

Osprey Medical Inc. - Revised CEO Capital Raising Presentation

Minnesota, United States and Melbourne, Australia – August 4, 2016 – Osprey Medical Inc. (ASX:OSP) has replaced slide 7 of the CEO Capital Raising Presentation to correct the formatting of the two charts on that slide. The revised presentation is enclosed. There were no other changes.

About Osprey

Osprey Medical is focused on protecting patients from the harmful effects of X-ray dye (contrast) used during commonly performed angiographic imaging procedures. The Company's core technologies originated from research conducted by Dr David Kaye at Melbourne's Baker IDI Heart and Diabetes Institute. Its proprietary dye reduction and monitoring technologies are designed to help physicians minimize dye usage. The Company's DyeVert[™] System is a next-generation product that reduces contrast while maintaining image quality in a self-adjusting easy-to-use design. Osprey Medical's Board and Management are comprised of experienced and successful personnel with established track records covering medical device development, regulatory approvals, sales and marketing, and mergers-acquisitions. Osprey Medical's advisory board comprises world-recognised experts in heart and kidney diseases.

Forward-Looking Statements

This announcement contains or may contain forward-looking statements that are based on management's beliefs, assumptions and expectations and on information currently available to management. All statements that address operating performance, events or developments that we expect or anticipate will occur in the future are forward-looking statements, including without limitation our expectations with respect to our ability to commercialize our products including our estimates of potential revenues, costs, profitability and financial performance; our ability to develop and commercialize new products including our ability to obtain reimbursement for our products; our expectations with respect to our clinical trials, including enrolment in or completion of our clinical trials and our associated regulatory submissions and approvals; our expectations with respect to the integrity or capabilities of our intellectual property position. Management believes that these forward-looking statements are reasonable as and when made. You should not place undue reliance on forward-looking statements because they speak only as of the date when made. Osprey does not assume any obligation to publicly update or revise any forward-looking statements, whether as a result of new information, future events or otherwise. Osprey may not actually achieve the plans, projections or expectations disclosed in forward-looking statements, and actual results, developments or events could differ materially from those disclosed in the forward-looking statements.

Media	Investors	Company		
Gavin Lower	Rebecca Wilson	Doug Schoenberg		
Buchan Consulting	Buchan Consulting	VP of Marketing		
T: (613) 8866 1215	M: (61) 417 382 391	T: (952) 955 8230		
glower@buchanwe.com.au	rwilson@buchanwe.com.au	dschoenberg@ospreymed.com		



Capital Raising Presentation

August 2016



Investment highlights



Osprey is aggressively commercialising its dye savings product portfolio

- 7 consecutive quarters of growth in units sold and samples, with 45% growth in unit sales in 2Q16, as compared to 1Q16
- Original sales territory in San Antonio, Texas was profitable in June 2016 with other territories following a similar sales trajectory
- Over 65% of San Antonio hospitals have approved and purchased Osprey's products, which demonstrates the clear potential for market adoption in other sales territories
- Accelerating commercialization with 9 sales representatives operating in unique territories and on track to reach up to 20 reps by end of CY16
- US\$1.1+ billion addressable market targeting 3.2M procedures per year
- Top tier Board and management team, invested in Osprey's success

Capital raising overview



Osprey has raised A\$28m via a two tranche placement with a concurrent security purchase plan to follow to raise a further A\$1m

- Two tranche placement to sophisticated and institutional investors
 - Tranche 1 raising A\$10.8m under Osprey's existing placement capacity pursuant to ASX Listing Rule 7.1 and
 7.1A
 - Tranche 2 raising A\$17.2m conditional on stockholder approval
 - New CHESS Depositary Interests (CDIs) will rank equally with existing CDIs on issue (where 2 CDIs are equivalent to 1 fully paid ordinary share in Osprey common stock)
- Brandon Capital Partners (16.1% stockholder) has committed A\$10m to the Placement (to be invested in Tranche 2)
- A security purchase plan (SPP) will be offered to existing stockholders, capped at approximately A\$1m
- Use of funds
 - Expand commercialization of the DyeVert System
 - Continued research and development of the product portfolio, including DyeVert Plus
 - Ongoing clinical evaluations for scientific presentation and publication; and
 - Provide expansion capital to accelerate growth

Capital raising overview (cont.)



