

Inari Medical Investor Update

May 3, 2023

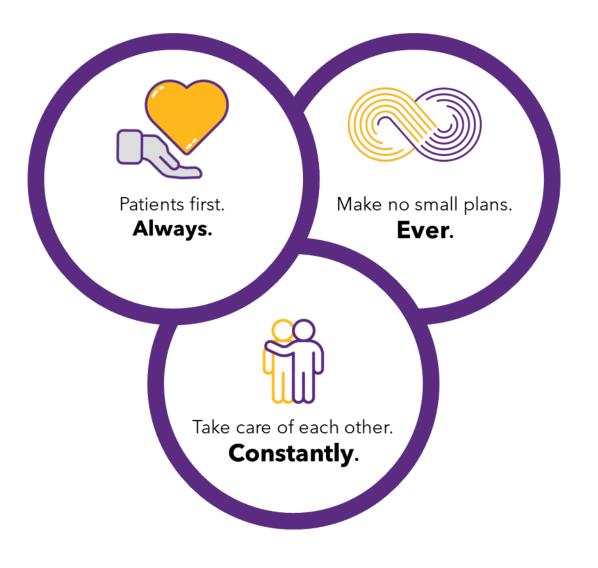
Larry | Newport Beach, CA FlowTriever Patient This presentation (together with any other statements or information that we may make in connection therewith) may contain are forward-looking statements. All statements other than statements of historical fact could be deemed forward-looking, including any estimates of fourth quarter revenue and total procedures, the potential impact of COVID-19 on the business, total addressable market, future results of operations, financial position, research and development costs, capital requirements and our needs for additional financing; our business model and strategic plans for our products, technologies and business, including our implementation thereof; competitive companies and technologies and our industry; the impact on our business, financial condition and results of operation from the ongoing and global COVID-19 pandemic, or any other pandemic, epidemic or outbreak of an infectious disease in the United States or worldwide; our ability to commercialize, manage and grow our business by expanding our sales and marketing organization and increasing our sales to existing and new customers; third-party payor reimbursement and coverage decisions; commercial success and market acceptance of our products; our ability to accurately forecast customer demand for our products and manage our inventory; our ability to establish and maintain intellectual property protection for our products or avoid claims of infringement; FDA or other U.S. or foreign regulatory actions affecting us or the healthcare industry generally, including healthcare reform measures in the United States; the timing or likelihood of regulatory filings and approvals; our ability to hire and retain key personnel; our ability to obtain additional financing; and our expectations about market trends. Without limiting the foregoing, the words "may," "will," "should," "expect," "plan," "anticipate," "could," "intend," "target," "project," "contemplate," "believe," "estimate," "predict," "potential" or "continue" or the negative of these terms and other similar

Forward-looking statements are based on and reflect management's current expectations, assumptions, estimates and projections that may or may not prove to be correct. These forward-looking statements are subject to a number of known and unknown risks, uncertainties, assumptions and other factors, many of which are beyond our control. Moreover, we operate in a very competitive and rapidly changing environment. New risks emerge from time to time. It is not possible for our management to predict all risks, nor can we assess the impact of all factors on our business or the extent to which any factor, or combination of factors, may cause actual results to differ materially from those contained in any forward-looking statement. In light of these risks, uncertainties, and assumptions, the future events and trends discussed in this presentation may not occur and our actual results, levels of activity, performance or achievements could differ materially and adversely from those anticipated or implied by any forward-looking statements. These and other known risks, uncertainties and factors are described in detail under the caption "Risk Factors" and elsewhere in our filings with the Securities and Exchange Commission ("SEC"), including our most recent Annual Report on Form 10-K and Quarterly Report on Form 10-Q. These filings are available in the Investor Relations section of our website at https://ir.inarimedical.com/ or at www.sec.gov.

The forward-looking statements in this presentation are made only as of the date hereof. Except to the extent required by law, we assume no obligation and do not intend to update any of these forward-looking statements after the date of this presentation or to conform these statements to actual results or revised expectations. All forward-looking statements are expressly qualified in their entirety by the foregoing cautionary statements. You are cautioned not to place undue reliance on these forward-looking statements.

This presentation is not an offer to sell securities of Inari Medical and it is not soliciting offers to buy securities of Inari Medical nor will there be any sales of securities of Inari Medical in any state or jurisdiction where the offer, solicitation or sale would be unlawful prior to registration or qualification under the securities laws of any such state or jurisdiction.





We've made improving lives our responsibility. And that drives our passion and success

A mission, a plan, and crisp execution producing sustained growth



- **Purpose Built, Highly Differentiated Solutions** designed to solve specific problems
- **BIG, Growing, and Efficient Commercial Team** of over 280 territories
- Large Markets, Lot of Runway (\$5.8B in US VTE alone, <6% penetrated)*
- Leading, Differentiated Data (6 major studies, including 2 ongoing RCTs)**
- **Robust Product Pipeline** (10 products launched 2021-2022)***
- **Efficient Procedures, Favorable Economics,** driven by limited hospital resources, avoiding ICU stay, and reducing total length of stay

** FLARE, FLASH, FLAME, CLOUT, PEERLESS. DEFIANCE

[•] Based on third party data and Inari management estimates

^{***} Products launched: Triever20 Curve catheter, FlowTriever2 catheter, FlowStasis, FlowSaver, Triever24 Flex catheter; Products launched 2022: ClotTriever BOLD catheter, Intri24 sheath, Protrieve sheath, InThrill system, Triever Gen 4 catheters

