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 13, 2016

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 Drive Latham, New York		12110
(Address of Principal Executive Offices)		(Zip Code)
(Re Check the appropriate box below if the Form 8-K filing is inte	(518) 795-1400 egistrant's telephone number, including area code) ended to simultaneously satisfy the filing obligation	of the registrant under any of the following provisions:
☐ Written communications pursuant to Rule 425 under the 5	Securities Act (17 CFR 230.425)	
Soliciting material pursuant to Rule 14a-12 under the Exc	change Act (17 CFR 240.14a-12)	
Pre-commencement communications pursuant to Rule 14	dd-2(b) under the Exchange Act (17 CFR 240.14d-2	(b))
Pre-commencement communications pursuant to Rule 13	se-4(c) under the Exchange Act (17 CFR 240.13e-4 ((c))

Item 7.01 – Regulation FD Disclosure.

On January 13, 2016, Joseph M. DeVivo, President and Chief Executive Officer of AngioDynamics, Inc. (the "Company"), will present to certain investors at the JP Morgan 34th Annual Healthcare Conference. The conference slides are furnished herewith as Exhibit 99.1.

The slides are being furnished pursuant to Item 7.01 and the information contained therein shall not be deemed "filed" for purposes of Section 18 of the Securities Exchange Act of 1934, as amended, or otherwise subject to the liabilities under that Section. Furthermore, the information in Exhibit 99.1 shall not be deemed to be incorporated by reference into the filings of the Company under the Securities Act of 1933, as amended.

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," 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, 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, the effects on pric

Item 9.01 - Financial Statements and Exhibits.

(d) Exhibits.

Exhibit No. Description

99.1 Presentation slides for the JP Morgan Healthcare Conference on January 13, 2016.

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)

By: /s/ Stephen A. Trowbridge

Date: January 13, 2016

Stephen A. Trowbridge Senior Vice President and General Counsel

EXHIBIT INDEX

Exhibit No. Description

Presentation slides for the JP Morgan Healthcare Conference on January 13, 2016.

99.1



JP Morgan Healthcare Conference

Joseph M. DeVivo, President and CEO January 13, 2016

ADVANCING CARE. REDUCING COMPLICATIONS. MINIMIZING COSTS.

Forward-Looking Statements

Notice Regarding Forward Looking Statements

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," 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, technological advances and patents attained by competitors, infringement of AngioDynamics' technology or 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, the effects on pricing from group purchasing organizations and competition, the ability of AngioDynamics to integrate purchased businesses,

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.

EmboMedics microsphere products have not been reviewed by the U.S. Food and Drug Administration or any other international regulatory body at this time; as such they are currently not available for sale by AngioDynamics.

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 news release, AngioDynamics has reported net sales excluding a supply agreement; adjusted sales growth; EBITDA (income before interest, taxes, depreciation and amortization); adjusted EBITDA; adjusted gross profit; adjusted net income and adjusted earnings per share. Additionally, this press release evaluates results on a constant currency basis. As a non-GAAP measure, constant currency excludes the impact of foreign currency exchange rate fluctuations. 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.

angiodynamics

2

Profile of AngioDynamics

Global, leading provider of innovative, image-guided, minimally invasive solutions.

Founded: 1998 IPO: May 2004 ANGO (NASDAQ) **Employee Base:** 1,300 employees 7 operating locations

Peripheral Vascular Vascular Access Oncology/Surgery

Franchises:

Global Footprint:

210+ global sales team in U.S., Australia, Canada, France, Germany, Netherlands and U.K. Present in 50+ markets through 110+ distributors

















IR Heritage Drives Future Growth

In 1990, AngioDynamics began manufacturing diagnostic catheters to serve the emerging practice of the interventional radiologist. Every development since then has resulted from the company's commitment to serving interventional radiologists and their ability to offer minimally invasive, innovative treatment options to patients worldwide.

AngioDynamics will continue to leverage that legacy to enable future growth.





