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): January 10, 2022

AngioDynamics, Inc. (Exact Name of Registrant as Specified in Charter)

Delaware	000-50761		11-3146460
(State or Other Jurisdiction of Incorporation)	(Commission File Number)		(IRS Employer Identification No.)
14 Plaza Driv	ve Latham, New York	12110	
(Address of Prin	ncipal Executive Offices)	(Zip Code)	
	(518) 795-1400		
(Registrant)	's telephone number, including ar	rea code)	
Check the appropriate box below if the Form 8-K filing is intended	d to simultaneously satisfy the fili	ng obligation of the reg	istrant under any of the following provisions:
$\hfill\Box$ Written communications pursuant to Rule 425 under the Security	rities Act (17 CFR 230.425)		
$\hfill\Box$ Soliciting material pursuant to Rule 14a-12 under the Exchange	ge Act (17 CFR 240.14a-12)		
$\hfill\Box$ Pre-commencement communications pursuant to Rule 14d-2(l	b) under the Exchange Act (17 C	FR 240.14d-2 (b))	
$\hfill\Box$ Pre-commencement communications pursuant to Rule 13e-4(e	c) under the Exchange Act (17 C	FR 240.13e-4 (c))	
Securities registered pursuant to Section 12(b) of the Act:			
Title of each class	Trading Symbol(s)	<u>Name of ea</u> registered	ch exchange on which
Common Stock, par value \$0.01 per share	ANGO	NASDAQ G	lobal Select Market
Indicate by check mark whether the registrant is an emerging grow 2 of the Securities Exchange Act of 1934 ($\$240.12b-2$ of this charge)	, ,	405 of the Securities A	ct of 1933 (§230.405 of this chapter) or Rule 12
Emerging growth company \square			
If an emerging growth company, indicate by check mark if the regimencial accounting standards provided pursuant to Section 13(a) $^{\circ}$		extended transition pe	eriod for complying with any new or revised

Item 7.01 - Regulation FD Disclosure.

On January 10, 2022, James Clemmer, President and Chief Executive Officer of AngioDynamics, Inc. ("AngioDynamics"), and Stephen Trowbridge, Executive Vice President and Chief Financial Officer of AngioDynamics, will present at the 24th Annual Needham Virtual Growth 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," "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 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 and labor shortages), 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 a

Item 9.01 - Financial Statements and Exhibits.

(d) Exhibits.

Exhibit No. Description

99.1 Presentation slides for the 24th Annual Needham Virtual Growth Conference, dated January 10, 2022

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: January 10, 2022

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



Notice Regarding Forward-Looking Statements

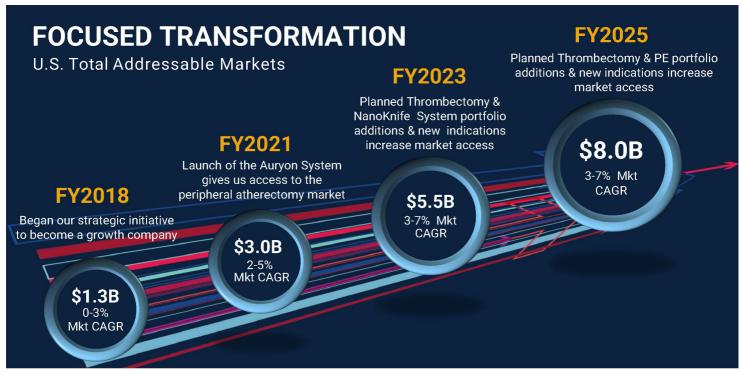
This presentation contains forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995. All statements regarding AngioDynamics' expected future financial position, results of operations, cash flows, business strategy, budgets, projected costs, capital expenditures, 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 (including inflation and labor shortages), 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

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 EBITDA, adjusted net income (loss), 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.





