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): November 6, 2017

MRI INTERVENTIONS, INC.

(Exact name of registrant as specified in its charter)

DELAWARE (State or other jurisdiction of incorporation) 001-34822 (Commission File Number) 58-2394628 (I.R.S. Employer Identification Number)

5 Musick Irvine, Ca. 92618 (Address of principal executive offices, zip code)

(949) 900-6833 (Registrant's telephone number, including area code)

the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under the following provisions:
Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)
Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)
Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))
Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))
e by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§240.12b-2 of this chapter). ⊠
nerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying y new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act.

Item 2.02. Results of Operations and Financial Condition.

On November 6, 2017, MRI Interventions, Inc. (the "Company") issued a press release announcing its financial performance for the third fiscal quarter and nine months ended September 30, 2017. A copy of the press release is furnished herewith as Exhibit 99.1.

The information in Item 2.02 of this Form 8-K, as well as Exhibit 99.1 attached hereto, shall not be deemed "filed" for purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the "Exchange Act"), nor shall it be deemed incorporated by reference in any filing under the Securities Act of 1933, as amended (the "Securities Act"), or the Exchange Act, except as shall be expressly set forth by specific reference in such a filing.

Item 7.01. Regulation FD Disclosure

On November 6, 2017, MRI Interventions, Inc. posted an updated investor presentation to its website at http://ir.stockpr.com/mriinterventions/investor-presentation. A copy of the investor presentation is being furnished herewith as Exhibit 99.2. The Company may use the investor presentation from time to time in conversations with analysts, investors and others.

The information in Item 7.01 of this Form 8-K, as well as Exhibit 99.2 attached hereto, shall not be deemed "filed" for the purposes of Section 18 of the Exchange Act, nor shall it be deemed incorporated by reference in any filing under the Securities Act, or the Exchange Act, except as shall be expressly set forth by specific reference in such a filing.

The information contained in Exhibit 99.2 is summary information that is intended to be considered in the context of the Company's filings with the Securities and Exchange Commission ("SEC") and other public announcements that the Company may make from time to time, by press release or otherwise. The Company undertakes no duty or obligation to publicly update or revise the information contained in this investor presentation, although it may do so from time to time as its management believes is warranted. Any such updating may be made through the filing or other reports or documents with the SEC, through press releases or other public disclosure.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits.

The following exhibits are furnished herewith:

Exhibit 99.1 Press Release dated November 6, 2017.

Exhibit 99.2 Investor Presentation dated November 2017.

SIGNATURES

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.

Date: November 6, 2017 MRI INTERVENTIONS, INC.

By: /s/ Harold A. Hurwitz

Harold A. Hurwitz Chief Financial Officer

MRI Interventions, Inc. Reports 26% Year-Over-Year Increase in Third Quarter Procedures

Gross Margin Increases to 60% Record 469 ClearPoint Procedures Performed Year-to-Date

IRVINE, CA, November 6, 2017 – MRI Interventions, Inc. (OTCQB: MRIC) (the "Company") today announced financial results for the third quarter ended September 30, 2017.

2017 Third Quarter and Recent Highlights

- Total revenue increased 6% to \$1.7 million, from \$1.6 million in the same period of 2016, reflecting a 12% increase in disposable sales, offset by lower capital equipment sales;
- Revenue year-to-date increased 39% to \$5.7 million, from \$4.1 million in the same period of 2016;
- Gross margin increased to 60%, compared with 54% in the same period of 2016;
- Net loss declined 44% year-over-year to \$1.4 million, as compared with \$2.6 million for the same period in 2016;
- Completed 161 procedures using the ClearPoint[®] Neuro Navigational System, a quarterly increase of 26%, and completed a record 469 procedures year-to-date;
- Sold one ClearPoint system, and initiated two new system evaluations;
- Grew the installed base of ClearPoint Systems to 52 centers in the U.S., including the Company's seventh top-ten children's hospital;
- Furthered research and development efforts with partners toward the development of new therapeutic applications for intra-cranial hemorrhage and pancreatic cancer;
- Participated in a One Room-One Procedure laser ablation Practical Clinic during the Congress of Neurological Surgeons; and
- Announced the appointment of Joseph Burnett as President and Chief Executive Officer effective November 7, 2017.

Frank Grillo, current President and Chief Executive Officer of MRI Interventions, Inc., said: "ClearPoint treatments increased 26% year over year in the third quarter, demonstrating increased market adoption of our platform. Treatment volume was slower at the beginning of the quarter in what we believe was a seasonal effect, and then accelerated at the end of the quarter, an increased pace that has continued into October and November. We remain focused on expanding utilization at our clinical sites across the U.S., as well as securing additional centers to further our neurosurgical market share. Several evaluation sites have successfully adopted ClearPoint, and we are focused on converting these locations into sales in the 2017 fourth quarter and beyond. Neurosurgeons are increasingly recognizing the value of real-time intraoperative MRI guidance in high-growth areas such as deep brain stimulation and laser ablation, as well as the compelling safety and patient comfort benefits inherent to the ClearPoint platform. Our hospital partners benefit as well, since patients are more comfortable with ClearPoint procedures than traditional approaches which may require the patient to be awake during surgery."

