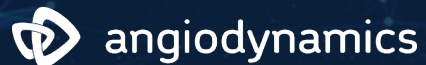


08 | 11 | 2021

ANGIODYNAMICS

Canaccord Genuity Annual Growth Conference 2021

Jim Clemmer, President & CEO
Stephen Trowbridge, EVP & CFO



Notice Regarding Forward-Looking Statements

This presentation contains forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995. All statements regarding AngioDynamics' expected future financial position, results of operations, cash flows, business strategy, budgets, projected costs, capital expenditures, products, competitive positions, growth opportunities, plans and objectives of management for future operations, as well as statements that include the words such as "expects," "reaffirms," "intends," "anticipates," "plans," "projects," "believes," "seeks," "estimates," "optimistic," or variations of such words and similar expressions, are forward-looking statements. These forward-looking statements are not guarantees of future performance and are subject to risks and uncertainties. Investors are cautioned that actual events or results may differ materially from AngioDynamics' expectations, expressed or implied. Factors that may affect the actual results achieved by AngioDynamics include, without limitation, the scale and scope of the COVID-19 global pandemic, the ability of AngioDynamics to develop its existing and new products, technological advances and patents attained by competitors, infringement of AngioDynamics' technology or assertions that AngioDynamics' technology infringes the technology of third parties, the ability of AngioDynamics to effectively compete against competitors that have substantially greater resources, future actions by the FDA or other regulatory agencies, domestic and foreign health care reforms and government regulations, results of pending or future clinical trials, overall economic conditions, the results of on-going litigation, challenges with respect to third-party distributors or joint venture partners or collaborators, the results of sales efforts, the effects of product recalls and product liability claims, changes in key personnel, the ability of AngioDynamics to execute on strategic initiatives, the effects of economic, credit and capital market conditions, general market conditions, market acceptance, foreign currency exchange rate fluctuations, the effects on pricing from group purchasing organizations and competition, the ability of AngioDynamics to obtain regulatory clearances or approval of its products, or integrate acquired businesses, as well as the risk factors listed from time to time in AngioDynamics' SEC filings, including but not limited to its Annual Report on Form 10-K for the year ended May 31, 2021. AngioDynamics does not assume any obligation to publicly update or revise any forward-looking statements for any reason.

In the United States, the NanoKnife System has received a 510(k) clearance by the Food and Drug Administration for use in the surgical ablation of soft tissue, and is similarly approved for commercialization in Canada, the European Union and Australia. The NanoKnife System has not been cleared for the treatment or therapy of a specific disease or condition.

Notice Regarding Non-GAAP Financial Measures

Management uses non-GAAP measures to establish operational goals and believes that non-GAAP measures may assist investors in analyzing the underlying trends in AngioDynamics' business over time. Investors should consider these non-GAAP measures in addition to, not as a substitute for or as superior to, financial reporting measures prepared in accordance with GAAP. In this presentation, AngioDynamics has included adjusted earnings per share. Management uses these measures in its internal analysis and review of operational performance. Management believes that these measures provide investors with useful information in comparing AngioDynamics' performance over different periods. By using these non-GAAP measures, management believes that investors get a better picture of the performance of AngioDynamics' underlying business. Management encourages investors to review AngioDynamics' financial results prepared in accordance with GAAP to understand AngioDynamics' performance taking into account all relevant factors, including those that may only occur from time to time but have a material impact on AngioDynamics' financial results.

