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 5, 2012

AngioDynamics, Inc.

(Exact Name of Registrant as Specified in Charter)

Delaware

(State or Other Jurisdiction of Incorporation)

000-50761 (Commission File Number) 11-3146460 (IRS Employer Identification No.)

603 Queensbury Avenue, Queensbury, New York 12804

(Address of Principal Executive Offices)

(Zip Code)

(518) 798-1215

(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:

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

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

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

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

Item 2.02 – Results of Operations and Financial Condition.

On January 5, 2012, AngioDynamics, Inc. (the "Company") issued a press release announcing financial results for the fiscal second quarter ended November 30, 2011.

The information set forth in Item 2.02 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 incorporated by reference in any filing under the Securities Act of 1933, as amended, or the Exchange Act, except as shall be expressly set forth by specific reference in such a filing.

Forward-Looking Statements

This document and its attachments include "forward-looking statements" intended to qualify for the safe harbor from liability established by the Private Securities Litigation Reform Act of 1995. Investors can identify these statements by the fact that they do not relate strictly to historical or current facts. These statements contain words such as "expect," "reaffirm," "anticipate," "plan," "believe," "estimate," "may," "will," "predict," "project," "might," "intend," "potential," "could," "would," "should," "optimistic," "seek," "continue," "pursue," or "our future success depends," or the negative or other variations thereof or comparable terminology, are intended to identify such forward-looking statements. In particular, they include statements relating to, among other things, future actions, strategies, future performance and future financial results of the Company. These forward-looking statements are based on current expectations and projections about future events.

Investors are cautioned that forward-looking statements are not guarantees of future performance or results and involve risks and uncertainties that cannot be predicted or quantified and, consequently, the actual performance or results of the Company may differ materially from those expressed or implied by such forward-looking statements. Such risks and uncertainties include, but are not limited to, the factors described from time to time in the Company's reports filed with the SEC, including the Company's Form 10-K for the fiscal year ended May 31, 2011, financial community and rating agency perceptions of the Company; the effects of economic, credit and capital market conditions on the economy in general, and on medical device companies in particular; domestic and foreign health care reforms and governmental laws and regulations; third-party relations and approvals, technological advances and patents attained by competitors; and challenges inherent in new product development, including obtaining regulatory approvals. In addition to the matters described above, the ability of the Company to develop its products, future actions by the FDA or other regulatory agencies, results of pending or future clinical trials, the outcome of pending litigation, overall economic conditions, general market conditions, market acceptance, foreign currency exchange rate fluctuations, and the effects on pricing from group purchasing organizations and competition and the ability of the Company to integrate purchased businesses, may affect the actual results achieved by the Company.

Any forward-looking statements are made pursuant to the Private Securities Litigation Reform Act of 1995 and, as such, speak only as of the date made. The Company disclaims any obligation to update the forward-looking statements. Investors are cautioned not to place undue reliance on these forward-looking statements which speak only as of the date stated, or if no date is stated, as of the date of this document.

Item 9.01 – Financial Statements and Exhibits.

(d) *Exhibits*.

Exhibit No.	Description
99.1	Press Release dated January 5, 2012.

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 5, 2012

By: /s/ D. Joseph Gersuk

D. Joseph Gersuk Chief Financial Officer

Exhibit No.	Description	

99.1 Press Release dated January 5, 2012.

ANGIODYNAMICS[®]

FOR IMMEDIATE RELEASE

Company Contact: <u>AngioDynamics, Inc.</u> D. Joseph Gersuk, CFO (800) 772-6446 x1608 jgersuk@AngioDynamics.com Investor Relations Contacts: <u>EVC Group, Inc.</u> Greg Gin/Doug Sherk (646) 445-4801; (415) 652-9100 <u>ggin@evcgroup.com</u>; <u>dsherk@evcgroup.com</u> Media Contact: <u>EVC Group, Inc.</u> Chris Gale (646) 201-5431 <u>cgale@evcgroup.com</u>

AngioDynamics Reports Fiscal 2012 Second Quarter Financial Results

ALBANY, N.Y. January 5, 2012 – AngioDynamics (NASDAQ: ANGO), a leading provider of innovative, minimally invasive medical devices for vascular access, surgery, peripheral vascular disease and oncology, today reported financial results for the fiscal 2012 second quarter ended November 30, 2011.