Strong leadership team to capitalize on our opportunity





Drew Hykes Chief Executive Officer





Tom Tu, M.D. Chief Medical Officer

Angela Ahmad

General Counsel

Brian Strauss

SVP Engineering

John Borrell SVP Sales

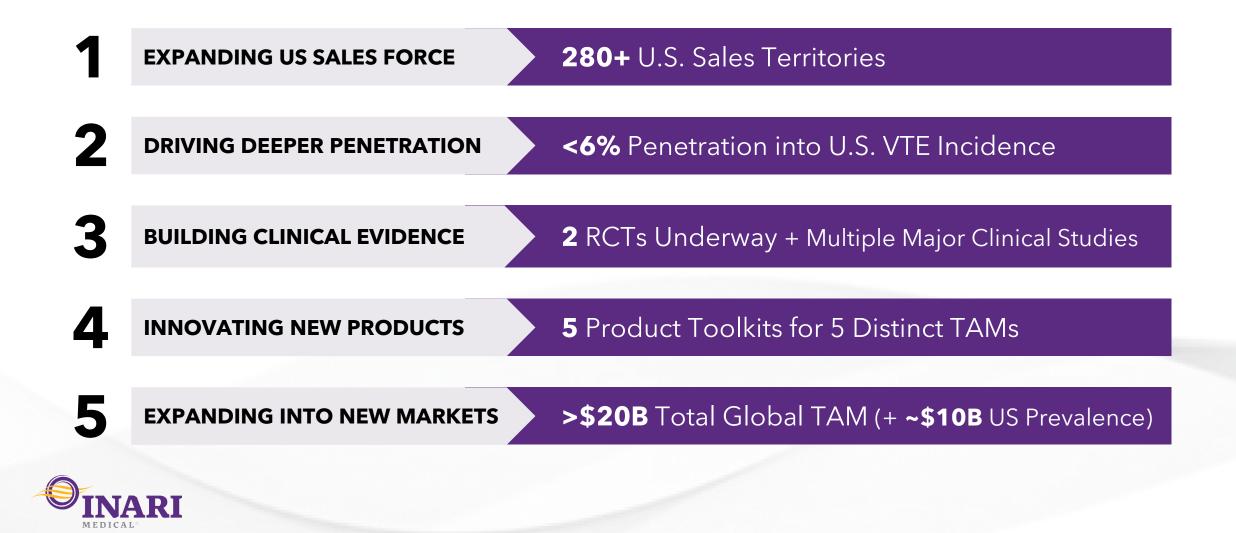
Eric Khairy SVP Marketing &

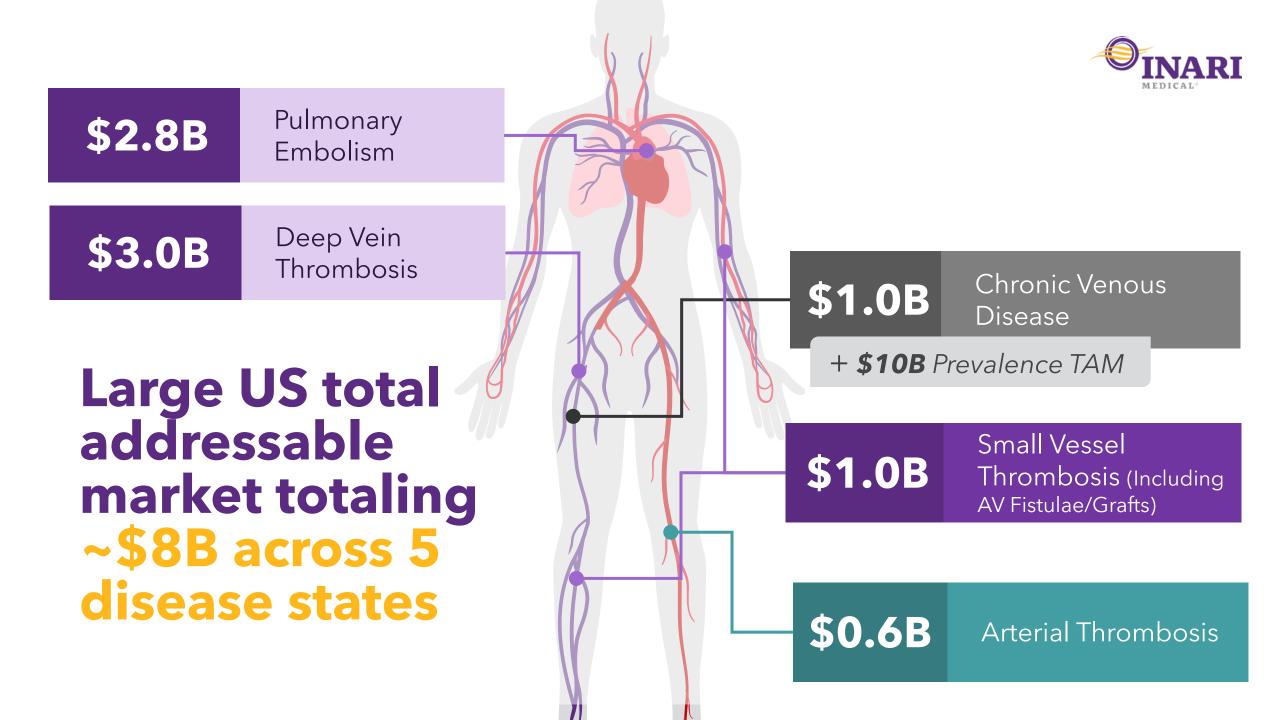
Commercial Operations

Paul Koehn SVP Operations

Tara Dunn SVP Clinical Affairs & Market Development

Our five growth drivers remain the roadmap







Venous Thromboembolism

Pulmonary Embolism and Deep Vein Thrombosis

Audry | Detroit, MI FlowTriever Patient

Venous thrombus requires purpose-built solutions



	Arterial System	Venous System	
Hemodynamics:	High flow, high pressure	Low flow, low pressure	
Vessel morphology:	Small vessels that taper in direction of flow	Large vessels that enlarge in direction of flow	
Clot morphology:	Small amounts of soft clot, "floating" in the vessel	Large amounts of firm/hard clot, adhered to vessel wall	

Only purposebuilt solutions can address the challenges of venous thrombus

Repurposing Arterial Thrombectomy Systems for venous clot results in inadequate safety, performance, and the need for thrombolytics



Inadequate thrombectomy options lead to use of thrombolytics, an ineffective option for venous clot

For Venous Clots, Thrombolytics are Generally:

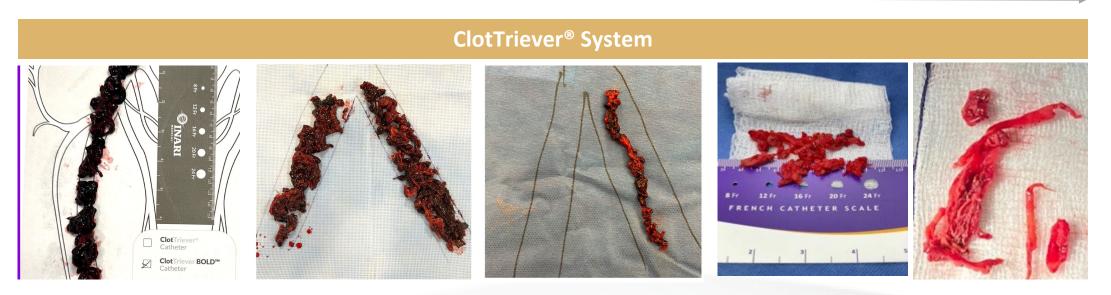
Ineffective	 Symptoms often appear gradually, and venous clot can become large/hardened Clot morphology changes over time The older the clot, the fewer "targets" of thrombolytics remain
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High Risk	 Can carry significant rates of bleeding complications Conservative patient selection and low dosage do not always eliminate bleeding risks Up to 50% of patients with VTE are relatively or absolutely contraindicated
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 Can be highly costly Requires multiple procedures and prolonged hospital stays Bleeding risks necessitate ICU stay (the most expensive bed in the Reimbursement is relegated to low-paying, medically-orientated I 	1 •
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Most venous clot does not respond to thrombolytics

