UNITED STATES SECURITIES AND EXCHANGE COMMISSION WASHINGTON, D.C. 20549

FORM 8-K

CURRENT REPORT Pursuant to Section 13 or 15(d) of the Securities Exchange Act of 1934

Date of Report (Date of earliest event reported): April 12, 2021

AngioDynamics, Inc.

(Exact Name of Registrant as Specified in Charter)

Delaware

(Commission File

11-3146460 (IRS Employer

Identification No.)

Name of each exchange on which registered

The Nasdaq Global Select Market

(State or Other Jurisdiction of Incorporation)

Number)

000-50761

12110

(Address of Principal Executive Offices) (Zip Code)

14 Plaza Drive Latham, New York

(518) 795-1400

(Registrant's telephone number, including area code)

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions:

Trading Symbol(s) ANGO

□ Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)

□ Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)

□ Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2 (b))

□ Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4 (c))

Securities registered pursuant to Section 12(b) of the Act:

Title of each class

Common Stock, par value \$0.01 per share

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (§230.405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§240.12b-2 of this chapter).

Emerging growth company \Box

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act. \Box

Item 7.01 - Regulation FD Disclosure.

On April 12, 2021, James C. Clemmer, President and Chief Executive Officer of AngioDynamics, Inc. ("AngioDynamics"), and Stephen A. Trowbridge, Executive Vice President and Chief Financial Officer of AngioDynamics, will present at the Needham Virtual Healthcare Conference. The presentation slides are furnished herewith as Exhibit 99.1.

The presentation slides furnished pursuant to Item 7.01 of this Form 8-K (including Exhibit 99.1) shall not be deemed "filed" for purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the "Exchange Act"), or otherwise subject to the liabilities under that Section. Furthermore, the presentation slides shall not be deemed to be incorporated by reference into any filing under the Securities Act of 1933, as amended, or the Exchange Act.

Forward-Looking Statements

This document and its attachments contain 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," "believes," "seeks," estimates," "project", "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, express 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' technology infringes the technology of third parties, 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 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 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 pur

(d)	Exhibits.	
Exhibit No	<u>.</u>	Description
<u>99.1</u>		Presentation slides for the 20th Annual Needham Virtual Healthcare Conference, dated April 12, 2021

SIGNATURE

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

ANGIODYNAMICS, INC. (Registrant)

Date: April 12, 2021

By: /s/ Richard C. Rosenzweig Name: Richard C. Rosenzweig Title: Senior Vice President, General Counsel and Secretary

04 | 12 - 15 | 2021

ANGIODYNAMICS

The 20th Annual Needham Virtual Healthcare Conference

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

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Forward-Looking Statement

Notice Regarding Forward-Looking Statements

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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 usesnon-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' 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 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 incomparing AngioDynamics' performance over different periods. By using thesenon-GAAP measures, management believes that investors get a better 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 tablesthat follow for a reconciliation of non-GAAP measures to measures prepared in accordance with GAAP.



ANGIODYNAMICS

Strategic Transformation



PURSUIT OF LARGER MARKETS

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

FOCUSED RESOURCE DEVELOPMENT

Resource deployment is focused in areas that offer best opportunities for success

