

Switzer Investment Conference

ASX:OSP October 2020



Investment Highlights



Clear and large problem: Contrast-Induced Acute Kidney Injury (CI-AKI) is increasingly associated with poor patient outcomes and costs hospitals over US\$900m a year in the USA alone¹



Our technology is the solution: DyeVert has a ~\$1.1B addressable market² and is clinically proven to reduce the risk of CI-AKI through dye minimization and monitoring in angiographic procedures



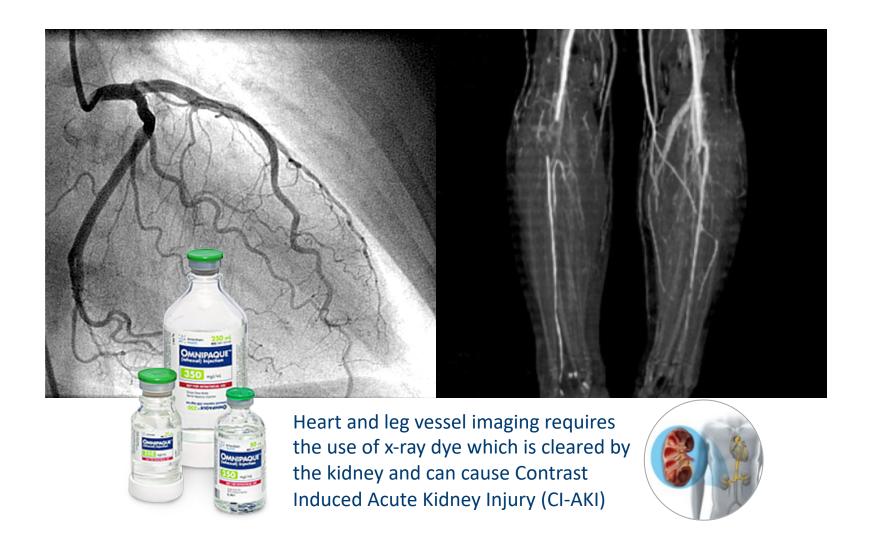
Clear US growth plan and distribution strategy outside US: Core focus to increase market penetration through US GPO strategy; alongside GE Healthcare distribution agreement in Europe, Middle East and Asia and recently signed agreement in Australia and New Zealand



A great value opportunity: Continued strong year on year revenue growth of 84% CAGR CY16-19 has not translated to share price growth



Clear and large problem | Making angiography safer for Chronic Kidney Disease patients





Clear and large problem | Osprey's DyeVert technology represents a significant market opportunity

Opportunity of 3.2M procedures per year in the USA and Western EU¹

CKD

1.3M procedures per year

Diabetes

1.0M procedures per year

Peripheral

450K procedures per year

STEMI

440K procedures per year





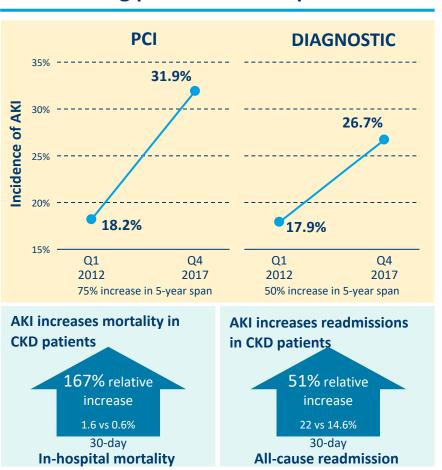
Clear and large problem | The Burden of Illness study¹ highlights the costs of CI-AKI to both patients and hospitals



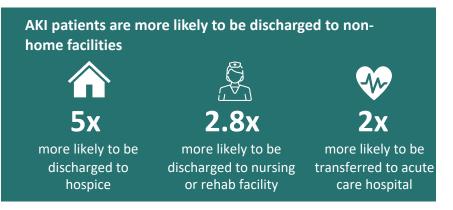
A study of 749 hospitals with 2.8m angiography patients with CKD

A rising problem in CKD patients

AKI increases hospital costs¹









Clear and large problem | There is a concerted and growing focus on AKI avoidance

Physician consensus position on CI-AKI

Avoidance guidelines for at-risk patients







Class 1 Level B recommendation for CI-AKI reduction



Patients with an eGFR < 60 ml/min are at high risk for AKI events



Adequate preparatory hydration should be given to at-risk patients



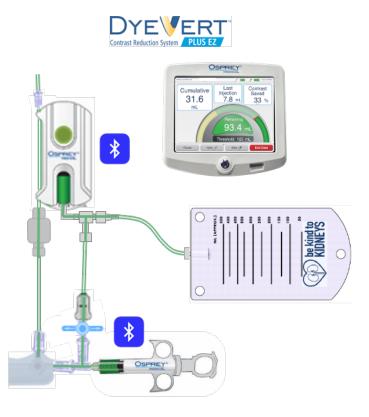
Minimise contrast dosage to high risk patients



Our technology is the solution | Osprey's proprietary technology is patent protected

Osprey's proprietary solution

Clinically proven results



On average 40% reduction in Dye1



Reduce dye in angiographic procedures¹



No compromise to image quality



Allow for real-time contrast monitoring of maximum allowable dose



Commercial approach | Key commercial highlights demonstrate strong customer adoption