The placement was oversubscribed reflecting a strong endorsement from the investment community

• Funds will be raised via the issue of 100m CHESS Depositary Interests (CDIs) (equivalent to 50m shares of common stock) The issue price is A\$0.28 per New CDI, representing a 10.3% discount to the 15-day volume weighted average price (VWAP) Private Placement will be conducted in two tranches: **Placement** - Tranche 1 will be an unconditional placement of approximately 38.5m CDIs issued under the Company's existing placement capacity pursuant to ASX Listing Rule 7.1 and 7.1A (to raise A\$10.8m); - Tranche 2 will be a conditional placement of approximately 61.5m CDIs (to raise A\$17.2m), subject to stockholder approval at a special meeting (Special Meeting) • An SPP will be offered to eligible CDI holders registered on the record date as at 7pm AEST on 30 August 2016 Security Purchase Plan Capped at A\$1m and up to a maximum of A\$15,000 per eligible CDI holder SPP to be offered at the same issue price as the placement (A\$0.28 per New CDI) New CDIs will rank equally with existing CDIs on issue Ranking and Indicative date for the Special Meeting is Tuesday, 30 August 2016, a Notice of Meeting will be **Timing** dispatched shortly

Capital raising timetable



The indicative timetable for the capital raising is provided below

Indicative capital raising timetable

Trading halt		Tuesday, 2 August 2016
Record date for SPP		Wednesday, 3 August 2016
Placement and SPP announced and Company resumes trading		Thursday, 4 August 2016
Settlement of issue of Placement CDIs under Tranche 1		Wednesday, 10 August 2016
Allotment of issue of Placement CDIs under Tranche 1		Thursday, 11 August 2016
Special Meeting for approval of issue of Placement CDIs under Tranche 2		On or around Tuesday, 30 August 2016
Settlement of Placement CDIs under Tranche 2 (subject to approval)		Monday, 5 September 2016
Allotment of Placement CDIs under Tranche 2 (subject to approval)		Tuesday, 6 September 2016

^{*}The timetable above is indicative only and may be varied subject to the ASX Listing Rules

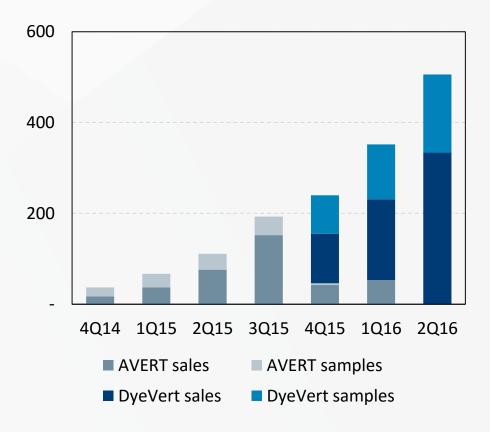
Strong sales growth



Sales traction evidenced by clear product market fit and strong customer growth

- 7 consecutive quarters of growth in units sold and sampled since first sales
- 88% growth in DyeVert units sales and 45% growth in DyeVert and AVERT unit sales combined in 2Q16, as compared to 1Q16
- Cash receipts from customers increased by 66% to US\$113k in 2Q16
- Average selling price of DyeVert is US\$355 (unchanged from 1Q16)
- All sales units transitioned to the upgraded DyeVert System in 2Q16

Quarterly product unit sales & samples since inception

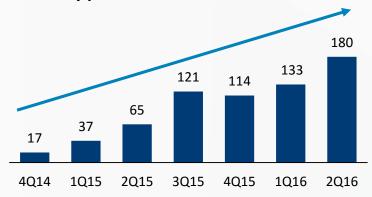


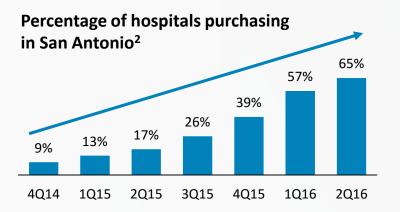
San Antonio sales territory statistics



Original sales territory in San Antonio, Texas was profitable in June 2016 with other territories following a similar sales trajectory

Quarterly product unit sales in San Antonio¹





- San Antonio was profitable in June 2016 with 100 units sold, compared to cashflow breakeven requirement of 75 units per month
- Over 65% (15 of 23) of San Antonio hospitals have approved and purchased Osprey's products
- Physicians have now used DyeVert or AVERT on 667 CKD patients, representing 40% of all CKD patients treated in these hospitals
- San Antonio is demonstrating the clear potential for market adoption in other territories
- Sales traction within the 2 largest hospital systems,
 Methodist and Christus, is particularly encouraging

