

Dimerix

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

Investor Presentation

June 2025



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, along with those factors outlined in the most recent Dimerix Limited Annual Report.



Overview

Phase 3 Global Opportunity

4

DMX-200 licensing partners across key territories¹

FSGS Indication

a **rare disease** that causes scar tissue of kidneys, which leads to irreversible kidney damage² ~\$1.4 billion

in total upfront &
potential development
and sales milestone
payments plus
royalties¹

>\$65 million

in total payments
received to date¹

No approved treatments

available to treat FSGS

FSGS kidney damage can lead to dialysis, kidney transplants or death²

Successful Phase 3 interim analysis:

analysis showed DMX-200 had **performed better than placebo** in reducing proteinuria³ **Lead Drug Candidate**

DMX-200 in a **Phase 3 clinical trial** for focal
segmental
glomerulosclerosis
(FSGS)

Orphan drug designation

regulatory, marketing exclusivity and pricing **benefits** in key territories



Strong body of evidence with significant progress

| | 3 | |
|------------|--|---|
| \bigcirc | Mechanism of Action | ➤ Precision therapy to disrupt inflammatory feedback loops in the kidney of patients with FSGS¹ |
| | Pre-Clinical | ➤ FDA confirmed proposed pre-clinical safety package sufficient to support marketing submission ² |
| | Manufacturing | Commercial scale up in place – manufacturing sites in USA ³ |
| | Phase 1 / Phase 2 clinical trials | Encouraging efficacy and positive safety signals across Phase 1 & Phase 2 studies (n=>100), including demonstrating a reduction in proteinuria and inflammatory markers in FSGS patients ⁴ |
| | ACTION3 Phase 3: Part 1 interim analysis | Interim analysis (n=72 @ 35 weeks) showed DMX-200 performing better than placebo in reducing proteinuria ⁵ |
| \bigcirc | FDA and Project PARASOL | Alignment on proteinuria as primary endpoint for final approval⁶ |
| \bigcirc | 3rd Party Validation | 4 licensing deals executed for various key territories, all of whom conducted independent, extensive due diligence ⁷ |
| | ACTION3 Phase 3: Part 2 interim analysis | ▶ Blinded data collection and analysis expected after PARASOL project outcomes and FDA feedback ⁶ |
| | ACTION3 Phase 3: Part 3 final analysis | 2-year proteinuria (potential primary endpoint) and eGFR (primary and/or secondary endpoint) data |

serves as basis for full approval (n=~286)



ACTION3 Phase 3: Part 3 final analysis

Cycle of damage:

What is FSGS?

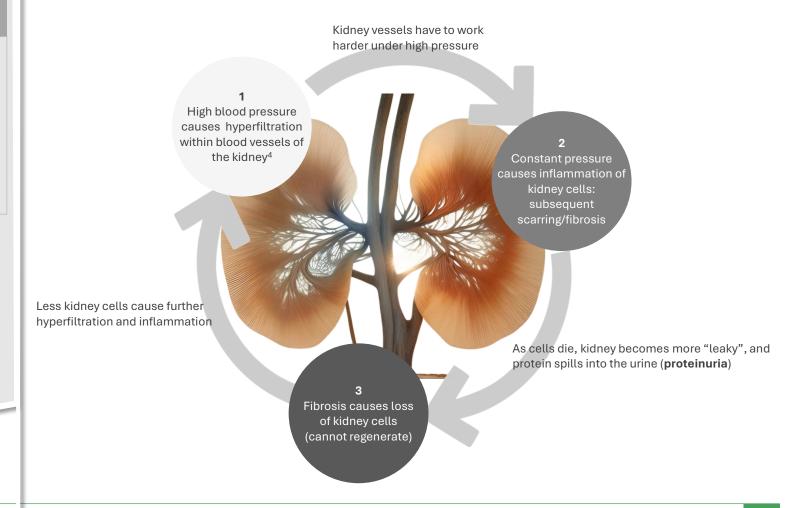
Focal = some

Segmental = sections

Glomerulo = of the kidney filtering units

Sclerosis = are scarred

in glomerular diseases





Cycle of damage:

What is FSGS?

