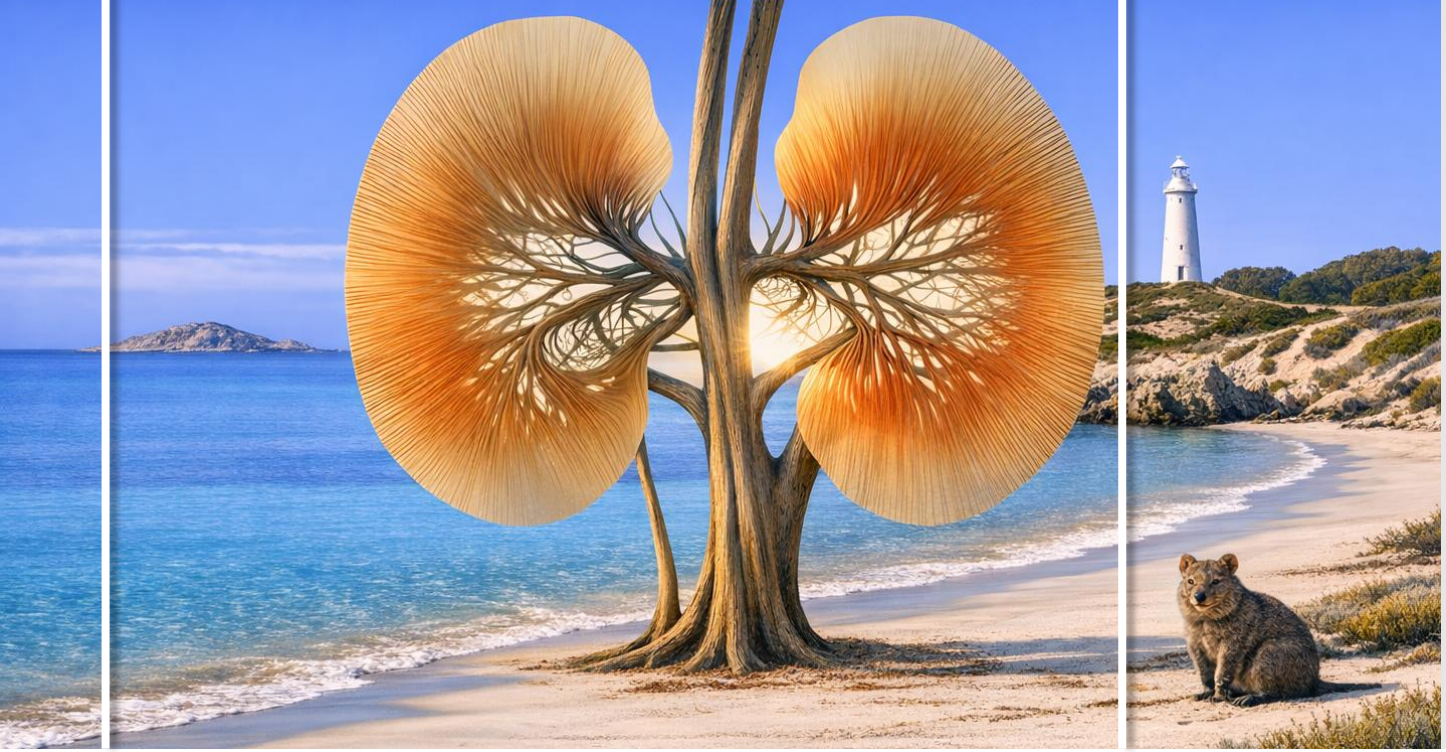


Dimerix

(ASX:DXB)

Developing new therapies to treat inflammatory causes of kidney disease with unmet clinical needs

Euroz Hartleys Institutional Conference

Rottnest Island – 17-19 March 2026

Authorised for lodgement by the Board of the Company

Forward looking statements

This presentation includes forward-looking statements that are subject to risks and uncertainties. Although we believe that the expectations reflected in the forward looking statements are reasonable at this time, Dimerix can give no assurance that these expectations will prove to be correct. Readers are cautioned not to place undue reliance on forward-looking statements.

Actual results could differ materially from those anticipated. Reasons may include risks associated with drug development and manufacture, risks inherent in the regulatory processes, delays in clinical trials, results of clinical trials, contractual risks, risks associated with patent protection, future capital needs or other general risks or factors, including but not limited to those factors outlined in the most recent Dimerix Limited Annual Report.

Overview | Phase 3 Global Opportunity

Phase 3 trial **recruitment complete** in trial of DMX-200 in focal segmental glomerulosclerosis (FSGS)

Reduced risk Proteinuria endpoint **passed blinded interim** (futility) assessment (March 2024)[1]

FSGS indication is a **rare disease** that causes scarring of the kidney, leading to irreversible damage[2]

No approved treatments specifically for FSGS: damage can lead to **dialysis, transplant or death**[2]

Orphan drug designations regulatory, marketing exclusivity and pricing **benefits** in key territories[3]

4 commercial partners **DMX-200 licensed** in USA, Europe, Canada, Australia, NZ, Japan and GCC[4]

up to $1.4 billion in total development and sales milestone payments **plus** royalties[3]

Dimerix

1. ASX release ASX release 11 March 2024 2. Guruswamy Sangameswaran KD, Baradhi KM. Focal Segmental Glomerulosclerosis (July 2021), online: https://www.ncbi.nlm.nih.gov/books/NBK532272/; 3. ASX releases: 14 December 2015, 21 November 2018, 07 June 2021; 4. ASX release 05 October 2023, 27 May 2024, 07 January 2025 and 01 May 2025

Cycle of damage : in glomerular diseases

What is FSGS?

Focal	**= some**
Segmental	**= sections**
Glomerulo	**= of the kidney filtering units**
Sclerosis	**= are scarred**

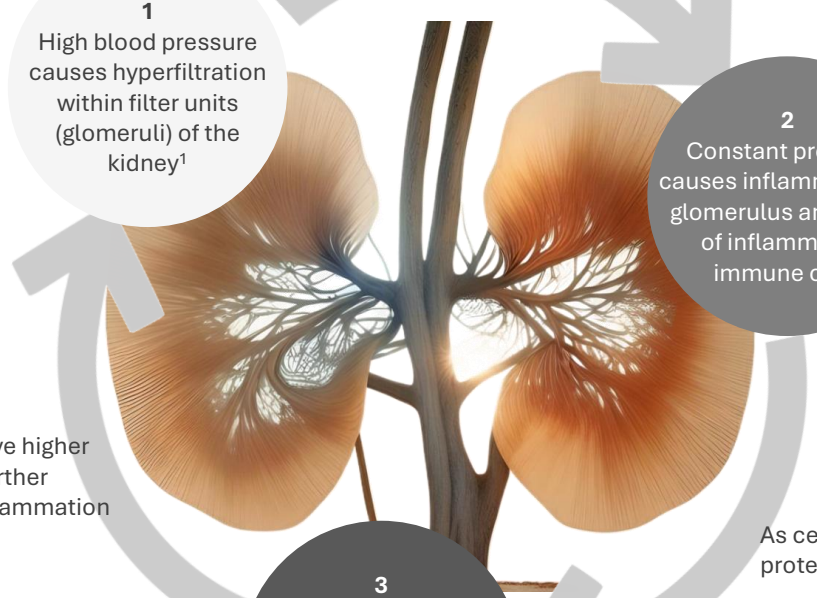
Existing blood pressure medication

DMX-200

Glomeruli exposed to stress from high blood pressure

1
High blood pressure causes hyperfiltration within filter units (glomeruli) of the kidney[1]

2
Constant pressure causes inflammation of glomerulus and influx of inflammatory immune cells



Fewer kidney cells drive higher blood pressure and further hyperfiltration and inflammation