Net sales in the second quarter were \$58.1 million, a 9% increase over the \$53.4 million reported in the prior year period. U.S. sales were \$49.7 million and increased 6% compared to the second quarter of fiscal 2011. International sales of \$8.4 million rose 27% from the prior year period. Oncology/Surgery sales increased 25% to \$19.8 million from the prior year and included \$3.2 million in NanoKnife[®] System sales, a 100% year-over-year increase. Vascular sales were \$38.3 million, an increase of 2% from the second quarter a year ago.

Second quarter gross profit was reduced by \$1.5 million due to costs attributable to the voluntary Class II recall of NeverTouch[®] procedure kits used with the VenaCure EVLT[®] System, Morpheus[®] Smart PICC CT PICCs and DuraMax[®] Chronic Hemodialysis Catheters. The recalls stemmed from nonconforming parts manufactured by suppliers. Shipments of all three products have resumed, however, management believes that a backlog of orders for NeverTouch procedure kits may exist until the end of February 2012.

Gross margin in the second quarter was 57.2%, or 59.8% excluding product recall costs, compared with 59.1% in the prior year second quarter and the first quarter of fiscal 2012. Excluding recall costs, the increase in gross margin reflects the impact of ongoing programs to reduce material costs and increase manufacturing efficiency. Operating income in the second quarter was \$3.9 million, or \$5.3 million excluding \$1.4 million of restructuring and other items, compared with \$5.4 million of operating income a year ago. The restructuring and other items primarily include the CEO transition and the beginning of a program to close a facility in the UK. Net income in the second quarter was \$2.3 million, or \$0.09 per share, compared with \$3.3 million, or \$0.13 per share, a year ago. Excluding restructuring and other items, second quarter net income was \$3.3 million, or \$0.13 per share.

During the second quarter, AngioDynamics generated net cash flow from operations of \$2.7 million and purchased 142,000 shares of its common stock at a cost of \$2.1 million under its share repurchase program. Approximately \$17.9 million remains under the current authorization. Cash and investments totaled \$136.3 million at November 30, 2011.

"Our sales and marketing teams' execution and focus led to a strong quarter," said Joseph DeVivo, President and CEO. "NanoKnife[®] System sales rose 100% year-over-year, Vascular and Oncology/Surgery grew in the U.S., and our International efforts delivered another 20 percent-plus growth quarter. We did identify areas needing improvement in our quality management system, which are being implemented by our team. After a comprehensive review, we initiated a voluntary recall of the NeverTouch[®] laser fiber and other catheter products. While these actions reduced our financial results in the second quarter, we believe they will benefit our Company longer term. Additionally, the voluntary recalls will impact operating results in the third quarter, and we have updated guidance accordingly.

"We expect that beginning in the first calendar quarter of 2012 we will begin to see clinicians present their experiences and patient outcomes with the NanoKnife System at medical conferences around the world," Mr. DeVivo continued. "We have decided to focus our NanoKnife System global clinical and regulatory efforts, hoping to establish it as a standard of care for the ablation of pancreatic tumors since we see the significant clinical unmet need as the top market priority for this technology. At the same time, we continue our corporate development effort to pursue acquisitions that can benefit from our strong Oncology/Surgery and Vascular sales organizations."

Highlights of the quarter, and more recent activities, include the following:

- Continued commercial use of AngioDynamics' NanoKnife System, with 158 patients treated in the second quarter. To date, more than 1,000 patients have been treated with the NanoKnife System.
- Continued progress on NanoKnife System clinical programs, with completion of enrollment in the NanoPanc European Study (ONC-208) evaluating its use to treat pancreatic cancer in 10 patients.
- Receipt of U.S. FDA 510(k) Market Clearance for the NanoKnife System that provides system enhancements such as implementation of RFID, touch screen capabilities and tightened system parameters.
- Continued double digit growth of VenaCure EVLT[®] System sales driven by the recent market introduction of the 90 cm NeverTouch[®] procedure kit and the VenaCure[®] 1470nm laser.
- Expansion of the leadership team with the appointments of Donna Haire, Senior Vice President of Regulatory, Quality, Clinical and Medical Affairs; Alan Panzer, Senior Vice President and General Manager, Vascular

Division; and Matthew Kapusta, Senior Vice President of Business Development.