Focal = some

Segmental = sections

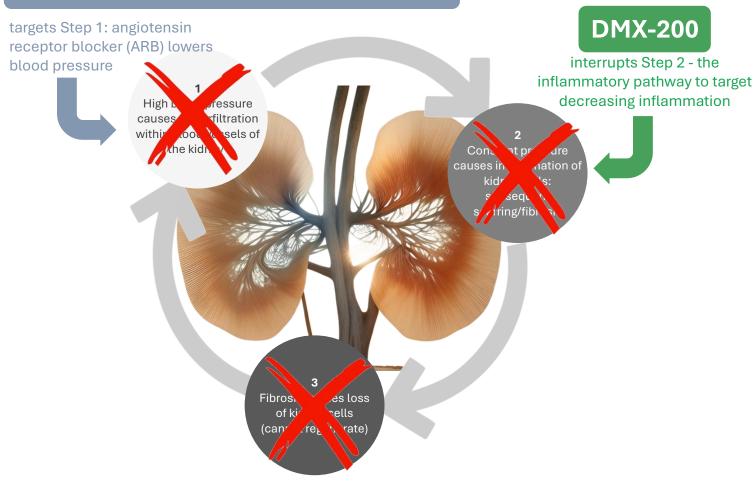
Glomerulo = of the kidney filtering units