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

MED TECH Invest for Growth

Thrombus Management

AngioVac Uni-Fuse+



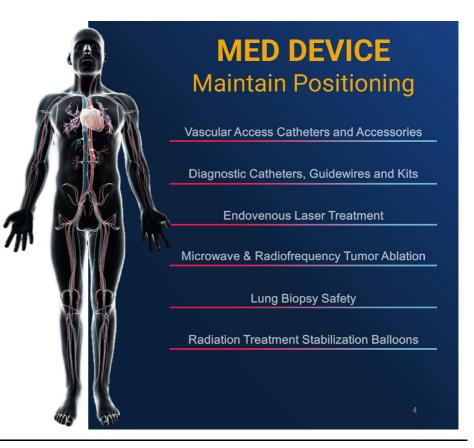
Peripheral Atherectomy

AURYON

Irreversible Electroporation







MED TECH

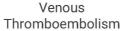
THROMBUS MANAGEMENT





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Deep Vein **Thrombosis** **Pulmonary Embolism**





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

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

DVT and PE are collectively referred to as VTE

208,000 Iliofemoral Cases¹

171,000 High-risk & intermediate-risk PE Cases¹ 100,000 VTE-Related Deaths in the USA Annually2









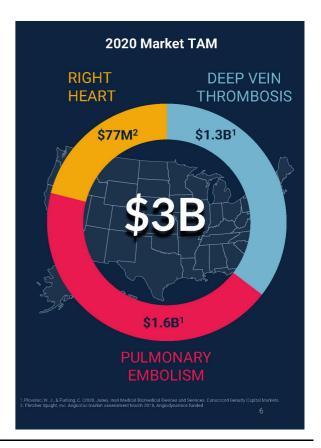


Stages of Clot

Clot in Transit (traveling through the heart)

Clot in Pulmonary Arteries (PE)







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

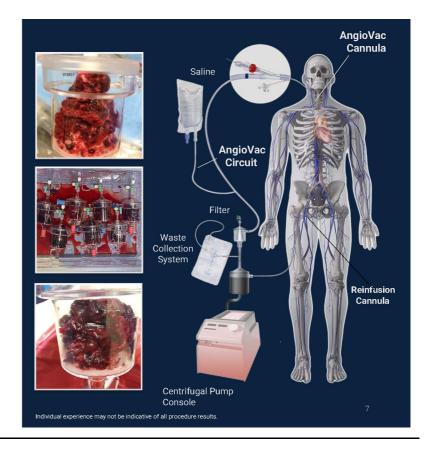
Simultaneously reinfusing the patient's own filtered blood to limit procedural blood loss







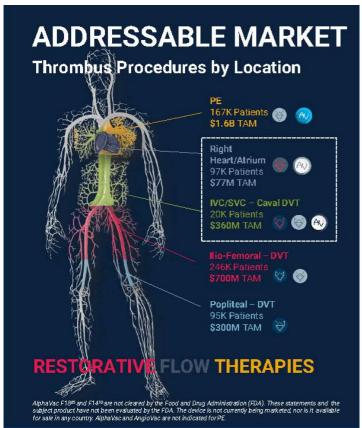


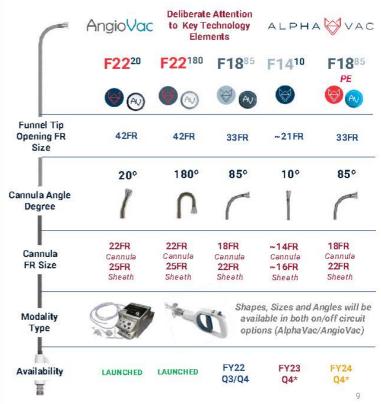




Control Features for Handle & Cannula

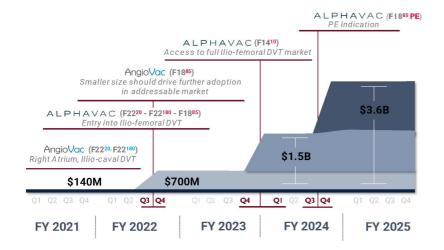






THROMBUS MANAGEMENT

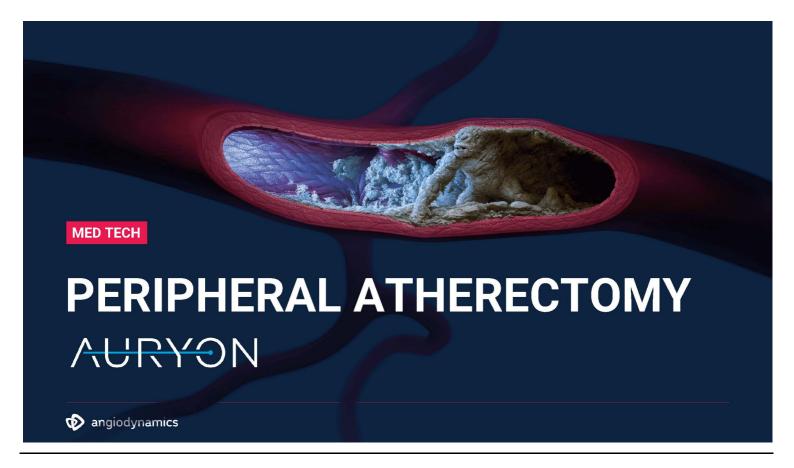
Planned Portfolio Additions & U.S. Addressable Markets Expansion