Our Strategic Foundation

Improve patient outcomes while lowering overall costs within the healthcare system and maximizing shareholder value

1

Invest in growth drivers

2

Focus on R&D to position for future growth opportunities

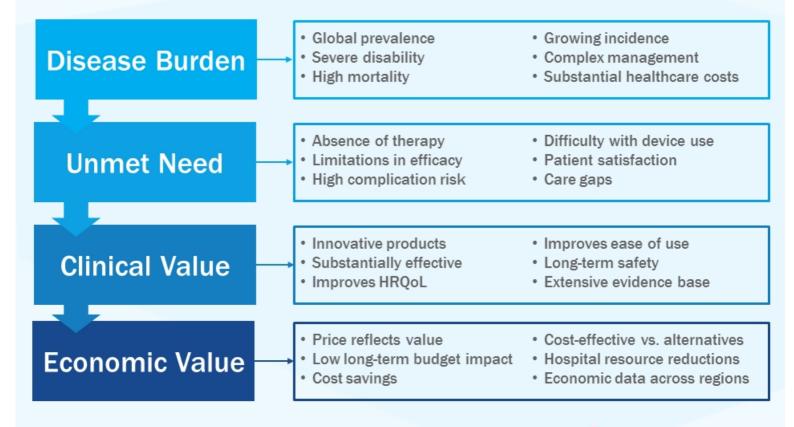
3

Transform through operational excellence



5

Economic and Quality Measures



angiodynamics



1

IMPROVING PATIENT OUTCOMES BY INVESTING IN GROWTH DRIVERS



Peripheral Vascular Overview

Thrombus Management



Venous Insufficiency



Fluid Management



Drainage



Angiographic Catheters

66(

FY2015 Revenue: \$193 million



8

Clinical Unmet Need: VTE

VENOUS THROMBOEMBOLISM (VTE) INCIDENCE

Additional: Overall ~1 Million VTE Events per year in U.S. **Annual Incidence** Overall 300,000 VTE Deaths per year in U.S. - #3 cause Catheter/Lead RA 200,000 1,000,000 595,000 Heart 800,000 **RA Mass** Cases Per Year 47,000 600,000 574,000 Cancer 400,000 **IVCF** Thrombosis 296,000 VTE 200.000 14,000 0 COPD 138,000 TOTAL DVT PE VTE TV Endocarditis ■FATAL 2,000 294,000 296,000 2,500 Stroke 129,000 ■NON-FATAL 237,000 613,000 376,000 Mechanical / Current Surgical Oral Systemic and **Embolectomy PharmacoMechanical** Anticoagulation **Treatment** Catheter Directed **Options Thrombolysis**

Sources:

Heit JA, et al. Blood. 2005;106:267A.

Murphy SL, et al. Deaths: Preliminary Data for 2010. National Vital Statistics Reports; 2012.

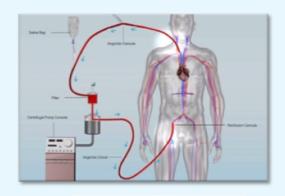


The Thrombus Management Solution

A COMPREHENSIVE SOLUTION FOR VENOUS THROMBOEMBOLISM

ANGIOVAC

Designed for the removal of soft, fresh thrombi or emboli, AngioVac venous drainage system includes the Venous Drainage Cannula and the Extracorporeal Circuit



UNI*FUSE INFUSION CATHETER

Comprehensive line of catheter directed thrombolysis products offering increased flexibility for a wide array of clinical applications





10

Vascular Access Overview







FY2015 Revenue: \$108 million



11

Clinical Need: Addressable PICC Complications

\$450M —	— 6x —	→ \$2.65B	AVG. RATE	AVG. TREATMENT COST
2.7M PICCs		\$1.5B*	Upper Extremity – Deep Vein Thrombosis: 2.0-7.8% ⁴	\$15,973 ¹
Placed		\$107M	Occlusions 25%2**	\$123.77
Annually		\$1.07B	Catheter-Related Bloodstream Infections: 0.4-0.8 per 1,000 catheter days ⁴ OR 2.4% ⁵	
PICC Line Mark		Cost to Treat	ciated	