Sclerosis = are scarred

This synergistic activity of both agents disrupts the cycle of damage in FSGS

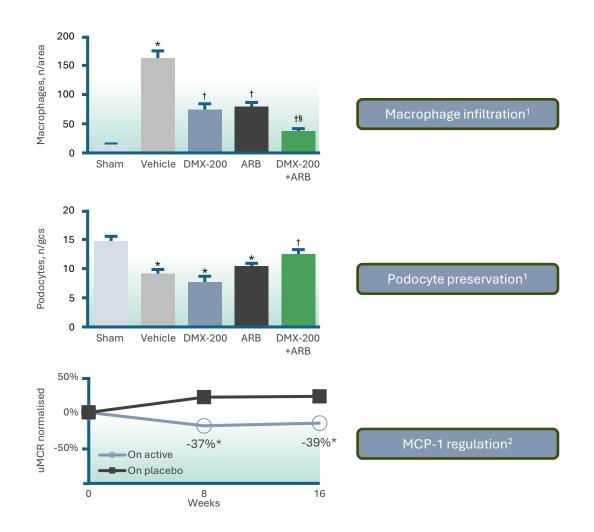
in glomerular diseases

Existing blood pressure medication





DMX-200: mechanism of action



 CCR2 is required for recruitment of inflammatory cells to the kidney



DMX-200 blocks CCR21

Macrophage/monocyte: regulate inflammatory cells



DMX-200 **reduces** inflammatory cells^{1,2,3}

Podocytes: specialist filtration cells in the kidney



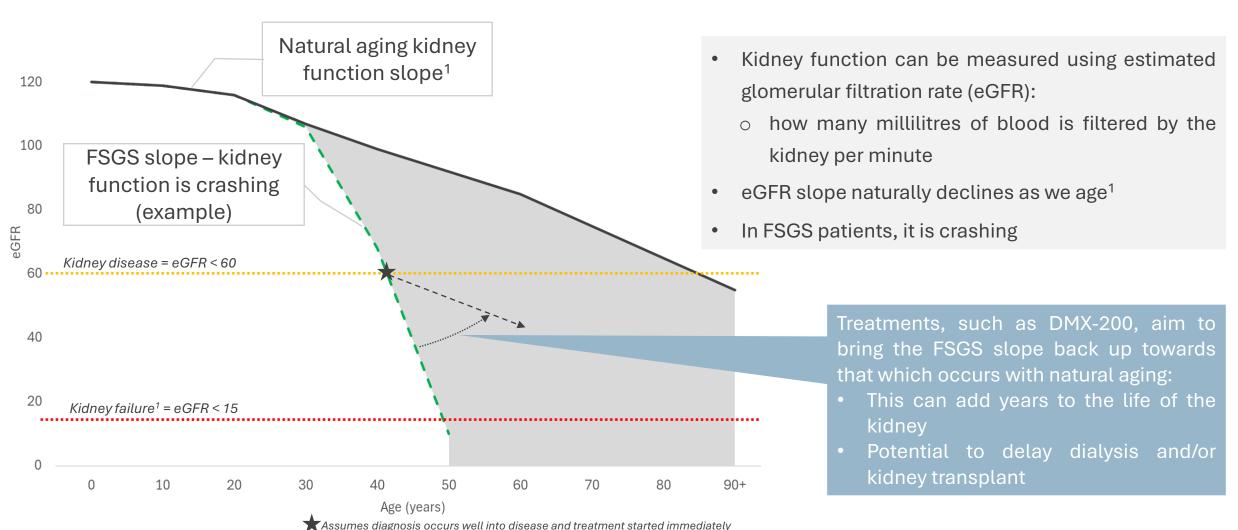
DMX-200 preserves podocytes¹



Damaged podocyte⁴



Significance of stabilising eGFR slope: clinical endpoint

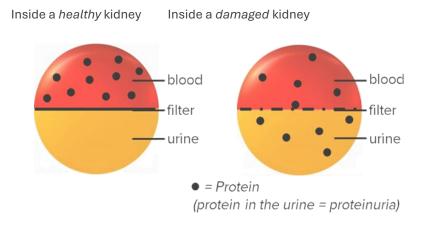




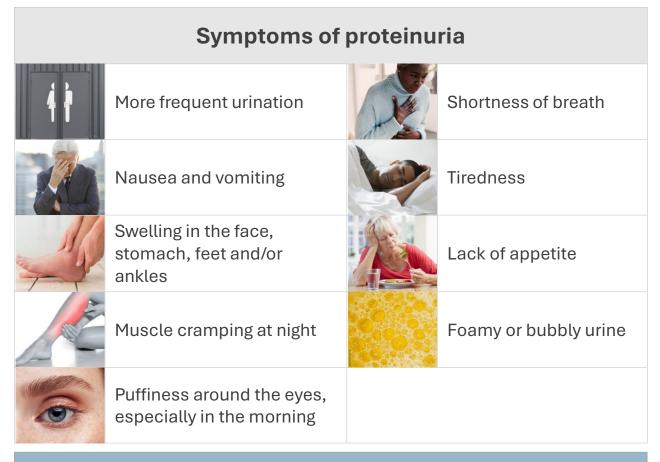
Significance of decreasing proteinuria: clinical endpoint

Why are kidneys important?

 A healthy kidney is a good filter and allows little to no protein in the urine¹



- When kidneys are damaged, protein can leak into the urine causing proteinuria
- Proteinuria represents an important early marker of kidney function²



DMX-200 aims to reduce the inflammation of the kidneys:

if DMX-200 reduces inflammation, the amount of proteinuria should decrease



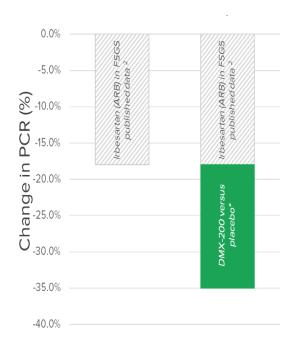
DMX-200: Phase 2 met primary and secondary endpoints



Clinically meaningful outcomes achieved for patients,^{2,3} with no safety issues



Average reduction of **17**% in proteinuria after 16 weeks treatment on DMX-200 versus placebo¹



