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): **March 6, 2023**

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)

(518) 795-1400

(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:

☐ 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))

Securities registered pursuant to Section 12(b) of the Act:

Title of each class	Trading Symbol(s)	Name of each exchange on which registered
Common Stock, par value $0.01 per share	ANGO	NASDAQ Global Select Market

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (§230.405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§240.12b-2 of this chapter).

Emerging growth company ☐

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act. ☐

Item 7.01 – Regulation FD Disclosure.

On March 6, 2023, James Clemmer, President and Chief Executive Officer of AngioDynamics, Inc. ("AngioDynamics"), and Stephen Trowbridge, Executive Vice President and Chief Financial Officer of AngioDynamics, will present at the Raymond James 44th Annual Institutional Investors Conference. The presentation slides are furnished herewith as Exhibit 99.1.

The presentation slides furnished pursuant to Item 7.01 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 otherwise subject to the liabilities under that Section. Furthermore, the presentation slides shall not be deemed to be incorporated by reference into any filing under the Securities Act of 1933, as amended, or the Exchange Act.

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," "projects", "optimistic," or variations of such words and similar expressions, are forward-looking statements. These forward-looking statements are not guarantees of future performance and are subject to risks and uncertainties. Investors are cautioned that actual events or results may differ materially from AngioDynamics' expectations, expressed or implied. Factors that may affect the actual results achieved by AngioDynamics include, without limitation, the scale and scope of the COVID-19 global pandemic, the ability of AngioDynamics to develop its existing and new products, technological advances and patents attained by competitors, infringement of AngioDynamics' technology or assertions that AngioDynamics' technology infringes the technology of third parties, the ability of AngioDynamics to effectively compete against competitors that have substantially greater resources, future actions by the FDA or other regulatory agencies, domestic and foreign health care reforms and government regulations, results of pending or future clinical trials, overall economic conditions (including inflation, labor shortages and supply chain challenges including the cost and availability of raw materials), 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 obtain regulatory clearances or approval of its products, or to integrate acquired 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, 2022. AngioDynamics does not assume any obligation to publicly update or revise any forward-looking statements for any reason.

Item 9.01 – Financial Statements and Exhibits.

(d) *Exhibits*.

Exhibit No.	Description
99.1	Presentation slides for the Raymond James 44th Annual Institutional Investors Conference, dated March 6, 2023

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: March 6, 2023

By: /s/ Stephen A. Trowbridge
Name: Stephen A. Trowbridge
Title: Executive Vice President and
Chief Financial Officer

Exhibit 99.1


angiodynamics
Raymond James
44th Annual Institutional Investors Conference
March 6, 2023
Jim Clemmer, President & CEO
Stephen Trowbridge, Executive Vice President & CFO

Notice Regarding Forward-Looking Statements

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In the United States, the NanoKnife System has received a 510(k) clearance by the Food and Drug Administration for use in the surgical ablation of soft tissue, and is similarly approved for commercialization in Canada, the European Union and Australia. The NanoKnife System has not been cleared for the treatment or therapy of a specific disease or condition.

Notice Regarding Non-GAAP Financial Measures

Management uses non-GAAP measures to establish operational goals and believes that non-GAAP measures may assist investors in analyzing the underlying trends in AngioDynamics' business over time. Investors should consider these non-GAAP measures in addition to, not as a substitute for or as superior to, financial reporting measures prepared in accordance with GAAP. In this presentation, AngioDynamics has reported adjusted EBITDA (income before interest, taxes, depreciation and amortization and stock-based compensation); adjusted net income and adjusted earnings per share. Management uses these measures in its internal analysis and review of operational performance. Management believes that these measures provide investors with useful information in comparing AngioDynamics' performance over different periods. By using these non-GAAP measures, management believes that investors get a better picture of the performance of AngioDynamics' underlying business. Management encourages investors to review AngioDynamics' financial results prepared in accordance with GAAP to understand AngioDynamics' performance taking into account all relevant factors, including those that may only occur from time to time but have a material impact on AngioDynamics' financial results. Please see the tables that follow for a reconciliation of non-GAAP measures to measures prepared in accordance with GAAP.

AngioDynamics

A medical technology platform company focused on a select group of large, high growth markets where meaningful treatment gaps exist in current standard of care. Our technologies positively impact treatment options and patients' quality of life.

AngioDynamics

Focused on making a positive patient impact in large markets

Cardiovascular disease and cancer have the highest morbidity and mortality worldwide

Strong Synergies[1]

These diseases are closely related with highly overlapping risk factors, disease development, and illness from the side effects of treatment.