Acute



FlowTriever® System

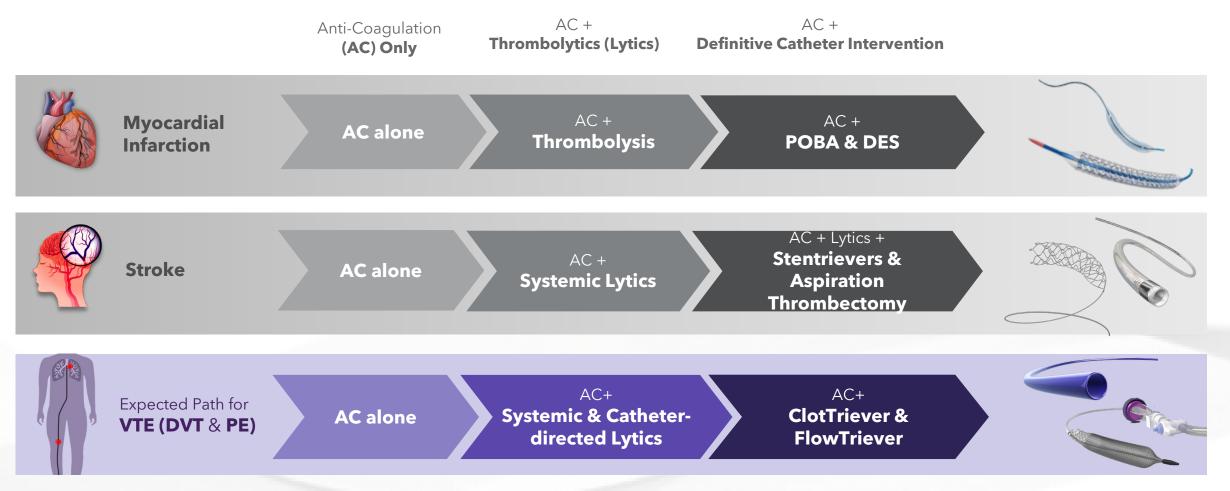




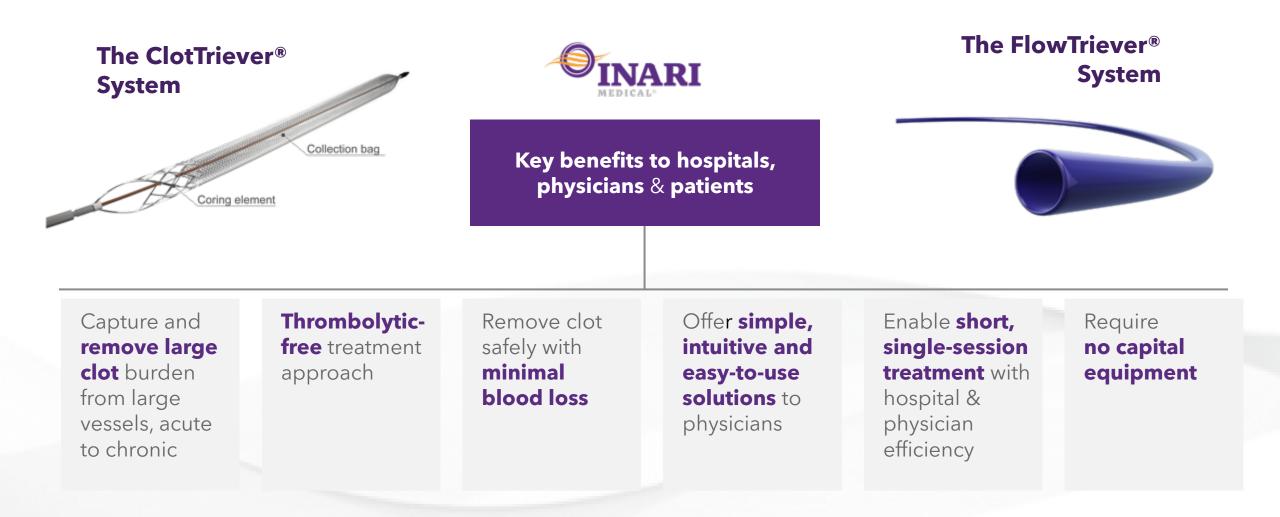
Chronic



Treatment of thrombotic diseases consistently evolves to definitive mechanical catheter intervention



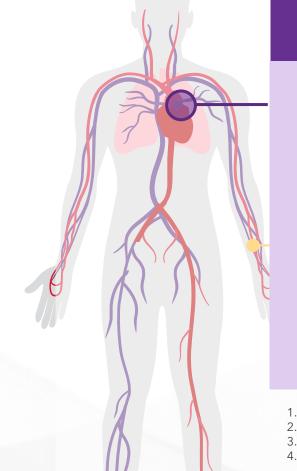
Our solutions are designed to offer significant benefits to patients, physicians, and hospitals



Pulmonary Embolism (PE)



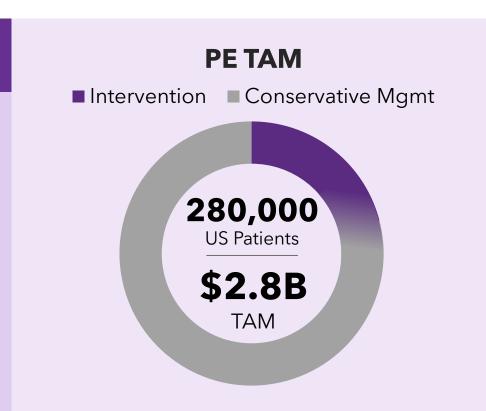
Transforming the lives of patients suffering from PE



PULMONARY EMBOLISM (PE)

3rd leading cause of cardiovascular death¹

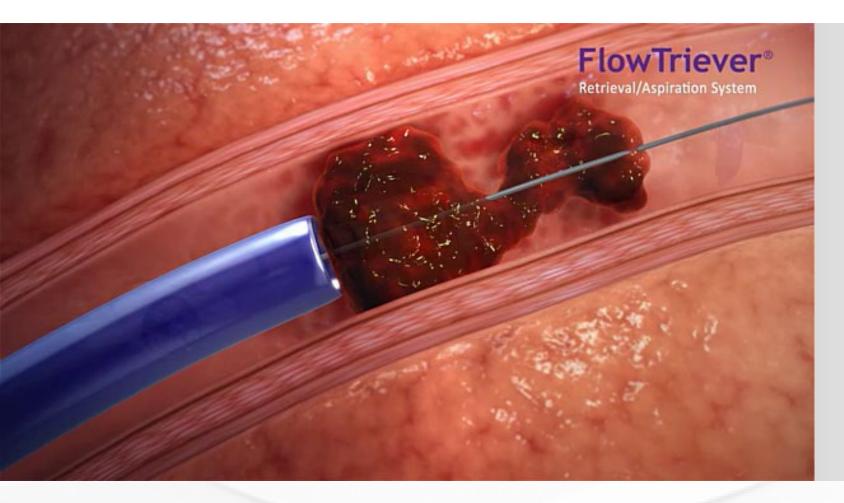
- A/C alone leaves clot behind in up to half of patients^{2,3}
- Long-term complications are common⁴



"Pulmonary Embolism in 2017: Increasing Options for Increasing Incidence", National Center for Biotechnology Information, May 2017.
 Picart, et al. Predictors of residual pulmonary vascular obstruction after pulmonary embolism: Results from a prospective cohort study. Thrombosis Research. 2020.
 Dzikowska-Diduch, et al. The post-pulmonary syndrome - results of echocardiographic driven follow up after acute pulmonary embolism. Thrombosis Research. 2020.
 Sista AK, et al. Vasc Med. 2017 Feb;22(1):37-43

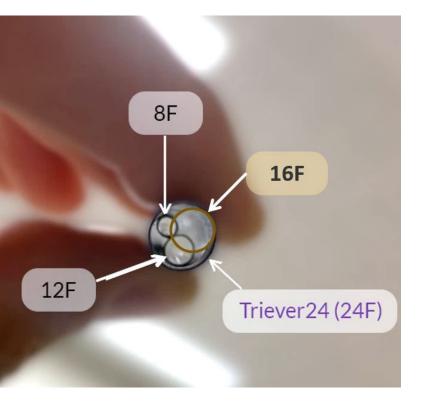
FlowTriever: Large bore catheters for large clot hauls

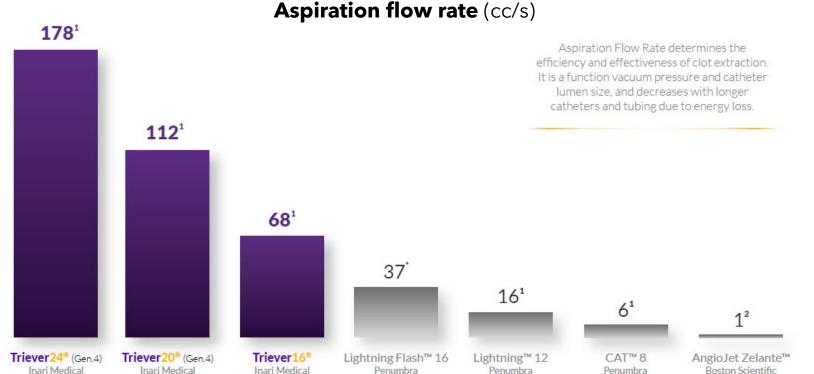




- Designed to extract large volumes of clot
- Blood can be returned with FlowSaver[®]
- Single session
- Lytic-free approach
- Avoid lytic-based ICU stay
- Rapid symptom relief