PORTFOLIO TRANSFORMATION

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

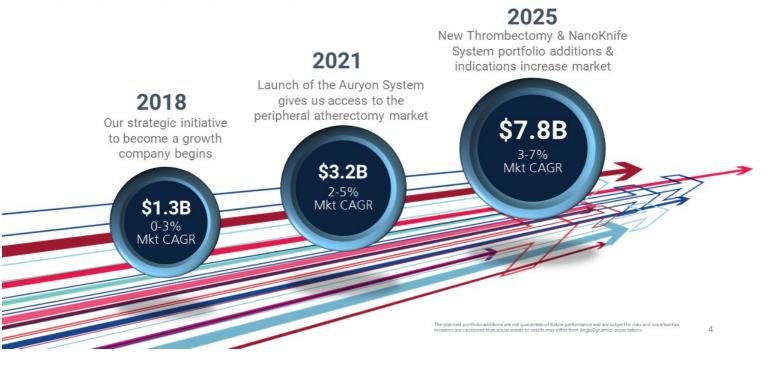
TOP TALENT

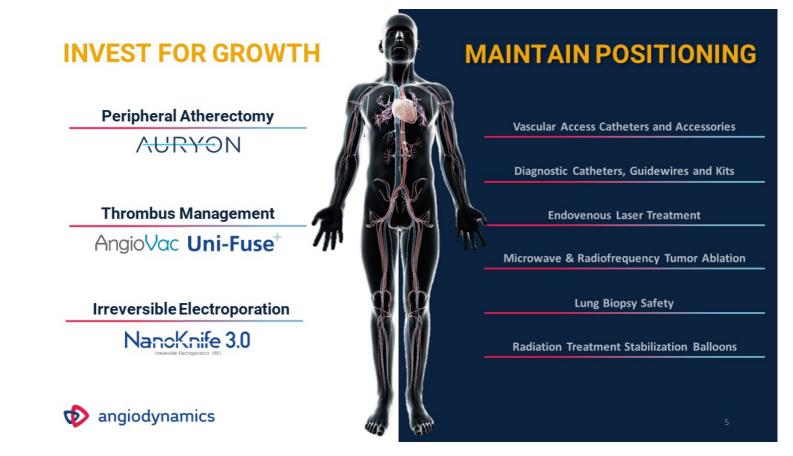
Portfolio combined with talent drives value

3

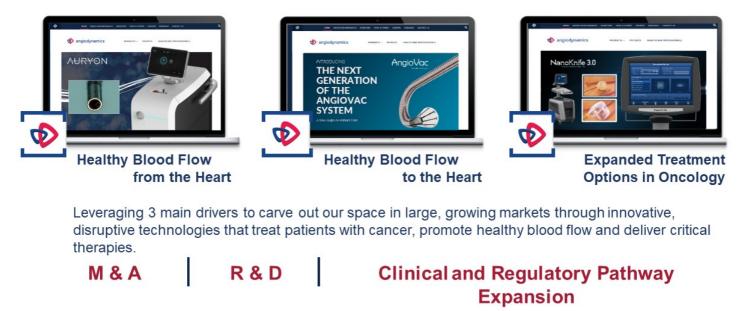
FOCUSED TRANSFORMATION

U.S. Total Addressable Markets





FOCUSED INVESTMENT



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6

04 | 12 - 15 | 2021

PERIPHERAL ATHERECTOMY

ngiodynamics

auryon

Pulse



2.35 mm Aspiration capability and indicated Aspiration and Off-Center capabilities^{for} Peripheral Atherectomy and ISR and indicated for Peripheral





1.5 mm

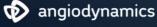
0.9 mm Indicated for Peripheral Atherectomy

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. ^{b, c}

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. ^a



a. Auryon. Instructions for use. AngioDynamics; 2019. b. Herzog A. Bogdan S. Gilkson M. Ishaaya AA, Love C. Selective tissue ablation using laser radiation at 355 nm in lead extraction by a hybrid catheter, a preliminary report. Lasers Surg Med. 2016;44(3):281-287. c. Spectrametics Corporation. CVX-300 Excimer Laser System: Operator's Manual. Version 28. 2019:1-56.

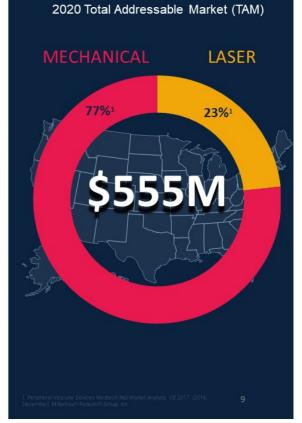
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PERIPHERAL ATHERECTOMY

US Addressable Markets & Competitive Landscape

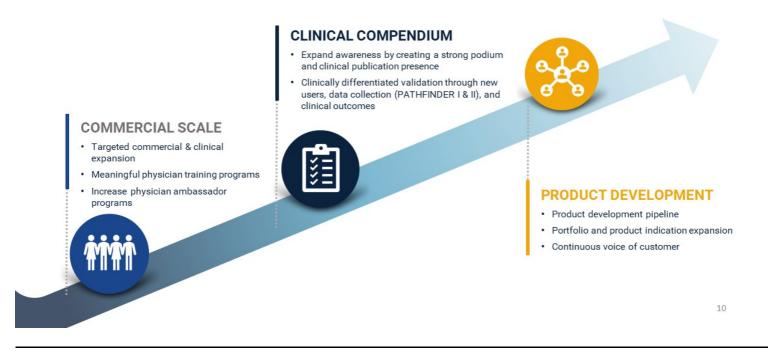
Company	Product	Thrombus	Plaque	Fibrotic	ISR	Calcified
ANGO	Auryon	***	***	**	***	**
Philips	Excimer Laser	**	**		***	*
MDT	TurboHawk		***	*		*
BSC	Jetstream	**	*	*		
CSI	Diamonback 360		**	*		***