Risk Factors

- Ageing
- Lifestyle
- Environmental factors
- Genetics
- Chronic conditions
- Access to healthcare

Global Cardiovascular Disease Burden[2]

523M diagnosed in 2020
~19 million deaths

Cardiovascular Disease causes 1 in 3 deaths globally

Cancer causes 1 in 6 deaths globally

19.3M diagnosed in 2020
~10 million deaths

Global Cancer Burden[3]

1. https://www.ncbi.nlm.nih.gov/pmc/articles/PMC5233740/#:~:text=After%20a%20first%20VTE%20recurrence,subsequent%20cancer%20diagnosis%20%5B10%5D.
2. https://professional.heart.org/-/media/PHD-Files-2/Science-News/2/2022-Heart-and-Stroke-Stat-Update/2022-Stat-Update-factsheet-Global-Burden-of-Disease.pdf https://www.cdc.gov/pcd/issues/2022/22_0347.htm
3. https://acsjournals.onlinelibrary.wiley.com/doi/10.3322/caac.21660#:~:text=Worldwide%2C%20an%20estimated%2019.3%20million,skin%20cancer)%20occurred%20in%202020.

AngioDynamics

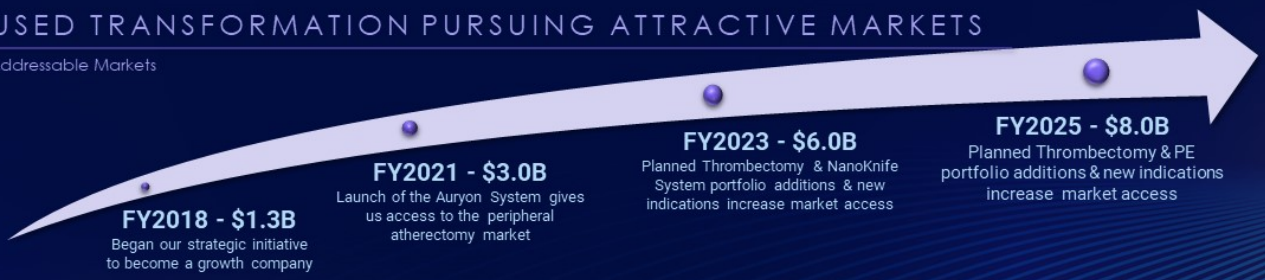
Investments in our Med Tech platforms are funded by operating cash flows from our Med Device portfolio

Med Tech: Invest for Growth
Peripheral Arterial Disease
Venous Thromboembolism
Cardiac Thrombus & Emboli
Solid Tumor

Med Device: Maintain Positioning	
Vascular Access Catheters & Accessories	Microwave & Radiofrequency Ablation
Diagnostic Catheters, Guidewires & Kits	Lung Biopsy Safety
Endovenous Laser Treatment	Radiation Treatment Stabilization Balloons

FOCUSED TRANSFORMATION PURSUING ATTRACTIVE MARKETS

U.S. Total Addressable Markets



FY2018 - $1.3B
Began our strategic initiative to become a growth company

FY2021 - $3.0B
Launch of the Auryon System gives us access to the peripheral atherectomy market

FY2023 - $6.0B
Planned Thrombectomy & NanoKnife System portfolio additions & new indications increase market access

FY2025 - $8.0B
Planned Thrombectomy & PE portfolio additions & new indications increase market access

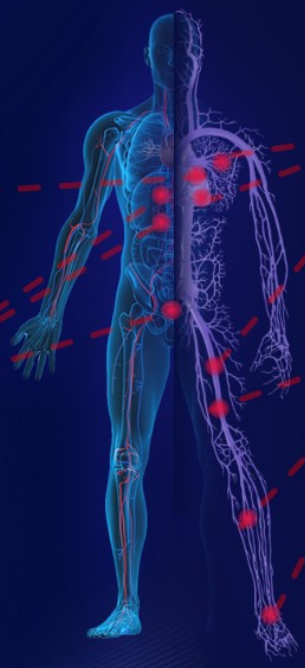
The planned portfolio additions and new indications are based on management estimates and industry sources as of July 2022 and are not guarantees of future performance and are subject to risks and uncertainties including FDA clearance. Investors are cautioned that actual events or results may differ from AngioDynamics' expectations.