TREATING THE COMPLICATIONS CAUSED BY PICCs IS

6X

with PICC Lines

GREATER THAN THE TOTAL ACQUISITION COST OF THE DEVICE

- * DVT cost was calculated on a mean DVT rate of 3.5%
- ** Assumption is single-dose efficacy
- 1. Evans S. et al. Chest (2013);143(3)
- 2. Deitcher SR, Fesen MR, Kiproff PM, et al. J Clin Oncol. 12 2002;20(1):317-324
- 3. Centers for Disease Control and Prevention (CDC). MMWR Morb Martal Wkly Rep. 2011;60(8):243-248
- Chopra V et al. The American Journal of Medicine. 2012; 125(8)
- 5. Maki, D. et al. Mayo Clinic Proceedings. 2006;81(9)



The BIOFLO Solution

A REVOLUTIONARY PICC TECHNOLOGY

NO HEPARIN

Minimizes complications associated w/ heparin

NO ANTIBIOTICS

Reduces risks associated w/ bacterial resistance

NOT A COATING

Present throughout entire catheter

NOT ELUTING

Present for life of device

UNLIKE OTHER TECHNOLOGIES THAT ARE SUPERFICIAL AND/OR TRANSIENT, THE BIOFLO MATERIAL IS DESIGNED TO BE PRESENT THROUGHOUT THE CATHETER AND PERMANENT

Coatings

(On the surface)



- ←Coating (Blue)
- ← Catheter Wall





Impregnated Agent (Gold)

← Catheter Wall

BioFlo Technology

(Throughout catheter material)



Catheter Wall

angiodynamics

13

Performance in Clinical Evaluations

FACILITY NAME	NUMBER OF PICCs PLACED	DVT REDUCTION
Academic Medical Center	1,251	85%
Community Hospital	533	25%
Community Hospital	1,212	36%
Hospital Group	252	79%
International Hospital	60	46%
Health System	3,891	32%
Academic Medical Center	52	55%
TOTAL	7,251	51%
TOTAL FACILITY NAME	7,251 NUMBER OF PICCs PLACED	
FACILITY NAME	NUMBER OF PICCs PLACED	TPA REDUCTION
FACILITY NAME Children's Hospital	NUMBER OF PICCs PLACED 272	TPA REDUCTION 42%
FACILITY NAME Children's Hospital Children's Hospital	NUMBER OF PICCs PLACED 272 477	TPA REDUCTION 42% 62%
FACILITY NAME Children's Hospital Children's Hospital Community Hospital	NUMBER OF PICCs PLACED 272 477 533	TPA REDUCTION 42% 62% 75%

MORE THAN 8,000 BIOFLO PICCS HAVE BEEN EVALUATED

51% DVTs
63% tPA

Data on file

Data was obtained during hospital product evaluations; several of these sites are working toward publications

14 Based on data collected at individual institutions; results may not be indicative of clinical experiences at other institutions



Opportunity: BioFlo for Contracted Business



Opportunity: BioFlo Midline

NEW BIOFLO MARKET OPPORTUNITIES

Midline product is drawing not only from the current PICC market, but also from the peripheral IV market

CLINICAL ADVANTAGES

- Ease of insertion for the clinician compared to other competitive devices
- 2. Consistently, effectively draws blood over duration of the treatment
- 3. Treat patients for the entire length of therapy





Opportunity: Tip Location Services

Tip location, coupled with our innovative BioFlo Technology, will improve our competitive position in the PICC market



- March 6, 2014: Acquired regulatory control over Celerity platform and rights to next generation technology
- Mid-summer 2014: U.S. Clearance
- ✓ Winter 2014/15: U.S. No X-ray Clearance
- Fall 2015: EU No X-ray Clearance
- Spring 2016: Navigation
- Fall 2017: FireFly