Joe Burnett, incoming President and Chief Executive Officer of MRI Interventions, Inc., remarked: "I look forward to capitalizing on the existing installed base of ClearPoint Systems to further increase procedure growth, as well as converting our large and growing pipeline of prospective sites into ClearPoint centers of excellence. During Frank's tenure as CEO, the Company has successfully commercialized its platform, brought ClearPoint into the mainstream of neurosurgery procedures and obtained funding to allow it to set its sights on strategic objectives. This is an exciting time at the Company, and I look forward to working closely with our team to create further shareholder value through continued growth."

Financial Results - Quarter Ended September 30, 2017

Total revenues were \$1.7 million for the three months ended September 30, 2017, an increase of \$101,000, or 6%, compared with \$1.6 million for the same period in 2016.

ClearPoint disposable product sales increased \$149,000, or 12%, to \$1.4 million for the three months ended September 30, 2017, compared with \$1.3 million for the same period in 2016. This growth in disposable sales reflected 161 ClearPoint procedures performed in the 2017 third quarter.

ClearPoint reusable product sales were \$208,000 for the three months ended September 30, 2017, compared with \$309,000 for the same period in 2016. Reusable products consist primarily of computer hardware and software bearing sales prices that are appreciably higher than those for disposable products and fluctuate from quarter to quarter.

Gross margin for the three months ended September 30, 2017 was 60%, compared to gross margin of 54% for the same period in 2016. The increase in gross margin primarily reflected decreased charges for inventory obsolescence and a favorable mix of product sold, comprised of a greater share of disposable products during the three months ended September 30, 2017, relative to the same period in 2016.

Research and development costs were \$590,000 during the three months ended September 30, 2017, compared to \$691,000 during the same period in 2016, a decrease of \$101,000, or 15%. The decrease was due primarily to reductions in software development and intellectual property costs, partially offset by an increase in new product development costs.

Selling, general and administrative expenses were \$1.8 million for the three months ended September 30, 2017, as compared to \$1.9 million for the same period in 2016, a decrease of \$120,000, or 6%. The decrease was due primarily to reduced financing costs and stock compensation expense, which were partially offset by increased recruiting expenses during the three months ended September 30, 2017, relative to the same period in 2016.

The Company's operating loss for the three months ended September 30, 2017 declined \$382,000, or 22%, to \$1.3 million, as compared with \$1.7 million for the same period in 2016.

In August 2016, the Company recorded a debt restructuring loss of \$933,000 resulting from amendments entered into with two holders of the 2014 junior secured notes payable (the "2014 Note Holders") who then converted \$1.75 million of aggregate principal balance of their notes into equity in connection with the Company's private placement of equity securities in September 2016.

During the three months ended September 30, 2017 and 2016, the Company recorded gains of \$110,000 and \$324,000, respectively, resulting from changes in the fair value of derivative liabilities. For the three months ended September 30, 2017, such derivative liabilities related to: (a) the issuance of warrants in connection with a 2013 private placement transaction; (b) a note amendment entered into with Brainlab AG ("Brainlab") in June 2016; and (c) the amendments entered into with the 2014 Note Holders discussed above. For the three months ended September 30, 2016, derivative liabilities included the foregoing and warrants issued with down-round price protection provisions in connection with a 2012 private placement transaction.

Net interest expense during the three months ended September 30, 2017 and 2016 was \$211,000 and \$240,000, respectively, a decrease of \$29,000, or 12%. This decrease was due to the reduction of principal balances resulting from the conversion into equity of an aggregate \$1.75 million principal balance of the notes discussed above.

Reflecting the effects of these non-operational items, net loss for the three months ended September 30, 2017 was \$1.4 million, as compared with \$2.6 million for the same period in 2016.

Cash used in operating activities for the three months ended September 30, 2017 was \$1.7 million, as compared with \$1.4 million of for the same period in 2016. The increase was due primarily to growth in accounts receivable, consistent with growth in total revenues during the three months ended September 30, 2017, relative to the same period in 2016, and to a planned increase in inventory safety stock levels during the three months ended September 30, 2017.

Financial Results - Nine Months Ended September 30, 2017

Total revenues were \$5.7 million for the nine months ended September 30, 2017, an increase of \$1.6 million, or 39%, compared with \$4.1 million for the same period in 2016. This increase was due primarily to an increase in the Company's disposable and reusable product sales.

ClearPoint disposable product sales increased \$1.1 million, or 33%, to \$4.5 million for the nine months ended September 30, 2017, compared with \$3.4 million for the same period in 2016. This growth in disposable sales reflected a record 469 ClearPoint procedures performed during the nine months ended September 30, 2017. ClearPoint reusable product sales were \$923,000 for the nine months ended September 30, 2017, compared with \$608,000 for the same period in 2016.