As cells die, glomerular become scarred and protein leaks into the urine (**proteinuria**)

3
Pro-inflammatory environment drives sclerosis and fibrosis that is permanent and non-reversable

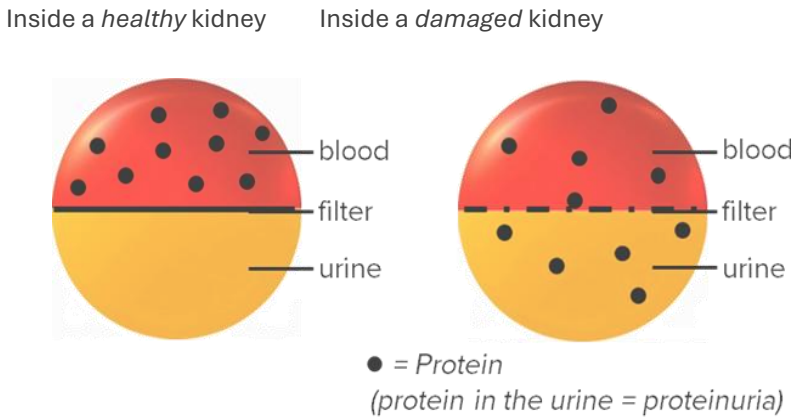
Dimerix

1. Lewis, E. J. et al. (2001), New Engl J Medicine 345, 851–860

Interpreting proteinuria as a surrogate endpoint

Proteinuria is the quantity of protein in the urine

A healthy kidney is a good filter and allows little to no protein into the urine[1]



- When kidneys are damaged, protein can leak into the urine causing proteinuria
- Proteinuria represents an important early marker of kidney function[2]

Proteinuria as a predictor of kidney disease[3]

Dip test	*uPCR g/g*
Neg	<0.13
Trace	0.13
+	0.62
	0.88
++	1.24
+++	2.30
++++	>3.09

NORMAL
MICROALBUMINURIA
HEAVY ALBUMINURIA

normal
proteinuric
nephrotic

Minimal uPCR for ACTION3 inclusion

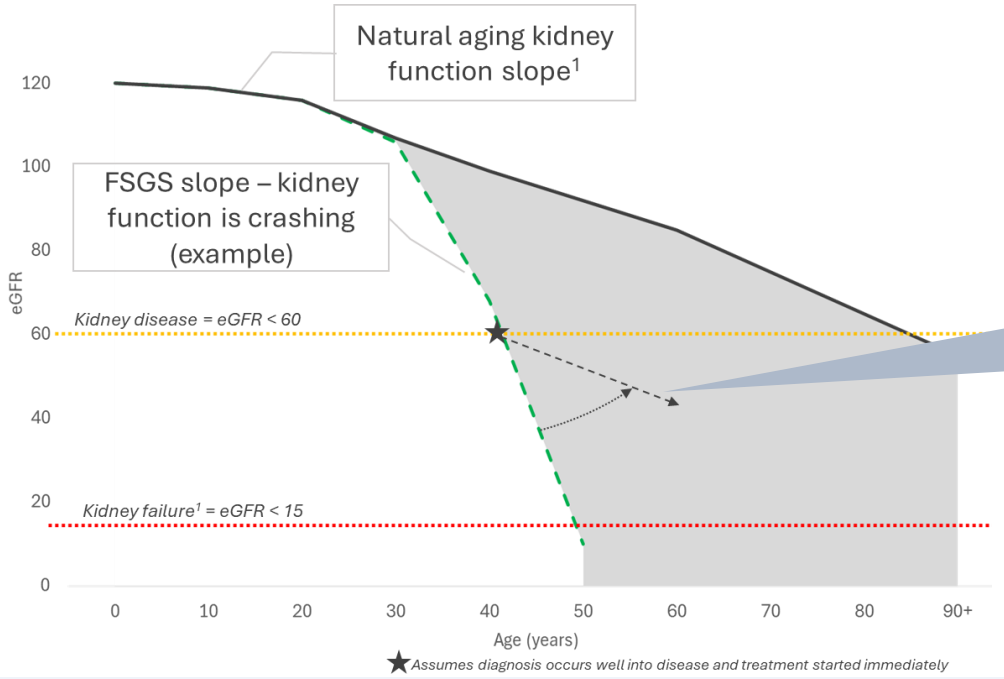
Proteinuria is typically less variable and easier to measure than eGFR[4]

Dimerix

1.Guruswamy Sangameswaran KD, Baradhi KM. Focal Segmental Glomerulosclerosis (July 2021), online: https://www.ncbi.nlm.nih.gov/books/NBK532272/; 2. Nephcure FSGS living with the disease (2024) at https://nephcure.org/livingwithkidneydisease/ns-and-other-glomerular-diseases/understanding-fsgs/ ; 3. Adapted from graphics prepared by Renal Unit at the Royal Infirmary of Edinburgh and the University of Edinburgh; 4. PARASOL outcomes 2024, a working group made up of: NephCure, ISGD, NKF, KHI are co-sponsors of the project, FDA and EMA involvement, University of Michigan is data coordinating center, Industry invited as participants

Measuring eGFR as a surrogate endpoint

Estimated glomerular filtration rate (eGFR) is the measurement of the kidney filtration rate



Treatments, such as DMX-200, aim to bring the FSGS slope back up:

- can add years to the life of the kidney
- potential to delay dialysis and/or kidney transplant

- Kidney function is measured using the estimated rate of blood filtered by the kidney per minute (millilitres per minute)
- eGFR slope naturally declines as we age[1]
- In FSGS patients, kidney function is decreasing rapidly

1. National Kidney Foundation: Estimated Glomerular Filtration Rate (eGFR): https://www.kidney.org/atoz/content/gfr

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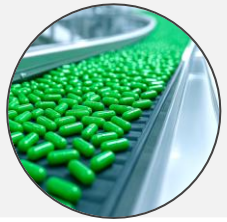
DMX-200 – inflammatory modulator

A CCR2 inhibitor working synergistically alongside the current standard of care (AT1R blocker): G protein-coupled receptor (GPCR)

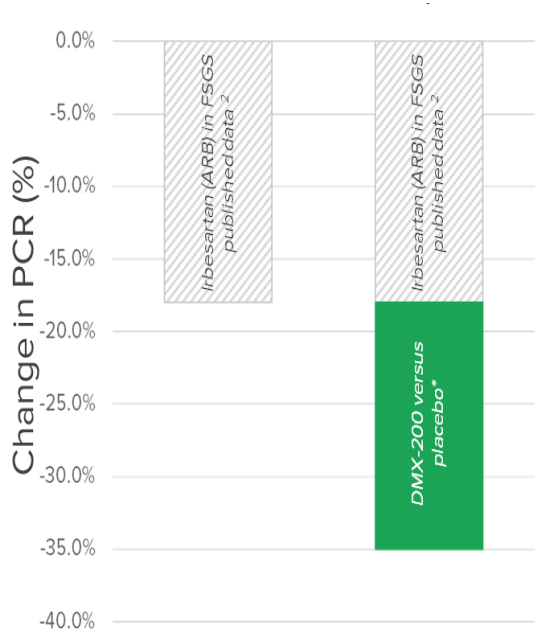
Commercial manufacturing sites established in USA[3]	1 x 120mg capsule BID	Well tolerated with no safety concerns to date (>400 patients dosed)[3]	4 clinical studies completed to date: positive safety and efficacy signals across studies[3]
			
Small molecule	**Easy & convenient dosing**	**No safety concerns noted[3]**	**Positive efficacy signals[3]**

