before or after exposure to the virus

ISLA-101 reduces viral load¹



Above: dose response showing ISLA-101's ability to protect against dengue infection

- In freshly isolated human cells, ISLA-101 was shown to potently reduce viral infection with a sub micromolar EC₅₀
- Island's SAD study was designed to also investigate the ability to achieve appropriate blood concentrations in healthy human volunteers (slide 16)

1. Fraser et al. J. Infect. Dis 2014

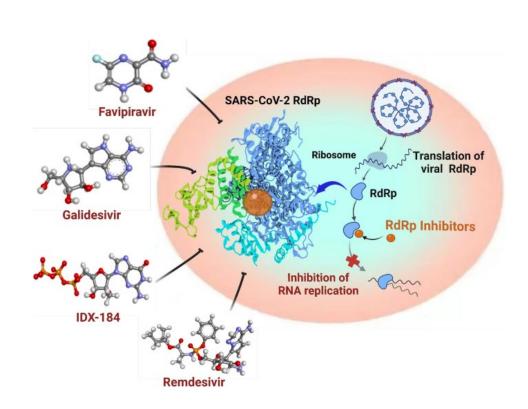
GALIDESIVIR PREVENTS VIRAL REPLICATION



Galidesivir inhibits propagation of RNA viruses

- RNA-dependent RNA polymerase (RdRp) is crucial in viral replication and transcription
- Catalyses viral RNA synthesis and is the therapeutic target of

https://link.springer.com/article/10.1007/s41061-023-00432-x



PLAYERS IN THE MEDICAL COUNTERMEASURES SPACE



2020

Programs backed by BARDA and/or NIAID

- BioFactura
- BioFire Defense
- Certara
- Emergent BioSolutions
- Evotec SE
- IDbyDNA (DIGET program)
- Juxtopia
- DARPA Panacea Program
- ARMR
- JnJ Blue Knight Consortium



https://globalbiodefense.com