Platform Technology Focused

On the global threat from cancer and cardiovascular disease

Cardiac
Thrombus & Emboli

Solid Tumors
Prostate, Pancreas, Liver



VTE
Venous Thromboembolism
DVT + PE = VTE

PAD
Peripheral Arterial Disease
Above the Knee, Below the Knee, In-Stent Restenosis

AngioDynamics

Focused technology platforms targeting attractive markets with meaningful treatment gaps, where our differentiated technologies can address unmet needs

Disease State	Platform	Treatment	Status
PAD Peripheral Arterial Disease	AURYON	Atherectomy	Launched
VTE Venous Thromboembolism	ANGIOVAC	Large Vessel Thrombectomy	Launched
	ALPHAVAC		Launched
	AURYON	Small Vessel Thrombectomy*	In development with targeted launch end of calendar 2024
	ALPHAVAC	Pulmonary Embolism*	APEX study currently enrolling Targeted launch early calendar 2025
Cardiac Thrombus & Emboli	ANGIOVAC	Right Heart	Launched
		Left Heart*	In development
	ALPHAVAC	Clot in Transit	Launched
Solid Tumor	NanoKnife	Prostate Tissue*	PRESERVE study >50% enrolled Launch targeted end of calendar 2024

*AlphaVac PE, Auryon Venous Thrombectomy/DVT, AngioVac Left Heart & NanoKnife Prostate are not cleared by the US Food and Drug Administration (FDA) for these indications.

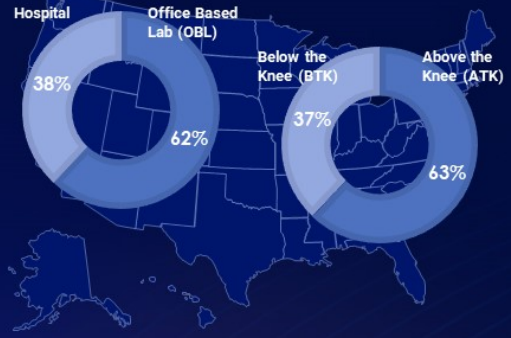
PAD

With over 25,000 cases performed, the Auryon Atherectomy System is the only atherectomy solution with the safety profile and versatility to treat every lesion location and morphology

THE MARKET

2022 Served

$760M



OUR SOLUTION

AURYON

- Peripheral Atherectomy

WHY IT MATTERS

Treat all levels of calcification [a-c]

- Indicated for in-stent restenosis*
- Treats above and below the knee (inc. below the ankle)

*2.0mm and 2.35mm catheters are indicated for ISR.

Protective of vessel wall [c-e]

- Targeted biological reactions to address risk of perforations
- Built-in aspiration to address risk of embolization†

†Built-in aspiration available with the 2.0-and 2.35-mm catheters.

Designed for hospital and lab [a-c, f]

- Portable, 110V outlet, low noise, touch screen
- Debulk in fewer passes

"We've always known that Auryon's technology is one-of-a-kind and unmatched. With the new [hydrophilic coating], we should be able to prove this – case after case after case"

– Dr. Curtis Anderson, Vascular & Interventional Radiologist

Source: Peripheral Vascular Devices Medtech 360 Market Analysis US December, 2021. Millennium Research Group, Inc.

a-f See reference page

VTE

Our differentiated technology platforms offer potential treatment solutions across the entire disease state

THE MARKET

Deep Vein Thrombosis **DVT** + Pulmonary Embolism **PE** = Venous Thromboembolism **VTE**

A blood clot that forms in a deep vein, usually the leg, groin or arm

A DVT breaks free and travels to the lungs blocking some or all of the blood supply

DVT and PE are collectively referred to as VTE

2022 TAM

$3.9B

OUR SOLUTION

ANGIOVAC

- Large Vessel Venous Thrombectomy/DVT

ALPHAVAC

- Large Vessel Venous Thrombectomy/DVT
- Pulmonary Embolism*

AURYON

- Small Vessel Venous Thrombectomy/DVT*

WHY IT MATTERS

- Only solution on the market with continuous aspiration and simultaneous reinfusion of filtered blood
- Aspirates large clot burden

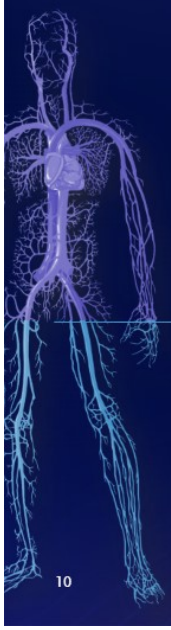
- Controlled aspiration
- Aspirates large clot burden
- APEX-AV study for PE

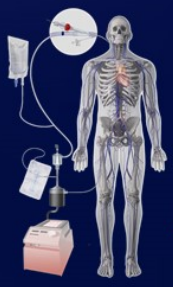
- Auryon's combination of laser technology and aspiration restores flow in occluded vessels

Source: Management estimate & industry sources as of July 2022.
*AlphaVac PE and Auryon Venous Thrombectomy/DVT are not cleared by the US FDA for these indications.