MED TECH

Invest for Growth

Thrombus Management

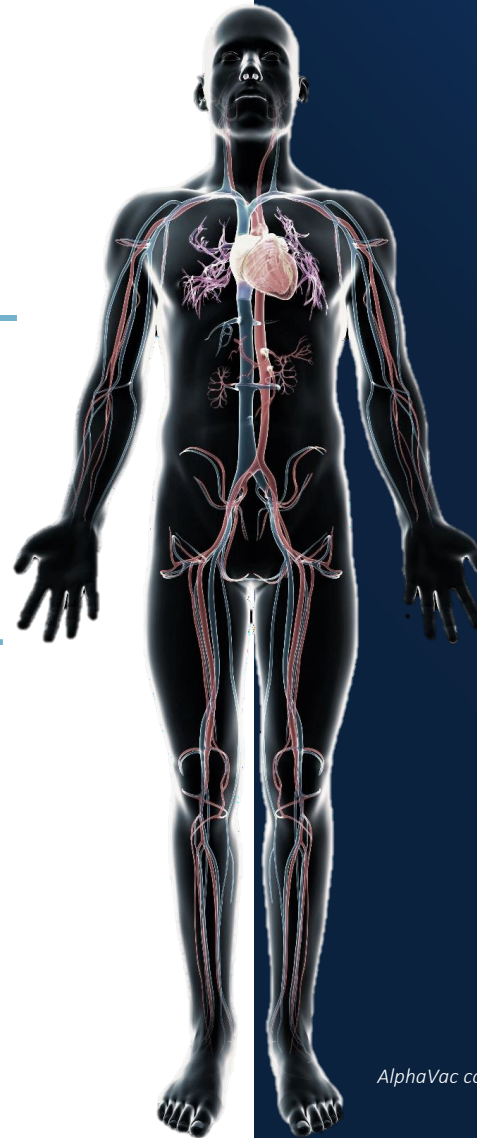
AngioVac Uni-Fuse⁺



Peripheral Atherectomy

AURYON

Irreversible Electroporation



MED DEVICE

Maintain Positioning

Vascular Access Catheters and Accessories

Diagnostic Catheters, Guidewires and Kits

Endovenous Laser Treatment

Microwave & Radiofrequency Tumor Ablation

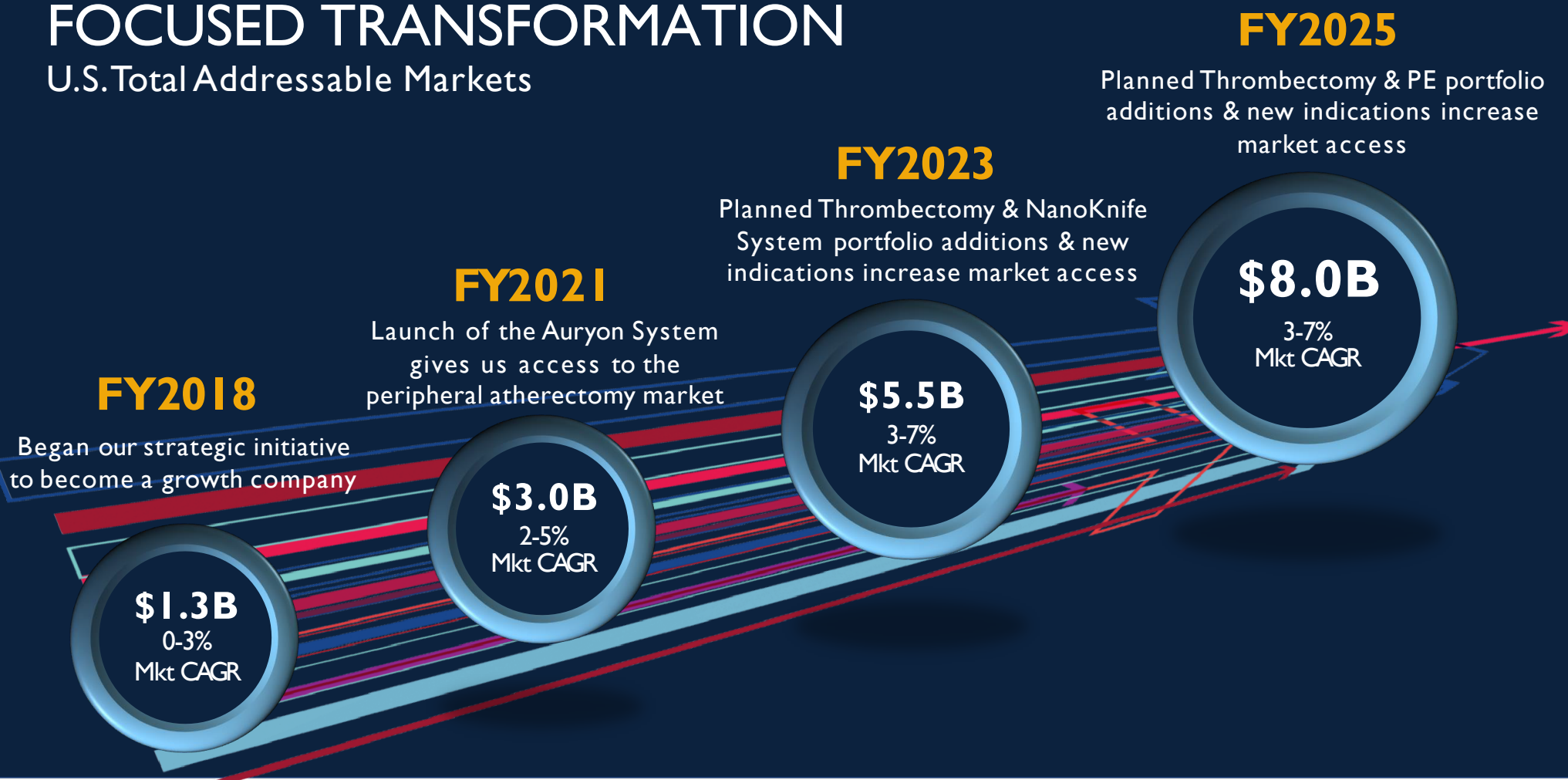
Lung Biopsy Safety

Radiation Treatment Stabilization Balloons

AlphaVac commercial launch planned for 4th quarter calendar year 2021.

FOCUSED TRANSFORMATION

U.S.Total Addressable Markets



VTE Represents 390k Cases Annually

Deep Vein
Thrombosis

DVT

A blood clot that forms in a deep vein, usually the leg, groin or arm

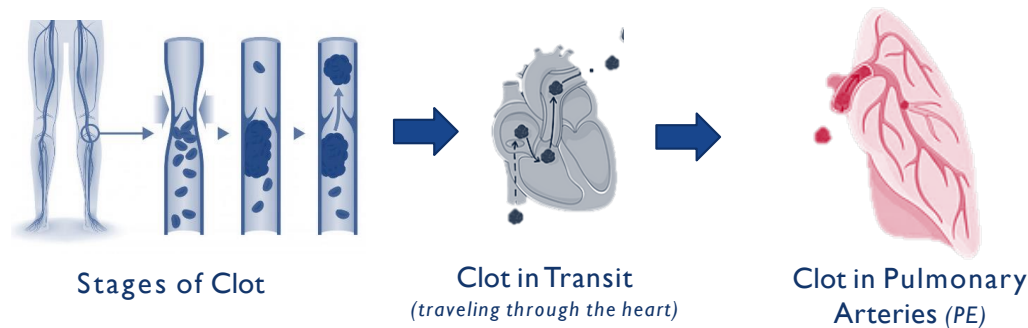
208,000 Iliofemoral Cases¹

Pulmonary
Embolism

PE

A DVT breaks free from a vein wall and travels to the lungs blocking some or all of the blood supply