Gross margin for the nine months ended September 30, 2017 was 61%, compared to gross margin of 52% for the same period in 2016.

Research and development costs were \$2.2 million during the nine months ended September 30, 2017, compared to \$2.1 million during the same period in 2016, an increase of \$133,000, or 6%. The increase was due to the upfront payments required under the previously announced development agreements entered into in April 2017 with the Mayo Clinic and Acoustic MedSystems, Inc., which were partially offset by reductions in software development and intellectual property costs, and compensation expenses.

Selling, general and administrative expenses were \$5.7 million during each of the nine months ended September 30, 2017 and 2016. Increases in personnel-related costs due to headcount increases in the Company's commercial team were offset by decreases in professional fees and stock-based compensation costs.

The Company's operating loss for the nine months ended September 30, 2017 declined \$1.2 million, or 21%, to \$4.5 million, as compared with \$5.7 million for the same period in 2016.

During the nine months ended September 30, 2016, the Company recorded a net loss from debt restructuring of \$812,000, the components of which were: (a) a gain of \$941,000 resulting from the restructuring of the note payable to Brainlab in April 2016; (b) a loss of \$820,000 resulting from amendments made in June 2016 to: (i) the note payable to Brainlab; and (ii) the notes held by the 2014 Note Holders; and (c) a loss of \$933,000 resulting from amendments entered into with the 2014 Note Holders in August 2016 as described above.

During the nine months ended September 30, 2017 and 2016, the Company recorded gains of \$48,000 and \$748,000, respectively, resulting from changes in the fair value of the derivative liabilities existing at those respective dates as described above.

During the nine months ended September 30, 2017, the Company recorded other income of \$7,000, as compared with other income of \$210,000 recorded during the same period in 2016, representing a decrease of \$203,000, or 97%. This decrease was due primarily to grant income from a U.S. federal agency related to a project in process during the nine months ended September 30, 2016, which was discontinued by the agency later in 2016. The Company has not since undertaken any additional such projects.

Net interest expense during the nine months ended September 30, 2017 and 2016 was \$637,000 and \$836,000, respectively, a decrease of \$199,000, or 24%. This decrease was due to the reduction of principal balances as described above.

Reflecting the effects of these non-operational items, net loss for the nine months ended September 30, 2017 was \$5.1 million, as compared with \$6.4 million for the same period in 2016.

Cash used in operating activities for the nine months ended September 30, 2017 was \$4.2 million, as compared with \$4.7 million of for the same period in 2016. This improvement was due primarily to the reduction in the Company's net loss for the nine months ended September 30, 2017, relative to the same period in 2016, which was partially offset by the planned increase in inventory safety stock levels during the three months ended September 30, 2017.

Private Placement

As previously announced, on May 26, 2017, the Company completed a private placement of equity units, which resulted in gross proceeds of \$13.25 million, before deducting placement agents' fees and offering expenses.

Conference Call and Webcast

Investors and analysts who would like to participate in a conference call today at 4:30 p.m. Eastern Time (1:30 p.m. Pacific Time) to review the Company's financial results may do so via telephone at (877) 407-9034, or at (201) 493-6737 if calling from outside the U.S. or Canada. Callers should dial in at least 5 minutes prior to the call start time.

A live and archived webcast may be accessed by visiting the Company's website at www.mriinterventions.com, by selecting "Investors" / "News" / "IR Calendar." The conference call may also be accessed at http://mriinterventions.equisolvewebcast.com/q3-2017. A replay of the conference call will be available shortly after completion of the call until November 13, 2017 by calling (877) 660-6853, or (201) 612-7415 if calling from outside the U.S. or Canada, and then entering conference ID number 413671.

About MRI Interventions, Inc.

Building on the imaging power of magnetic resonance imaging ("MRI"), MRI Interventions is creating innovative platforms for performing the next generation of minimally invasive surgical procedures in the brain. The ClearPoint Neuro Navigation System, which has received 510(k) clearance and is CE marked, utilizes a hospital's existing diagnostic or intraoperative MRI suite to enable a range of minimally invasive procedures in the brain. For more information, please visit www.mriinterventions.com.

Forward-Looking Statements

Statements herein concerning the Company's plans, growth and strategies may include forward-looking statements within the context of the federal securities laws. Statements regarding the Company's future events, developments and future performance, as well as management's expectations, beliefs, plans, estimates or projections relating to the future, are forward-looking statements within the meaning of these laws. Uncertainties and risks may cause the Company's actual results to differ materially from those expressed in or implied by forward-looking statements. Particular uncertainties and risks include those relating to: future revenues from sales of the Company's ClearPoint Neuro Navigation System products; the Company's ability to market, commercialize and achieve broader market acceptance for the Company's ClearPoint Neuro Navigation System products; and estimates regarding the sufficiency of the Company's cash resources. More detailed information on these and additional factors that could affect the Company's actual results are described in the "Risk Factors" section of the Company's Annual Report on Form 10-K for the year ended December 31, 2016, as well as the Company's Quarterly Report on Form 10-Q for the quarter ended September 30, 2017, both of which have been filed with the Securities and Exchange Commission.

