

J.P. Morgan

43rd Annual Healthcare Conference January 16, 2025

Jim Clemmer, President & CEO

Forward looking statements



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 AnaioDynamics' 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 (including inflation, labor shortages and supply chain challenges including the cost and availability of raw materials), 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 to 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, 2024. AngioDynamics does not assume any obligation to publicly update or revise any forward-looking statements for any reason.

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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 reported pro forma results, adjusted EBITDA (income before interest, taxes, depreciation and amortization and stock-based compensation); adjusted net income and 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. Please see the tables that follow for a reconciliation of non-GAAP measures to measures prepared in accordance with GAAP.

Angio Dynamics Summary



Our Med Tech segment operates in key markets with unique technologies that deliver proven clinical outcomes

We have a strong track record in portfolio management, R&D, clinical and regulatory expansion, and customer-centric sales & marketing

Our Med Device segment funds investments driving Med Tech growth

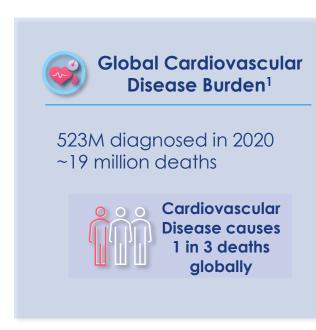
We maintain a debt-free, strong balance sheet

We expect to be Adj. EBITDA positive by FY2025 and cash flow positive by FY2026

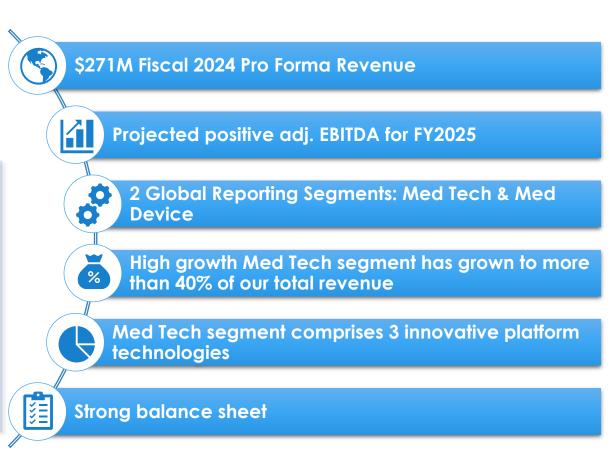
Our company is positioned for sustainable revenue growth and profit for years to come



We are a dynamic, diversified medical technology company committed to expanding treatment options and improving patient outcomes and quality of life by focusing on cardiovascular disease and cancer. Our execution strategy is built on innovative R&D, clinical and regulatory pathway expansion, and customer centric sales performance.









Our broad based clinically focused portfolio targets treating two of the largest global healthcare markets

Cardiovascular Disease

Cancer

Med Tech Venous
Thromboembolism (VTE)
\$5.3B US TAM*



Peripheral Artery
Disease (PAD)
\$760M US TAM*



Prostate Care \$780M US TAM*



Med Device

Uni-Fuse

















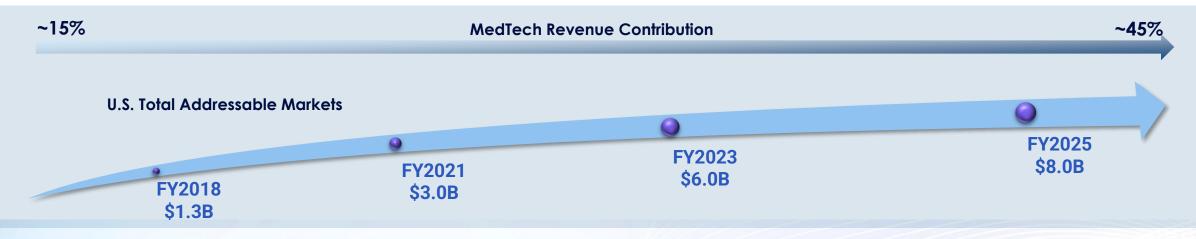
Mabib 4x





Our demonstrated successful execution in portfolio optimization, clinical and regulatory expansion, and product introductions...