"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 2020³



EFFICACY

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



 No safety concerns – reduced development risk



ACTION3 phase 3 clinical trial – next steps



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

Total number of patients required - anticipated H2 20251

202

Patients recruited, randomised and dosed²



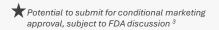
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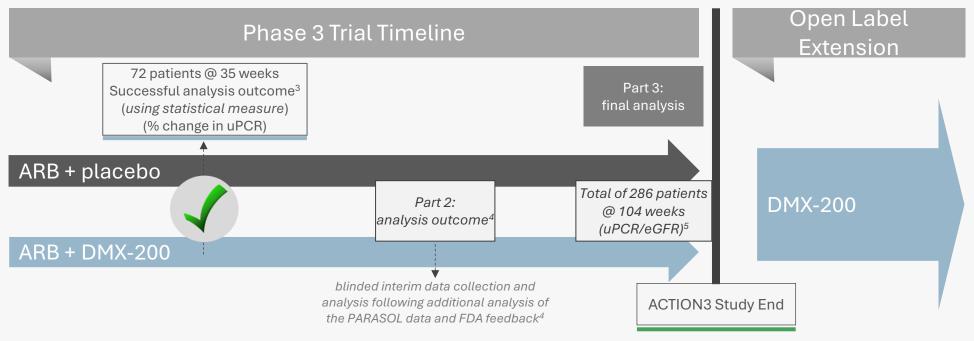
Patients enrolled over into Open Label Extension Study²



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









Positive Type C meeting held in March 2025 with US Food & Drug Administration (FDA) on proteinuria trial endpoints, and potential for accelerated approval for DMX-200¹

FDA confirmed that a proteinuria-based endpoint for full marketing approval in the US, which may be either:

1

- the proportion of patients meeting the proteinuria responder definition; or
- percentage change in proteinuria from baseline

Dimerix is working with the PARASOL working group on additional analysis of existing PARASOL data to further assess what may represent an appropriate and meaningful endpoint for accelerated approval in FSGS¹

• This additional work may provide the justification to support an accelerated approval endpoint, which will be discussed with the FDA prior to any potential submission

3-6 months¹



Competitive landscape in FSGS



No approved therapies for FSGS



Low competition



DMX-200 is the only inflammatory modulator in development

| | Phase 1 | Phase 2 | Phase 3 | Company |
|----------------------------------|--------------------------------------|-------------------------------------|-----------------------------|------------------------|
| DMX-200 QYTOVRA® [REPAGERMANIUM] | | | ACTION3 FSGS CLINICAL STUDY | t Dimerix |
| Sparsentan | AT₁R/ET₄R dual inhibitor – Failed P | Ph3 eGFR endpoint: resubmitted to F | DA on proteinuria endpoints | Travere Therapeutics |
| VX-147 | APOL1 inhibitor – specific type of g | genetic FSGS | | Vertex Pharmaceuticals |
| BI-764198 | TRPC inhibitor | | | Boehringer Ingelheim |
| Atrasentan | AT_1R/ET_A antagonist | | | Chinook |
| R3R01 | Lipid modifying | | | River 3 Renal |

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FSGS market – potential for growth

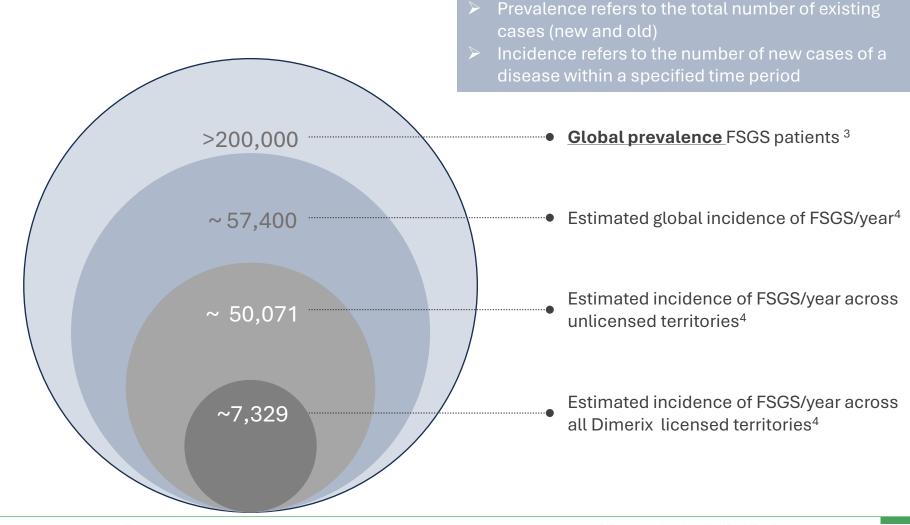
Biopsy

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

7 per 1,000,000

Global <u>incidence</u> rate of FSGS per capita per year¹

FSGS is the most frequent primary glomerular disease that reaches end-stage renal failure in the US²