For the six months ended November 30, 2011, net sales were \$112.5 million, a 7% increase over the \$104.9 million reported for the prior year period. Gross margin for the six months ended November 30, 2011, was 58.1%, or 59.4% excluding product recall costs, compared with 58.7% for the six months ended November 30, 2010. For the six months ended November 30, 2011, operating income was \$6.6 million, compared to \$8.9 million for the year ago period. Net income of \$3.7 million, or \$0.15 per share, compared to \$5.2 million, or \$0.21 per share, as reported for the prior year period. Excluding restructuring and other items, first half net income was \$5.2 million, or \$0.21 per share.

Fiscal 2012 Guidance

As shown in the tables below, the Company updated its guidance for fiscal 2012 to reflect second quarter operating results and its outlook for the balance of the year. The guidance reflects the impact of the recent recalls, the expiration of the Company's contract to distribute LC Beads on December 31, 2011, and \$1.1 million in restructuring costs in the second half of the fiscal year, primarily for the closure of a facility in the UK. Finally, the guidance for the fiscal year includes approximately \$21 million in R&D spending and a \$0.14 loss per share impact from the NanoKnife[®] System program.

FY 2012 GUIDANCE, INCLUDING ITEMS (GAAP) (\$ in mil's, except EPS)							
	Q3	Q4	FY 2012				
Sales (\$)	52.0 - 54.0	53.5 – 55.5	218.0 - 222.0				
Sales Growth (%)	(5)% - (1)%	(5)% - (2)%	1% - 3%				
Gross Margin (%)	58.5% - 59.5%	59.5% - 60.5%	58.5% - 59.5%				
Operating Income (\$)	2.7 - 3.7	4.6 - 5.6	13.9 - 15.9				
EBITDA (\$)	6.2 - 7.2	8.1 - 9.1	27.5 – 29.5				
EPS (\$)	0.07 - 0.09	0.11 - 0.13	0.32 - 0.37				

FY 2012 GUIDANCE, EXCLUDING ITEMS (Non-GAAP)* (\$ in mil's, except EPS)							
	Q3	Q4	FY 2012				
Sales (\$)	52.0 - 54.0	53.5 - 55.5	218.0 - 222.0				
Sales Growth (%)	(5)% - (1)%	(5)% - (2)%	1% - 3%				
Pro Forma Sales Growth (%)**	0% - 4%	10% - 15%	5% - 7%				
Gross Margin (%)	58.5% - 59.5%	59.5% - 60.5%	58.5% - 59.5%				
Operating Income (\$)	3.5 - 4.5	4.9 - 5.9	17.4 - 19.4				
EBITDA (\$)	7.0 - 8.0	8.4 - 9.4	31.0 - 33.0				
EPS (\$)	0.09 - 0.11	0.11 - 0.13	0.41 - 0.45				

* Excludes CEO transition, the closure of a facility in the U.K., and certain other items

** Pro Forma Sales Growth excludes LC Beads in all periods

Conference Call

AngioDynamics management will host a conference call to discuss its second quarter results today beginning at 4:30 p.m. Eastern Time. To participate in the live call by telephone, please dial 1 (877) 941-6009. In addition, individuals can listen to the live call and the replay on the Internet by visiting the investor relations portion of the AngioDynamics Web site at <u>http://investors.angiodynamics.com</u>. To listen to the live call, please go to the Web site 15 minutes prior to its start to register, download and install the necessary audio software.

Use of Non-GAAP 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 news release, AngioDynamics has reported and provided projections for non-GAAP gross margin, non-GAAP operating income, non-GAAP net income, non-GAAP net income per share, non-GAAP EBITDA (income before interest, taxes, depreciation, amortization and impairment charges) and non-GAAP 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. Management believes the presentation of these measures is relevant and useful for investors because they allow investors to view performance in a manner similar to the method used by management, help improve their ability to understand the Company's operating performance and make it easier to compare the Company's results with other companies that have different financing and capital structures or tax rates. In addition, these measures are among the primary measures used externally by the Company's investors, analysts and peers in its industry for purposes of valuation and comparing the operating performance of the Company to other companies in the industry. 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 Operating Income to non-GAAP measures.