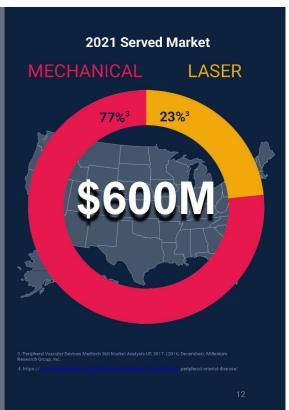


The planned portfolio additions are not guarantees of future performance and are subject to risks and uncertainties, including clearance by the FDA. Investors are cautioned that actual events or results may differ from AngioDynamics' expectations.

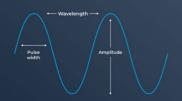














2.35 mm

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 Atherectom



0.9 mm

Indicated for Peripheral Atherectom

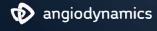
Why wavelength matters

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

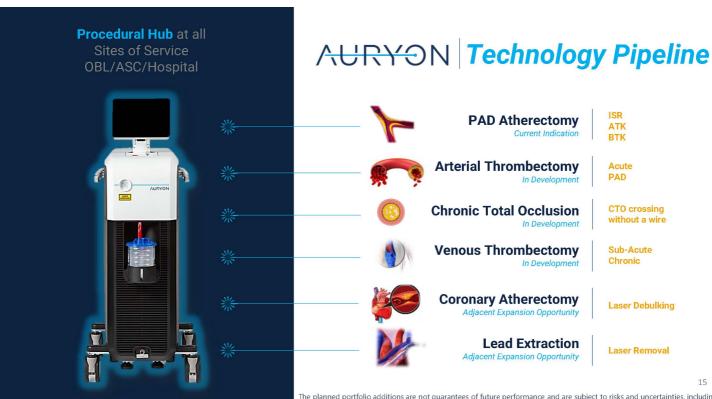


5. Auryce. Instructions for use. AngioDynamics; 2019. 6. Herzog A Bogdan S, Olikson M, Ishaaya AA, Love C. Selective tissue ablation using laser radiation at 355 nm in lead extraction by a hybrid catheler; a preliminary spect. Lasers Surg Med. 2016;48(5):281-287.
7. Spectrametric Corporation, CVX-300 occimient user system: Operator's Manual, Version 28, 2019;1-56.





* As of 01/06/2022 **Perinberal Vascular Devices, Millennium Research Data 2020 based on 288K Atherectomy Procedures Annual



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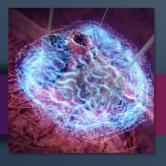


INNOVATION DOCTORS NEED

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



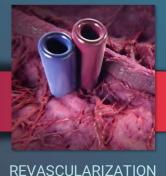
PROBE PLACEMENT NanoKnife can be confidently used in all segments of an organ.10.11



DECELLULARIZATION Destroys targeted tissue with precise treatment margins.10,11



NON-THERMAL Spares vital structures by retaining the structural integrity of tissue.12,13



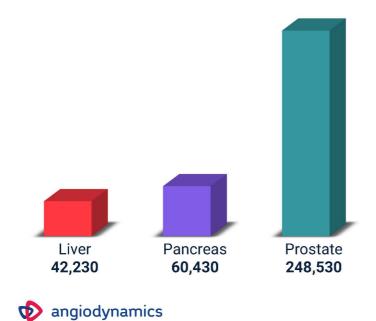
Facilitates functional tissue regeneration post-ablation. 12,13