angiodynamics

Oncology / Surgery Overview

Non-Thermal Ablation



Microwave Ablation



Radiofrequency Ablation





Surgical Resection





FY2015 Revenue: \$52 million



18

Global Cancer Incidence

GLOBAL STATISTICS

Cancer is the leading cause of death worldwide1

14 million new cases of cancer each year1

8.2 million cancer-related deaths in 2012¹ and has not changed in over 50 years²

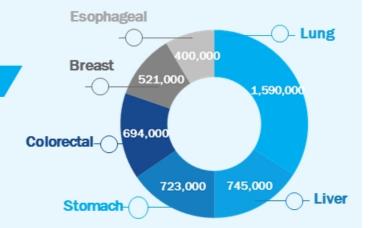
TREATMENT OPTIONS

Surgical Ablation

Radiotherapy

chemotherapy

Palliative Treatment



Most Common Causes of Cancer Death1

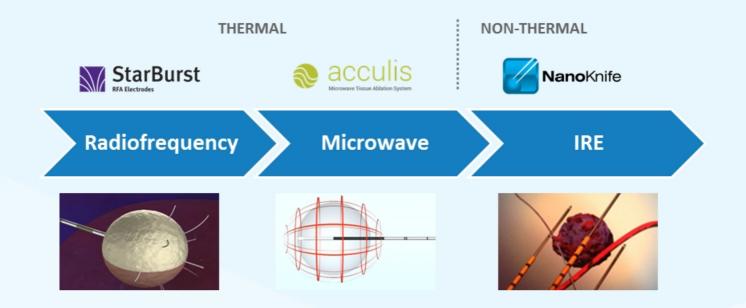


19

1. http://www.who.int/mediacentre/factsheets/fs297/en/

AngioDynamics Leadership in Surgical Ablation

Angio Dynamics offers a full complement of tissue ablation products that provide clinicians maximum choice in treating patients.





The NanoKnife Solution

A REVOLUTIONARY SURGICAL TECHNOLOGY

A new treatment modality for locally advanced pancreatic and hepatic tumors

NOVEL TECHNOLOGY FOR THE ABLATION OF SOFT TISSUE

- Series of short, low energy direct current electrical pulses
- Does not rely on heat to ablate tissue
- Creates defects (pores) in cell membranes
- · Cell death occurs by apoptosis
- This immune mediated cell death allows cellular clearance of debris and creates minimal tissue distortion



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. In the United States, the NanoKnife System has not received clearance for the therapy or treatment of any specific disease or condition.

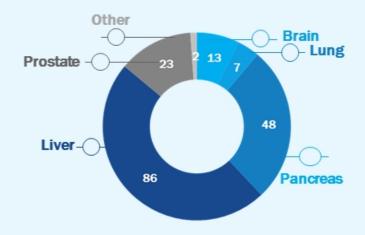


Building Clinical Evidence for NanoKnife



Mounting Clinical Data

More than 150 peer reviewed articles published





Opportunity: Asia-Pacific





2 INVESTING IN THE NEXT GENERATION OF GROWTH



Strong Product Development Pipeline*

	Inflation Device	FY16		
	Drainage Line Extension	FY16		
	AngioVac Line Extension	FY16		
PV	Non-Vascular Introducer	FY16		
	Pain-Free Laser Procedure		FY17	
	New Thrombus Management Product		FY17	
	Fluid Management Line Extension		FY17	
	Celerity Navigation	FY16		
	BioFlo Port Line Extension	FY16		
VA	BioFlo PICC Line Extension	FY16		
	FireFly Tip Location			FY18
	Next Generation BioFlo Port			FY18
	Next Generation BioFlo PICC			FY18
	Next Generation Microwave Ablation		FY17	
0/S	NanoKnife Single Insertion Device		FY17	
,	Next Generation NanoKnife System			FY18
25 *Timelin	es are estimated and subject to change		angio	dynamics



TRANSFORM THROUGH OPERATIONAL EXCELLENCE



Operational Improvements Plan and Impact



Q2 Summary Financial Performance

\$ in millions, except per share	Q2 FY2016	Q2 FY2015	% Change	% Change FX Adjusted
Net sales	\$89.2	\$92.1	-3%	-1% ^(b)
GAAP Earnings per share (EPS)	\$(0.01)	\$0.04	-127%	-108%
Adjusted EPS ^(a)	\$0.14	\$0.17	-19%	-15%
Free Cash Flow	\$9.2	\$(4.6)	-	-

FDA lifts final warning letters for **AngioDynamics**

DECEMBER 3, 2015 BY BRAD PERRIELLO

AngioDy of warning

angiodynamics

The feder

Funding pact includes pause for medical device tax

DECEMBER 16, 2015 BY BRAD PERRIELLO — LEAVE A COMMENT



The \$1.6 trillion deal struck last night on government includes a provision that we Obamacare's medical device tax.