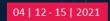
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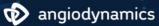
PERIPHERAL ATHERECTOMY

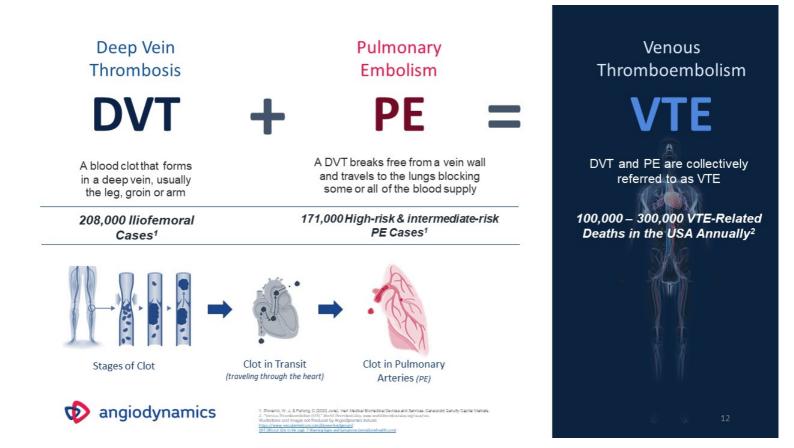
Continuing our momentum of growth





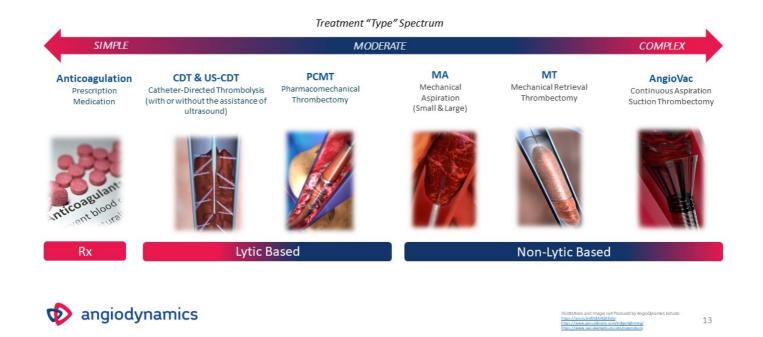
THROMBUS MANAGEMENTAngioVac**Uni-Fuse**⁺

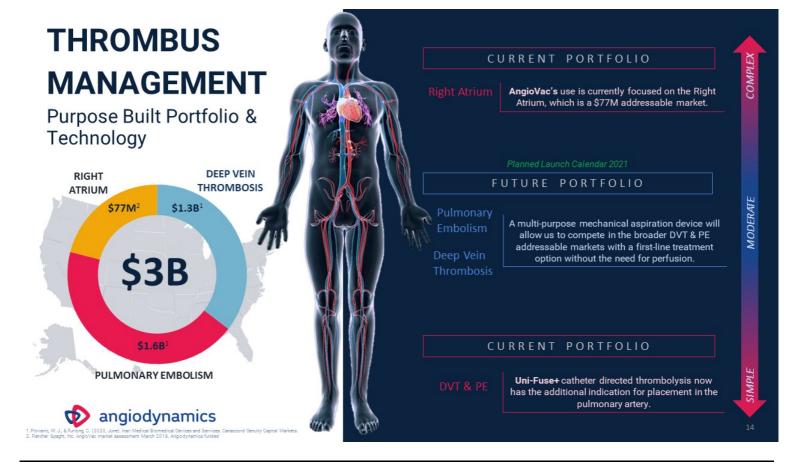




DVT & PE TREATMENT OPTIONS

Percutaneous Thrombectomy







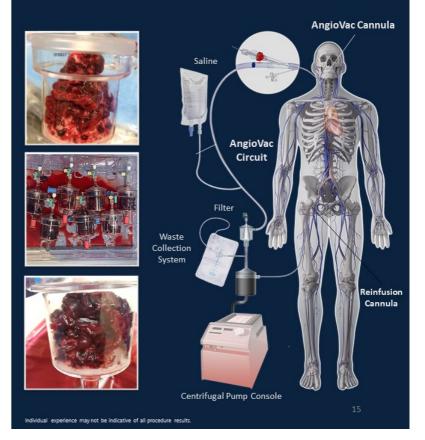
The AngioVac System allows for the **continuous aspiration** of embolic material such as fresh, soft thrombi or vegetation from the venous system