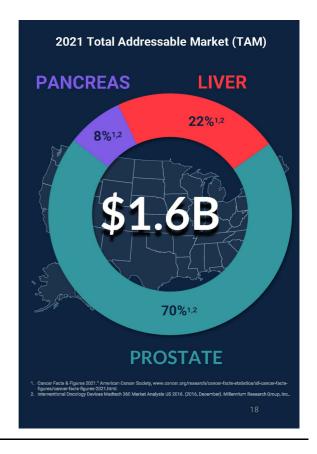


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THE NANOKNIFE SYSTEM

Estimated # of US Patients Diagnosed in 20211





PROSTATE INITIATIVE

Prostate cancer is the only solid tumor without a standardized local treatment option1

Advances in genetics, imaging, and methodology support a local treatment option for PCa2

But existing technology has less than ideal outcomes for cancer control and quality of life^{3,4}

Effectively destroys targeted tissue with precise treatment margins⁶ Spares vital structures within the ablation zone⁷ Enables treatment to be performed

in all segments of an organ⁸



Bridge the gap between an active surveillance strategy and whole-gland treatment

Approximately men are estimated to be candidates for focal therapy each year⁵

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 Sites selected, focused on geographic and
- demographic diversity, high-volume focal therapy institutions
- 118 Intermediate-risk patients to be enrolled through 1-year follow up

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

First Enrollment: Expected Q3

Enrollment Period: 12 months

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Date of last update: Jan 7, 2022

FINANCIALS



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SECOND QUARTER AND YTD HIGHLIGHTS

Financial Performance

\$ in thousands (except per share data)

	Q2 FY2022	Q2 FY2021	Change	YTD FY2022	YTD FY2021	Change
Revenue	\$78,280	\$72,770	7.6%	\$155,251	\$142,986	8.6%
Gross Margin	51.8%	55.2%	(340 bps)	52.0%	53.1%	(110 bps)
Net Loss	(\$8,351)	(\$4,268)	(\$4,083)	(\$15,323)	(\$8,536)	(\$6,787)
GAAP EPS	(\$0.21)	(\$0.11)	(\$0.10)	(\$0.39)	(\$0.22)	(\$0.17)
Adjusted EPS	(\$0.02)	\$0.01	(\$0.03)	(\$0.04)	\$0.03	(\$0.07)
Adjusted EBITDA	\$4,421	\$5,158	(\$737)	\$7,992	\$9,625	(\$1,633)

Updated FY22 Guidance

Adjusted EPS
Previous Guidance
\$0.00 - \$0.05

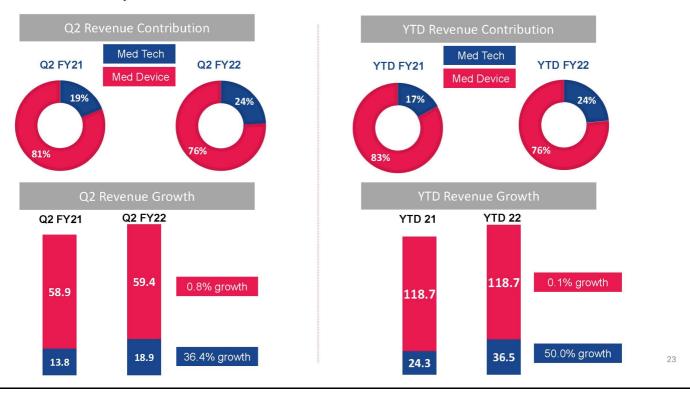
Adjusted EPS
Updated Guidance
(\$0.02) - \$0.02

Revenue (unchanged) \$310 - \$315 million Gross Margin

Previous Guidance Updated Guidance

~55% 52% - 54%

SECOND QUARTER AND YTD HIGHLIGHTS



SECOND QUARTER AND YTD FY2022 RESULTS (unaudited)