1. ASX release and investor presentation 29Jul20 and ASX release 11Mar24 BID: bis in die/ twice a day

DMX-200: Phase 2 met primary endpoint

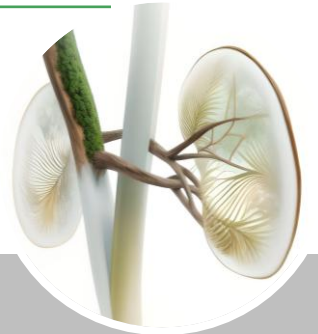
Clinically encouraging outcomes achieved for patients,[1,2] with no safety concerns noted[3]

Average reduction of **17%** in proteinuria after 16 weeks treatment on DMX-200 versus placebo[3]



"Any reduction in proteinuria could yield years of preserved native kidney function and delay the onset of kidney failure and its attendant morbidity and mortality"

Kidney survival study – Troost et al, August 2020[2]



EFFICACY

- **86%** of patients demonstrated reduced proteinuria
- DMX-200 reduced inflammatory biomarker by **39%** vs placebo

SAFETY

- No safety concerns – reduced development risk

Dimerix

PCR = protein creatinine ratio; ARB = angiotensin receptor blocker; 1. Trachtman, et al., 2018. J Amer Soc Nephrology 29(11):2745-2754; 2. Troost JP et al (August 2020); doi.org/10.1053/j.ajkd.2020.04.014; 3. Repeated measures mixed model analysis per protocol; top line data was reported as grouped analysis on 29 July 2020, study not designed for statistical significance;

ACTION3 phase 3 clinical trial

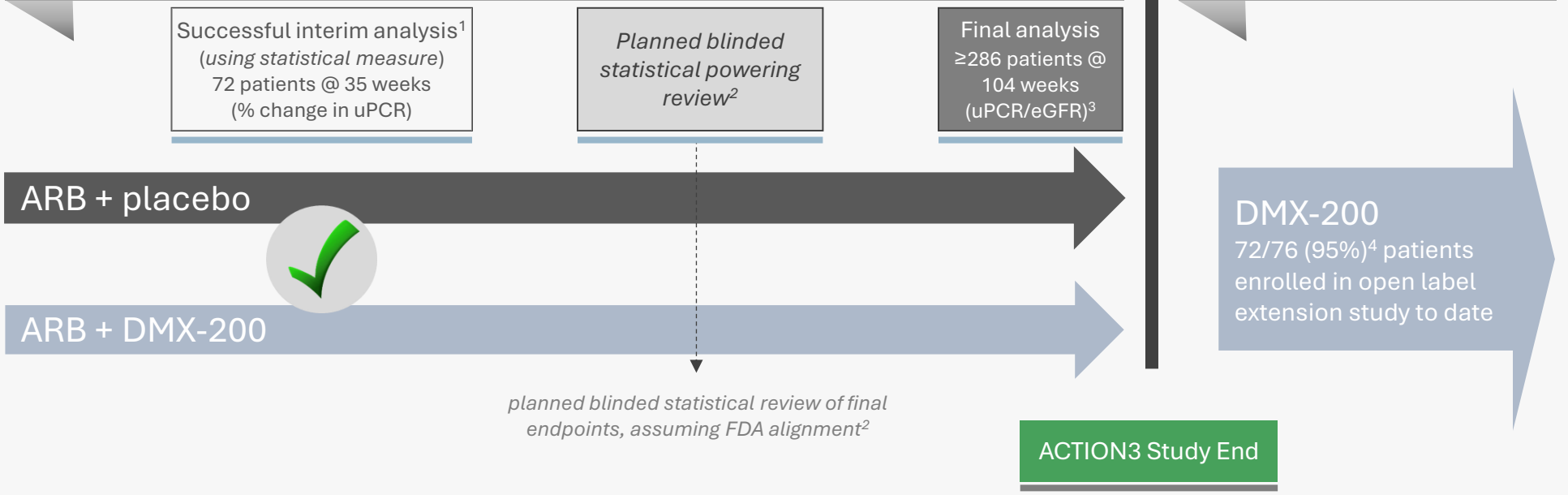
FSGS CLINICAL STUDY

A randomised, double-blind, multi-centre, placebo-controlled study of renal outcomes of DMX-200 in patients with FSGS receiving an ARB (n=≥286)

Background

- Patients recruited, then screened and stabilised on background medications
- Patients randomised to receive drug or placebo
- DXB remains blinded at all times during study