Quarterly unit sales fell in 4Q15 following the introduction of the DyeVert System, as hospitals sampled the new product

^{2.} This chart reflects the percentage of the 23 hospitals in San Antonio that have approved and purchased Osprey's products, since the beginning of 4Q14

Commercialization strategy



Osprey's growing team of sales representatives are targeting areas in the US with the highest incidence of kidney failure

Sales expansion

- Sales reps incentivised to achieve month on month sales growth and 100 unit sales per month (cashflow breakeven requirement is 75 per month)
- 9 existing sales reps, on track to reach up to 20 sales reps by end of CY16
- Factors driving selection of new territories include number of CKD procedures and prevalence of kidney failure

Marketing

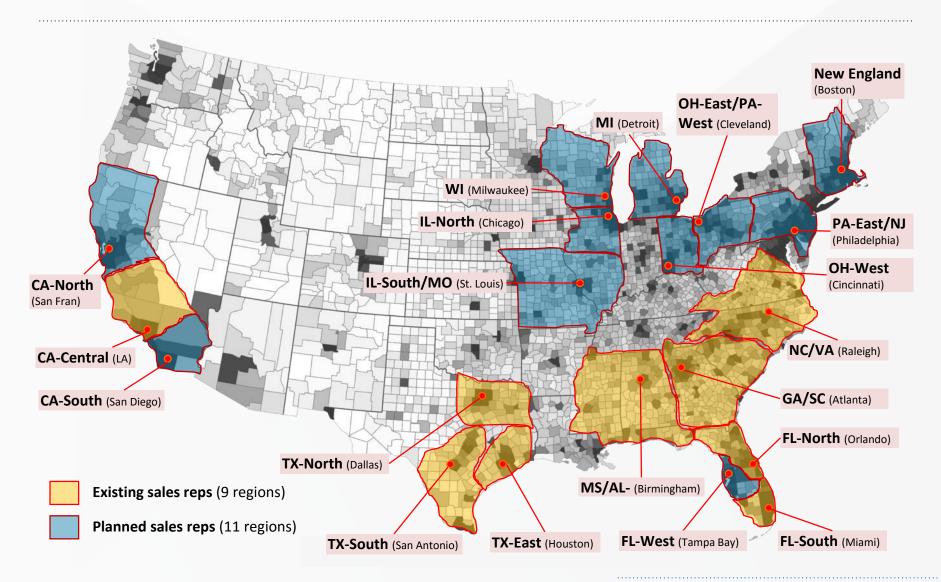
- Osprey is taking advantage of US healthcare reform and stricter national guidelines
- Increasing awareness around kidney protection and the economic arguments for Osprey's products also drives adoption

Key opinion leader advocates

 Developing strong relationships with key opinion leading physicians who are advocating for the benefits of Osprey's products at public forums

Sales territories – planned for CY16





Economic effectiveness



US healthcare payment reform has created an economic incentive for the use of Osprey's products



 The Patient Protection & Affordable Care Act 2010 ("ObamaCare") has changed the economic landscape for medical care in the US and created a favorable economic climate for Osprey's technology



 ObamaCare has redefined hospital and physician payment structures so that payment will be based on quality metrics rather than volume



 Dye volume and CI-AKI are key quality metrics under ObamaCare according to the National Cardiovascular Data Registry



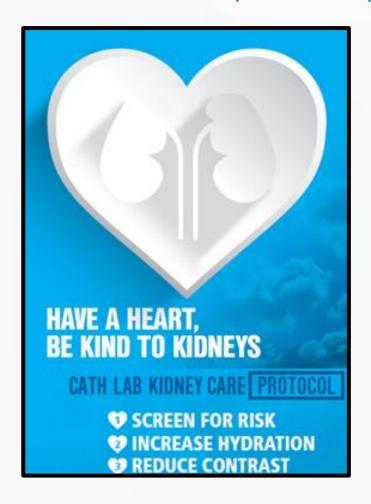
Economic penalties can apply to hospitals that do not comply with ObamaCare

