AngioDynamics

First Quarter 2021 Earnings Presentation September 29, 2020



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," "believes," "seeks," "estimates," "optimistic," or variations of such words and similar expressions, are 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 from AngioDynamics' expectations. 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, t

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; adjusted earnings per share and free cash flow. 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.



First Quarter FY2021 Highlights

Financial Performance					
(in millions)	Q1 FY21	Q1 FY20	YOY Change		
Revenue	\$70.2	\$66.0	6.3%		
Gross Margin	50.9%	57.9%	(700 bps)		
Adjusted EPS	\$0.02	\$0.08	(\$0.06)		
Adjusted EBITDA	\$4.5	\$7.3	(\$2.8)		
Cash Used In Operations	\$5.4	\$6.5			
Free Cash Flow	(\$7.2)	(\$7.9)			

Product Family Year-over-Year Sales Growth				
Vascular Interventions and Therapies				
AngioVac®	46%			
Auryon	NA			
Thrombolytic	(8%)			
Core Peripheral	0%			
Venous Insufficiency	(16%)			
Vascular Access				
Midlines	86%			
C3	NA			
PICCs	45%			
Ports	(5%)			
Dialysis	(2%)			
Oncology				
NanoKnife® Capital	(76%)			
NanoKnife® Disposables	(5%)			
Solero® Microwave	7%			
BioSentry	29%			
Alatus and IsoLoc Balloons	(31%)			
RadioFrequency Ablation	(29%)			



First Quarter FY2021 Results (unaudited)

\$ in thousands (except per share data)	Q1 FY2021	Q1 FY2020	YOY Change
Revenue Vascular Interventions and Therapies Vascular Access Oncology	\$70,216 29,857 28,105 12,254	\$66,042 28,913 23,159 13,970	6.3% 3.3% 21.4% (12.3%)
United States International	54,108 16,108	52,937 13,105	2.2% 22.9%
Net Loss Adjusted Net Income	(\$4,268) \$618	(\$1,275) \$3,174	(\$2,993) (\$2,556)
GAAP EPS Non-GAAP Adjusted EPS	(\$0.11) \$0.02	(\$0.03) \$0.08	(\$0.08) (\$0.06)
Gross Margin	50.9%	57.9%	700 bps
Adjusted EBITDA	\$4,466	\$7,280	(\$2,814)
Free Cash Flow	(\$7,249)	(\$7,925)	\$676

	Q1 FY2021	Q4 FY2020	Change
Cash	\$47,929	\$54,435	(\$6,506)
Debt	\$40,000	\$40,000	-



Corporate Developments

- Previously disclosed order to National Health Services in the UK for \$5.2 million in the first quarter of fiscal year 2021.
- Gross margin was 50.9% and was primarily impacted by COVID-related operating protocols designed to ensure supply-chain security and employee safety.
 - Inventory was reduced by \$7.2 million in the first quarter.
- Procedural volumes began to rebound in June with steady improvement throughout the first quarter but not yet back to pre-COVID levels.
- Continued investment in three key technology platforms: NanoKnife, AngioVac, and Auryon.
- NanoKnife disposable growth of 7% in the United States was more than offset by softness in China due to the ongoing impacts of COVID-19.
- NanoKnife DIRECT study: 23 sites have secured IRB approval, compared to 21 at the end of the fourth quarter.
- Official launch of Auryon on September 21, 2020.

FY2021 Guidance					
Revenue	\$278 - \$284 million				
Adjusted EPS	\$0.00 - \$0.05				



GAAP to Non-GAAP Reconciliation



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

A		Three Months Ended			
Amounts in thousands	Aug 31, 2020		Aug 31, 2019		
		(una	idited)		
Net loss	\$	(4,268)	\$	(1,275)	
Amortization of intangibles		4,953		3,868	
Change in fair value of contingent consideration		(657)		(448)	
Acquisition, restructuring and other items, net (1)		1,319		1,500	
Write-off of deferred financing fees (2)		_		593	
Tax effect of non-GAAP items (3)		(729)		(1,064)	
Adjusted net income	\$	618	\$	3,174	
		Three Mo	nths End	led	
	Aug	g 31, 2020	Au	g 31, 2019	

	Aug	Aug 31, 2020 A		Aug 31, 2019	
		(unaudited)			
Diluted loss per share	\$	(0.11)	\$	(0.03)	
Amortization of intangibles		0.13		0.10	
Change in fair value of contingent consideration		(0.02)		(0.01)	
Acquisition, restructuring and other items, net (1)		0.03		0.04	
Write-off of deferred financing fees (2)		_		0.02	
Tax effect of non-GAAP items (3)		(0.01)		(0.04)	
Adjusted diluted earnings per share	\$	0.02	\$	0.08	
A.P. (1.19)		20.101		20.150	
Adjusted diluted sharecount		38,191		38,158	

- (1) 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 first 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 August 31, 2020 and 2019.



Reconciliation of Net Loss to Adjusted EBITDA

		Three Months Ended		
Amounts in thousands	Aug 31, 2020		Aug 31, 2019	
	(unaudited)			
Net loss	\$	(4,268)	\$	(1,275)
Income tax benefit		(545)		(116)
Interest expense, net		215		465
Depreciation and amortization		6,538		5,170
Change in fair value of contingent consideration		(657)		(448)
Stock based compensation		1,864		1,984
Acquisition, restructuring and other items, net (1)		1,319		1,500
Adjusted EBITDA	\$	4,466	\$	7,280
Per diluted share:				
Adjusted EBITDA	\$	0.12	\$	0.19

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