Contact: Harold A. Hurwitz, Chief Financial Officer

(949) 900-6833

Matt Kreps, Darrow Associates Investor Relations

(512) 696-6401; mkreps@darrowir.com

(tables follow)

MRI INTERVENTIONS, INC. Condensed Consolidated Statements of Operations (Unaudited)

For The Three M	Ionths Ended	nded
Sentemb	er 30	

	 Septen	nber 30,	
	2017		2016
Revenues:			
Product revenues	\$ 1,628,435	\$	1,580,826
Service and other revenues	88,635		35,507
Total revenues	1,717,070		1,616,333
Cost of product revenues	688,847		748,305
Research and development costs	589,716		691,330
Selling, general, and administrative expenses	1,765,830		1,886,220
Operating loss	(1,327,323)		(1,709,522)
Other income (expense):			
Loss from debt restructuring	_		(933,134)
Gain from change in fair value of derivative liabilities	109,803		324,035
Other income (expense), net	3,363		(4,877)
Interest expense, net	 (211,362)		(239,733)
Net loss	\$ (1,425,519)	\$	(2,563,231)
Net loss per share attributable to common stockholders:			
Basic and diluted	\$ (0.14)	\$	(0.92)
Weighted average shares outstanding:			
Basic and diluted	10,339,210		2,779,803

MRI INTERVENTIONS, INC. Condensed Consolidated Statement of Operations (Unaudited)

For Th	e Nine	Months	Ended
	Cantar	nhor 20	

	 Septen	1ber 30,	
	2017		2016
Revenues:			
Product revenues	\$ 5,443,287	\$	4,013,531
Service and other revenues	256,860		100,818
Total revenues	 5,700,147	•	4,114,349
Cost of product revenues	2,239,808		1,965,839
Research and development costs	2,231,616		2,098,465
Selling, general, and administrative expenses	5,731,961		5,748,524
Operating loss	 (4,503,238)		(5,698,479)
Other income (expense):			
Loss from debt restructuring	_		(811,909)
Gain from change in fair value of derivative liabilities	48,064		748,080
Other income, net	6,774		209,504
Interest expense, net	 (637,270)		(836,208)
Net loss	\$ (5,085,670)	\$	(6,389,012)
Net loss per share attributable to common stockholders:	 		
Basic and diluted	\$ (0.75)	\$	(2.59)
Weighted average shares outstanding:			
Basic and diluted	6,783,605		2,467,437

MRI INTERVENTIONS, INC. Condensed Consolidated Balance Sheets (Unaudited)

September 30, 2017		_	D	December 31, 2016	
ASSETS					
Current Assets:					
Cash and cash equivalents	\$	11,028,973	\$	3,315,774	
Accounts receivable, net of allowance for doubtful accounts of \$26,714 and \$25,000					
at September 30, 2017 and December 31, 2016, respectively		863,758		865,943	
Inventory, net		2,181,245		1,768,382	
Prepaid expenses and other current assets		239,326		134,996	
Total current assets		14,313,302		6,085,095	
Property and equipment, net		246,381		328,249	
Software license inventory		889,400		976,900	
Other assets		10,641		10,641	
Total assets	\$	15,459,724	\$	7,400,885	
LIABILITIES AND STOCKHOLDERS' EQUITY (DEFICIT)					
Current liabilities:					
Accounts payable	\$	957,660	\$	1,546,926	
Accrued compensation	Ψ	809,323	Ψ	666,060	
Other accrued liabilities		529,942		450,424	
Derivative liabilities		72,450		131,173	
Deferred product and service revenues		281,826		223,117	
Total current liabilities		2,651,201		3,017,700	
Accrued interest		734,625		647,500	
Senior secured note payable		2,000,000		2,000,000	
2014 junior secured notes payable, net of unamortized discount and deferred issuance		2,000,000		2,000,000	
costs of \$120,516 and \$180,774 at September 30, 2017 and December 31, 2016,					
respectively		1,854,484		1,794,226	
2010 junior secured notes payable, net of unamortized discount of \$2,060,865 and					
\$2,302,472 at September 30, 2017 and December 31, 2016, respectively		939,135		697,528	
Total liabilities		8,179,445		8,156,954	
Commitments and contingencies	_				
Stockholders' equity (deficit):					
Preferred stock, \$0.01 par value; 25,000,000 shares authorized at September 30,					
2017 and December 31, 2016; none issued and outstanding at September 30, 2017					
and December 31, 2016		_		_	
Common stock, \$0.01 par value; 200,000,000 shares authorized; 10,339,210 shares					
issued and outstanding at September 30, 2017; and 3,622,032 issued and					
outstanding at December 31, 2016		103,391		36,220	
Additional paid-in capital		106,131,322		93,076,475	
Accumulated deficit		(98,954,434)		(93,868,764)	
Total stockholders' equity (deficit)		7,280,279		(756,069)	
Total liabilities and stockholders' equity (deficit)	\$	15,459,724	\$	7,400,885	
Tour institutes and stookholders equity (deficit)	4	10,.00,121	Ψ	,,.50,005	