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



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

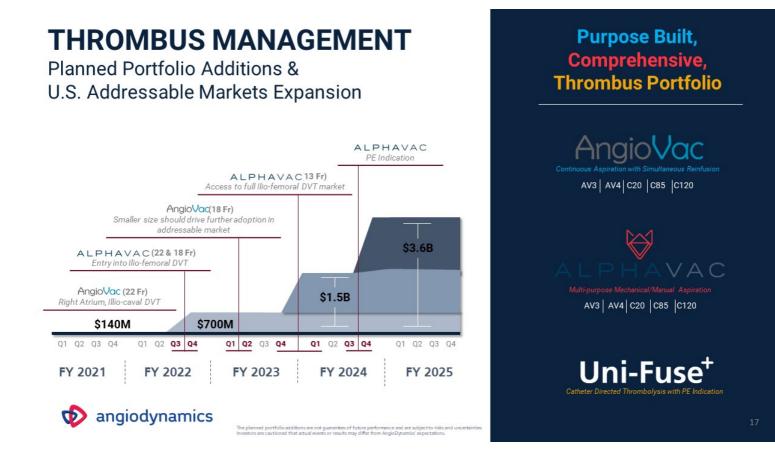
Versatile

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

THE NEXT PORTFOLIO INNOVATION

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

Not approved by the Food and Drug Administration (FDA). These statements and the subject product have not been evaluated by the Food and Drug Administration (FDA). This product is not intended to diagnose, treat, cure, or prevent any disease. The device is not currently being marketed, nor is it available for sale in any country.



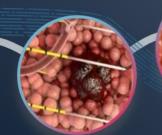
04 | 12 - 15 | 2021

IRREVERSIBLE ELECTROPORATION

ngiodynamics

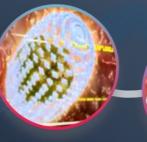
IRREVERSIBLE ELECTROPORATION

Non-thermal energy destroys cells while preserving critical structures



Probe placement

Placement of probes around a target area provides effective treatment coverage a-d



Electrical field

Electrical pulses cause pores to form within the cellular membrane leading to nonthermal cell death a-d



Decellularization

Target tissue undergoes complete decellularization while preserving the underlying structure of blood vessels, nerves, ducts, and tissue a-d



Non-thermal

By preserving those underlying structures the potential for revascularization of treated tissue is maintained^{a-d}



Revascularization

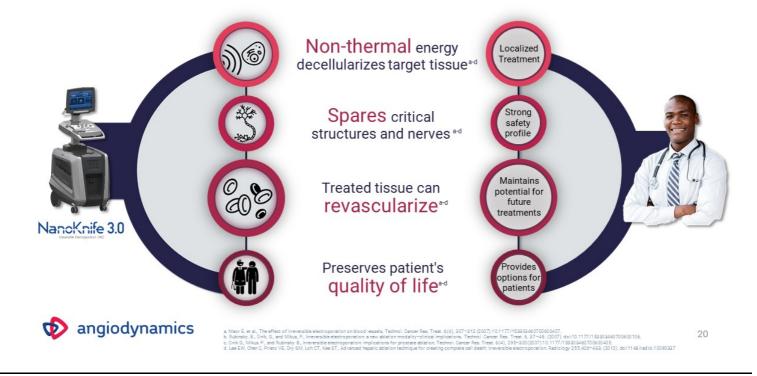
The NanoKnife System's unique technology enables physicians to provide localized treatments in locations and at times previously unavailable to them a-d