FlowTriever: Large bore catheters for large clot hauls





*Calculated using Poiseuille's Law

References:

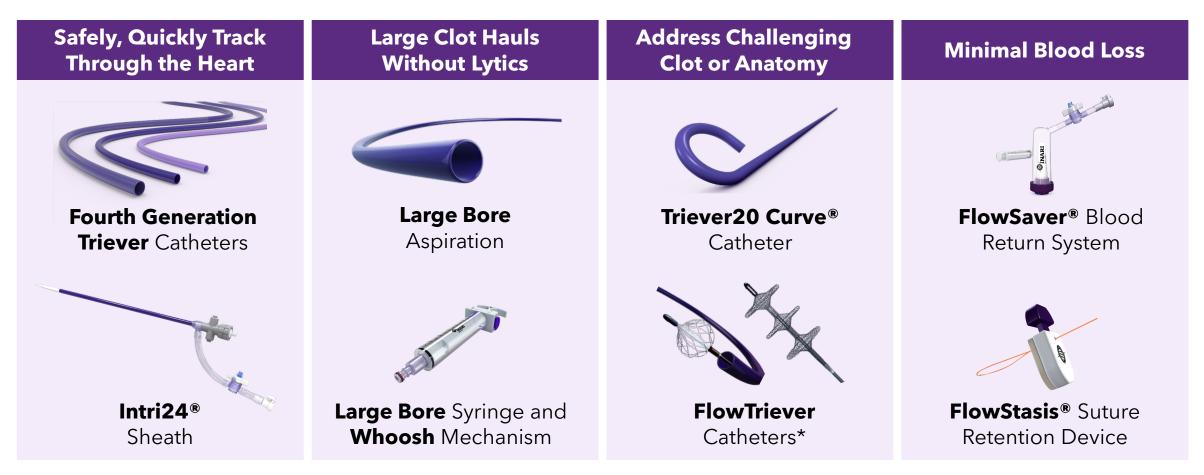
1. Experimental data on file.

2. AngioJet product brochure. https://www.bostonscientific.com/content/dam/bostonscientific/pi/portfolio-group/ham-portal-emea/resources/PI-750306-

AA%20EMEA%20AngioJet%20Interactive%20Brochure-%20FINAL.pdf

The FlowTriever® System: A full toolkit approach to PE





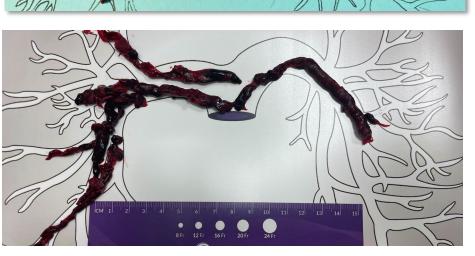
*The FlowTriever 2 catheter is not indicated for the treatment of PE

INDICATIONS FOR USE: **The FlowTriever Retrieval/Aspiration System** is indicated for: (1) The non-surgical removal of emboli and thrombi from blood vessels, and (2) The injection, infusion, and/or aspiration of contrast media and other fluids into or from a blood vessel. The FlowTriever Retrieval/Aspiration System is intended for use in the right atrium, but not in conjunction with FlowTriever Catheters. The FlowTriever Catheter is indicated for: the non-surgical removal of emboli and thrombi from blood vessels. The right atrium, but not in conjunction with FlowTriever Catheters. The FlowTriever Catheter is indicated for: the non-surgical removal of emboli and thrombi from blood vessels. The right atrium, but not in conjunction with FlowTriever Catheter is indicated for: the non-surgical removal of emboli and thrombi from peripheral blood vessels. Injection, infusion, and/or aspiration of contrast media and other fluids into or from a blood vessel. The FlowTriever Catheter is intended for use in the peripheral vasculature. The FlowStasis device is intended for temporary suture retention following a percutaneous venous procedure The FlowSaver Blood Return System is used with Triever Catheters for autologous blood transfusion

FlowTriever removes significant clot burden



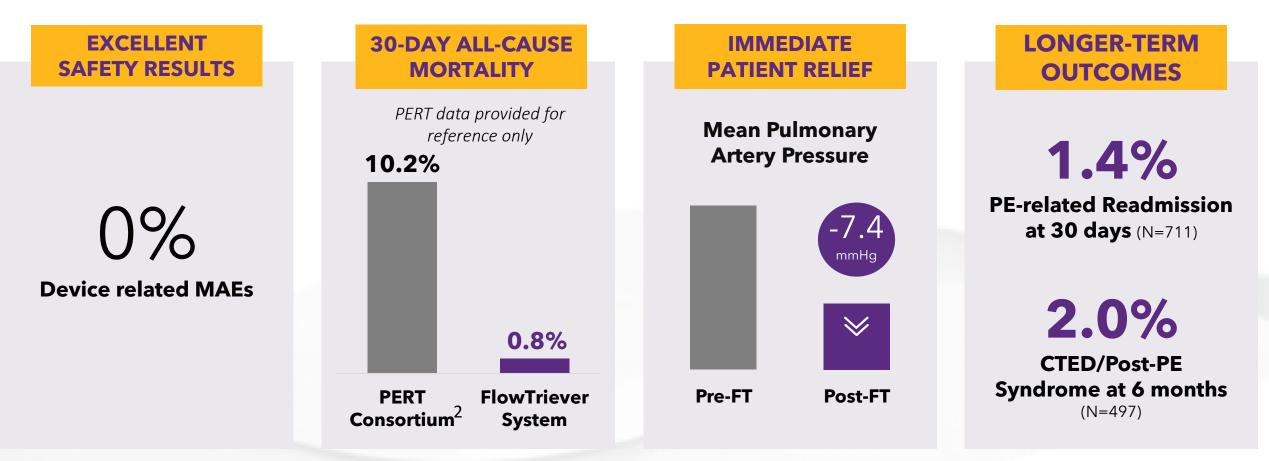




FLASH is the largest prospective registry in PE with exceptional results¹



800 patients, 50 sites, 32% were contraindicated to lytics



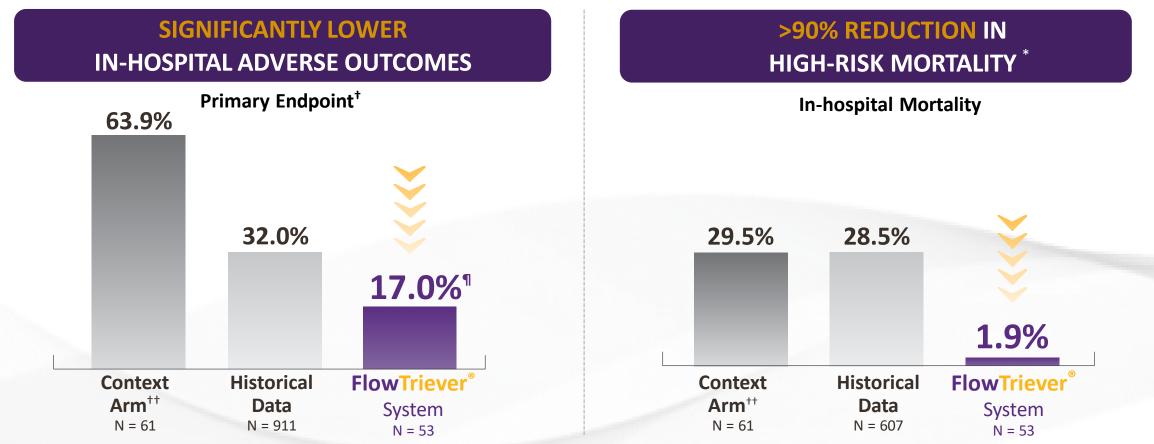
1. "FLASH data Presented by Dr. Catalin Toma, TCT 2022