MRI INTERVENTIONS, INC. Condensed Consolidated Statements of Cash Flows (Unaudited)

For T	The Nine	Months	Ended
	Santa	mbor 30	

	September 50,			
		2017		2016
Cash flows from operating activities:				
Net loss	\$	(5,085,670)	\$	(6,389,012)
Adjustments to reconcile net loss to net cash flows from operating activities:				
Depreciation and amortization		92,656		125,076
Share-based compensation		616,007		736,982
Expenses paid through the issuance of common stock		502,032		259,898
Gain from change in fair value of derivative liabilities		(48,064)		(748,080)
Amortization of debt issuance costs and original issue discounts		301,865		323,016
Loss from retirement of fixed assets		_		1,689
Loss from debt restructuring		_		811,909
Increase (decrease) in cash resulting from changes in:				
Accounts receivable		2,185		414,506
Inventory, net		(311,637)		(33,958)
Prepaid expenses and other current assets		(104,329)		(96,358)
Accounts payable and accrued expenses		(270,535)		(220,304)
Deferred revenue		58,709		106,479
Net cash flows from operating activities		(4,246,781)		(4,708,157)
Cash flows from investing activities:			· · · · · ·	
Purchases of property and equipment		(24,515)		(100,324)
Net cash flows from investing activities		(24,515)		(100,324)
Cash flows from financing activities:				
Proceeds from private equity offering		13,250,000		4,255,000
Offering costs		(1,265,505)		(417,865)
Other		_		(4,756)
Net cash flows from financing activities		11,984,495		3,832,379
Net change in cash and cash equivalents		7,713,199		(976,102)
Cash and cash equivalents, beginning of period		3,315,774		5,408,523
Cash and cash equivalents, end of period	\$	11,028,973	\$	4,432,421
SUPPLEMENTAL CASH FLOW INFORMATION				
Cash paid for:				
Income taxes	\$		\$	<u> </u>
Interest	\$	146,611	\$	976,295



Investor Presentation

November 2017



Forward Looking Statements

Statements herein concerning MRI Interventions, Inc. (the "Company") plans, growth and strategies may include forward-looking statements within the context of the federal securities laws. Statements regarding the Company's future events, developments and future performance, as well as management's expectations, beliefs, plans, estimates or projections relating to the future, are forward-looking statements within the meaning of these laws. Uncertainties and risks may cause the Company's actual results to differ materially from those expressed in or implied by forward-looking statements. Particular uncertainties and risks include those relating to: the Company's ability to obtain additional financing; estimates regarding the sufficiency of the Company's cash resources; future revenues from sales of the Company's ClearPoint System products; and the Company's ability to market, commercialize and achieve broader market acceptance for the Company's ClearPoint System products. More detailed information on these and additional factors that could affect the Company's actual results are described in the "Risk Factors" section of the Company's Annual Report on Form 10-K for the year ended December 31, 2016, which has been filed with the Securities and Exchange Commission, and our most recently filed Quarterly Report on Form 10-Q for the quarter ended June 30, 2017, which will be filed with the Securities and Exchange Commission on or before August 15, 2017.



MRIC: The Platform Company for MRI-Guided Therapies

- MRI Interventions: Leader in the Delivery of MRI-Guided Therapies
 - Primary Innovator in the Field, grown out of the Advanced MR department at Johns Hopkins
 - Extensive Intellectual Property position: 70+ issued US patents, 45 international
 - · Proven track record for conceiving, developing, commercializing and securing clinical adoption for our MRI-guided therapy platform
- Rapidly Growing Commercial Business; Established the Industry Platform for MRI-guided, Minimally-Invasive Neurosurgery
 - · Strong and growing clinical footprint: 48 hospitals in the US
 - Accelerating adoption, revenue growth: 8 successive quarters of record procedures
 - Current procedures include DBS electrode placement, laser ablation, biopsy, drug delivery
 - MRIC's platform integrates with products from multiple companies (med device companies, imaging companies, biotechs)
- Expanding Platform into Adjacent Areas to Address Additional Unmet Medical Needs
 - Expanding platform into the stroke market; Joint Development Agmt with Mayo Clinic
 - Expanding our capabilities, adding novel ultrasound ablation technology via Co-Dev Agmt with Acoustic MedSystems; initial market is pancreatic cancer
 - Leveraging our existing platform, install base and technologies to enable a rapid, cost-effective path to additional markets



Our MRI-Guided Therapy Platform is Currently Being Used to:

Implant Neuro Stimulation Products from:

- · Medtronic
- · St. Jude Medical
- NeuroPace







Place Laser Ablation Probes from:

- · Medtronic-Visualase
- · Monteris Medical





Deliver Drugs and Biologics from:

- Voyager
- · Medicenna
- · Oxford Biomedica
- · International Stem Cell Corporation









MRIC Platform Runs on All Major Scanners:

- · Siemens
- GE
- · Philips
- IMRIS











Broad and Growing User Base of Leading Neurosurgeons

ClearPoint® Is Installed in 52 Top US Hospitals and Growing

UC San Francisco San Francisco VA Stanford Univ **UCSF Benioff Childrens** USC UC San Diego Univ of Colo Univ of Utah Univ of Arizona Cook Children's MD Anderson Methodist Hosp Texas Children's Hosp Riverside Nationwide Children's Children's Mercy Kansas Univ Med Center Univ of Wisconsin Spectrum Health Ohio State Univ Univ of Cincinnati Univ of Michigan



Brigham & Women's Boston Children's Yale Univ Univ of Pitt Med Center Memorial Sloan Kettering Hackensack Univ Med Center Cornell Central Du Page Nat. Institutes of Health Nat Children's Hospital Children's Hosp of Philadelphia Univ of Virginia **Emory University** Carillion **Duke University** Children's of Alabama **CHOA Scottish Rite** Willis Knighton Mayo Clinic Jacksonville Miami Children's

Strong Commercial Sales and Clinical Support Teams in Place



Foundation of Our Platform:

ClearPoint Neuro Navigation System



Without ClearPoint, Minimally Invasive Neuro Procedures Are Performed "Blind"

Conventional Stereotactic Procedure



No real time images – images from earlier in the day or week

ClearPoint Neuro Procedure





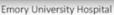
ClearPoint Neuro Navigation System: MRI-Guided NeuroSurgical Platform

ClearPoint Components:











Univ. of California San Francisco Medical Center



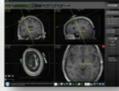
Univ. of Pittsburgh Medical Center

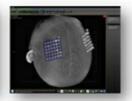


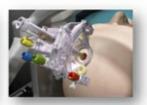
Details on Our ClearPoint Neuronavigation System Platform

ClearPoint Procedure Overview:

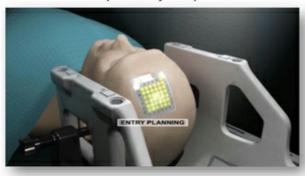








ClearPoint Video **Entry and Trajectory**



ClearPoint Video Alignment and Insertion

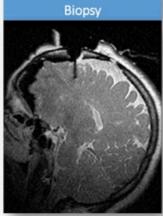


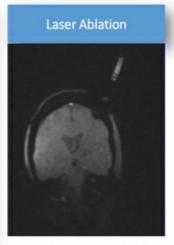


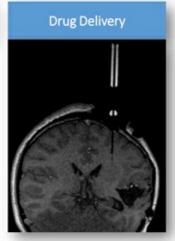
Multiple Clinical Applications for Our ClearPoint System

MRI-Guided Therapy Platform









Delivering Therapies to Address Significant Unmet Medical Needs:

- Parkinson's disease, Epilepsy, Brain Tumor, Dystonia

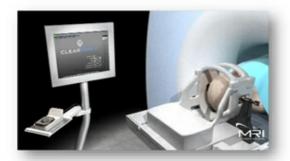
Enabling Multiple Therapies from Multiple Companies:

- Electrode Placement: Medtronic, St. Jude, Neuropace Laser Ablation: MDT/Visualase, Monteris Drug Delivery: Voyager, Medicenna, Oxford Biomedica, Int'l Stem Cell
- Laser Ablation MR Thermometry is an MRI-based functionality available on most MR scanner platforms and it is a feature built into products from several third party vendors. The ClearPoint system enables MRI-guided procedures and allows physicians to use this inherent MR capability during a procedure.
 Drug Delivery The SmartFlow* cannula received 510(k) clearance for injection of cytarabine, a chemotherapy drug, to the ventricles or removal of CSF from the ventricles during intracranial procedures. Delivery of other therapeutic agents, and delivery of agents to other areas of the brain, using the SmartFlow cannula is investigational.



Strong Business Model for Our Core MRI-Guided Navigation Platform

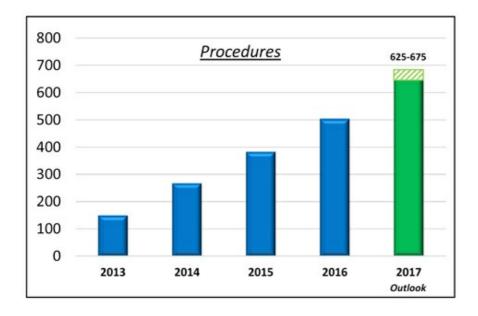
- ClearPoint Hardware/Software: \$100,000 \$150,000 ASP
- ClearPoint Disposables: ~\$7,500 ASP per procedure with strong margins
- Recurring revenue from the sale of disposables
- Procedures covered by existing inpatient DRG reimbursement codes
- Same razor/razorblade model applies to future product sales







ClearPoint Neurosurgical Procedure Growth



CAGR, 2013 - 2016: 46%

- Q1, 2017: 146 Procedures
- Q2, 2017: 162 Procedures
- Q3, 2017: 161 Procedures



Expanding Our Platform:

New Procedural Applications



2017 MRI INTERVENTIONS, INC.

Expanding MRIC's MRI-Guided Therapy Platform Benefits Larger Patient Populations

First, MRIC established the Capability of the Platform...

