ANGIODYNAMICS

Third Quarter 2021 Earnings Presentation March 30, 2021



Forward-Looking Statement

Notice Regarding Forward-LookingStatements

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 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 integrate acquired businesses, as well as the risk fact

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 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 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 prepared in accordance with GAAP.



Corporate Developments – Q3 Highlights



- Continued focused investment in our 3 key technology platforms: Auryon, AngioVac & NanoKnife
 - Procedural volumes negatively impacted in January and first half of February
 - Began to rebound in the second half of February
 - AngioVac and Auryon procedure volume exhibited resiliency in Q3
 - NanoKnife disposable growth: US 12% YOY; worldwide decline of 7% YOY
 - NanoKnife DIRECT study: 22 sites have secured IRB approval, down 4 from Q2
 - Decommissioned 4 sites to continue to refine and optimize the process
 - We are encouraged by the overall execution of the study in the current environment
 - Completed enrollment of PATHFINDER 1 pilot registry
 - · Shifting our focus to the pivotal study phase
 - Recognized a \$1.9 million benefit to net income in Q3 from the CARES Act



Reduced debt outstanding under revolver by \$20 million as of March 2021

Revised FY2021 Guidance

Adjusted EPS Revenue **Revised Guidance Previous Guidance Revised Guidance** \$0.04 - \$0.06 \$0.00 - \$0.05 \$285 - \$288 mil

Third Quarter FY2021 Highlights

Financial Performance

\$ in thousands (except per share data)	Q3 FY2021	Q3 FY2020	YOY Change
Revenue	\$71,182	\$69,780	2.0%
Gross Margin	54.1%	57.8%	(370 bps)
Net Loss	(\$3,544)	(\$5,709)	\$2,165
GAAP EPS	(\$0.09)	(\$0.15)	\$0.06
Adjusted EPS	\$0.02	\$0.01	\$0.01
Adjusted EBITDA	\$5,379	\$3,790	\$1,589
Cash Provided by (Used in) Operations	\$5,871	(\$17,837)	\$23,708

Product Family Sales Growth Over Prior Year Periods

Q3 FY2021	YTD FY2021
27%	31%
NA*	NA*
2%	(1%)
(5%)	(1%)
(16%)	(14%)
Q3 FY2021	YTD FY2021
16%	40%
10%	NA**
(3%)	14%
(2%)	(1%)
2%	2%
Q3 FY2021	YTD FY2021
(5%)	(40%)
(7%)	4%
(4%)	3%
17%	24%
(18%)	(24%)
(24%)	(24%)
	 27% NA* 2% (5%) (16%) Q3 FY2021 16% 10% (3%) (2%) 2% Q3 FY2021 (5%) (5%) (7%) (4%) 17% (18%)

* The Auryon full market launch took place in the second quarter of fiscal year 2021.

** The C3 Wave acquisition took place in December 2019.

Third Quarter FY2021 Results (unaudited)

\$ in thousands (except per share data)	Q3 FY2021	Q3 FY2020	Change	YTD FY2021	YTD FY2020	Change
Revenue	\$71,182	\$69,780	2.0%	\$214,168	\$205,825	4.1%
Vascular Interventions and Therapies	\$33,251	\$30,552	8.8%	\$97,008	\$90,616	7.1%
Vascular Access	\$24,813	\$24,642	0.7%	\$76,848	\$70,585	8.9%
Oncology	\$13,118	\$14,586	(10.1%)	\$40,312	\$44,624	(9.7%)
United States	\$58,654	\$54,889	6.9%	\$173,446	\$163,381	6.2%
International Net Loss Non-GAAP Adjusted Net Income	\$12,528	\$14,891	(15.9%)	\$40,722	\$42,444	(4.1%)
	(\$3,544)	(\$5,709)	\$2,165	(\$12,080)	(\$9,720)	(\$2,360)
	\$738	\$362	\$376	\$1,919	\$5,687	(\$3,768)
GAAP EPS	(\$0.09)	(\$0.15)	\$0.06	(\$0.32)	(\$0.26)	(\$0.06)
Non-GAAP Adjusted EPS	\$0.02	\$0.01	\$0.01	\$0.05	\$0.15	(\$0.10)
Gross Margin	54.1%	57.8%	(370 bps)	53.4%	58.3%	(490 bps)
Adjusted EBITDA	\$5,379	\$3,790	\$1,589	\$15,004	\$17,480	(\$2,476)

\$ in thousands (except per share data)	Q3 FY2021	Q4 FY2020	Change
Cash	\$54,469	\$54,435	\$34
Debt	\$30,000*	\$40,000	(\$10,000)

* In March 2021, another \$10 million was paid on the revolver. Total debt outstanding is now \$20 million.



GAAP to Non-GAAP Reconciliation



Reconciliation of GAAP to Non-GAAP Net Income and EPS

	Three Months Ended					Nine Months Ended				
(in thousands, except per share data)		28, 2021	Fel	Feb 29, 2020		b 28, 2021	Feb 29, 2020			
	(unaudited)					(unaudited)				
Net loss	\$	(3,544)	\$	(5,709)	\$	(12,080)	\$	(9,720)		
Amortization of intangibles		4,292		5,019		13,838		13,417		
Change in fair value of contingent consideration		183		419		(290)		116		
Acquisition, restructuring and other items, net (1)		610		1,565		3,057		4,486		
Write-off of deferred financing fees (2)				_				593		
Tax effect of non-GAAP items (3)		(803)		(932)		(2,606)		(3,205)		
Adjusted net income	\$	738	\$	362	\$	1,919	\$	5,687		

	Three Months Ended				Nine Months Ended				
	Feb 28, 2021		Feb 29, 2020		Feb 28, 2021		Feb 29, 2020		
Diluted loss per share		(unau	idited)	2	(unaudited)				
	\$	(0.09)	\$	(0.15)	\$	(0.32)	\$	(0.26)	
Amortization of intangibles		0.11		0.13		0.36		0.35	
Change in fair value of contingent consideration		-		0.01		(0.01)		_	
Acquisition, restructuring and other items, net (1)		0.02		0.04		0.08		0.12	
Write-off of deferred financing fees (2)				_		_		0.02	
Tax effect of non-GAAP items (3)		(0.02)		(0.02)		(0.06)		(0.08)	
Adjusted diluted earnings per share	\$	0.02	\$	0.01	\$	0.05	\$	0.15	
Adjusted diluted sharecount (4)		39,271		38,094		38,770		38,111	

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

(2) Deferred financing fees related to the old credit agreement were written off during the second quarter of fiscal year 2020.

(3) 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.

(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

	Three Months Ended					Nine Months Ended			
(in thousands)		Feb 28, 2021		Feb 29, 2020		Feb 28, 2021		Feb 29, 2020	
	(unaudited)			(unaudited)					
Net loss	\$	(3,544)	\$	(5,709)	\$	(12,080)	\$	(9,720)	
Income tax benefit		(583)		(824)		(2,033)		(1,506)	
Interest expense, net		226		166		676		672	
Depreciation and amortization		6,340		6,401		19,276		17,434	
Change in fair value of contingent consideration		183		419		(290)		116	
Stock based compensation		2,147		1,772		6,398		5,998	
Acquisition, restructuring and other items, net (1)		610		1,565		3,057		4,486	
Adjusted EBITDA	\$	5,379	\$	3,790	\$	15,004	\$	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.