2. PERT Consortium Quality Database. October 2021 (Presented by Secemsky E); Darki A & Jaber WA. Endovascular Today. July 2022 Supplement (PERT Updates)

FLAME high-risk PE study shows very low rates of adverse events and mortality



High-risk PE. Non-randomized study with FlowTriever (N=53), Historical Reference, and Context Arm (N=61)



*>90% reduction in high-risk PE in-hospital all-cause mortality vs. other contemporary treatments in historical data

[†]Composite primary endpoint consisted of in-hospital all-cause mortality, bailout to an alternate thrombus removal strategy, clinical deterioration, and major bleeding

^{+†}Context arm patients were treated with systemic thrombolysis (68.9%), anticoagulation alone (23.0%), CDT (6.6%) or surgical thrombectomy (1.6%)

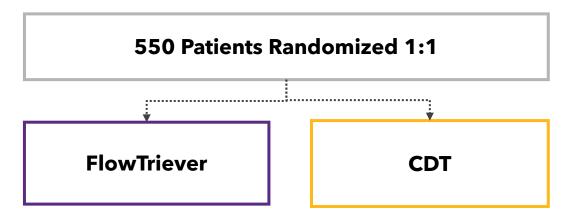
 $\P<\!0.01$ vs. performance goal based on historical data

Source: Outcomes In High-risk Pulmonary Embolism Patients Undergoing FlowTriever Mechanical Thrombectomy: Results From The FLAME Study presented at ACC March 2023 by Dr. Mitchell J. Silver



Superiority RCT of FlowTriever vs CDT in PE

Intermediate Risk PE



Patients Followed for 30 Days

HIGHLIGHTS

Currently, Catheter Directed Thrombolysis (CDT) is used in nearly half of interventions commercially*



Primary endpoint via win ratio:

- All-Cause Mortality
- Intracranial Hemorrhage
- ISTH Major Bleeding
- Clinical Deterioration/Bailout
- ICU Admission & ICU LOS



Enrollment ahead of schedule



Designed to transform standard of care away from CDT

*Based on third party data and Inari management estimates.

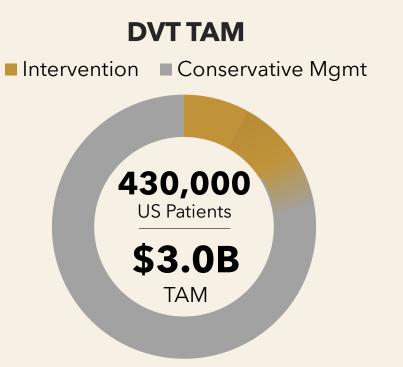
Deep Vein Thrombosis (DVT)

Transforming the lives of patients suffering from DVT



DEEP VEIN THROMBOSIS (DVT)

- Up to 50% develop Post-Thrombotic Syndrome (PTS)¹
- A/C alone leaves clot behind in up to half of patients²
- Lytics don't address chronic clot, and come with bleeding risk

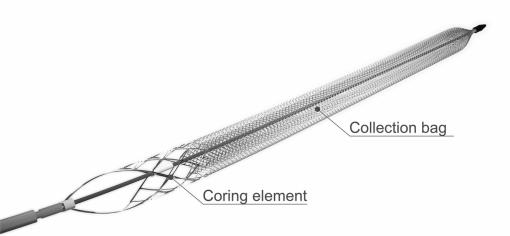


1. Kahn, Susan R. Hematology Am Soc Hematol Educ Program. 2016 Dec 2; 2016(1): 413-418

2. Young et al., Post-treatment residual thrombus increases the risk of recurrent deep vein thrombosis and mortality. J Thromb Haemost 2006; 4: 1919–24.

ClotTriever is effective on clot of all ages

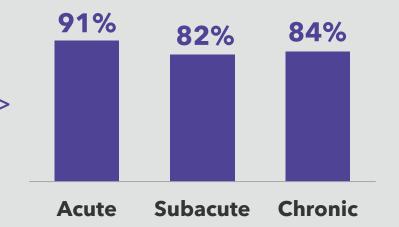




Effective on Clot of all Ages

% limbs with complete or near complete (≥75%) thrombus removal

(as assessed by Marder Score)¹





ClotTriever removes significant clot burden





20F sheath designed for right IJ access and **optimal positioning within the IVC**

Wall apposing funnel designed to trap emboli



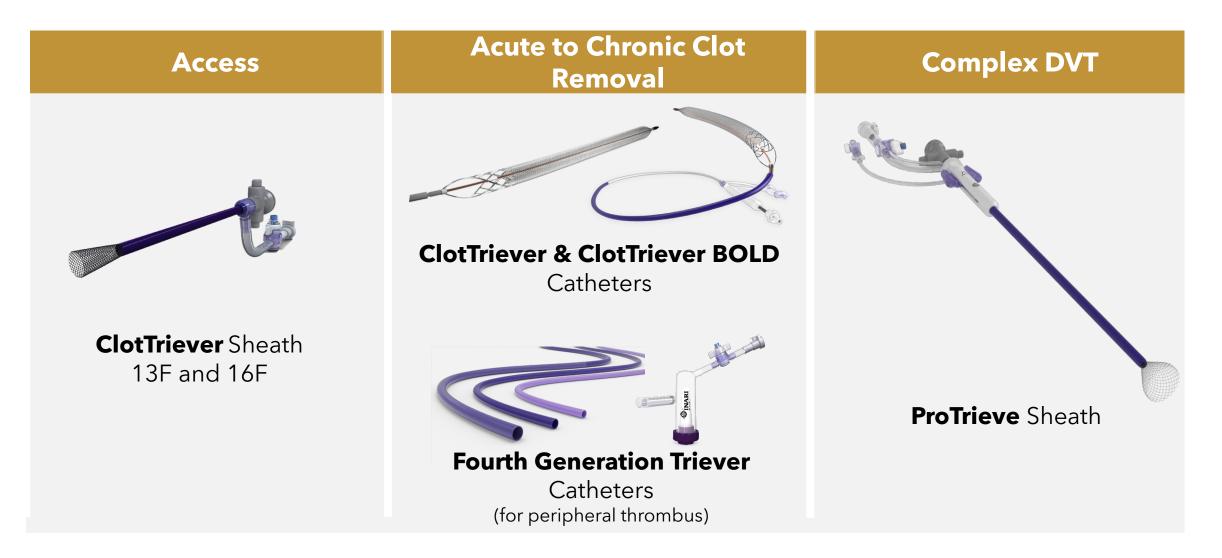
Compatible with **ClotTriever** and **FlowTriever** platforms