171,000 High-risk & intermediate-risk PE Cases¹



1. Plovnic, W.J., & Furlong, C. (2020, June). Inari Medical Biomedical Devices and Services. Canaccord Genuity Capital Markets.
2. "Venous Thromboembolism (VTE)." *World Thrombosis Day*. www.worldthrombosisday.org/issue/vte. Illustrations and Images not Produced by AngioDynamics Include: <https://www.vascularmedicare.com/disease-background-dvt-blood-clot-in-the-leg-7-warning-signs-and-symptoms> (emedicinehealth.com)

Venous Thromboembolism

VTE

DVT and PE are collectively referred to as VTE

- **VTE** Affects up to **900k** Americans each year
- **100,000** VTE-Related Deaths in the USA Annually²
- Roughly 30% of Americans who get a blood clot will have a recurrence in less than 10 years
- VTE Costs our US Healthcare system \$10 Billion a year

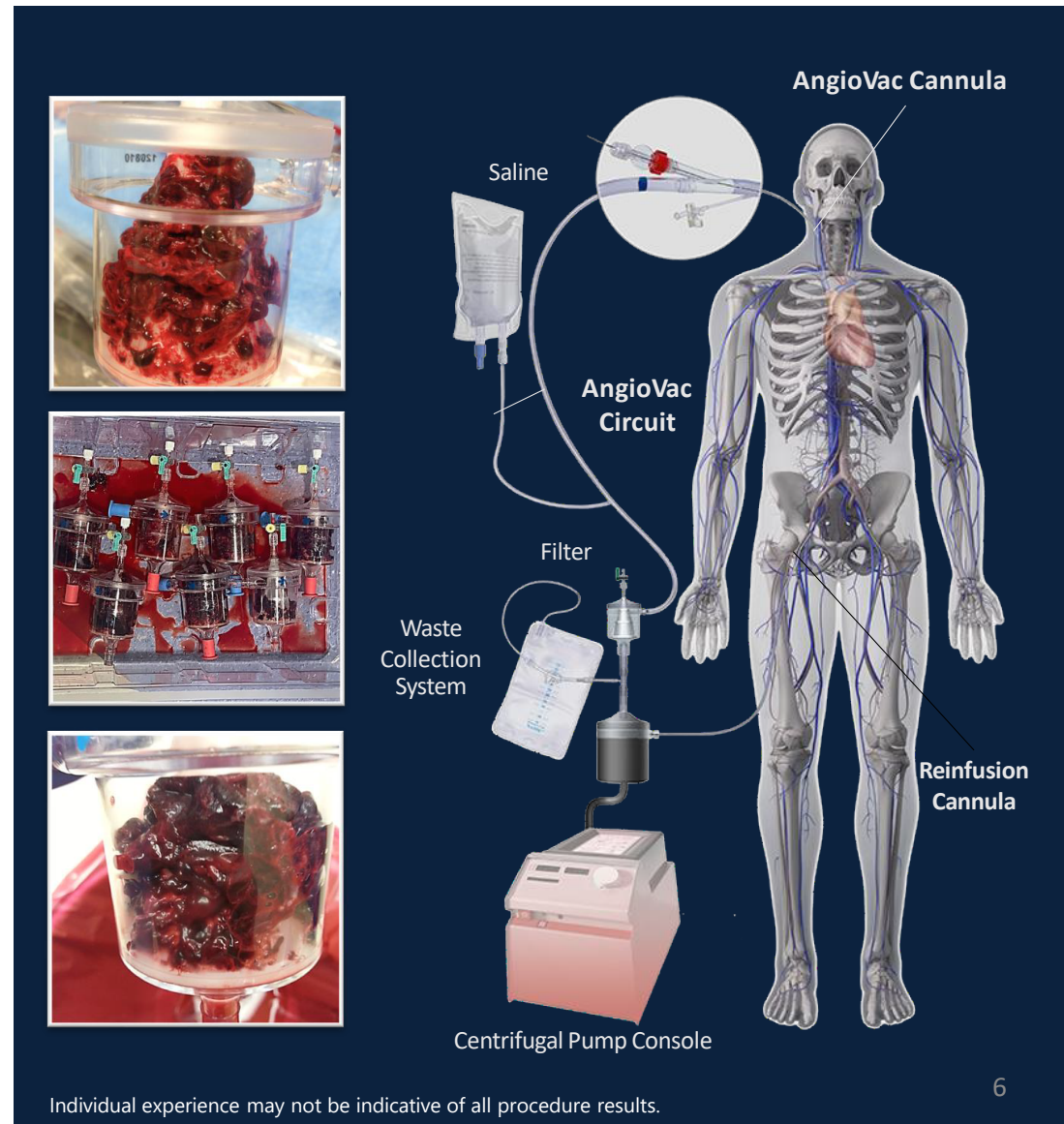
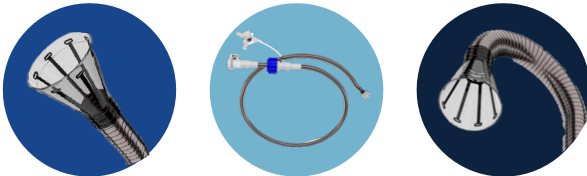
<https://www.cdc.gov/ncbddd/dvt/data.html>

The AngioVac Difference

The AngioVac System allows for the **continuous aspiration** of embolic material such as fresh, soft thrombi or vegetation from the venous system

Utilizing a self-expanding, nitinol reinforced **funnel tip**

Simultaneously reinfusing the patient's own filtered blood to limit procedural blood loss



THE NEXT GENERATION OF ANGIOVAC

Physician requests for use in DVT drive new product development



THE NEXT PORTFOLIO INNOVATION

A purpose-built, innovative product leveraging the strengths of the AngioVac cannula technology with *off-circuit* manual aspiration control