Rare kidney disease pricing examples



US Medicare Drug Price Negotiation Program¹

US Medicare Drug Price Negotiation Program **exempts orphan drugs** that treat "only one rare disease or condition" from drug
price negotiations



One Big Beautiful Bill Act²

proposes to **expand that exemption** by including drugs that treat "one <u>or more</u> rare diseases or conditions" from pricing negotiations

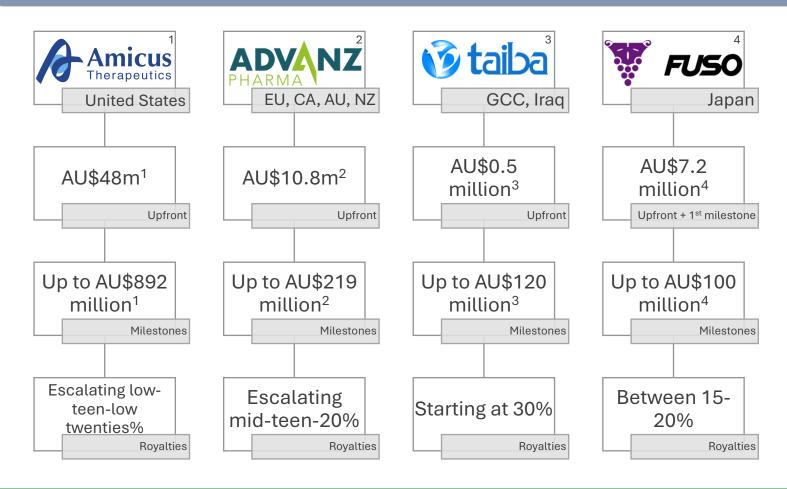
Example current pricing for other rare kidney disease drugs:

- in the US (i.e. Filspari in IgAN)³: **US\$9,900 p/month**
- in Europe/UK (i.e. Kinpeygo/Tarpeyo)⁴: **US\$8,267 p/month**