Can be used as a conduit for **compatible filter** retrieval sheaths

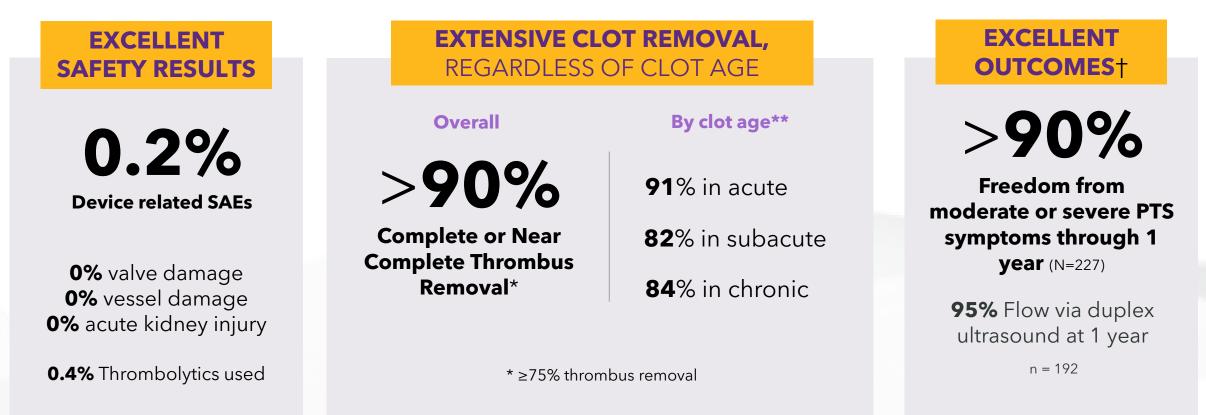
A comprehensive solution for DVT and peripheral thrombus





CLOUT is the largest mechanical thrombectomy dataset in DVT with exceptional results

500 patients | 47 sites | 70% subacute and/or chronic clot | 30% lytics contraindicated



500-patient results presented by Dr. David Dexter, VEINS 2022

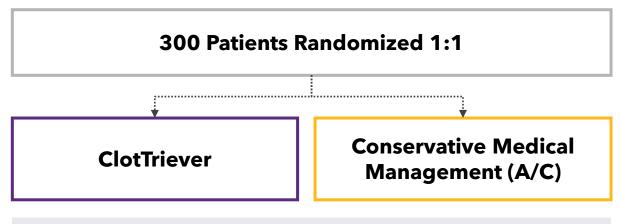
**Subset of 250 patients presented at AVF 2022

[†] One-year interim outcomes from the multicenter prospective CLOUT registry presented by Dr. David Dexter AVF 2023



Superiority RCT of ClotTriever vs Anticoagulation in DVT

Moderate-Severe Iliofemoral DVT



Patients Followed for 6 Months

HIGHLIGHTS



First global industrysponsored RCT for DVT



Primary endpoint via win ratio:

- Treatment failure or escalation of therapy
- Post-Thrombotic Syndrome severity at 6 months