Powerful

Proven funnel tip design allows efficient aspiration and compression of large clot burden



Controlled

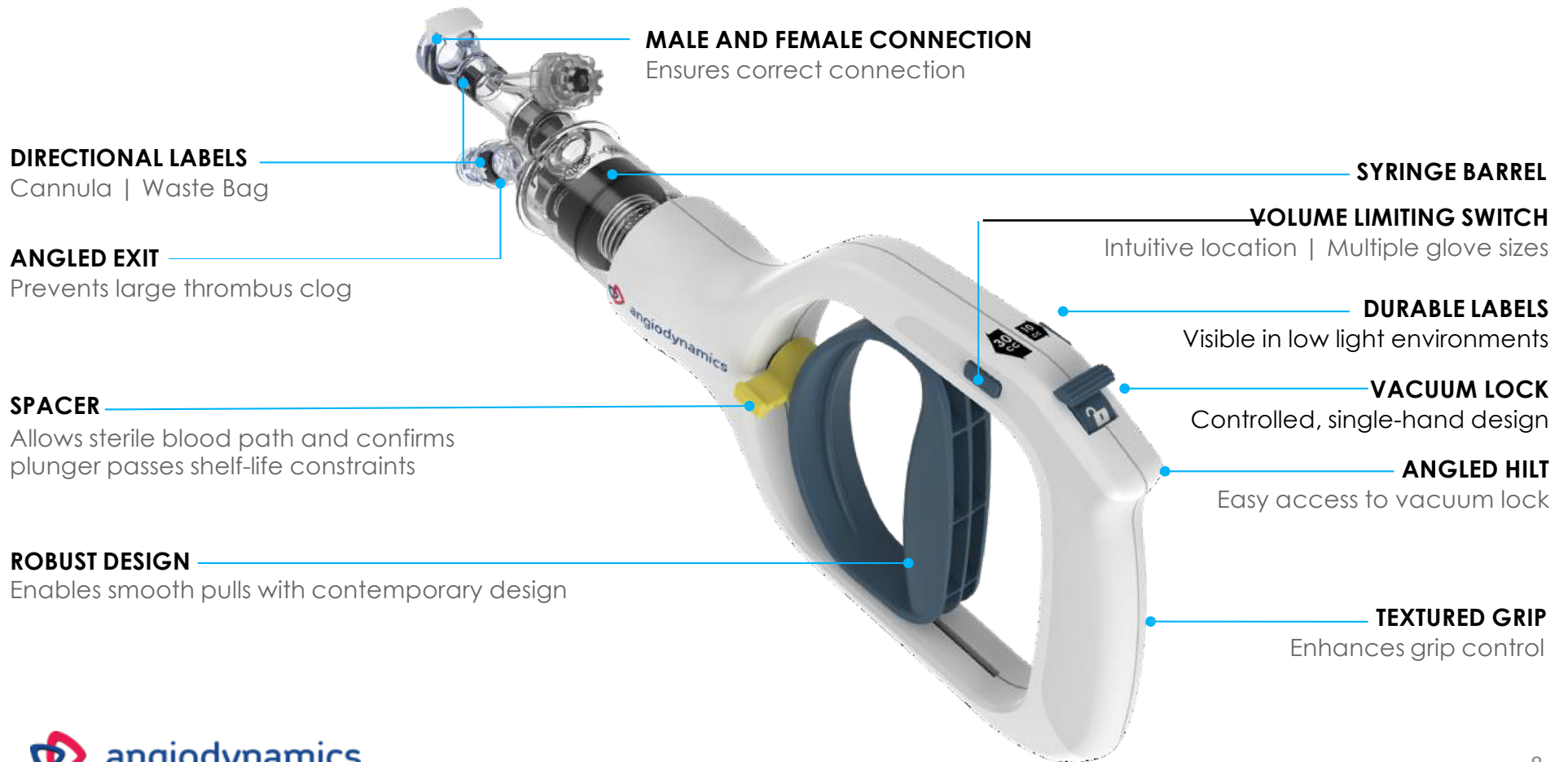
Designed to allow the end-user command and control of the mechanical aspiration



Versatile

Broadens Thrombus Management portfolio and is designed to provide an intuitive, first-line treatment option without the need for lytics and advanced procedural support

Handle | Control Features



F18^{85°} Cannula | Simple Design. Powerful Features.

OBTURATOR

Lubricious Shaft Material
Easy delivery through tortuous anatomy

Tapered Distal Tip
Enhanced navigation and safety

SHEATH

Quarter Turn Valve
Locks tip angle in place

Side Arm Flush Port
Remove air between sheath and cannula

Radiopaque Tapered Soft Tip
Enhanced visibility
Atraumatic transition to obturator

Funnel Shaped Handle
Guided device insertion

Hemostasis Valve
Prevents blood loss during device exchange

CANNULA

Handle Alignment Rib
Indicates cannula curve

Triple Durometer Braided Shaft
Stiffness for powerful push, terminates with a more flexible atraumatic distal end with 1:1 torque

Nitinol Reinforced Funnel Tip
Reliable clot entrapment and removal

Quick Connect

85° Cannula Bend
Enhanced direct-ability

ADDRESSABLE MARKET

Thrombus Procedures by Location



PE
167K Patients
\$1.6B TAM



Right Heart/Atrium
97K Patients
\$77M TAM



IVC/SVC – Caval DVT
20K Patients
\$360M TAM



Ilio-Femoral – DVT
246K Patients
\$700M TAM



Popliteal – DVT
95K Patients
\$300M TAM



RESTORATIVE FLOW THERAPIES

AlphaVac F18⁸⁵ and F13¹⁰ are not cleared by the Food and Drug Administration (FDA). These statements and the subject product have not been evaluated by the FDA. The device is not currently being marketed, nor is it available for sale in any country. AlphaVac and AngioVac are not indicated for PE.