...Positions us for sustained growth and profitability in our focus markets









Platform Technologies







 $ilde{ riangle}$ Exploring new indications

Dynamic Markets



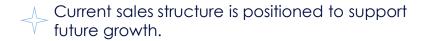


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New dynamics in Percutaneous Coronary
Interventions (PCI) may significantly increase
use of laser atherectomy

Structural heart procedure growth may open new opportunities for AngioVac

Leverageable Infrastructure



Margins and profits continue to expand steadily.

Current G&A structure supports our strategic plan

Manufacturing footprint restructuring estimated to save \$15M in FY 2027



NanoKnife is a unique technology poised to drive change in the standard of care for focal therapy in Prostate, the most diagnosed cancer in men



Cancer

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Thromboembolism (VTE)
\$5.3B US TAM*



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Med Device

Uni-Fuse



















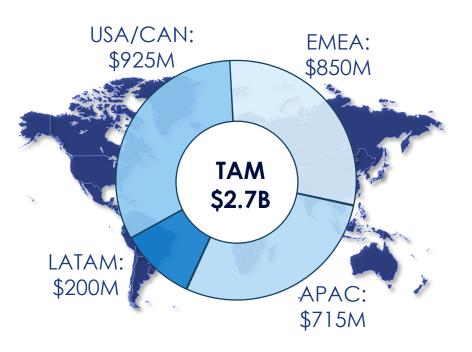


NanoKnife Prostate Care



Men are being forced to compromise between their quality of life or controlling their cancer

MARKET DYNAMICS



Prostate Cancer is Prevalent, On the Rise & Debilitating

The most diagnosed male cancer in 112 countries, including the U.S.⁵

Incidence projected to double by 2040⁵

Current Standard Treatment Options Remain Suboptimal

Active Surveillance³

Radical Surgery⁴

Radiation Therapy⁴

Frectile Dysfunction

Syears 50.2%
10 years 38.7%
15 years 33.7%

Baseline 34%
1-year 85%

Radiation Therapy⁴

Brectile Dysfunction

Frectile Dysfunction

Baseline 30%
1-year 85%

Baseline 30%
1-year 62%

Baseline 32%
1-year 62%

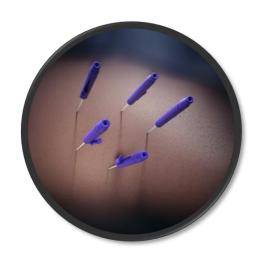
Clinical Research Favors Focal Therapy As "A Middle Ground"

Focal therapy aims to destroy the index lesion while preserving the natural anatomy, continence and erectile function of the man.⁶

NanoKnife Technology



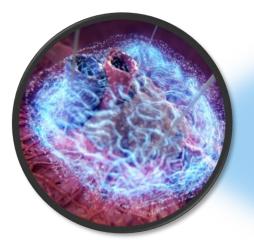
Preserves the underlying structure of tissue giving physicians a more precise tool...



Probe Placement

Up to 6 probes placed per procedure



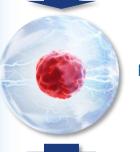


Non-Thermal Decellularization

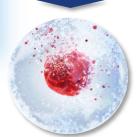
Destroys targeted tissue with precise margins, preserving vital structures and tissue integrity



IRE selectively targets cell membranes⁷



Sufficient voltage permanently opens the ion channels of the cell⁷



Cell loses homeostasis leading to cell death⁷



PRESERVE Pivotal Study



...to minimize quality of life side effects by helping to maintain both sexual and urinary function for patients