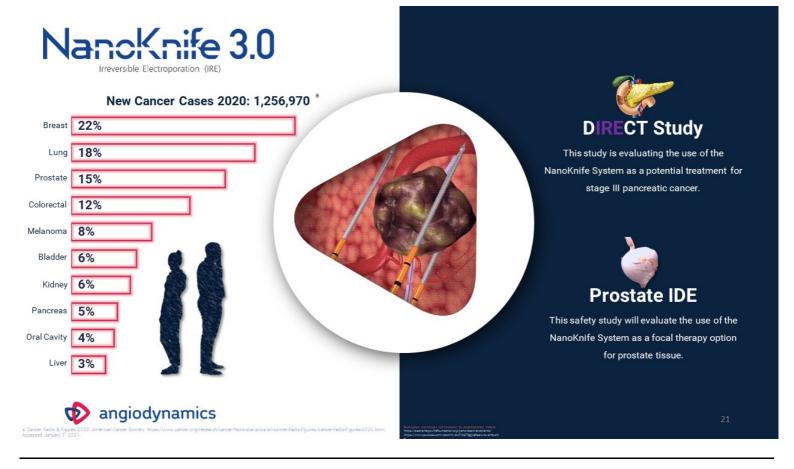


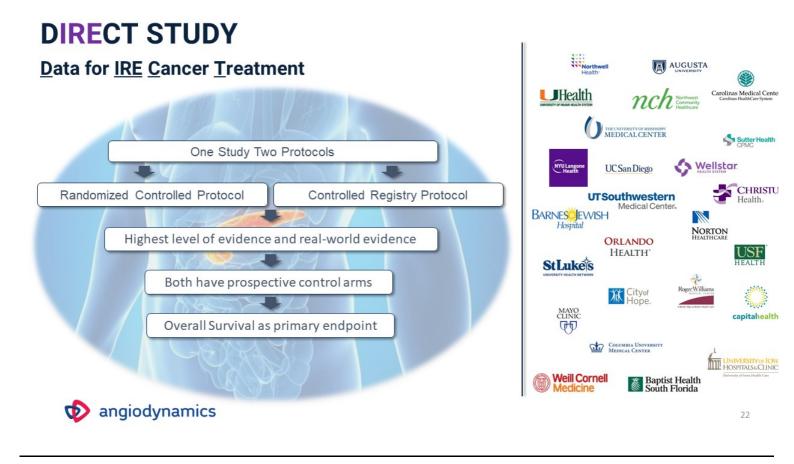
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PROVIDES THE INNOVATION DOCTORS NEED

Expands treatment options and helps preserve patient's quality of life^{®d}







CLINICAL EXPANSION

Within the U.S.

191,930 men will

27% will undergo radiotherapy of which 66% report E.D. at 36 months

31% will undergo a radical prostatectomy of which **79%** report E.D. at 36 months _{b,c}

42% will undergo active surveillance

An estimated **38%** of these patients are ideal candidates for focal therapy_d



Focal therapy

bridges the gap between whole gland treatment and active surveillance in an attempt to maximize quality of life

by avoiding the effects of whole gland radiation or surgery.

However... adoption remains low (<5%) due to gaps within existing technology.

be diagnosed with prostate cancer in 2020.

An estimated **38%** of

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NANOKNIFE PLATFORM



Expanding Indications

Pancreas pivotal study underway (DIRECT)

Prostate safety study underway

FDA Pre-Sub meeting regarding prostate tissue indication complete



Expanded Reimbursement

Inpatient ICD-10 Code (2018) DRGs (2019)

Organ Agnostic CPT Code (2020)

Outpatient (2021) ASC (2021)



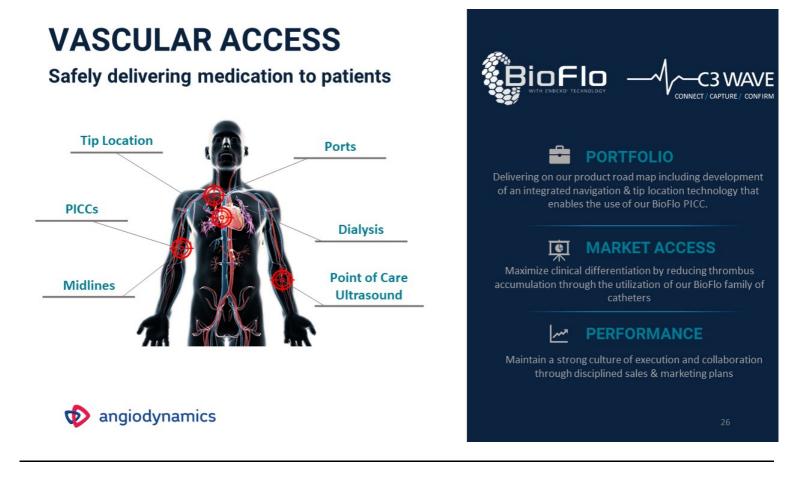


24

04 | 12 - 15 | 2021

VASCULAR ACCESS

🔊 angiodynamics



Q3 YTD FY2021 Highlights

Financial Performance

in millions (except per share data)	YTD FY2021	YTD FY2020	YOY Change
Revenue	\$214.2	\$205.8	4.1%
Gross Margin	53.4%	58.3%	(490 bps)
Net Loss	(\$12.1)	(\$9.7)	(\$2.4)
GAAP EPS	(\$0.32)	(\$0.26)	(\$0.06)
Adjusted EPS	\$0.05	\$0.15	(\$0.10)
Adjusted EBITDA	\$15.0	\$17.5	(\$2.5)
Free Cash Flow	\$7.3	(\$24.2)	\$31.5
Please see Appendix A for reconciliation of GAAP to Non-GAAP.			