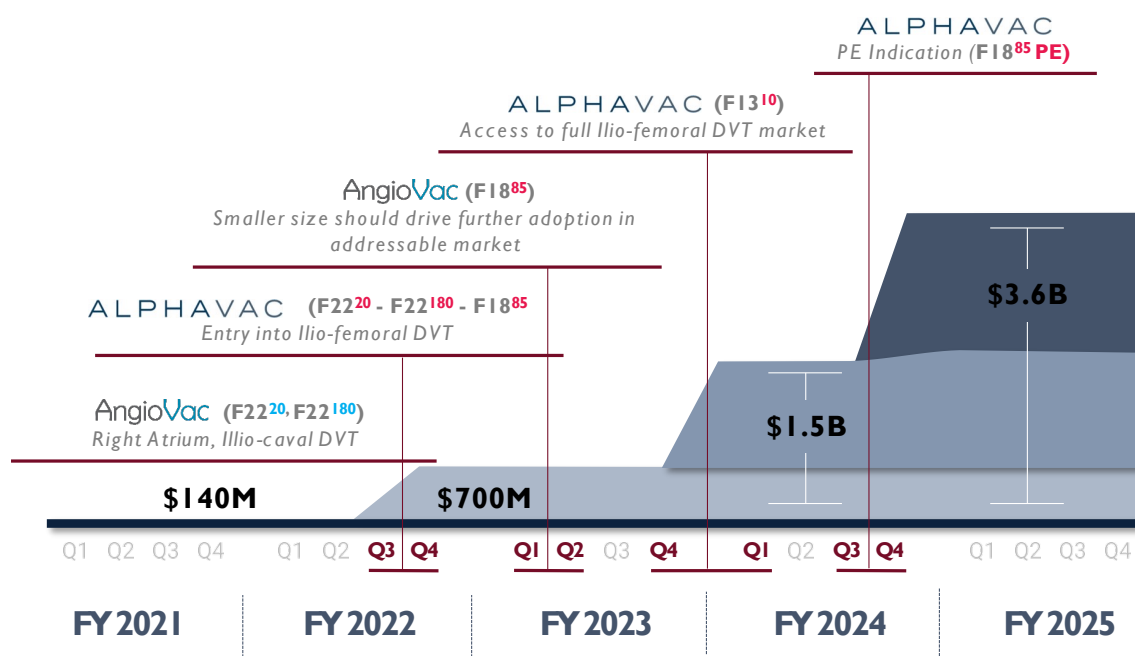
Deliberate Attention to Key Technology Elements

	AngioVac F22 ²⁰	AngioVac F22 ¹⁸⁰	AlphaVac F18 ⁸⁵	AlphaVac F13 ¹⁰	AlphaVac F18 ⁸⁵ PE
Funnel Tip Opening FR Size	42FR	42FR	33FR	~16FR	33FR
Cannula Angle Degree	20° 	180° 	85° 	10° 	85°
Cannula FR Size	22FR Cannula 25FR Sheath	22FR Cannula 25FR Sheath	18FR Cannula 22FR Sheath	~13FR Cannula ~16FR Sheath	18FR Cannula 22FR Sheath
Modality Type	 Shapes, Sizes and Angles will be available in both on/off circuit options (AlphaVac/AngioVac)				
Availability	FY22 Q2/3	FY22 Q2/3	FY22 Q3*	FY23 Q4*	FY24 Q4*

AlphaVac commercial launch planned for 4th quarter calendar year 2021.

THROMBUS MANAGEMENT

Planned Portfolio Additions & U.S. Addressable Markets Expansion



The planned portfolio additions are not guarantees of future performance and are subject to risks and uncertainties, including clearance by the FDA. Investors are cautioned that actual events or results may differ from AngioDynamics' expectations.

Purpose Built, Comprehensive, Thrombus Portfolio

\$1.3B¹

AngioVac

Continuous Aspiration with Simultaneous Reinfusion

F22²⁰ | F22¹⁸⁰ | F18⁸⁵ | 18⁸⁵ PE | F13¹⁰



ALPHA VAC

Multi-purpose Mechanical/Manual Aspiration

F22²⁰ | F22¹⁸⁰ | F18⁸⁵ | 18⁸⁵ PE | F13¹⁰

Uni-Fuse⁺

Catheter Directed Thrombolysis with PE Indication

PERIPHERAL ATHERECTOMY

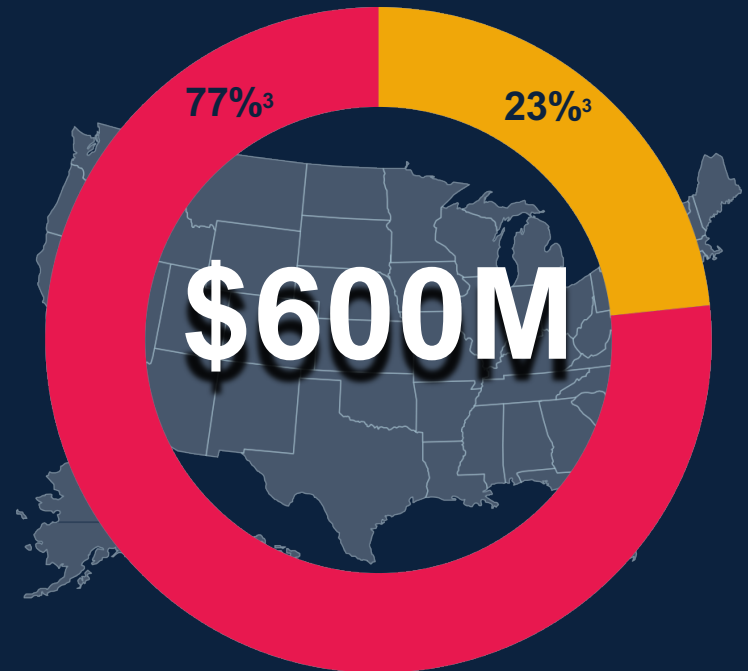
US Addressable Markets & Competitive Landscape