PRESERVE

Number of Sites

17 US Sites

Number of Patients

121

Follow Up

12 Months

Efficacy⁸

Safety⁸

84% of patients were free from clinically significant infield disease

3.3% of patients had a device related SAE, all of which resolved

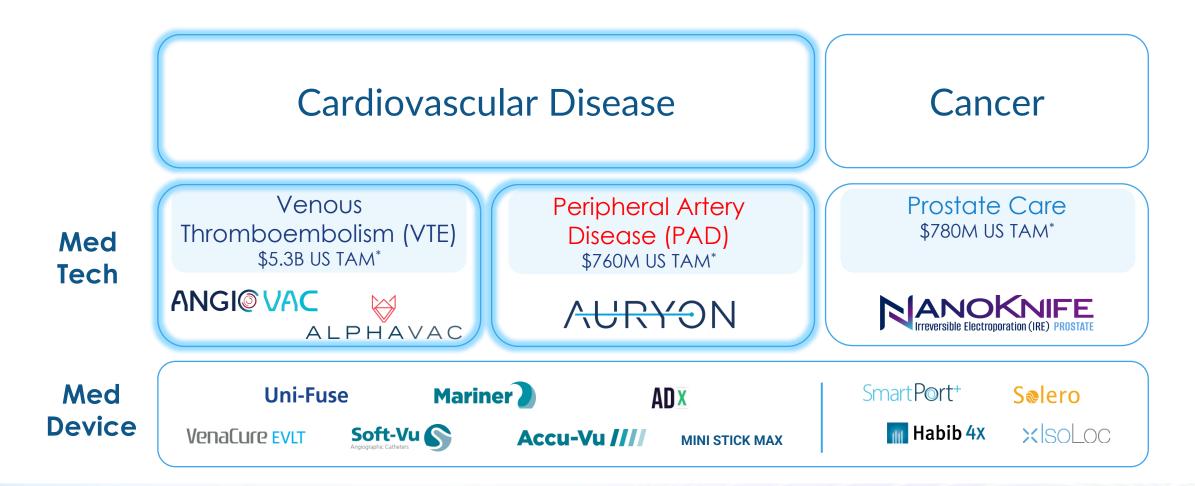
Change from baseline at 1 year^{8, 9}

	IRE	Radical Surgery	Radiation Therapy
Erectile Function	(9%)	(51%)	(30%)
Urinary Continence	(1.2%)	(41%)	(8%)

NanoKnife safely & effectively treats prostate tumors while avoiding the high incidence of erectile dysfunction and incontinence associated with radical surgery and radiation



Differentiated technologies and a comprehensive portfolio enable us to be a strong competitor within high growth Cardiovascular Disease market

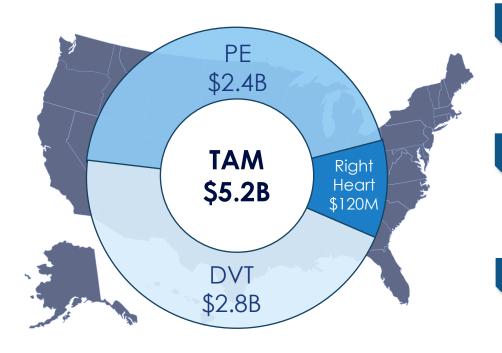


Cardiovascular Venous



Venous Thromboembolism (VTE) represents an attractive, high growth, underpenetrated market opportunity

MARKET DYNAMICS



Pulmonary Embolism (PE)

- Mechanical Thrombectomy is estimated to have penetrated ~25% of the TAM 11,12
- Mechanical Thrombectomy is expected to grow ~16% in 2025 10

Deep Vein Thrombosis (DVT)

- Mechanical Thrombectomy is estimated to have penetrated ~15% of the TAM ^{11,12}
- Mechanical Thrombectomy is expected to grow ~20% in 2025¹⁰

Right Heart

- Growth is driven by rising intravenous drug abuse, an aging population, and increased pacemaker prevalence
- More interventionalists embrace percutaneous techniques and the structural heart market is expanding

AngioVac Technology



Only system allowing for the continuous aspiration and simultaneous reinfusion of blood allowing physicians to treat more complex cases minimally invasively

Aspiration & Simultaneous Reinfusion



On-circuit aspiration provides surgical results via percutaneous access by simultaneously reinfusing blood back in the patient's body, minimizing blood loss.