Phase 3 Trial Timeline



Open Label Extension

DMX-200

72/76 (95%)[4] patients enrolled in open label extension study to date

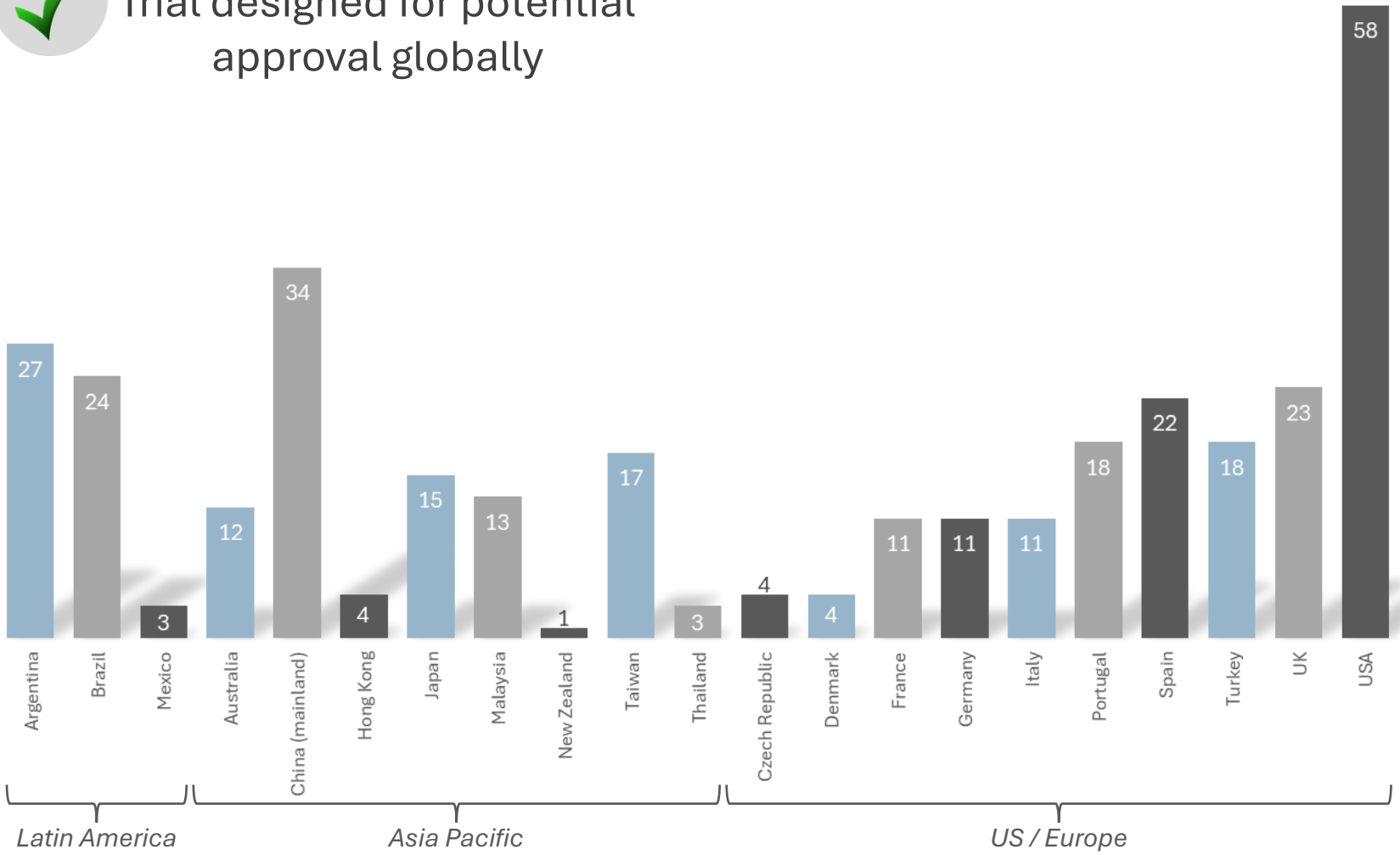
ACTION3 Study End

Dimerix

1. Blinded interim Phase 3 analysis data does not guarantee a statistically significant outcome at the end of the trial, ASX release 11 March 2024; 2. The potential for accelerated (or conditional) approval submissions, following the second blinded review and any potential unblinding, will be assessed based on discussions with the appropriate regulatory authorities such as the FDA in the US and blinded statistical powering outcomes; 3. Regardless of any accelerated (conditional) approval potential, ACTION3 study will complete full 2 year analysis and regulatory submission for potential traditional (full) approval; 4. number and % of eligible patients who have completed 2 years treatment as at 13 March 2026; ARB = angiotensin receptor blocker; uPCR = urinary proteinuria; eGFR = estimated glomerular filtration rate (kidney function);

Adult patient recruitment by territory

Trial designed for potential approval globally



Recruitment completed

(adult population)[1]

333

Adult patients recruited, randomised and dosed (target ≥286)[2]

6

Paediatric patients recruited, randomised and dosed[2]

1. ASX release 15 December 2025; 2. Final numbers of adult patients, ASX release 10 March 2026; paediatric patients will continue to recruit, and will not impact final analysis timelines

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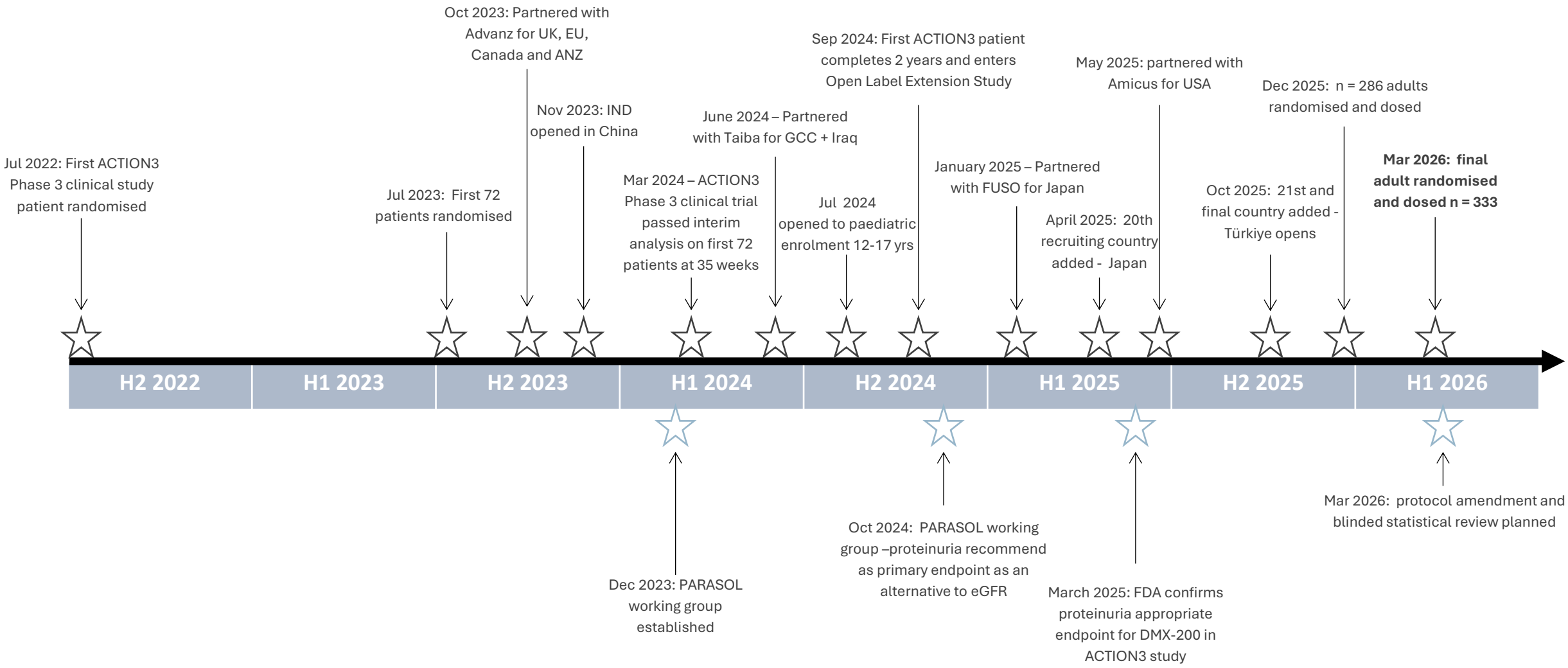
Evolution of FSGS landscape & ACTION3 study



ACTION3 next steps

FSGS CLINICAL STUDY

PARASOL working group conducted “ACTION3-like” population analysis of larger PARASOL observation dataset[1]

- Results of this analysis are generally consistent with the broader PARASOL analysis conducted in 2024
- Potential relationship between proteinuria at 12 months and subsequent risk of kidney failure observed that may support proteinuria as a more powerful endpoint alternative to eGFR to detect a treatment benefit of new therapies, such as DMX-200

Step 1

PARASOL FSGS working group 12-month data analysis received[1]

Step 2

Seek FDA alignment on endpoints and process for blinded review, ACTION3 clinical study protocol updates[2]

Assumes FDA aligned with endpoints and review process

Step 3

Blinded review to assess endpoint power

~late March 2026[2]

Timing of analysis based on protocol update in each territory

Step 4

Decision on next steps towards marketing approval submission, in line with commercial partners[2,3]