- Over 8 Million² Americans Suffer from PAD
- Over 150,000 Limbs⁴ are Lost Every Year because of PAD
- 50% Mortality Rate⁴ Associated with PAD after Limb Loss

2021 Served Market

MECHANICAL

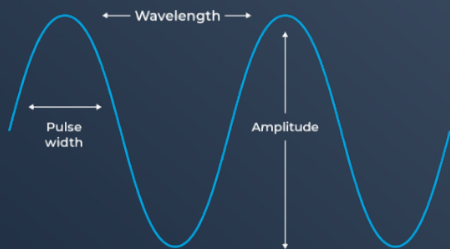
LASER



3. Peripheral Vascular Devices Medtech 360 Market Analysis US 2017. (2016, December). Millenium Research Group, Inc.

4. <https://www.cookmedical.com/peripheral-intervention/10-facts-about-peripheral-arterial-disease/>

AURYON



2.35 mm

Aspiration and Off-Center capabilities and indicated for Peripheral Atherectomy and In-Stent Restenosis (ISR)



2.0 mm

Aspiration capability and indicated for Peripheral Atherectomy and ISR



1.5 mm

Indicated for Peripheral Atherectomy



0.9 mm

Indicated for Peripheral Atherectomy

Why wavelength matters

Each type of tissue interacts differently with a given wavelength

The Auryon System produces a photon energy of 3.5 eV, which is low enough to be nonreactive to vessel endothelium, but high enough to vaporize calcium.^{6,7}

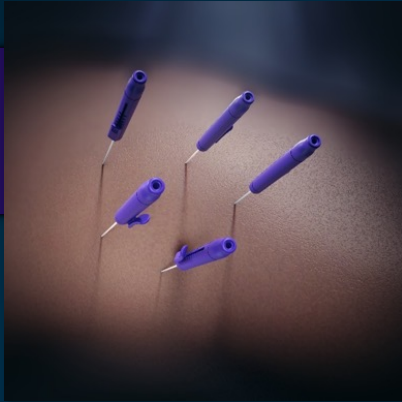
Why pulse width and amplitude matter

Greater amplitude is achieved with shorter pulses, which can deposit energy before thermal diffusion occurs

The Auryon System has a pulse width of 10 to 25 ns, ensuring enough power to target the lesion and spare the vessel.⁵

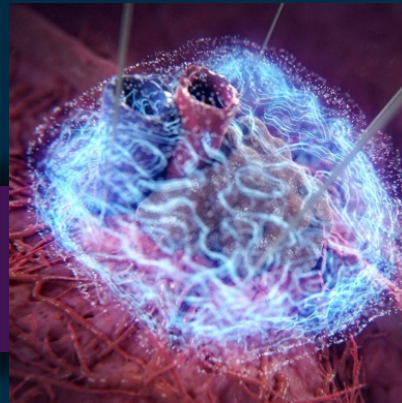
INNOVATION DOCTORS NEED

Expands treatment options and help preserve patient's quality of life



PROBE PLACEMENT

NanoKnife can be confidently used in all segments of an organ.^{10,11}



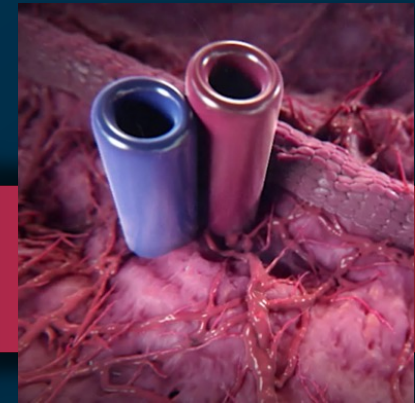
DECELLULARIZATION

Destroys targeted tissue with precise treatment margins.^{10,11}



NON-THERMAL

Spares vital structures by retaining the structural integrity of tissue.^{12,13}



REVASCULARIZATION

Facilitates functional tissue regeneration post-ablation.^{12,13}

10. Lee EW, Thai S, Kee ST. Irreversible electroporation: a novel image-guided cancer therapy. *Gut Liver*. (2010);4(SUPPL. 1):99–104. doi: 10.5009/gnl.2010.4.S1.S99