About AngioDynamics

AngioDynamics, Inc. is a leading provider of innovative, minimally invasive medical devices used by professional healthcare providers for vascular access, surgery, peripheral vascular disease and oncology. AngioDynamics' diverse product lines include market-leading ablation systems, vascular access products, angiographic products and accessories, angioplasty products, drainage products, thrombolytic products and venous products. More information is available at www.AngioDynamics.com.

Safe Harbor

This release 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. 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 from AngioDynamics' expectations. Factors that may affect the actual results achieved by AngioDynamics include, without limitation, the ability of AngioDynamics to develop its existing and new products, future actions by the FDA or other regulatory agencies, results of pending or future clinical trials, overall economic conditions, the results of on-going litigation, 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 purchased 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, 2011, and quarterly report on Form 10-Q for the fiscal quarter ended November 30, 2011. AngioDynamics does not assume any obligation to publicly update or revise any forward-looking statements for any reason.

In the United States, NanoKnife has been cleared by the FDA for use in the surgical ablation of soft tissue. This document may discuss the use of NanoKnife for specific clinical indications for which it is not cleared in the United States at this time.

ANGIODYNAMICS, INC. AND SUBSIDIARIES CONSOLIDATED INCOME STATEMENTS

(in thousands, except per share data)

		Three months ended			Six months ended				
	Nov	November 30, November 30, 2011 2010 (unaudited)				vember 30,	No	vember 30,	
					2011		2010		
		(unau	lanea)			(unau	unea)	inteu)	
Net sales	\$	58,099	\$	53,372	\$	112,530	\$	104,879	
Cost of sales		24,868		21,836		47,154		43,323	
Gross profit		33,231		31,536	_	65,376		61,556	
% of net sales		57.2%	, D	59.1%	,	58.1%		58.7%	
Operating expenses									
Research and development		5,125		5,259		10,715		10,501	
Sales and marketing		15,847		13,793		32,156		28,237	
General and administrative		4,625		4,173		8,937		8,759	
Amortization of intangibles		2,300		2,142		4,594		4,409	
Restructuring and other costs		1,408		772		2,331		772	
Total operating expenses		29,305		26,139		58,733		52,678	
Operating income		3,926		5,397		6,643		8,878	
Other income (expense), net		(357)		(262)		(971)		(790)	
Income before income taxes		3,569		5,135	-	5,672		8,088	
Provision for income taxes		1,240		1,856		1,970		2,921	
Net income	\$	2,329	\$	3,279	\$	3,702	\$	5,167	
Earnings per common share									
Basic	\$	0.09	\$	0.13	\$	0.15	\$	0.21	
Diluted	\$	0.09	\$	0.13	\$	0.15	\$	0.21	
Weighted average common shares									
Basic		25,190		24,845		25,107		24,799	
Diluted		25,340		25,094		25,278		25,067	

ANGIODYNAMICS, INC. AND SUBSIDIARIES CONSOLIDATED INCOME STATEMENTS (in thousands, except per share data)

Reconciliation of Operating Income to non-GAAP EBITDA:

	Th	Three months ended				led				
		November 30, November 30, 2011 2010 (unaudited)				,		November 30, 2011		vember 30, 2010
				(unaudited) (unaudit			dited)			
Operating income	\$	3,926	\$	5,397	\$	6,643	\$	8,878		
Amortization of intangibles		2,300		2,142		4,594		4,409		
Depreciation		850		813	13 1,67		9 1,			
EBITDA	\$	7,076	\$	8,352	\$	12,916	\$	14,862		
EBITDA per common share										
Basic	\$	0.28	\$	0.34	\$	0.51	\$	0.60		
Diluted	\$	0.28	\$	0.33	\$	0.51	\$	0.59		
Weighted average common shares										
Basic	-	25,190		24,845		25,107		24,799		
Diluted	2	25,340		25,094		25,278		25,067		