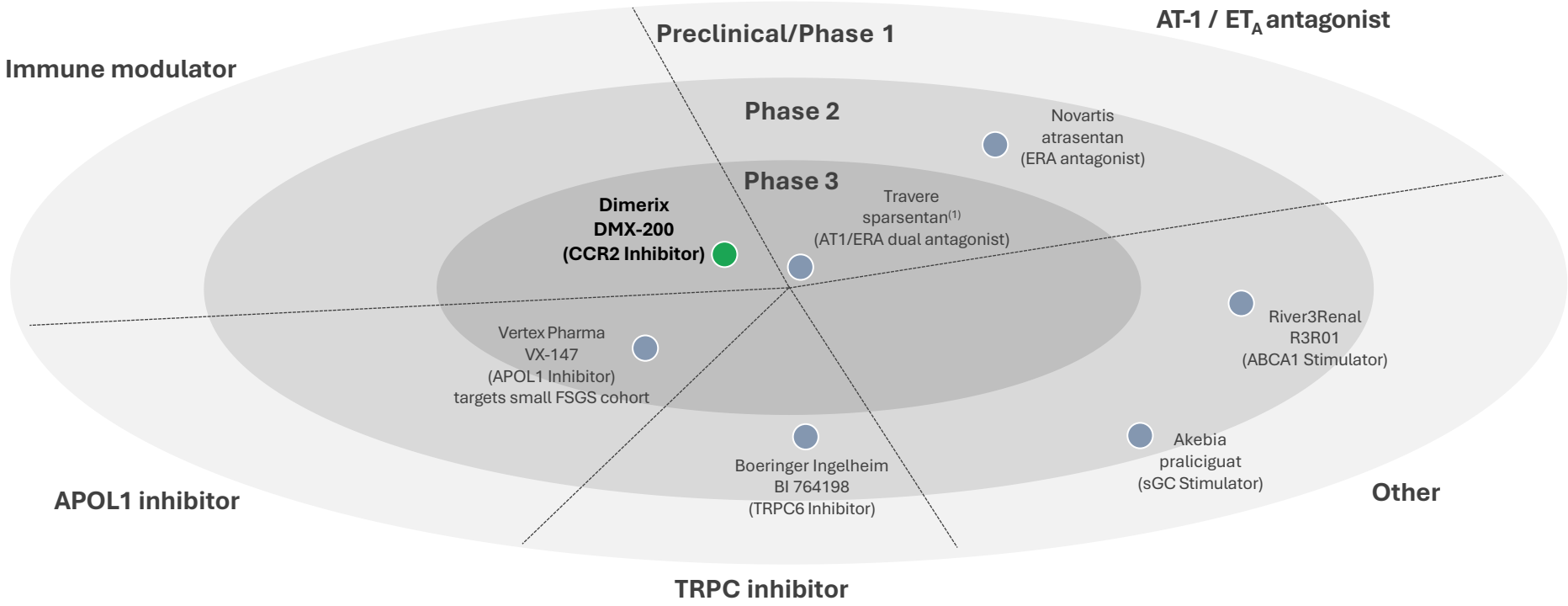
Regardless of outcome, ACTION3 study will complete full 2 year analysis for regulatory submission for potential traditional (full) approval

Dimerix

1. ASX release 08 October 2025; 2. ASX release 16 March 2026 ; 3. The potential for unblinding and any accelerated (or conditional) approval submissions will be assessed based on further discussion and agreement with the appropriate regulatory authorities such as the FDA in the US

Competitive/complementary landscape in FSGS

- Low competition in inflammatory treatment options, large unmet medical need
- DMX-200 is the only inflammatory modulator in development
- DMX-200 has potential for use in conjunction with other drugs in development if approved



Source: Company information and clinicaltrials.gov

Rare kidney disease – a potential growth market

Biopsy

FSGS diagnosis driven by rates of biopsy - growth potential as biopsy rates increase

7 per 1,000,000

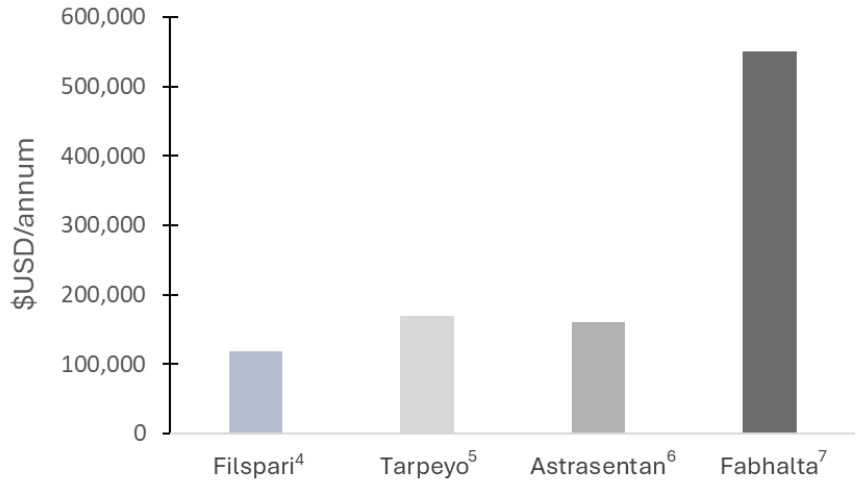
Global incidence rate of FSGS per capita, per year[1]

FSGS is the most frequent primary glomerular disease that reaches end-stage renal failure in the US[2]

DMX-200

Commercial manufacturing sites established in USA[3]

Example pricing: USA retail price for IgA Nephropathy products



Example price for other rare kidney disease drugs per patient:

- *in the US (i.e. Kinpeygo/Tarpeyo in IgAN)[8]:* ***US$15,123 per month***
- *in the UK (Kinpeygo/Tarpeyo in IgAN)[9]:* ***US$8,797 per month***

Other key territories, including Middle East and China, use US and/or Europe as pricing reference[10]

1. Kitiyakara et al Trends in the epidemiology of focal segmental glomerulosclerosis. Semin Nephrol. 2003 Mar;23(2):172-82. doi: 10.1053/snep.2003.50025. PMID: 12704577; 2. The United States Renal Data System (USRD), 2023 Annual Report, End Stage Renal Disease; 3. ASX investor presentation 09Mar20; 4. https://www.reuters.com/business/healthcare-pharmaceuticals/us-fda-approves-travere-therapeutics-kidney-disorder-drug-2023-02-17/; 5. http://www.vanrafia.com; 6. Ito, et al., 2025. https://doi.org/10.1182/blood.2024025176; 7. https://endpts.com/fda-clears-traveres-rare-kidney-disease-drug-will-come-with-rems-program; 8. Micromedex Red Book (2025) Comprehensive drug pricing and product information; 9. UK list price (2025) https://www.nice.org.uk/guidance/TA937/chapter/2-information-about-targeted-release-budesonide; 10. J Mark Access Health Policy. 2016 Mar 15;4:10.3402/jmahp.v4.30458. doi: 10.3402/jmahp.v4.30458

Summary of licensing deals for DMX-200 to date

Dimerix has successfully partnered DMX-200 across key markets

	1 Amicus Therapeutics	2 ADVANZ PHARMA	3 taiba	4 FUSO
	United States	EU, CA, AU, NZ	GCC, Iraq	Japan
Upfront	AU$48 million[1]	AU$10.8 million[2]	AU$0.5 million[3]	AU$7.2 million[4]
	Upfront	Upfront	Upfront	Upfront + 1st milestone
Milestones	Up to AU$892 million[1]	Up to AU$219 million[2]	Up to AU$120 million[3]	Up to AU$100 million[4]
	Milestones	Milestones	Milestones	Milestones
Royalties	Escalating low-teen-low twenties%	Escalating mid-teen-20%	Starting at 30%	Between 15-20%
	Royalties	Royalties	Royalties	Royalties

Licensing deals collectively valued up to

~AU$1.4 billion

in total upfront and potential milestone fees plus royalties[1]

Over

AU$65 million

in total payments received

Significant potential additional global **deal value remains**, as Dimerix pursues and progresses **licensing opportunities** with potential partners outside the licensed territories

1. ASX release 01 May 2025; 2. Based on Euro conversions & further terms outlined in ASX Announcement on 5 October 2023; 3. Based on US dollar conversions & further terms outlined in ASX Announcement on 27 May 2024; 4. Based on Japanese ¥ Yen conversions & further terms outlined in ASX Announcement on 7 January 2025

DMX-200 substantially de-risked Phase 3 renal asset

ACTION3 primary endpoint: proteinuria has potential for stronger statistical power to resolve DMX-200 effect[1]

Proteinuria endpoint reduced risk: passed blinded interim (futility) assessment (March 2024)[2]

Commercial validation: 4 commercial partners across key territories[3]

Study fully recruited: fixed timelines to full study completion in Q1 2028[4]

Passed 7 safety data monitoring meetings with no protocol changes requested[5]

Strong uptake into open label extension study: indicates patient willingness to continue treatment

Indication extension: DMX-200 proposed mechanism of action potentially suited to other inflammatory renal indications