Proprietary Funnel

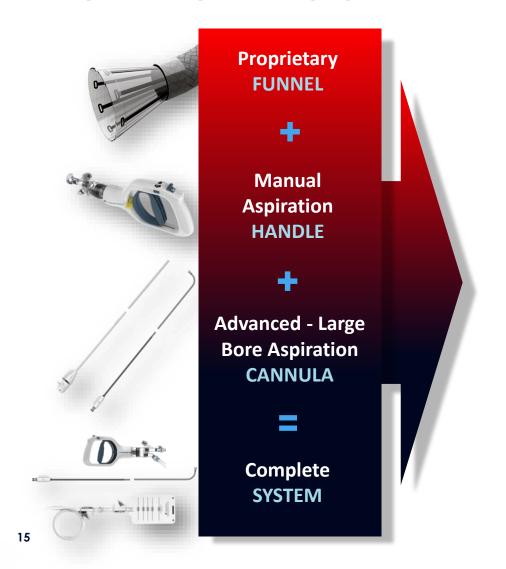
Funnel tip design enables efficient clot aspiration and compression

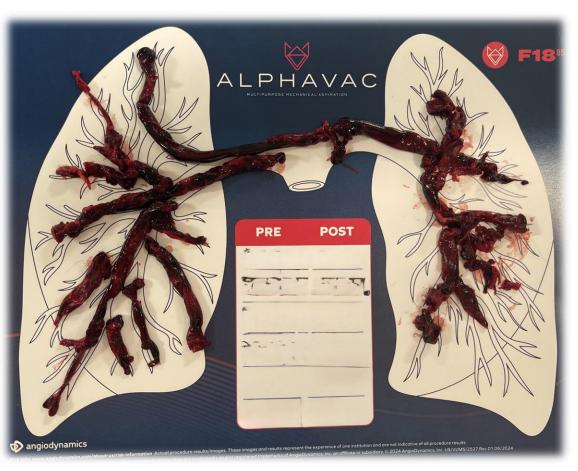
- More than 100 publications on the use of the AngioVac system¹³
- Use of the device has been published for caval thrombi, cardiac masses and thrombi¹⁴

AlphaVac Technology



AlphaVac combines AngioVac cannula technology with off-circuit manual aspiration control for superior aspiration, physician control and maneuverability, uniquely positioning it for PE





*Actual case result

APEX – AV Pivotal Study



AlphaVac received FDA clearance for PE following APEX-AV study demonstrating efficacy and significantly improved reduction in clot burden vs. competitive technologies

APEX-AV

Number of Sites 25 US Sites

Number of Patients 122

Timeline Oct. 2022 – Dec 2023

Significant reduction in the RV/LV ratio¹⁵

• Significant improvement in the RV function

Significant reduction in clot burden

• Large funnel size (33 Fr) may aid in reducing the clot burden

Procedure efficiency

- Atraumatic tip provides easy and efficient navigation in the Pas
- Ability to minimize blood loss

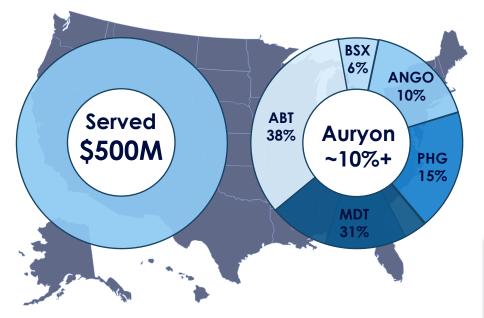
	APEX	FLARE	EXTRACT-PE
	(AlphaVac)	(FlowTriever)	(Indigo)
Reduction in Clot Burden ¹⁶⁻¹⁸	35.1%	9.3%	11.3%

Cardiovascular Arterial



With over 100,000 patients treated, the Auryon Atherectomy System has surpassed \$150M cumulative sales since its September 2020 launch

MARKET DYNAMICS









Treat all levels of calcification 19-21

- Indicated for in-stent restenosis
- Treats above and below the knee (inc. below the ankle)

Protective of vessel wall²¹⁻²³

- Targeted biological reactions to address risk of perforations
- Built-in aspiration to address risk of embolization