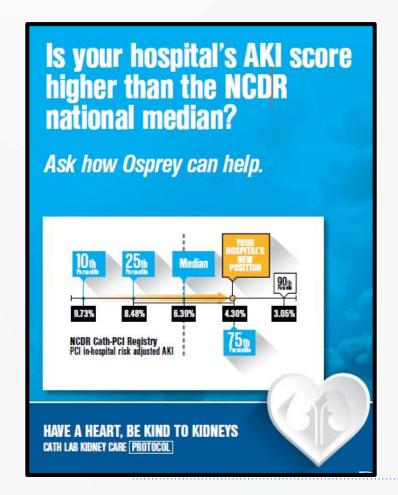


Marketing materials



Osprey is leveraging DyeVert's medical efficacy and economic benefits to drive market awareness and product adoption





Key opinion leader advocates



Key opinion leaders and prominent physicians help drive early adoption of Osprey's dye savings portfolio via podium presentations

Dr Anand Prasad

Assistant Professor of Medicine, University of Texas

"DyeVert demonstrated **dye savings of 49%** with uncompromised image quality. The fully disposable, hands free system is **easy to use and saves more dye**. The reaction from the physician community at SCAI was very positive"





Dr Roxana Mehran, MD

AVERT Principal Investigator, Director of Interventional Cardiovascular Research at Mount Sinai Hospital, NY

"Having a system like that in a cath lab is going to be extremely useful, especially for chronic total occlusion (CTO) and chronic kidney disease (CKD) patients. For these high risk individuals, this product is a great thing"

Large addressable market



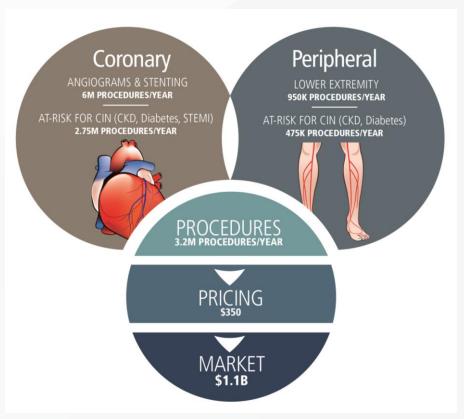
US\$1.1 to 1.4 billion addressable market with initial commercialization focus on the 1.3 million CKD procedures

Market opportunity:

3.2 million procedures per year in the USA and Western Europe which can benefit from DyeVert



- CKD: 1.3 million procedures per year with a 20% occurrence of CI-AKI
- Diabetes: 1.0 million procedures per year with a 15% occurrence of CI-AKI
- STEMI: 440K procedures per year with a 15% occurrence of CI-AKI
- Peripheral: 450K procedures per year with a 15% occurrence of CI-AKI



Ongoing news flow



Osprey remains firmly focused on sales to drive stockholder returns

SALES GROWTH Grow sales team and territories

- Ongoing quarter on quarter sales growth is expected with a larger sales team set to drive commercial success
- Marketing efforts expected to continue growing product sales

R&D Development of R&D portfolio

- Osprey is developing DyeVert Plus, which records dye savings in realtime to comply with new cardiology guidelines
- FDA approval for DyeVert Plus expected in 1H 2017

DYEVERT Development milestones

- Enrolment for DyeVert RCT is complete, findings are expected to be presented in 2016
- DyeVert Plus submitted for innovative new technology award at TCT conference in Q4 2016

ECONOMICS Capitalize on new legislation

- Capitalize on the shift of hospital/physician payments based on "procedure volume" to "improving quality"
- Take advantage of mandatory dye savings guidelines

Appendix

Osprey Medical 5600 Rowland Road | Suite 250 Minnetonka, MN 55343



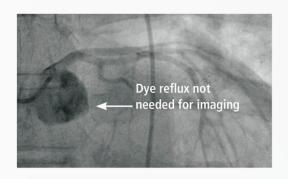
www.ospreymed.com

Osprey – protecting kidneys from dye



Osprey is a revenue stage company with proprietary medical device technology designed to protect kidneys from the harmful effects of dye

- Heart and leg vessels imaging (x-ray) for surgical procedures requires the injection of x-ray dye which is then cleared by the kidney
- The use of dye in these commonly performed procedures can cause damage called Contrast Induced Acute Kidney Damage (CI-AKI)
- Osprey is a medical devices company with proprietary technology designed to protect patients' kidneys by reducing the amount of contrast dye injected