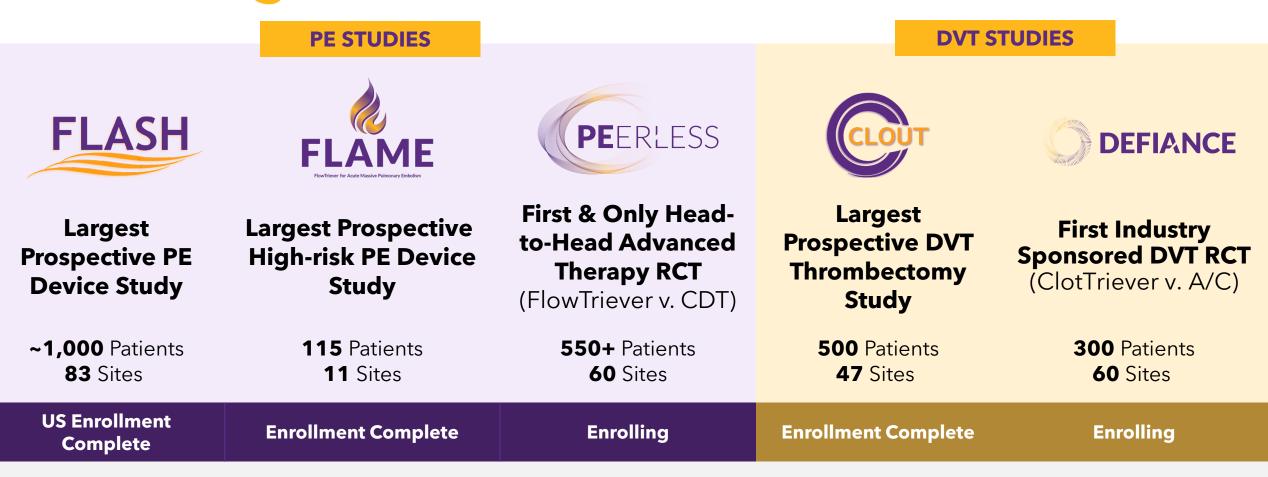
Enrollment started in January 2023



Designed to transform standard of care

Setting a high bar for VTE evidence to change standard of care





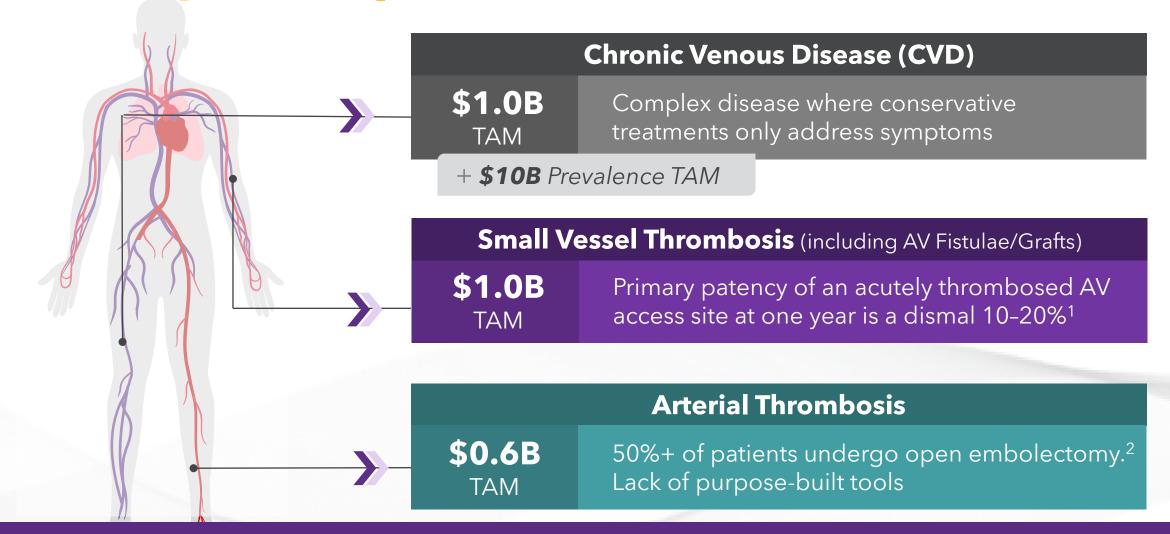
~2,500 patients across 5 studies

New Markets



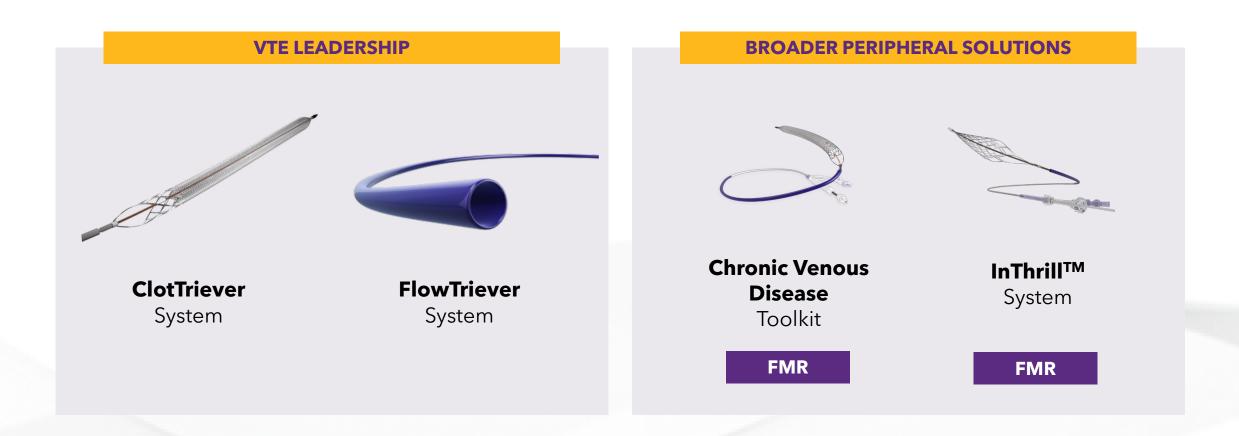
Continuing our mission to help more patients in need





1. Quencer KB, Oklu R. Hemodialysis access thrombosis. Cardiovasc Diagn Ther. 2017 Dec;7(Suppl 3):S299-S308. 2. Based on third party data and Inari management estimates.

Expanding beyond VTE to develop





ClotTriever BOLD was designed to **extract the full range of clot chronicity**



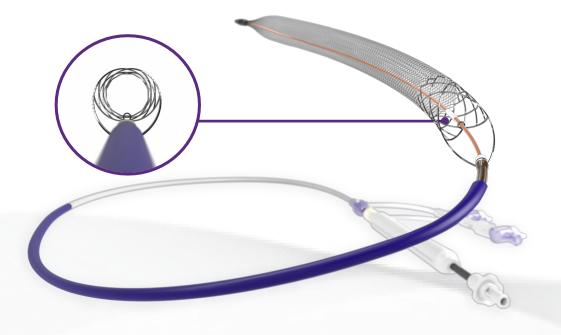
Clot is often older than symptoms suggest



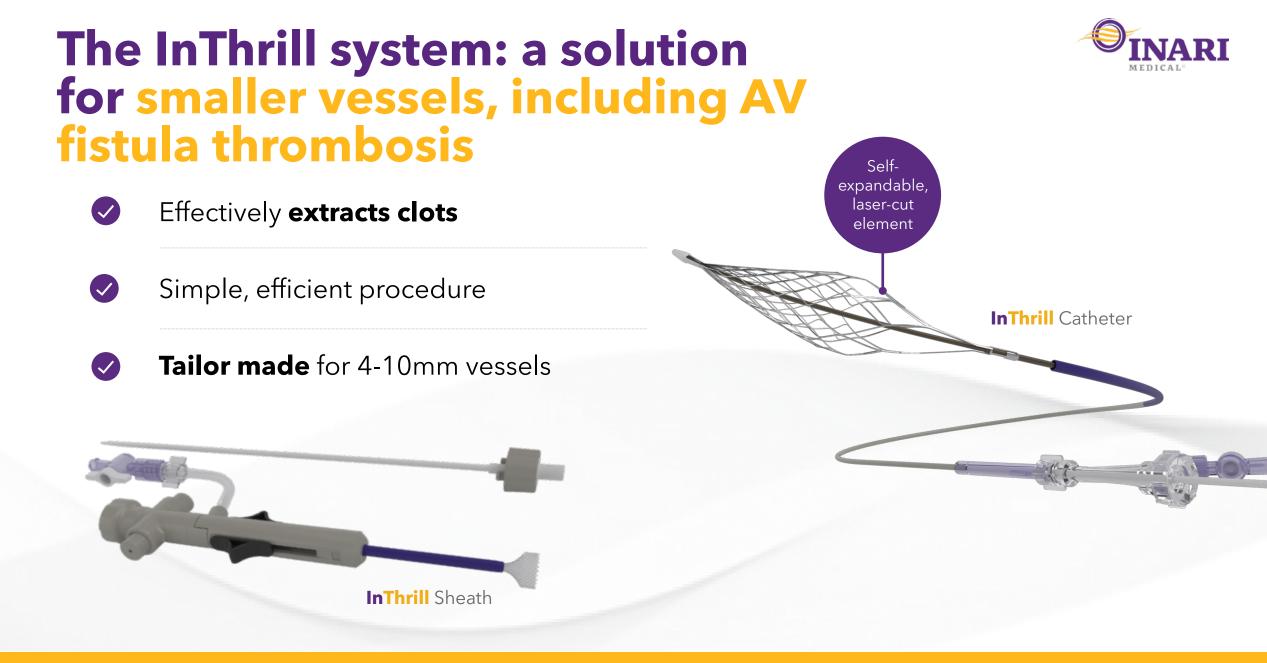
~30% greater radial force for **improved** wall apposition



Improved thrombus engagement to treat the full range of acute to chronic clot







Note: the InThrill device is indicated for use in the peripheral vasculature, including AV fistulae and grafts

US and Global Commercial Execution



Leveraging high-touch commercial system to solve patient needs

Single-tier sales team w/ ~90% case presence

Mining information across all sources, informing every decision we make

Solution-based toolkits, not widgets

Scalable across new markets with significant unmet needs

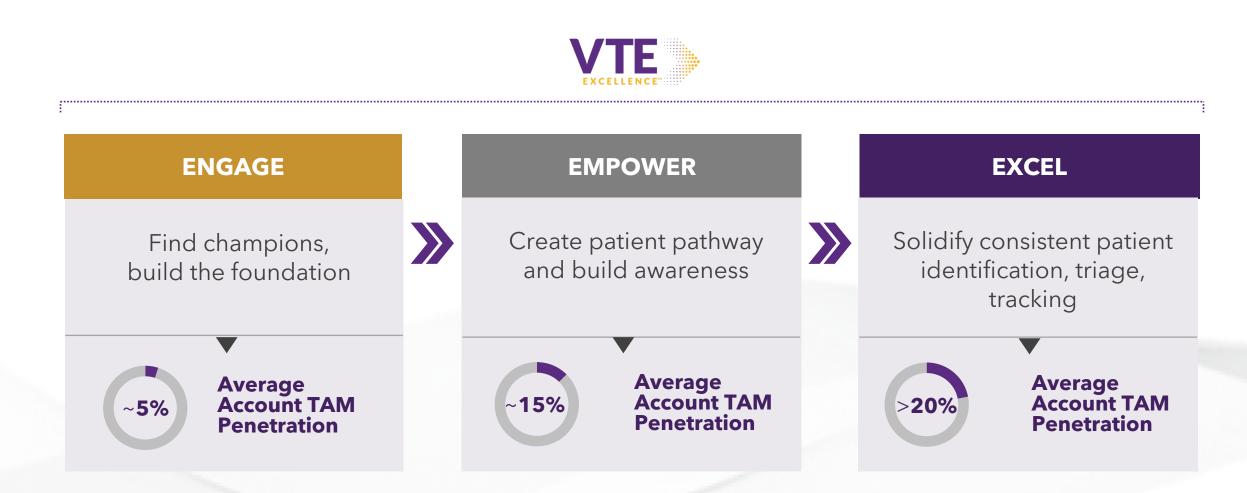
Supported by robust commercial market development team