- Began with Electrode Placement (DBS) Parkinson's disease, dystonia
- ✓ Expanded to Biopsy Brain Tumor
 - ✓ Expanded to Laser Ablation Probe Placement Epilepsy and Brain Tumor
 - ✓ Expanded to Drug Delivery Parkinson's Disease and Brain Tumor

...now, Expanding the Reach of the MRI-Guided Platform

- ✓ Expanding into the Stroke Market Intra-Cranial Hemorrhage
- ✓ Adding Novel Ablation Technology to Platform Initial focus in Pancreatic Cancer



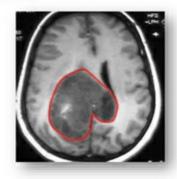
Expanding into the Stroke Market Leveraging Existing Capabilities and Install Base

Expanding the Reach of the MRIC Platform into the Stroke Market

- · Joint Development Agreement with Mayo Clinic
- · Dr. Bernard Bendok, Chair of Neurosurgery, Mayo Arizona

Large Stroke Market

- · 5th leading cause of death in US
- · Leading cause of permanent disability
- · Affects 800,000 people in the US every year



Initial Stroke Product (ClearAway™) Targets Intracerebral Hemorrhage (ICH)

- · Only major stroke subtype w/o clearly effective therapy major unmet medical need
- Affects 80,000 to 100,000 people in the US each year
- MRIC market opportunity is 12,000 to 15,000 cases/year



Our MRI-Guided Approach to Intra Cerebral Hemorrhagic Stroke

Current Approaches for Hemorrhage Removal and Decompression are Inadequate:

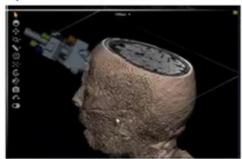
- · Open craniotomy provides visibility but is highly invasive, destroys brain tissue
- Minimally Invasive approach has very limited ability to quantify volume reduction of the hemorrhage or monitor subsequent bleeds

Our Unique ClearAway MRI-Guided Therapy Approach to ICH

- · Detailed, continuous, high resolution, 3 dimensional visibility
- · Minimally invasive approach

Expected Path to Market

- · Builds directly on our ClearPoint Platform
 - · ClearPoint software and hardware; SmartFrame
 - · Hemorrhage aspiration components
- · Potential 510(k) regulatory path
- · Projected market introduction as soon as 12-18 months



U.S. Market Opportunity = \$72 million to \$90 million

95% of Our Existing ClearPoint Adult Sites Have a Stroke Program!



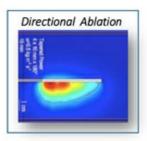
Expanding our Platform with a Novel Ablation Technology Co-Development Agreement Will Add Additional Capabilities

License and Co-Development Agreement with Acoustic MedSystems (AMS)

- · Leading developer of ultrasound ablation technologies and systems
- · Base technology has 510k clearance
- · Initial license areas: WW Excl. for Pancreatic Cancer, Excl. Option for Brain Applications

AMS Ultrasound Technology Provides Unique Capabilities over Other Ablative Energy Sources

- Precision ablation over small and large areas
- · Directional and depth control
- · "Protected Zone" between transducer and ablated tissue



AMS Ultrasound Technology Works with MRI Interventions' Technology

- · MRI-friendly components
- · Enhanced by real-time, MRI-guidance
- Can incorporate real-time MRI-thermometry



Strong Pipeline of New Products to Add to our Existing Products Leveraging our MRI-Guided Therapy Platform

Projected Development Pipeline		2017	40	2018			
A CONTRACTOR OF THE STATE OF TH	Q2	Q3	Q4	Q1	Q2	Q3	Q4
New Products in Development							
ClearPoint Software 2.0			\rightarrow			commercializing	
ClearPoint Drug Delivery Cannula – CE Mark					commercializi	ng	
ClearAway™ - Intracerebral Hemorrhage					Commercializing / F		
ClearAblate TM – Pancreas/Liver (Gen 1)							
ClearAblate™ – Pancreas (Gen 2)					de	velopment	
Drug Trials in Progress							
Voyager (Parkinson's disease)		T.				II.	
Medicenna (GBM Brain Tumors)				- 1			
Oxford Biomedica (Parkinson's disease)							
Lysogene (Sanfilippo A)							
International Stem Cell (for Parkinson's disease)				1			