Dimerix

1. ASX release 24 Dec 2025; 2. ASX release ASX release 11 March 2024 ; 3. ASX release 01 May 2025; 4. ASX release 15 Dec 2025; 5. ASX release 1 Nov 2025

Growth strategy

Deliver ACTION3 Phase 3 clinical trial

- Ensure drug supply continuity and patient visits for recruited patients
- Complete recruitment of paediatric patients
- Maintain regulatory engagement (FDA, EMA, PMDA, NMPA + others)
- With partners, prepare for potential market approval and launch readiness

Expand global commercial partnerships

- Build on existing licensing agreements and relationships
- Secure additional partnerships to expand and accelerate market access

Advance pipeline development

- Identify and progress new assets in renal and/or rare disease indications
- Leverage DMX-200 platform for additional indications

Grow sustainable shareholder value through clinical success, global partnerships, and pipeline diversification

Corporate overview

Ticker Symbol	ASX: DXB
Cash Balance (Dec25)	$38.5 million
Market Capitalisation[1]	$220 million
Share price[1]	$0.35
Total ordinary shares on issue[1]	600,396,776
Average Daily Liquidity by value for past 30 trading days[2]	$0.65 million

Research Coverage	Analyst
EUROZ HARTLEYS	Seth Lizee
PETRA CAPITAL	Tanushree Jain

SHARE PRICE

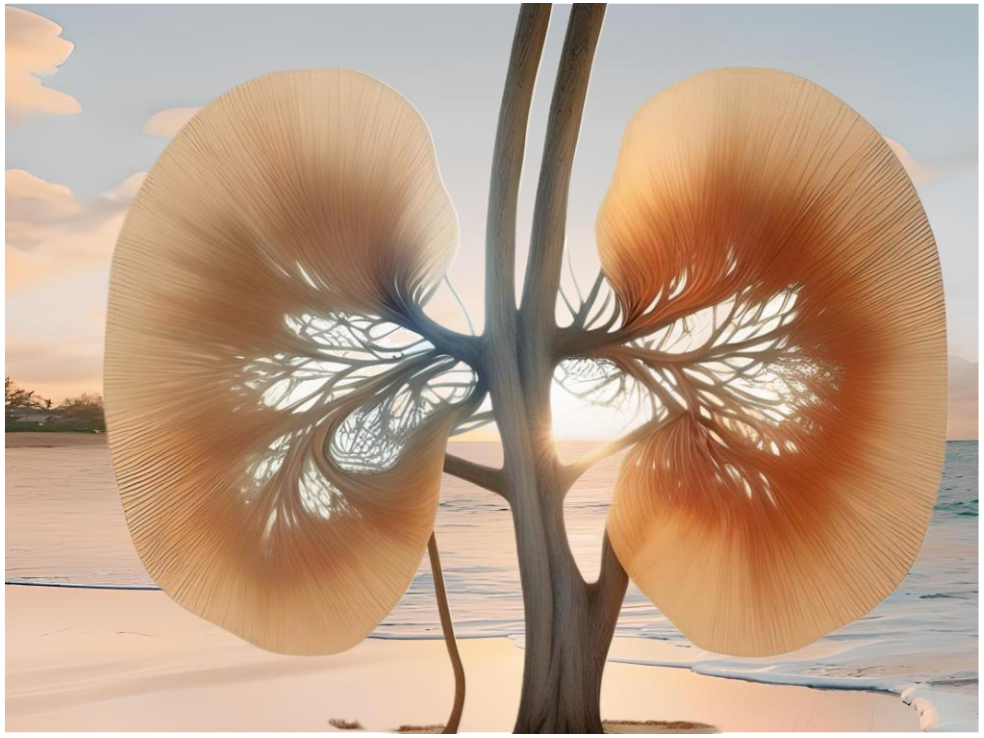
SUBSTANTIAL SHAREHOLDERS[3]

Position	Holder Name	Holding	% IC
1	Mr P Meurs	87,259,311	14.5%
TOTAL (TOP 5) Shareholders		**149,412,198**	**24.9%**

1. As at 13 March 2026; 2. Past 30 trading days liquidity as at 13 March 2026; 3. Shareholder register as at 13 March 2026

Dimerix

(ASX:DXB)



A biopharmaceutical company developing innovative new therapies in areas with unmet medical needs, with a core focus on inflammatory disease treatments such as kidney and respiratory diseases.

SCAN ME

WELL POSITIONED **TO DELIVER** AGAINST STRATEGIC PLAN

ESG Statement

Dimerix is committed to integrating Environmental, Social and Governance (ESG) considerations across the development cycle of its programs, processes and decision making. The Dimerix commitment to improve its ESG performance demonstrate a strong, well-informed management attitude and a values led culture that is both alert and responsive to the challenges and opportunities of doing business responsibly and sustainably.

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E. investor@dimerix.com

DMX-200: unique pharmacology

- CCR2 activation promotes recruitment of inflammatory monocytes to the kidney

  DMX-200 **inhibits** CCR2[1]

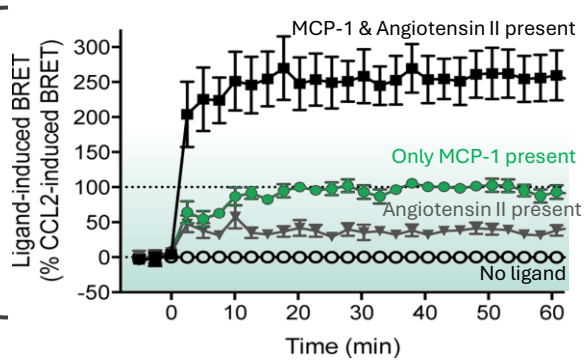
- Monocytes promote sclerosis and fibrosis of the kidney

  DMX-200 **reduces** inflammatory cells[1,2,3]

- Podocytes are the essential filter cells of the kidney

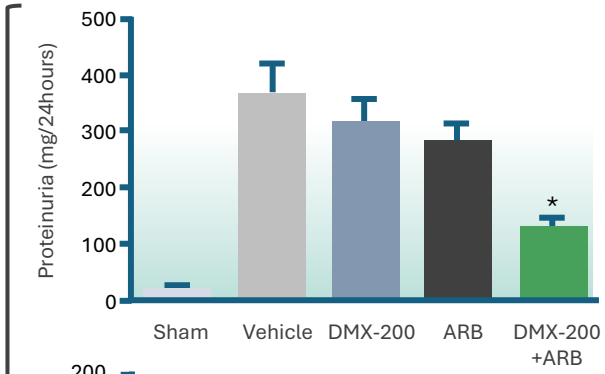
  DMX-200 **preserves** podocytes[1]

in-vitro

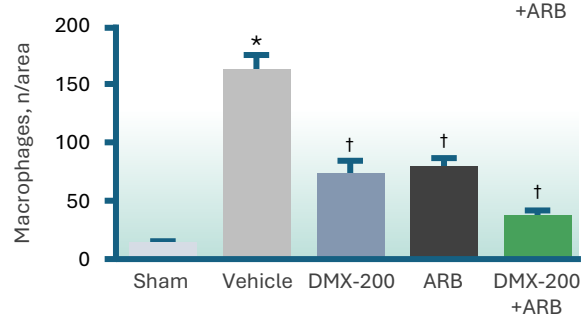


Complex of CCR2 and AT1R increases aberrant signaling when both receptors activated[1]