US Sales Territories in 2022

280 +



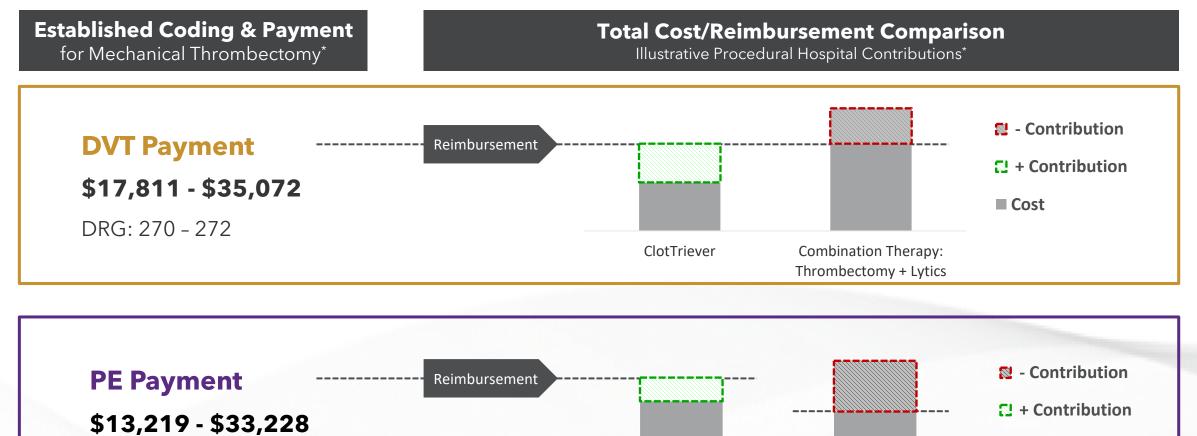




Cost

Ultrasonic Catheter Directed Lytics

Our products offer benefits and value to our hospital and physician customers



FlowTriever

DRG: 163 - 165

* Utilizes national average Medicare reimbursement rates for CY2023 IPPS and Inari management estimates around patients with and without MCC and CC.

Limited hospital resource use, excellent clinical and economic outcomes

Patients, physicians and hospitals all benefit from Inari products



Effective, short, single-session treatments with no capital equipment



Thrombolytic-free treatment approach



Avoid lytic-based ICU stay



Established **procedural** reimbursement



Short total hospital stay

Laying the foundation to treat patients globally

Cases Completed

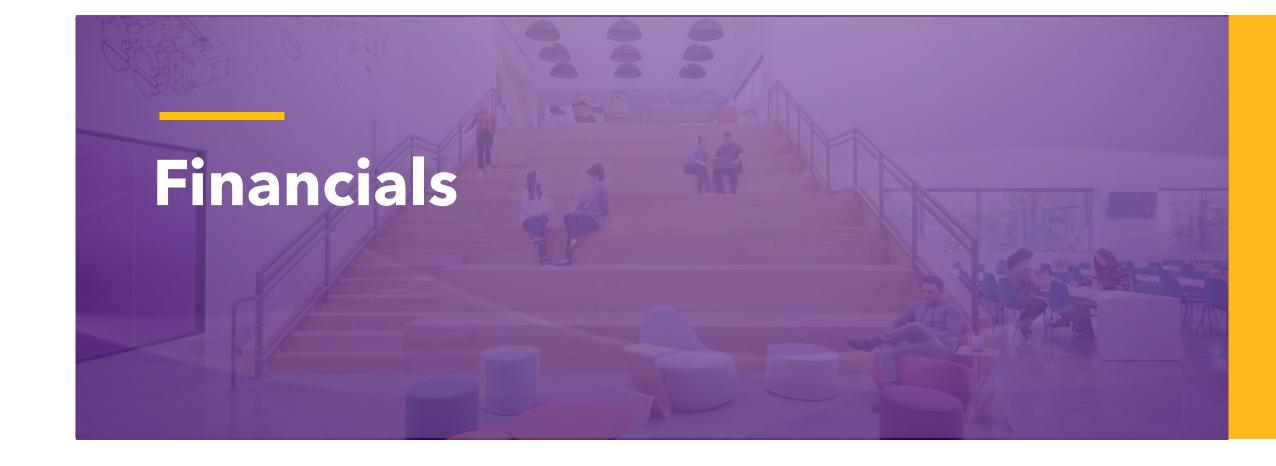
Pr-

Approval Received

Approvals In-Progress



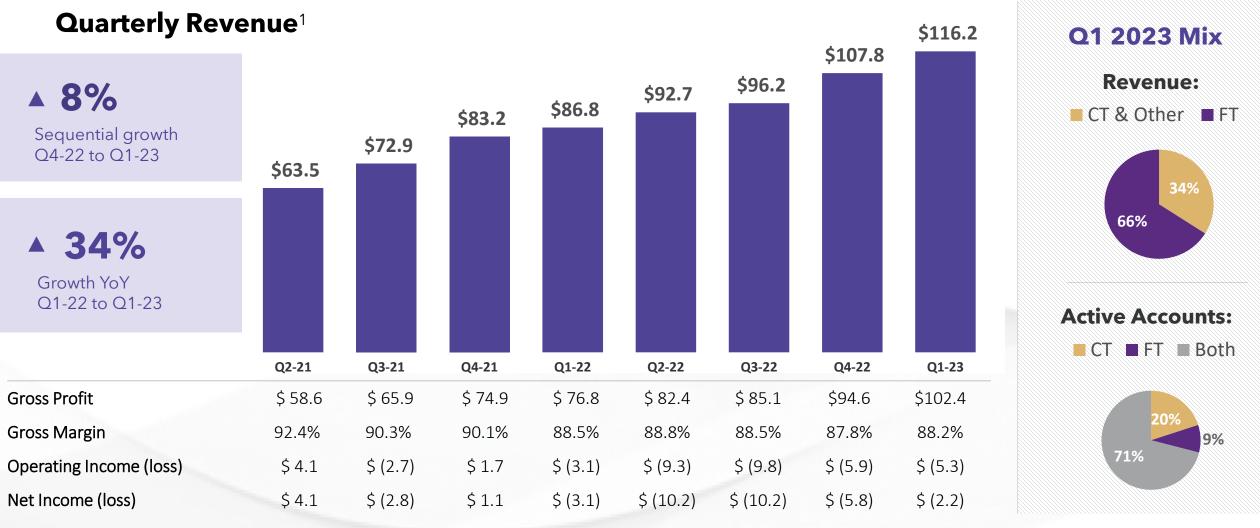
CASES COMPLETED IN 20+ COUNTRIES ACROSS THE GLOBE







Growing patient Impact reflected in financial performance



(1) Dollars are in millions.

Continued momentum into 2023 and beyond



2023 FY Guidance

\$478M - \$488M

25%-27% increase over full year 2022

Financial Profile

- Exceptional growth, significant runway
- Premium 85%+ gross margin profile
- Sustained operating profitability by 1H 2024
- Strong cash position



No small plans. And we're just getting started

1	EXPANDING US SALES FORCE	280+ U.S. Sales Territories
2	DRIVING DEEPER PENETRATION	<6% Penetration into U.S. VTE Incidence
3	BUILDING CLINICAL EVIDENCE	2 RCTs Underway + Multiple Major Clinical Studies
4	INNOVATING NEW PRODUCTS	5 Product Toolkits for 5 Distinct TAMs
5	EXPANDING INTO NEW MARKETS	>\$20B Total Global TAM (+ ~\$10B US Prevalence)



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