Summary Financial Performance

	Three months ended Nov 30, 2015		Six months end	ded Nov 30, 2015
\$ in millions, except per share amounts	Sales	YoY Growth	Sales	YoY Growth
WW	\$89.2	-3%	\$172.9	-4%
WW ^(a)	\$90.2	-2%	\$174.9	-3%
PV	\$51.1	3%	\$98.2	1%
VA	\$25.0	-11%	\$49.7	-9%
0/S	\$12.4	-9%	\$23.7	-9%
US ^(b)	\$71.4	-2%	\$140.4	-1%
Int'I	\$17.8	-6%	\$32.5	-12%
Int'l ^(a)	\$18.8	-1%	\$34.4	-6%
	Q2 2016	YoY Growth		
Adjusted EBITDA(c)	\$13.4	-16%		
Adjusted EPS(c)	\$0.14	-19%		
Adjusted EPS(a)(c)	\$0.15	-15%		

⁽a) Constant-ourrency basis
(b) Excludes impact of our supply agreement.
(c) Adjusted results exclude costs relating to acquisitions, debt financing, business restructuring, litigation, facility consolidations, amortization of basis step-up of acquired inventory, revaluation of contingent earn outs related to acquisitions, recalls, product discontinuations and amortization of intangible assets.



Balance Sheet and Cash Flow

\$ in millions	Nov30,2015	May 31, 2015
Cash & investments	\$20.6	\$20.1
Net working capital	\$91.5	\$90.3
Totalassets	\$760.4	\$773.6
Totaldebt	\$133.9	\$137.7
Total stockholder's equity	\$547.6	\$545.0
\$ in millions, except per share amounts	3 months ended Nov 30, 2015	3 months ended Nov 30, 2014
Cash Flow from operations	\$9.6	\$(2.1)
Free Cash Flow	\$9.2	\$(4.6)



Adjusted Income Statement(a)

\$ in millions, except per share amounts	Three months ended Nov 30, 2015	Three months ended Nov 30, 2014	Six months ended Nov 30, 2015	Six months ended Nov 30, 2014
Sales	\$89.2	\$92.1	\$172.9	\$179.5
Gross margins	51.4%	51.7%	51.5%	52.1%
Operating expenses	\$35.8	\$36.1	\$71.0	\$71.2
Operating income	\$10.0	\$11.6	\$18.0	\$22.3
Operating margin	11%	13%	10%	12%
Net Income (loss)	\$5.1	\$6.2	\$9.1	\$11.9
EPS	\$0.14	\$0.17	\$0.25	\$0.33
ЕВІТОА	\$13.4	\$15.9	\$25.3	\$30.4

 ⁽a) Adjusted results exclude costs relating to acquisitions, debtfinancing, business restructuring, litigation, facility consolidations, amortization of basis step-up of acquired inventory, revaluation of contingent earn outs related to acquisitions, recalls, product discontinuations and amortization of intangible assets.



AngioDynamics Transformation

Investor Profile

- Major acquisitions Navilyst (2012), Vortex Medical (2012), Microsulis (2013)
- Disruptive technology in PV, VA, and O/S markets with strong product pipeline
- · Significant international opportunity
- Operational Excellence Program in place to drive margin expansion
- · Emerging GPO presence

Fiscal Guidance* \$ in millions, except per share amounts Q3 Full-Year Sales(c) \$84-\$87 \$353-\$359 Adjusted EPS(d) \$0.10-\$0.14 \$0.59-\$0.63

Net Sales

	FY11	FY12	FY13	FY14	FY15
Net Sales	\$216M	\$222M	\$342M	\$354M	\$357M
Reported Growth	0%	3%	54%	4%	1%
Pro Forma	0%	3%	-1% ^(a)	4%	1%
Adjusted Growth	-	-	-	5% ^(b)	2% ^{(b)(c)}

(a) On a pro forma basis, prior year sales for AngioDynamics and Navilyst Medical were \$344.6 million

(b) Excludes impact of our supply agreement

32 (c) Constant-currency basis