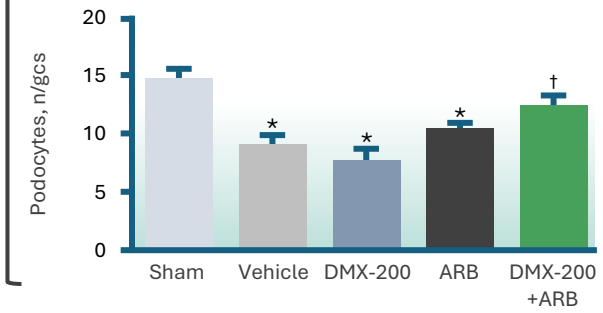
in-vivo[1]



Simultaneous inhibition of CCR2 and AT1R reduces proteinuria an important early marker of kidney function[1]



Simultaneous inhibition of CCR2 and AT1R reduces recruitment of monocytes to the kidney[1]



Simultaneous inhibition of CCR2 and AT1R preserves the number of essential filter cells (podocytes) in the kidney[1]

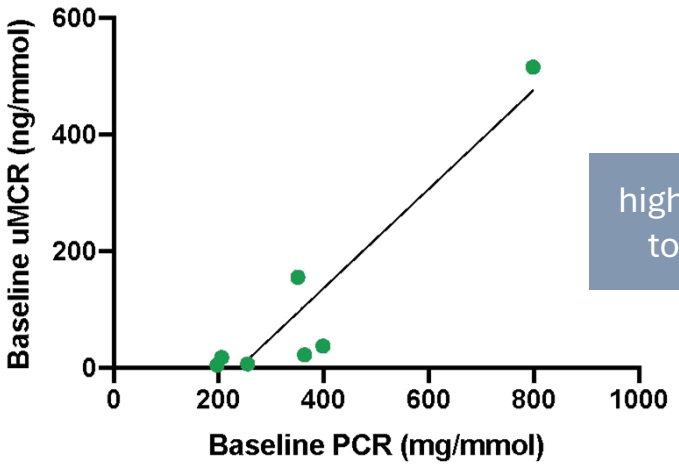
Dimerix

1. Ayoub MA, et al. (2015) PLoS One; doi.org/10:e0119803; 2. ASX presentation 27 October 2020; 3. Liu Y. et al (2024) Role of MCP-1 as an inflammatory biomarker in nephropathy, Front. Immunol., Sec. Inflammation doi.org/10.3389/fimmu.2023.1303076; CCR2: C-C chemokine receptor type 2; AT1R: Angiotensin II type 1 receptor; MCP1: Monocyte chemoattractant protein-1 (also known as CCL2, the ligand for CCR2); BRET: Bioluminescence Resonance Energy Transfer Assay; PCR = protein creatinine ratio; *P<0.05 vs sham STNx rats ; P<0.05 vs vehicle STNx rats;

DMX-200 Phase 2 effect on inflammatory biomarker[1]

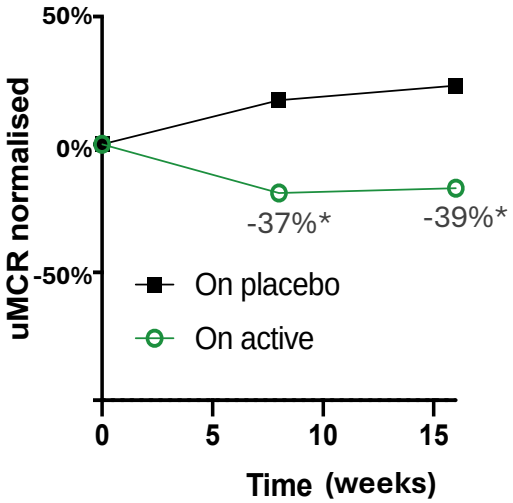
Unlike other CCR2 antagonists investigated to date, treatment with DMX-200 reduces the urine concentration of the pro-inflammatory ligand of CCR2 called MCP-1[2]

Average baseline MCP-1 versus average baseline proteinuria



high MCP-1 correlates to high proteinuria

Change in MCP-1 over time on DMX-200 versus placebo



MCP-1 levels reduced when on DMX- 200 treatment

- **16 weeks treatment with DMX-200 vs placebo reduced inflammatory biomarker by 39%:**
  - DMX-200 blocks receptor responsible for inflammation
  - Translates to reduced inflammation and subsequent fibrosis (scarring) in the kidney[2]

Dimerix

*uMCR = Urinary MCP-1 creatine ratio; PCR = protein creatinine ratio; *placebo adjusted difference*

1. ASX presentation 27 October 2020; 2. Liu Y. et al (2024) Role of MCP-1 as an inflammatory biomarker in nephropathy, Front. Immunol., Sec. Inflammation doi.org/10.3389/fimmu.2023.1303076

PARASOL: proteinuria as an endpoint for full FDA approval

1

PARASOL FSGS Working Group 2024

➢ PARASOL was formed in Dec-23 to address the need to **validate alternative surrogate endpoints** for FSGS, and is a coalition of nonprofit organizations, academia, registries, trials and Sponsors to share data to support analysis[1]

- PARASOL confirmed that eGFR slope is a valid endpoint for predicting progression of kidney disease
- It is recognised FSGS patients see higher proteinuria, even in remission, due to residual scaring of the glomeruli
- PARASOL data demonstrated the strong relationship between a reduction in proteinuria and a reduction in the progression of kidney disease in FSGS patients

- Reduction in proteinuria is now a validated endpoint for full FDA approval

2

Biological Plausibility

➢ The FDA has emphasised the need for programs wishing to use proteinuria endpoints to be able to justify the biological plausibility (scientific rationale of why or how the drug candidate is having the desired effect) of the drug on the endpoint chosen

- Dimerix has existing preclinical evidence on the preservation effect of DMX-200 on the specialist cells on the kidney – the podocytes[2]
- Dimerix has existing clinical and preclinical evidence of reduced recruitment of monocytes to the kidney and reduced MCP-1 levels

- PARASOL has increased the range of potential endpoints that may best show the treatment effect of DMX-200

3

ACTION3 capturing all proposed endpoint data: eGFR and proteinuria

Proteinuria

- Randomised, double blind PCR values over 24 months
- PCR captured across 4-week washout
- PCR measured over additional 24 month open-label period

eGFR slope

- Randomised, double blind eGFR values captured over 24 months, including raw values and total eGFR slope

Other endpoints

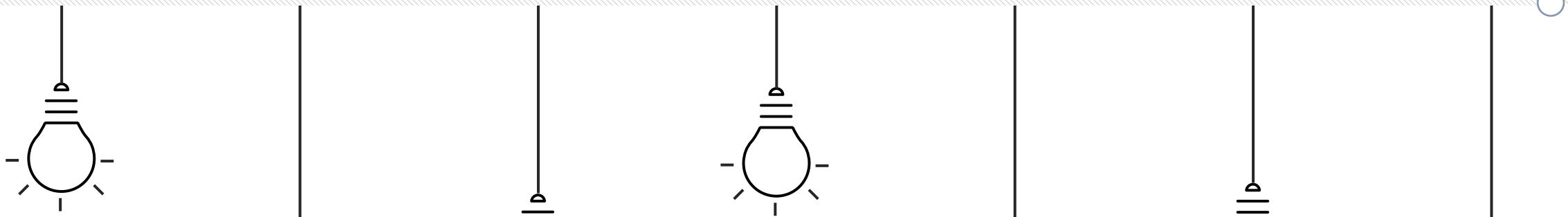
- Classical definitions of complete and partial remission
- PARASOL-informed response endpoints
- Hard-renal endpoints (where available)

Dimerix

1. Coalition made up of: NephCure, ISGD, NKF, KHI are co-sponsors of the project, FDA and EMA involvement, University of Michigan is data coordinating center, Industry invited as participants; 2. Ayoub MA, et al. (2015) PLoS One; doi.org/10:e0119803; 3. ASX presentation 27 October 2020