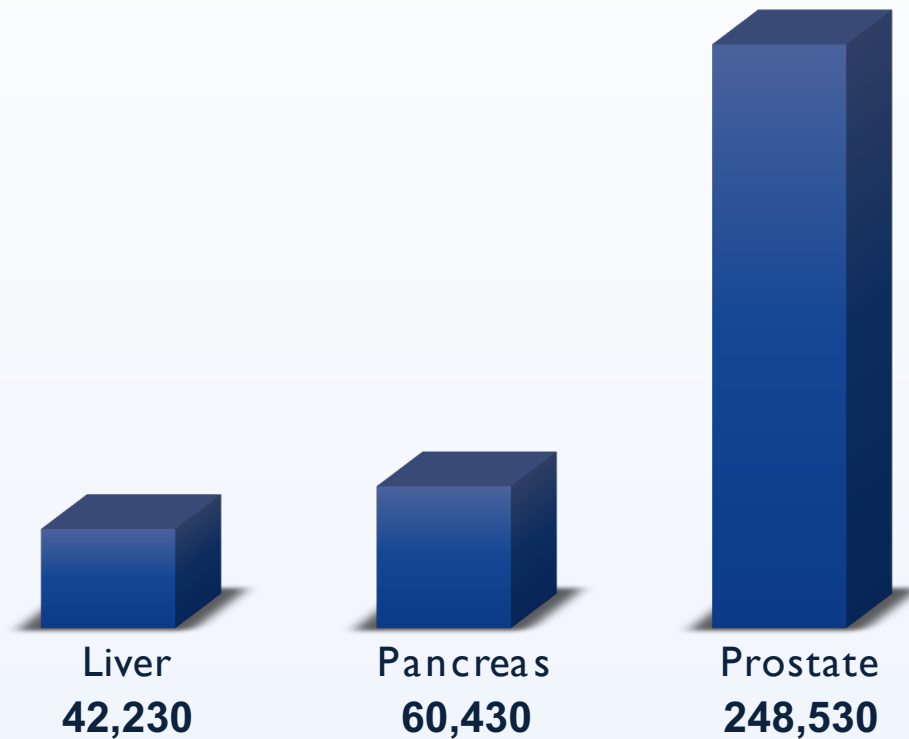
11. Guidance for Selection of NanoKnife Probe Array Configuration and Ablation parameters for the Treatment of Stage III Pancreatic Cancer.

12. Scheltema MJ, Chang JJ, van den Bos W, Gielchinsky I, Nguyen TV, Reijke TM, Siriwardana AR, Böhm M, de la Rosette JJ, Stricker PD. Impact on genitourinary function and quality of life following focal irreversible electroporation of different prostate segments. *Diagn Interv Radiol*. 2018 Sep;24(5):268-275. doi: 10.5152/dir.2018.17374. PMID: 30211680; PMCID: PMC6135060.

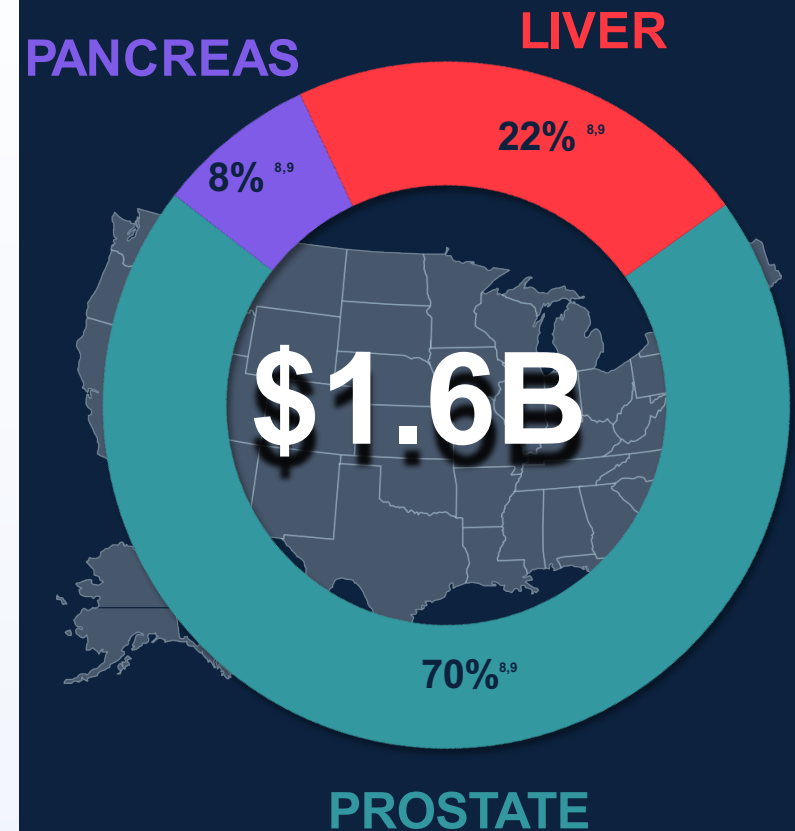
13. Li W, Fan Q, Ji Z, Qiu X, Li Z. The effects of irreversible electroporation (IRE) on nerves. *PLoS One*. 2011 Apr 14;6(4):e18831. doi: 10.1371/journal.pone.0018831. PMID: 21533143; PMCID: PMC3077412.