Ability to crack medial calcification²⁵

- Shock waves break severe calcification, including medial arterial calcification
- Contributes to plaque modification that may help improve vessel compliance

Designed for hospital and lab 19-21, 24

- Portable, 110V outlet, low noise, touch screen
- Debulk in fewer passes

Auryon Platform Technology



The Auryon System's technology has demonstrated safety and efficacy in PAD and presents an opportunity to expand into additional disease states

PERIPHERAL ARTERY DISEASE (PAD)

- PATHFINDER Registry 12 Mo Published Sept 2023
- Auryon BTK Prospective Study Published May 2024

Safety, Efficacy & Versatility

MicroCT Pre-Clinical Study Published Dec. 2023

Established Secondary Mechanism of Action

AMBITION BTK Enrollment Begins Q1 2025

- RCT: 200 Patients
- Registry: ~1,500 Patients

Auryon w/ Angioplasty vs. Angioplasty alone Below The Knee (BTK)

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CORONARY ARTERY DISEASE (CAD)

Initial European Experience in Coronary Artery Disease

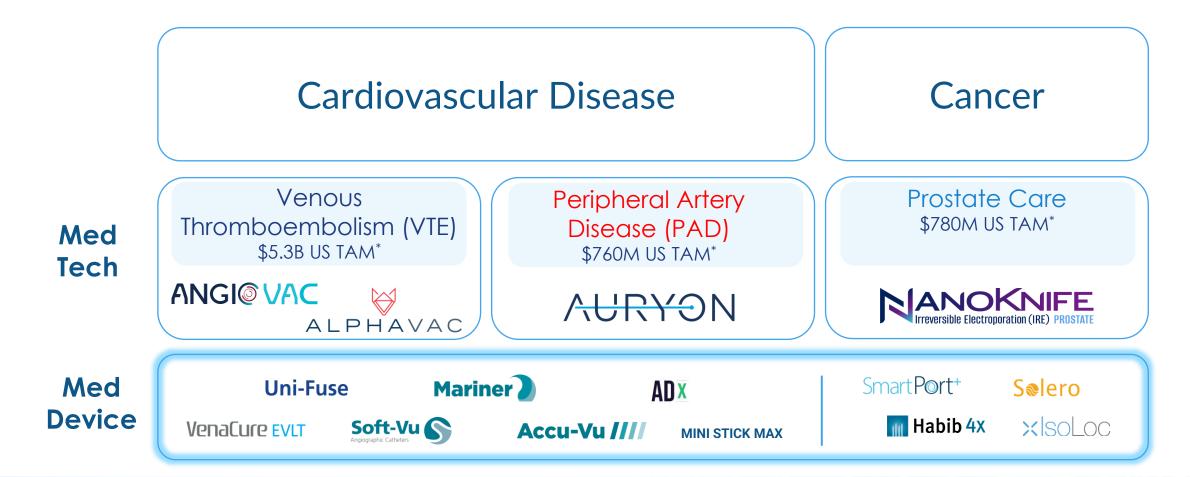
Promising Initial Study Results

~\$900M US TAM*

PMA Consideration



Med Device segment includes trusted brands serving both Cardiovascular and Cancer while providing stable earnings and cash flow enabling Med Tech investment



Fiscal Year 2025 Results and Guidance



Metric	Q2 2025	YTD 2025	Full Year FY2025 Guidance
Net Sales	\$73.0 million	\$140.5 million	\$282 - \$288 million
Med Tech Net Sales Med Tech Growth	\$31.5 million 25%	\$59.5 million 16.8%	12 – 15% YoY growth
Med Device Net Sales Med Device Growth	\$41.5 million (0.4%)	\$81.0 (2.0%)	Flat
Gross Margin	54.7%	54.6%	52 - 53%
Adjusted EBITDA	\$3.1 million	\$2.9 million	\$1.0 - \$3.0 million
Adjusted EPS	(\$0.04)	(\$0.15)	(\$0.34) – (\$0.38)

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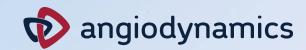
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