VTE

All-purpose technology platforms targeted at peripheral and cardiovascular thrombolytic events, including small and large vessels





Large Vessel

Small Vessel

RADIOPAQUE MARKERS
Better Tip Visibility

LARGE END HOLE ASPIRATION
42FR & 30FR Opening

PROPRIETARY FUNNEL DESIGN
Allows for Significant Clot Removal

MULTIPLE TIP ANGLES
20^0, 85^0, 180^0

ANGIOVAC

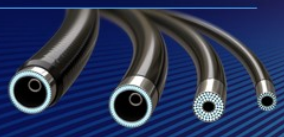
The **AngioVac** System allows for the **continuous aspiration** of embolic material such as thrombi and emboli from the venous system while **Simultaneously reinfusing** the patient's own filtered blood to limit procedural blood loss

ALPHAVAC

The **AlphaVac** System allows for the **controlled aspiration** of embolic material such as thrombi and emboli from the venous system

AURYON

POWERFUL
355 nm laser is designed to deliver an optimized wavelength, pulse width, and amplitude to restore flow in occluded vessels[c, d, g]



PRECISE
Protective of vessel wall[c-e]

ADAPTABLE
Potential to treat all types of small vessel DVT*

c-g See reference page
*Auryon Venous Thrombectomy/DVT is not cleared by the US FDA for this indication.

Cardiac Thrombus & Emboli

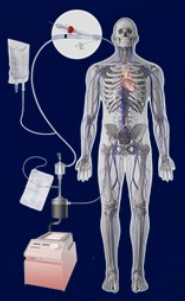
We are focused on offering percutaneous solutions for removing thrombus and emboli in the left and right heart

THE MARKET

2022 TAM

$530M

OUR SOLUTION



ANGIOVAC

- Right Heart
- Left Heart*

WHY IT MATTERS

- Continuous aspiration combined with the funnel tip, allows for the efficient removal of the targeted material while minimizing risk of blood loss
- Currently, there is no standard for right or left heart percutaneous approach

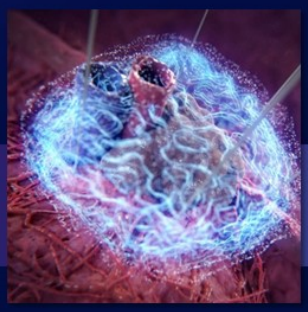
Source: Management estimate & industry sources as of July 2022.
*AngioVac Left Heart is not cleared by the US FDA for this indication.

NanoKnife Technology

Expands treatment options and help preserve patient's quality of life

PROBE PLACEMENT

NanoKnife can be confidently used in all segments of an organ.[1,2]



DECELLULARIZATION

Destroys targeted tissue with precise treatment margins.[1,2]

NON-THERMAL

Spares vital structures by retaining the structural integrity of tissue.[3,4]

REVASCULARIZATION

Facilitates functional tissue regeneration post-ablation.[3,4]

1. Lee EW, Thai S, Kee ST. Irreversible electroporation: a novel image-guided cancer therapy. Gut Liver. (2010);4(SUPPL. 1):99–104. doi: 10.5009/gnl.2010.4.S1.S99
2. Guidance for Selection of NanoKnife Probe Array Configuration and Ablation parameters for the Treatment of Stage III Pancreatic Cancer
3. Scheltema MJ, Chang JI, van den Bos W, Gielchinsky I, Nguyen TV, Reijke TM, Shivardana AR, Böhm M, de la Rosette JJ, Stricker PD. Impact on genitourinary function and quality of life following focal irreversible electroporation of different prostate segments. Diagn Interv Radiol. 2018 Sep;24(5):268-275. doi: 10.5152/dir.2018.17374. PMID: 30211680; PMCID: PMC6135060.
4. Li W, Fan Q, Ji Z, Qiu X, Li Z. The effects of irreversible electroporation (IRE) on nerves. PLoS One. 2011 Apr 14;6(4):e18831. doi: 10.1371/journal.pone.0018831. PMID: 21533143; PMCID: PMC3077412

Prostate Initiative*

Over 100,000 men with intermediate risk prostate cancer could be treated with this technology

THE MARKET

2022 TAM

$700M

OUR SOLUTION

NanoKnife

- Focal Therapy

WHY IT MATTERS

Targeted: Short electric pulses destroy cells without relying on extreme heat or cold and spare vital structures within the ablation zone

Quality of Life: Better preserves urinary control and erectile function

Versatile: Can be used in all segments of the prostate for primary and recurrent disease

Fast: Minimally invasive treatment that is delivered in a single session

Preserves future treatment options

*IDE Study in progress
Market Source: Management estimate & industry sources as of July 2022.