THE NANOKNIFE SYSTEM

Estimated # of U.S. Patients Diagnosed in 2021⁹



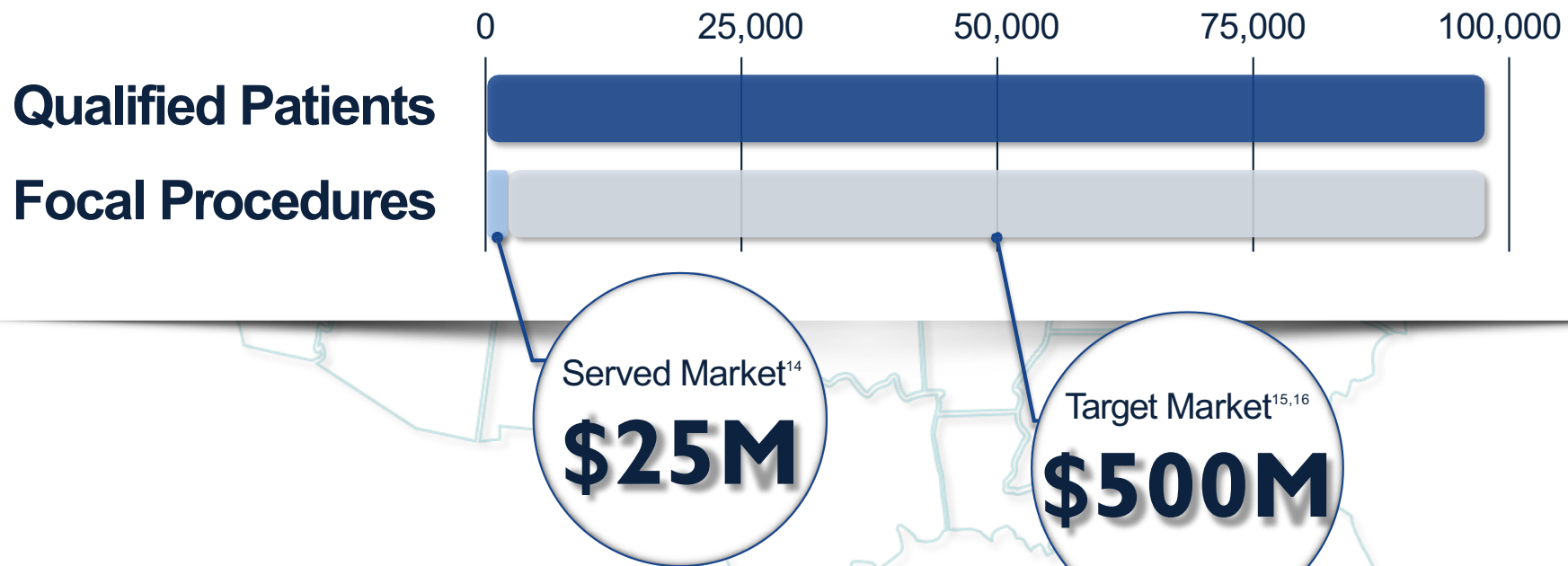
2021 Total Addressable Market (TAM)



8. Interventional Oncology Devices Medtech 360 Market Analysis US 2 (2016, December). Millennium Research Group, Inc.
9. "Cancer Facts & Figures 2021." American Cancer Society, www.cancer.org/research/cancer-facts-statistics/all-cancer-facts-figures/cancer-facts-figures-2021.html.

FOCAL THERAPY

U.S. Served and Target Markets



14. D" Cancer Facts & Figures 2021." American Cancer Society, www.cancer.org/research/cancer-facts-statistics/all-cancer-facts-figures/cancer-facts-figures-2021.html.

15. Definitive Healthcare All-Payor Hospital Outpatient Volume by CPT Code, 02/05/2021

16. Parry MG, Cowling TE, Sujenthiran A, et al. Risk stratification for prostate cancer management: value of the Cambridge Prognostic Group classification for assessing treatment allocation. BMC Medicine. 2020;18(1). doi:10.1186/s12916-020-01588-9

PRESERVE Prostate IDE

SUO-CTC is a clinical research investigator network of 500+ members from more than 250 clinical sites in the US and Canada.



37	SUO-CTC US sites responded to Call for Sites
Up to 20	Sites to be selected, focused on geographic and demographic diversity, high-volume focal therapy institutions
100	Intermediate-risk patients enrolled through 1-year follow up

Primary endpoint: Rate of negative in-field biopsy at 1 year

FINANCIAL GUIDANCE & GOALS

Transformation Toward Double Digit Revenue Growth

AngioDynamics in investment mode throughout the planning horizon

FY22

Revenue \$305M – \$310M
Growth 5% - 7%

Gross Margin ~55%

Adjusted EPS \$0.00 - \$0.05

FY23

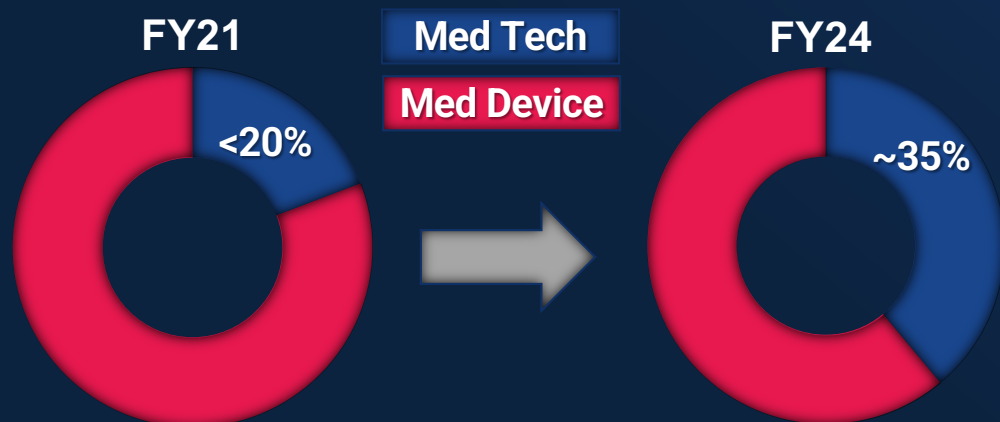
Revenue \$330M – \$336M
Growth 7% - 9%

- Planned significant investment in Med Tech platforms drives top line growth
- Bottom line leverage will ramp slower than top line growth

FY24

Revenue \$360M – \$375M
Growth 10% - 12%

Revenue Contribution



Revenue Growth CAGR



STRATEGIC TRANSFORMATION



PURSUE LARGER, FASTER GROWING MARKETS

Active portfolio management enables us to compete in larger, faster growing markets relying on technology & innovation to produce measurable patient outcomes

DEPLOY FOCUSED RESOURCE DEVELOPMENT

Resource deployment focused in areas that offer better opportunities for success

DRIVE PORTFOLIO TRANSFORMATION

Portfolio transformation & strength driven by R&D, M&A, and Clinical & Regulatory

ATTRACT AND RETAIN TOP TALENT

Strong and innovative portfolio combined with top talent drives value

THANK YOU