Intellectual property portfolio

DMX-200



Exclusivity
7/10 years
Orphan exclusivity from marketing approval date in many territories[1]
+
Paediatric
exclusivity period extension[1]

Portfolio 1
Exp.2032
(2033 in US)
Method of Use
Granted in key territories[2]

Portfolio 2
Exp.2042
(if granted)
Method of Use & Formulation
Global application[3]

Portfolio 3
Exp.2042
(if granted)
Formulation
Global application[3]

Portfolio 4
Exp.2044
(if granted)
Method of Use & Formulation
Global application[3]

Portfolio 5
Exp.2045/6
(if granted)
Method of Use
Provisional application[4]

Trademarks
Various trademarks
Global applications

1. DMX-200 is a "New Chemical Entity: an active moiety not previously approved, which can attract exclusivity periods in various territories; 2. Granted patents US9,314,450, US10,058,555, US10,525,038, US11,382,896, US12,083,102, CN2012800046165, CA2,821,985, EP12734251.7, HK 1404477.8, IL227414, JP2013-547780, SA2013/5897, AU2012206945; 3. Patent applications: PCT/AU2022/050013, PCT/AU2022/050249 and PCT/AU2024/050416; 4. US Provisional Application 63/887,984

Dimerix

Dimerix board

Mark Diamond
BSc, MBA
Non-Executive Chairman

Previous experience:

percheron THERAPEUTICS · Pfizer (Faulding)[1]

- Senior pharmaceutical executive with a demonstrated record of achievement and leadership over more than 30 years within the pharmaceutical and biotechnology industries
- Significant accomplishments in capital raising initiatives, pipeline development and licensing
  - ✓ BSc – Chemistry
  - ✓ MBA – Business

Nina Webster
PhD, MBA, M.IP.Law
CEO & Managing Director

Previous experience:

Wyeth[2] · ACRUX · immuron

- Experienced in product development, commercial strategy development & execution
- Successfully commercialized pharmaceutical products globally
  - ✓ BSc (Hons) – Pharmacology
  - ✓ PhD – Pharmaceutics
  - ✓ MBA – Business
  - ✓ M.IP.Law – Intellectual Property Law

Hugh Alsop
BSc (Hons), MBA
Non-Executive Director

Previous experience:

mayne pharma[1] · ACRUX · HatchTech · KINOXIS

- Extensive biotech drug development & commercial manufacturing experience
- Responsible for successful global commercialization programs & NDA registrations
  - ✓ BSc (Hons) – Chemistry
  - ✓ MBA – Business

Sonia Poli
PhD
Non-Executive Director

Previous experience:

Roche · addex · AC Immune · minoryx

- Experienced executive in pharmaceutical operations
- Background in small molecules development and analytical development
  - ✓ BSc (Hons) – Chemistry
  - ✓ PhD – Industrial Chemistry

Clinton Snow
BEng (Hons), BCom
Non-Executive Director

Previous experience:

Woodside Energy · PolyActiva · icetana[AI]

- Experienced technology and governance professional with a focus in operations, risk management, assurance, and AI
- Provides advisory services to a family office with multiple Australian biotech investments
  - ✓ BEng (Hons) – Chemical Engineering
  - ✓ BCom – Commerce

1. Acquired by Pfizer in 2015; 2. Acquired by Pfizer in 2009

Dimerix management

Nina Webster
PhD, MBA, M.IP.Law
CEO & Managing Director

Previous experience:

Wyeth[1] ACRUX immuron

- Experienced in product development, commercial strategy development & execution
- Successfully commercialised multiple pharmaceutical products
  - ✓ BSc (Hons) – Pharmacology
  - ✓ PhD – Pharmaceutics
  - ✓ MBA – Business
  - ✓ M.IP.Law – Intellectual Property Law

Mike Tonroe
BSc (Hons) FCA, MAICD
CFO & Company Secretary

Previous experience:

IMUGENE OPTHEA Genetic Technologies

- Experienced finance and governance executive with extensive experience of both ASX and NASDAQ-listed companies.
- Brings more than 30 years' international finance leadership experience across Australia, US, Canada, the UK and Hong Kong.
  - ✓ BSc (Hons) – Business Studies
  - ✓ MAICD
  - ✓ Chartered Accountant

David Fuller
B. Pharm (Hons), MBBS
CMO

Previous experience:

RACE ONCOLOGY Syneos Health genzyme

- 35 years international experience in drug development, commercialization and corporate leadership
- Planning, Financing, Pre-clinical, Clinical Development, Regulatory Approval, Product Launch, Pharmacovigilance, and Medical Affairs
  - ✓ B.Pharm (Hons) - Pharmacy
  - ✓ MBBS - Medicine and Surgery

Robert Shepherd
PhD, MBA,
COO

Previous experience:

MONASH University Medicines Development for Global Health

- Experienced pharmaceutical executive in project management, clinical development and research translation
- BD and strategic alliance leader
- Led multidisciplinary R&D&C teams for 13 years
  - ✓ BSc (Hons) – Genetics
  - ✓ PhD – Molecular Immunology
  - ✓ MBA – Business & Leadership

1. Acquired by Pfizer in 2009;

Medical Advisory Board

Professor
Hiddo Heerspink
PhD

Professor of Clinical Trials and Personalized Medicine: University Medical Center Groningen, the Netherlands. He specializes in the research of novel treatment approaches to slow the onset of diabetic cardiovascular and renal disease. Hiddo has been instrumental in interactions between industry, researchers and regulatory agencies in the validation of surrogate endpoints for renal trials.

Professor
Alessia Fornoni
MD, PhD, FASN

Professor of Medicine & Molecular & Cellular Pharmacology: University of Miami. Chief of the Katz Family Division of Nephrology and Hypertension. She has an extensive history of translational excellence for patients with renal disease and has uncovered novel pathogenetic mechanisms and therapeutic approaches for glomerular disorders.

Professor
Jonathan Barratt
MD, PhD, FRCP

Mayer Professor of Renal Medicine: Department of Cardiovascular Sciences; University of Leicester and Nephrologist. Jonathan is the IgA nephropathy Rare Disease Group lead for the UK National Registry of Rare Kidney Diseases (RaDaR) and a member of the steering committee for the International IgA Nephropathy Network.

Associate Professor
Lesley Inker
MD, MS, FRCPC

An attending physician and Director of the Kidney and Blood Pressure Center in the Division of Nephrology at Tufts Medical Center. Lesley's major research interest is in the estimation and measurement of glomerular filtration rate (GFR) and in defining alternative endpoints for CKD progression trials based on GFR decline and changes in albuminuria.

Dr Muh Geot Wong
MBBS, PhD, FRCP

Renal Physician and Head of the Renal Clinical trials at the Royal North Shore hospital, Sydney, Australia. Muh Geot's main areas of research are in understanding the mechanisms of kidney fibrosis, biomarkers research, and identifying strategies in delaying progressive kidney disease including glomerular diseases.

Professor
Howard Trachtman
MD, FASN

Graduated from Haverford College and the University of Pennsylvania School of Medicine. He has been a practicing pediatric nephrologist for 35 years. Has been the PI of NIDDK and industry sponsored clinical trials in glomerular disease and am a Co-Investigator in the NEPTUNE and CureGN observational cohort studies.

Associate Professor
Laura Mariani
MD, MSCE

Assistant Professor in the Division of Nephrology at the University of Michigan. Interest in observational studies in glomerular disease, including NEPTUNE and CureGN. Lead on PARASOL program to define FSGS endpoints with by applying statistical methods for clinical outcome definition and prediction of kidney disease progression.