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International Expansion Plan

Expanding our business reach in targeted regions & countries

Aligning our Go-to-Market strategy to the different regions and markets, utilizing new partnerships where appropriate to maximize growth

Preparing for EU and selected OUS launches of both the Auryon Atherectomy Product line, and the AlphaVac large bore Thrombectomy product Line

- Targeted launch date Auryon: 1H of calendar 2024
- Targeted launch date AlphaVac: 1H of calendar 2024

Continue to increase our global presence through our series of life symposiums which has attracted interest from global key opinion leaders who are gaining more access of our technologies

Medical Device

Med Device: Maintain Positioning

Vascular Access Catheters & Accessories	Microwave & Radiofrequency Ablation
Diagnostic Catheters, Guidewires & Kits	Lung Biopsy Safety
Endovenous Laser Treatment	Radiation Treatment Stabilization Balloons

PORTFOLIO

- Optimizing our commercial approach by re-aligning Core portfolio into new VA - Device centric commercial team

MARKET ACCESS

- Broader Med Device bag allows deeper customer engagement
- Maximize clinical differentiation & secure committed customers through targeted GPO/IDN contracting

PERFORMANCE

- Maintain a strong culture of execution and collaboration through disciplined sales & marketing plans
- Develop & export key talent throughout the organization

3 Year Transformational Plan

FY22	
Revenue	$305M – $310M
Growth	*5% - 7%*
Gross Margin	~55%
Adjusted EPS	$0.00 - $0.05

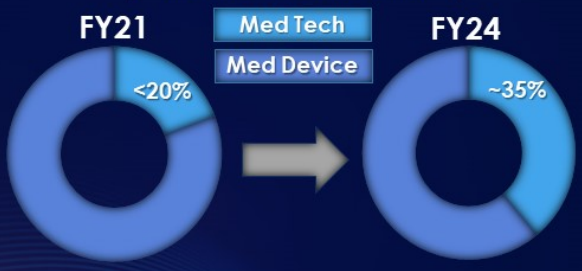
Actual $316M 9%

FY23	
Revenue	$330M – $336M
Growth	*7% - 9%*

FY24	
Revenue	$360M – $375M
Growth	*10% - 12%*

- Planned significant investment in Med Tech platforms drives top line growth
- Bottom line leverage will ramp slower than top line growth

Revenue Contribution



Revenue Growth CAGR

The projections and growth rates depicted on this slide are forward-looking statements. These forward-looking statements are not guarantees of future performance and subject to risks and uncertainties.

angiodynamics

Raymond James

44th Annual Institutional Investors Conference

March 6, 2023

Jim Clemmer, President & CEO

Stephen Trowbridge, Executive Vice President & CFO

Auryon References

a. Rundback J, Chandra P, Brodmann M, Weinstock B, Sedillo G, Cawich I, et al. Novel laser-based catheter for peripheral atherectomy: 6-month results from the Eximo Medical B-Laser™ IDE study. Catheter Cardiovasc Interv. 2019;1-8.

b. Shammas NW, Chandra P, Brodmann M, Weinstock B, Sedillo G, Cawich I, et al. Acute and 30-day safety and effectiveness evaluation of Eximo Medical's B-Laser™, a novel atherectomy device, in subjects affected with infrainguinal peripheral arterial disease: Results of the EX-PAD-03 trial. Cardiovas Revasc Med. 2020;21(1):86-92.

c. Auryon. Instructions for use. AngioDynamics; 2019.

d. Herzog A, Bogdan S, Glikson M, Ishaaya AA, Love C. Selective tissue ablation using laser radiation at 355 nm in lead extraction by a hybrid catheter; a preliminary report. Lasers Surg Med. 2016;48(3):281-287.

e. Herzog A, Steinberg I, Gaisenberg E, Nomberg R, Ishaaya AA. A route to laser angioplasty in the presence of fluoroscopy contrast media, using a nanosecond-pulsed 355-nm laser. IEEE J Sel Top Quantum Electron. 2016;22(3):342-347.

f. Kuczmik W, Kruszyna L, Stanisic MG, Dzieciuchowicz L, Ziaja K, Zelawski W, et al. Laser atherectomy using the novel B-Laser™ catheter, for the treatment of femoropopliteal lesions: twelve-month results from the EX-PAD-01 study. Not yet published.

g. Vogel A, Venugopalan V. Mechanisms of pulsed laser ablation of biological tissues. Chem Rev. 2003;103(2):577-644.