Addressable Markets for our MRI-Guided Therapy Platform

	Functiona	l Neurosurge	ry Market	Stroke	Oncology	Drug Deliv	ery Market
	Parkinsons	Epilepsy	Brain Tumor	ICH	Pancreas	Parkinson's	Brian Tumor
Prevalence	1,500,000	2,200,000	80,000	90,000	50,000	1,500,000	80,000
Annual Potential Procedures	12,500	28,000	15,000	13,500	12,500	25,000	26,000
Est. Average Selling Price	\$7,500	\$7,500	\$7,500	\$6,000	\$5,000	\$14,000	\$8,000
Annual Market Opportunity	\$93 Million	\$210 Million	\$112 Million	\$81 Million	\$62 Million	\$350 Million	\$208 Million

18-24 Months 12-18 Months 24+ Months **Growing Current Market** Same Hospital Same Hospital Continue Continue to Grow Install Base Same MRI Suite involvement in Same MRI Suite and Increase Utilization Neurosurgery New Physician **Current Trials New Application** New Application Add New Trials

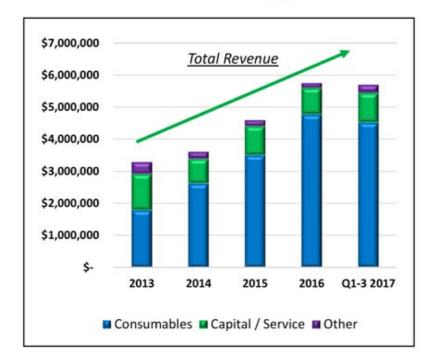
Note: Market sizes for brain tumor, ICH are Pancreatic cancer are incidence numbers because of the nature of those diseases. Prevalence numbers are based on either market research conducted by a third party on behalf of MRI Interventions or research conducted by MRI Interventions of publicly available sources.



Financials



Revenue Growth Accelerating...



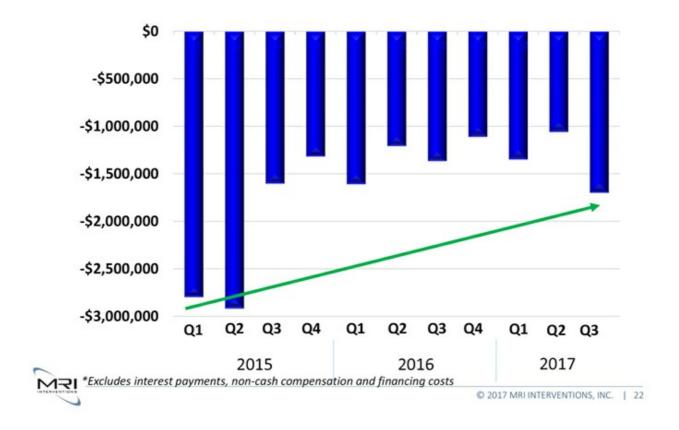
Disposable Product
Estimated CAGR: 30%

Other Results of Note:

- Installed at 52 of 250+ target US neurosurgery centers
- H1 2017 revenue of \$4.0 mm
- 2017 nine months of revenue are \$5.4million; 2016 full year revenue was \$5.6 million



Cash Flow From Operations*: 2015 – Q2, 2017



Income Statement, Last 5 Quarters

				Quar	ter Ende	d	
	Se	p-16	De	ec-16	Mar-17	Jun-17	Sep-17
Revenues:							
Disposable product revenues	\$	1,272	\$	1,363	\$ 1,663	\$ 1,436	\$ 1,421
Reusable product revenues	100 miles	309		224	259	457	208
Total product revenues		1,581		1,587	1,922	1,893	1,629
Service revenues		35		48	85	83	88
Total revenues		1,616		1,635	2,007	1,976	1,717
Cost of product revenues		748		677	752	798	689
Gross profit		868		958	1,255	1,178	1,028
Gross profit %		55%		60%	65%	62%	60%
Total operating expenses	<u> </u>	2,578		2,748	2,608	3,000	2,355
Operating loss	((1,710)		(1,790)	(1,354)	(1,822)	(1,327



Experienced Medical Device Management Team

Management Team

Joseph Burnett CEO





Peter Piferi COO

HeartWare Cordis

Wendelin Maners

VP Sales & Mrktg

Scientific



Hal Hurwitz **CFO**





Board of Directors















Summary

Significant Value in Owning the MRI-Guided Therapy Platform

Leader in this Field

- Primary Innovator, Established Clinical Footprint, Industry Integration, IP

World-Class Research Institutions Behind All Major Initiatives

Proven Ability to Develop, Commercialize and Secure Clinical Adoption of our Platform

Leveraging our Prior Investment to Cost-Effectively Expand into Stroke Market

Adding a Unique Ultrasound Ablation Capability to Broaden our Platform

Strong Revenue Growth and a Strong Product Pipeline