\$ in thousands (except per share data)	Q2 FY2022	Q2 FY2021	Change	YTD FY2022	YTD FY2021	Change
Revenue	\$78,280	\$72,770	7.6%	\$155,251	\$142,986	8.6%
Med Tech Med Device	\$18,886 \$59,394	\$13,849 \$58,921	36.4% 0.8%	\$36,504 \$118,747	\$24,335 \$118,651	50.0% 0.1%
Endovascular Therapies Vascular Access Oncology	\$39,660 \$25,070 \$13,550	\$33,900 \$23,930 \$14,940	17.0% 4.8% (9.3%)	\$77,718 \$50,026 \$27,507	\$63,757 \$52,035 \$27,194	21.9% (3.9%) 1.2%
United States International	\$65,350 \$12,930	\$60,684 \$12,086	7.7% 7.0%	\$129,814 \$25,437	\$114,792 \$28,194	13.1% (9.8%)
Net Loss Non-GAAP Adjusted Net Income (Loss)	(\$8,351) (\$856)	(\$4,268) \$564	(\$4,083) (\$1,420)	(\$15,323) (\$1,743)	(\$8,536) \$1,181	(\$6,787) (\$2,924)
GAAP EPS Non-GAAP Adjusted EPS	(\$0.21) (\$0.02)	(\$0.11) \$0.01	(\$0.10) (\$0.03)	(\$0.39) (\$0.04)	(\$0.22) \$0.03	(\$0.17) (\$0.07)
Gross Margin	51.8%	55.2%	(340 bps)	52.0%	53.1%	(110 bps)
Adjusted EBITDA	\$4,421	\$5,158	(\$737)	\$7,992	\$9,625	(\$1,633)



PANDEMIC LANDSCAPE

Taking Actions to Address Short-term Supply Chain Disruption & Long-term Ability to Supply Customers

Headwinds

- Tight Labor Market
 - Drives increasing labor costs
 - Impacts absorption
- Raw Materials Inflationary Pressure
- Increasing Freight Costs

Operational Focus

- Increasing and Qualifying Additional Manufacturing Capacity
- Improving Efficiencies

 - Robust supply chain planning
- Pricing and Shipping Adjustments

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APPENDIX

GAAP to Non-GAAP Reconciliation angiodynamics

Reconciliation of GAAP to Non-GAAP Net Income (Loss) and EPS

	Three Months Ended				Six Months Ended				
(in thousands, except per share data)		v 30, 2021	No	Nov 30, 2020		Nov 30, 2021		Nov 30, 2020	
, , , , , ,	(unaudited)				(unaudited)				
Net loss	\$	(8,351)	\$	(4,268)	\$	(15,323)	S	(8,536)	
Amortization of intangibles		4,889		4,593		9,710		9,546	
Change in fair value of contingent consideration		609		184		804		(473)	
Acquisition, restructuring and other items, net (1)		2,253		1,128		4,693		2,447	
Tax effect of non-GAAP items (2)		(256)		(1,073)		(1,627)		(1,803)	
Adjusted net income (loss)	\$	(856)	\$	564	\$	(1,743)	\$	1,181	
	No	v 30, 2021		v 30, 2020	No	ov 30, 2021		v 30, 2020	
	- No	Three Mo			No	Six Mon			
	(unaudited)			(unaudited)					
Diluted loss per share	\$	(0.21)	\$	(0.11)	\$	(0.39)	\$	(0.22)	
Amortization of intangibles		0.13		0.12		0.25		0.25	
Change in fair value of contingent consideration		0.02		_		0.02		(0.01)	
Acquisition, restructuring and other items, net (1)		0.05		0.03		0.12		0.06	
Tax effect of non-GAAP items (2)		(0.01)		(0.03)		(0.04)		(0.05)	
Adjusted diluted earnings (loss) per share	\$	(0.02)	\$	0.01	\$	(0.04)	\$	0.03	
Adjusted diluted sharecount (3)		39,053		38,473		38,893		38,50	

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

 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 November 30, 2021 and November 30, 2020. Diluted shares may differ for non-GAAP measures as compared to GAAP due to a GAAP loss.

Reconciliation of Net Loss to Adjusted EBITDA

(in thousands)	Three Months Ended				Six Months Ended			
	Nov 30, 2021		Nov 30, 2020		Nov 30, 2021		Nov 30, 2020	
	(unaudited)				(unaudited)			
Net loss	\$	(8,351)	\$	(4,268)	\$	(15,323)	\$	(8,536)
Income tax benefit		(512)		(905)		(2,148)		(1,450)
Interest expense, net		174		235		330		450
Depreciation and amortization		7,240		6,397		14,199		12,936
Change in fair value of contingent consideration		609		184		804		(473)
Stock based compensation		3,008		2,387		5,437		4,251
Acquisition, restructuring and other items, net (1)		2,253		1,128		4,693		2,447
Adjusted EBITDA	\$	4,421	\$	5,158	\$	7,992	\$	9,625

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