US Commercial strategy, direct sales model

DyeVert unit sales since 2015 (#)¹



NEW ACCOUNTS

Leverage GPO National Agreements



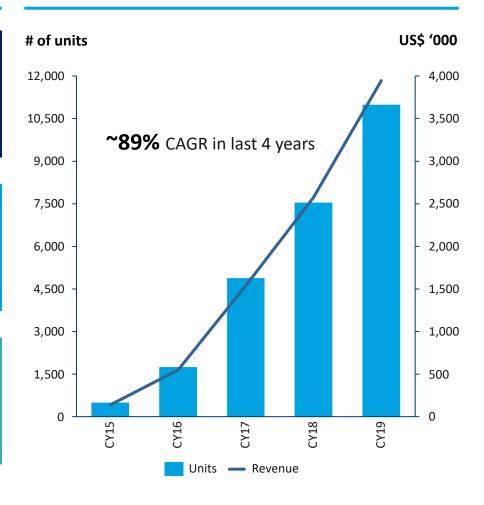
DRIVE PENETRATION

Increase penetration in accounts



PUBLISH CLINICAL SUCCESS

Reinforce protocol driven care with the DyeVert



Notes



Commercial approach | Real-world AKI prevention strategies that work

AKI reduction initiatives

St. Mary's Medical Center Huntington, WV

Presented at West Virginia ACC 2018 Annual Conference







Houston Methodist Sugarland Hospital

Sugarland, TX

Presented at ACC NCDR 2018 Annual Meeting











Presented at ACC Quality Summit 2019 Annual Meeting







Osprey Registry

up to 10K patients in 50 hospitals



Clear future growth strategy | Clear plan with significant whitespace for growth

Clear plan for accelerated future growth

Whitespace for commercial growth

1. Increased penetration existing US

- Protocols driven approach adds consistency
- · Tracking of AKI and Publication of results

2. GPO focus for opening new US customers

- Leverage 5 existing GPO contracts to expand to new hospitals
- Increase pace with AKI reduction publications

3. GE OUS market expansion

 Leverage GE's position as the largest global player in contrast media and molecular imaging agents (signed in Jul-20)

4. Australia & New Zealand distribution signed

 Leverage Regional Health Care Group's extensive experience in supplying medical equipment and contrast media (signed in Sep-20) 20% of US customers CKD patients were protected with DyeVert in 2019

50% coverage of hospitals under the GPO model

GE commercialisation team of over 120 in the contracted area who will be selling the Osprey portfolio of products

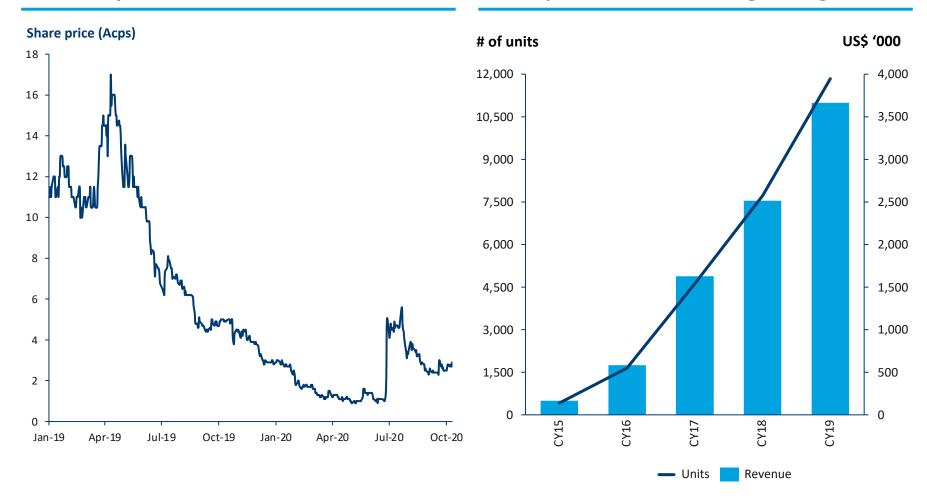
First entry into market where DyeVert technology was developed



Significant value upside | Strong revenue growth has not translated to share price growth

Share price remains at historic lows...

...despite continued strong sales growth¹





High calibre board and management team | Highly

experienced board and management team



Mike McCormick | President and CEO

- 30+ years medical device experience across private and public companies.
- Formerly CEO of Anulux and Centrepulse Spine Tech



John Erb | Non-Executive Chairman

 35+ years of medical device experience and also currently Chairman and CEO of CHF Solutions



Chris Nave | Non-Executive Director

 Founding partner of Brandon Capital and CEO of the Medical Research Commercialisation Fund



Sandra Lesenfants | Non-Executive Director

 Currently serves as Vice President & General Manager of endoVenous business in the Medtronic Cardiac & Vascular Group



Neville Mitchell | Non-Executive Director

 Formerly CFO and Company Secretary at Cochlear where he was for 20+ years and a board member at Sirtex Medical



Focus on driving shareholder value | Multi-pronged approach to drive near term sales growth



GPO Strategy

National contracts • and publications •

- Continue to build on GPO strategy within the US
- Use national contract to open new accounts
- Leverage published data from GPO hospitals to support growth



GE Partnership

A game changer for OUS

- GE agreement to drive sales in OUS regions
- Revenue certainty over the contract duration with prescribed minimum purchase levels with significant potential for upside
- Stable ASPs locking in margin



R&D

Continued investment in R&D

- DyeVert Power XT has CE Mark for EU commercialization by GE
- FDA clearance for the US is expected in early 2021



PODIUM

Scientific presentations

- DyeVert featured in the SCAI Scientific Session in 2020 with strong validation from several medical practitioners
- Continue to build brand awareness through presentations at various reputable conferences and support of key opinion leaders

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DyeVert[™], DyeVert Plus and DyeTect Systems Regulatory Status: Europe – CE Mark obtained; Australia – TGA approval obtained; United States – 510(k) cleared.