in millions	Q3 FY2021	Q4 FY2020	YOY Change
Cash	\$54.5	\$54.4	\$0.1
Debt	\$30.0	\$40.0	(\$10.0)



Growth From Key Products

Key Product Categories	Q1 FY2021	Q2 FY2021	Q3 FY2021	YTD FY2021
AngioVac®	46%	24%	27%	31%
NanoKnife [®] Disposables	(5%)	30%	(7%)	4%
Auryon*	\$1.1	\$2.1	\$3.3	\$6.5

* Auryon reflects revenue contribution in quarter vs growth rate due to acquisition date in FY20.

ANGIODYNAMICS

Strategic Transformation



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PORTFOLIO TRANSFORMATION

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

TOP TALENT

Portfolio combined with talent drives value

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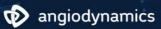
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Reconciliation of GAAP to Non-GAAP Net Income and EPS

		Nine Mor	ths Ended		
(in thousands, except per share data)	Fe	b 28, 2021	Feb 29, 2020		
		(unau	idited)		
Net loss	s	(12,080)	\$	(9,720)	
Amortization of intangibles		13,838		13,417	
Change in fair value of contingent consideration		(290)		116	
Acquisition, restructuring and other items, net (1)		3,057		4,486	
Write-off of deferred financing fees (2)		-		593	
Tax effect of non-GAAP items (3)		(2,606)		(3,205)	
Adjusted net income	\$	1,919	\$	5,687	
	Fe	b 28, 2021	Fel	5 29, 2020	
		(unau	idited)		
Diluted loss per share	\$	(0.32)	s	(0.26)	
Amortization of intangibles		0.36		0.35	
		0.36 (0.01)		0.35	
Change in fair value of contingent consideration				0100	
Change in fair value of contingent consideration Acquisition, restructuring and other items, net (1)		(0.01)		-	
Change in fair value of contingent consideration Acquisition, restructuring and other items, net (1) Write-off of deferred financing fees (2)	_	(0.01)		0.12	
Amortization of intangibles Change in fair value of contingent consideration Acquisition, restructuring and other items, net (1) Write-off of deferred financing fees (2) Tax effect of non-GAAP items (3) Adjusted diluted earnings per share	5	(0.01) 0.08	\$	0.12	

(1)

Includes costs related to merger and acquisition activities, restructurings, and unusual items, including asset impairments and write-offs, certain litigation, and other items. Deferred financing fees related to the old credit agreement were written offduring the second quarter of fiscal year 2020. Adjustment to reflect the income tax provision on a non-GAAP basis has been calculated assuming no valuation allowance on the Company's U.S. deferred tax assets and an effective tax rate of 23% for the periods ended February 28, 2021 and February 29, 2020. (2) (3)

(4) Diluted shares may differ for non-GAAP measures as compared to GAAP due to a GAAP loss.



Reconciliation of Net Loss to Adjusted EBITDA

	Nine Months Ended				
(in thousands)		b 28, 2021	Fel	Feb 29, 2020	
	(unaudited)				
Net loss	\$	(12,080)	\$	(9,720)	
Income tax benefit		(2,033)		(1,506)	
Interest expense, net		676		672	
Depreciation and amortization		19,276		17,434	
Change in fair value of contingent consideration		(290)		116	
Stock based compensation		6,398		5,998	
Acquisition, restructuring and other items, net (1)		3,057		4,486	
Adjusted EBITDA	\$	15,004	s	17,480	

Includes costs related to merger and acquisition activities, restructurings, and unusual items, including asset impairments and write-offs, certain litigation, and other items. (1)

Reconciliation of Free Cash Flow

Nine Months Ended (in thousands) Feb 28, 2021 Feb 29, 2020 (unaudited) 11,894 \$ (18,434) Net cash provided by (used in) operating activities \$ (4,567) (5,756) 7,327 \$ (24,190) Additions to property, plant and equipment s Free Cash Flow