Without Osprey dye reduction



With Osprey dye reduction









Company overview



Osprey's product portfolio comprises the only FDA and CE indicated technology for contrast volume reduction

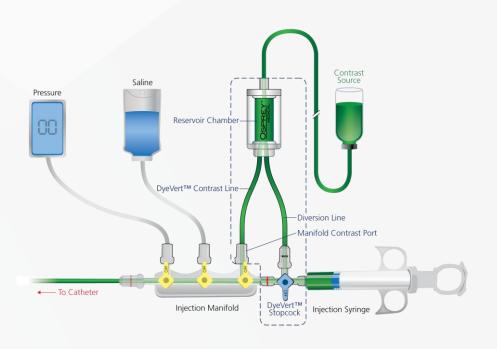
Financial information

CDI price (1-Aug-16)	A\$0.340	
52 week high / low	A\$0.20 / A\$0.85	
Number of CDIs (m)	154.2	
Market Capitalization	A\$52.4m	
Cash (30-Jun-16)	US\$5.8m / A\$7.7m	
Debt (30-Jun-16)	No debt	
Enterprise value	A\$44.7m	

Note: Assumes AUDUSD exchange rate of 0.75

Top stockholders	CDIs	%
Talu Ventures	34.0m	22.1%
Brandon Capital Partners	24.8m	16.1%
Kinetic Investment Partners	8.7m	7.1%

Note: Grey shading represents substantial holdings associated with Osprey Board members, Andy Jane and Chris Nave



Disclaimer



This presentation has been prepared by Osprey Medical, Inc. ("Osprey" or the "Company") for the sole purpose of providing general and background information on Osprey. This presentation does not contain all information necessary to make an investment decision.

This presentation does not constitute an offer, invitation, solicitation or recommendation by any person to sell or apply for securities in Osprey in any jurisdiction, and none of this presentation document or its contents shall form the basis of any contract or commitment. This presentation is not intended to constitute legal, tax or accounting advice or opinion, or financial product advice and should not be relied upon as a representation of any matter that a person should consider in evaluating Osprey. You must not rely on the presentation provided but make your own independent assessment of the presentation and seek and rely upon your own independent taxation, legal, financial or other professional advice in relation to the presentation. This presentation does not take into account an your investment objectives, taxation situation, financial situation or needs. Osprey is not licensed to provide financial product advice in respect of its securities or any other financial products. Cooling off rights do not apply to the acquisition of Osprey securities.

None of Osprey, its officers, directors, employees and agents, nor any other person makes any representation or warranty, express or implied, as to, or endorsement of, Osprey, the accuracy or completeness of any information, statements or representations contained in this presentation and none of them accepts any responsibility or liability for any errors or omissions in this presentation whatsoever.

The information in this presentation is subject to change without notice and Osprey does not have any responsibility or obligation to inform you of any matter arising or coming to their notice, after the date of this presentation, which may affect any matter referred to in this presentation. The distribution of this presentation may be restricted by law and you should observe any such restrictions.

This presentation contains certain forward looking statements which involve known and unknown risks, uncertainties, and other factors which may cause the actual results or performance of Osprey to be materially different from the results or performance expressed or implied by such forward looking statements. Past performance is not necessarily a guide to future performance and no representation or warranty is made as to the likelihood of achievement or reasonableness of any forward looking statements or other forecast.

All figures in the presentation are A\$ thousands on a constant currency basis based on an exchange rates of A\$1: US\$0.75 unless stated otherwise and all market shares are estimates only. The pro-forma historical financial information included in this presentation does not purport to be in compliance with Article 11 of Regulation S-X of the rules and regulations of the US Securities and Exchange Commission. This presentation may contain certain financial data that is "non-GAAP financial measures" under Regulation G under the U.S. Securities Exchange Act of 1934, as amended. The disclosure of such non-GAAP financial measures in the manner included in this presentation would not be permissible in a registration statement under the Securities Act. These non-GAAP financial measures do not have a standardised meaning prescribed by AIFRS and, therefore, may not be comparable to similarly titled measures presented by other entities, nor should they be construed as an alternative to other financial measures determined in accordance with AIFRS. Although we believe these non-GAAP financial measures provide useful information to users in measuring the financial performance and condition of our business for the reasons set out in this presentation, you are cautioned not to placed undue reliance on any non-GAAP financial measures and rations included in this presentation.

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