ANGIODYNAMICS, INC. AND SUBSIDIARIES NET SALES BY PRODUCT CATEGORY AND BY GEOGRAPHY (in thousands)

	Three months ended			led																						
		November 30, 2011		November 30, 2010																				vember 30, 2011	Nov	vember 30, 2010
		(unau	dited)			(unau	dited)																			
Net Sales by Product Category																										
Vascular																										
Peripheral Vascular	\$	23,078	\$	22,004	\$	44,046	\$	42,705																		
Access		15,204		15,516		30,801		30,729																		
Total Vascular		38,282		37,520		74,847		73,434																		
Oncology/Surgery		19,817		15,852		37,683		31,445																		
Total	\$	58,099	\$	53,372	\$	112,530	\$	104,879																		
Net Sales by Geography United States International	\$	49,653 8,446	\$	46,703 6,669	\$	96,958 15,572	\$	92,176 12,703																		
Total	\$	58,099	\$	53,372	\$	112,530	\$	104,879																		
Total	Ψ.	50,055	Ψ	55,572	Ψ	112,550	Ψ	104,075																		

ANGIODYNAMICS, INC. AND SUBSIDIARIES CONSOLIDATED BALANCE SHEETS

(in thousands)

		November 30, 2011 (unaudited)		2011 20		May 31, 2011 naudited)
Assets						
Current Assets						
Cash and cash equivalents	\$	42,955	\$	45,984		
Marketable securities		93,364		85,558		
Total cash and investments		136,319		131,542		
Receivables, net		31,451		27,141		
Inventories, net		29,427		28,126		
Deferred income taxes		2,851		2,821		
Prepaid income taxes		2,269		503		
Prepaid expenses and other		3,519		4,172		
Total current assets		205,836		194,305		
Property, plant and equipment, net		23,196		23,804		
Intangible assets, net		43,691		48,037		
Goodwill		161,951		161,951		
Deferred income taxes		4,870		5,835		
Other non-current assets		4,046		3,489		
Total Assets	\$	443,590	\$	437,421		
Liabilities and Stockholders' Equity						
Current portion of long-term debt	\$	290	\$	275		
Other current liabilities		26,330		25,232		
Long-term debt, net of current portion		6,125		6,275		
Total Liabilities		32,745		31,782		
Stockholders' equity		410,845		405,639		
Total Liabilities and Stockholders' Equity	\$	443,590	\$	437,421		
Shares outstanding		25,103		- 24,986		

ANGIODYNAMICS, INC. AND SUBSIDIARIES CONSOLIDATED STATEMENTS OF CASH FLOWS

(in thousands)

	Six	months ended
	November 3 2011	30, November 30, 2010
	(unaudited	l) (unaudited)
Cash flows from operating activities:		
Net income	\$ 3,7	702 \$ 5,167
Depreciation and amortization	6,2	5,984
Tax effect of exercise of stock options	(1	198) (29)
Deferred income taxes	1,0	2,285
Stock-based compensation	1,8	377 2,254
Other	1	168 71
Changes in operating assets and liabilities		
Receivables	(4,5	515) 4,157
Inventories	(1,5	546) (3,515)
Accounts payable and accrued liabilities	8	392 (5,944)
Other	(1,9	998) 1,195
Net cash provided by operating activities	5,7	713 11,626
Cash flows from investing activities:		
Additions to property, plant and equipment		058) (1,489)
Acquisition of intangible and other assets		
Proceeds from sales of intangible and other assets		- 000
Purchases, sales and maturities of marketable securities, net	(8,3	
Net cash provided by (used in) investing activities	(8,7	735) (43,883)
Cash flows from financing activities:		
Repayment of long-term debt	(1	135) (130)
Proceeds from exercise of stock options and ESPP	2,2	250 718
Repurchase and retirement of shares	(2,1	- 104)
Net cash provided by financing activities		11 588
Effect of exchange rate changes on cash		(18) 38
Decrease in cash and cash equivalents		029) (31,632)
Cash and each equivalents		
Cash and cash equivalents		984 58,763
Beginning of period	45,9	
End of period	\$ 42,9	955 \$ 27,131