Summary of licensing deals for DMX-200 to date

Dimerix has successfully partnered DMX-200 across key markets



Licensing deals collectively valued up to

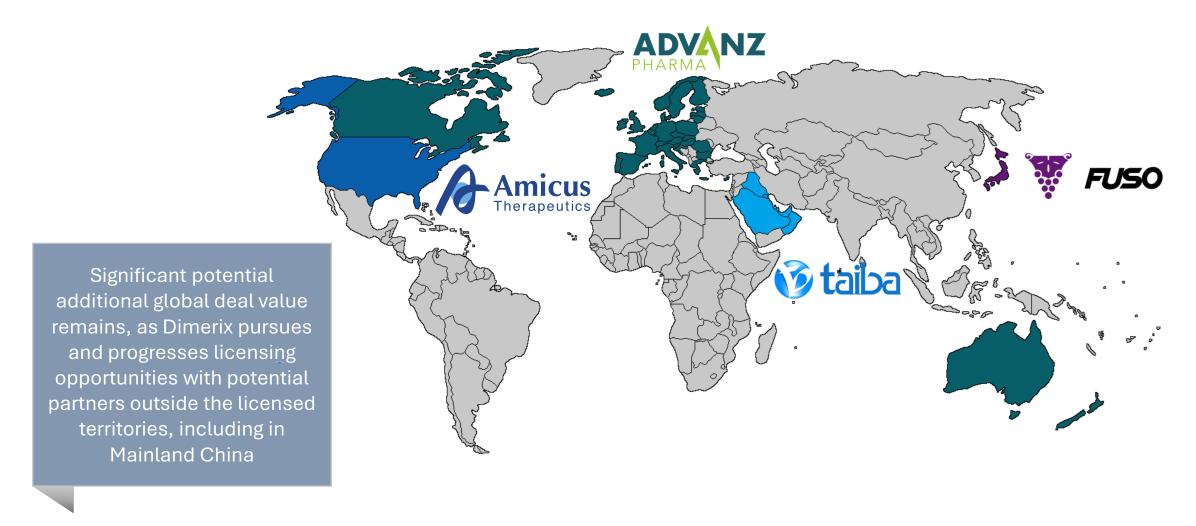
~AU\$1.4 billion

in total upfront and potential milestone fees <u>plus</u> royalties¹

AU\$65
million
in total payments received



Potential for additional partnering opportunities





Corporate overview

| Ticker Symbol | ASX: DXB | |
|--|---------------|--|
| Cash Balance (Mar25)* | \$17 million | |
| Market Capitalisation ¹ | \$295 million | |
| Share price ¹ | \$0.50 | |
| Total ordinary shares on issue ¹ | 590,068,104 | |
| Average Daily Liquidity by value for past 30 trading days ² | \$2.0 million | |



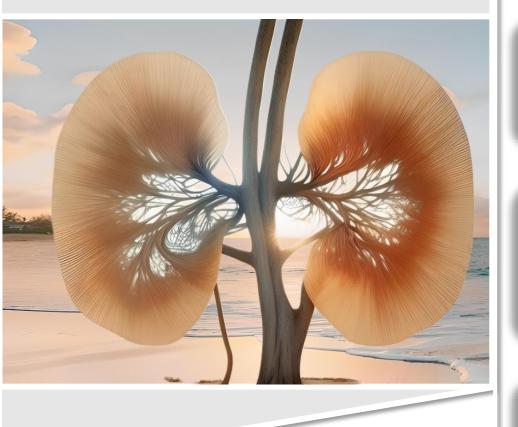
| SUBSTANTIAL SHAREHOLDERS ³ | | | | | | | |
|---------------------------------------|-----------------|-------------|-------|--|--|--|--|
| Position | Holder Name | Holding | %IC | | | | |
| 1 | Mr P Meurs | 87,259,311 | 14.8% | | | | |
| TOTAL (TOP | 5) Shareholders | 142,856,456 | 24.2% | | | | |

*Cash balance does not include:

- ~\$48 million upfront fee received from Amicus Therapeutics licensing agreement (ASX release 6 May 2025)
- ~\$4.3 million payment anticipated on 1st development milestone achievement in May 2025
- Up to \$6.3 million potential conversion of 41,026,596 DXB options (as at 31 March 2025) exercisable at 15.4c per share (expire 30 June 2025)

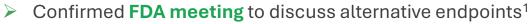


Potential catalysts



CY 2025

Q1/Q2 2025





> ~AU\$4.1 million² development milestone anticipated from FUSO¹

Q3/Q4 2025

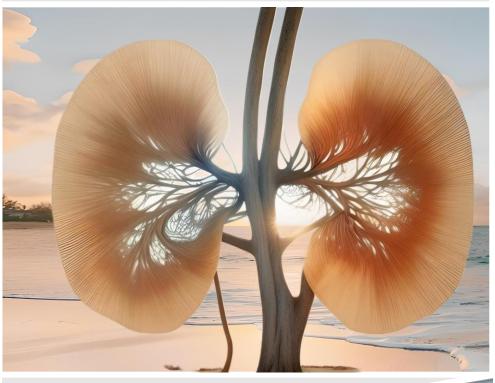
- > Planned blinded interim data collection anticipated in H2 2025³
 - Potential for accelerated (or conditional) approval submission, subject to PARASOL outcomes, FDA feedback and interim analysis outcomes^{1,3}
- Full study recruitment of 286 adult patients anticipated in H2 2025⁴

Potential upside - at any time

Additional **licensing partners** for DMX-200: Dimerix continues to pursue potential licensing opportunities in un-licensed territories, including China







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.



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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