

Inari Investor Day Agenda

Time	Session		
7:30 - 8:00 AM	Registration and Breakfast Bar		
7:55 - 8:00 AM	Welcome Remarks		
8:00 - 8:45 AM	Site Tour and VTE Tech Demos		
8:45 – 9:00 AM	Break		
9:00 - 11:30 AM	Management Presentation		
11:30AM - 12:00 PM	Informal lunch, New Product Tech Demos, and Meet the Management Team		
12:00 PM	Conclude		

Please check your badges



Product Demos





Brian Strauss SVP Engineering

Purple 1 - 3

Paul Koehn SVP Operations

Orange 1 + 2



Site Tour

Eric Louw VP Manufacturing

Orange 3 + 4





Financial Information

Unless otherwise indicated, all financial and operational information included herein is as of June 30, 2022.

Indications for Use and Publication References

Indications for Use and relevant labeling information for all Inari products included in this presentation are included in the appendix. References to clinical studies and other publications cited in this presentation are located in the appendix.

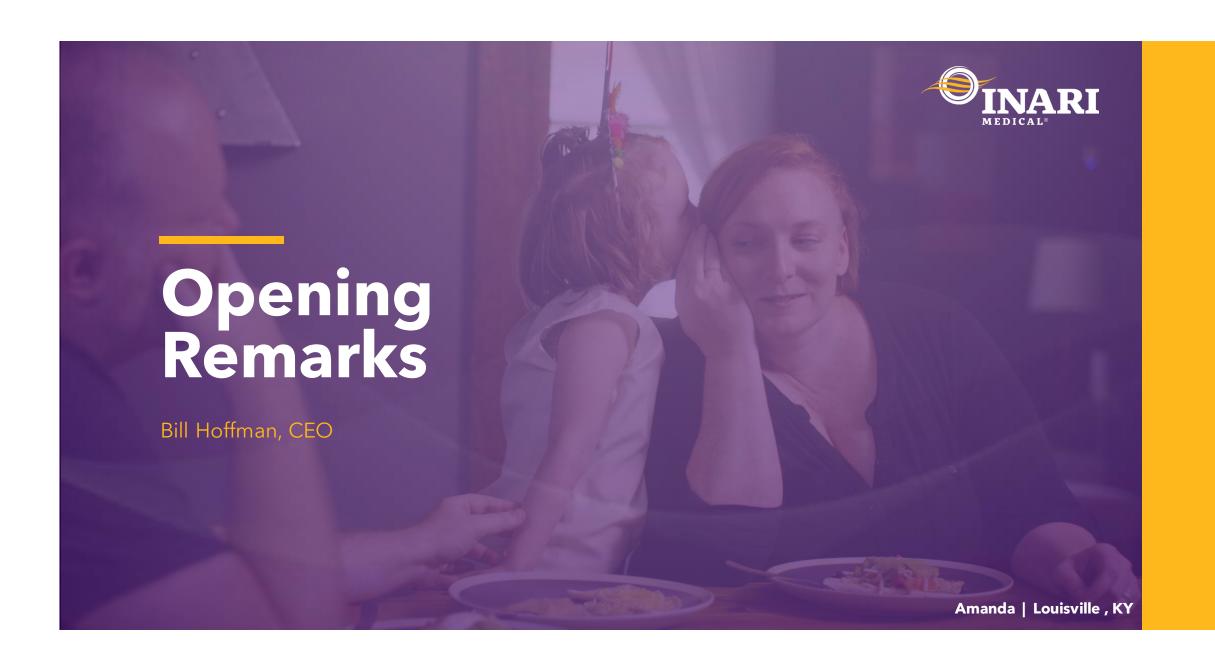
Forward Looking Statements

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 revenue and total procedures, 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; our ability to grow and maintain our US sales force; our ability to develop new tools and new markets; the results of our clinical studies; 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 expressions are intended to identify forward-looking statements, although not all forward-looking statements contain these words.

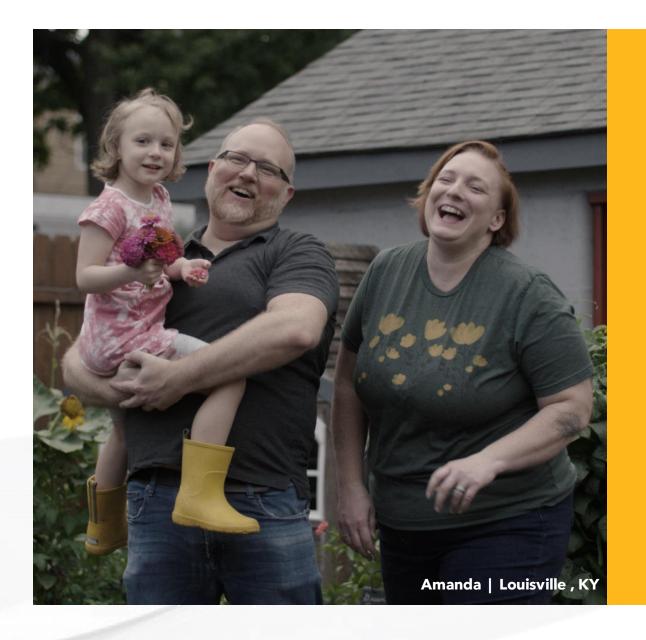
Forward-looking statements are based on and reflect management's current expectations, assumptions, estimates and projections that may or may not prove to be correct. These forward-looking statements are subject to a number of known and unknown risks, uncertainties, assumptions and other factors, many 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, 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 Reports 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.



Our dedication to changing lives drives every decision we make









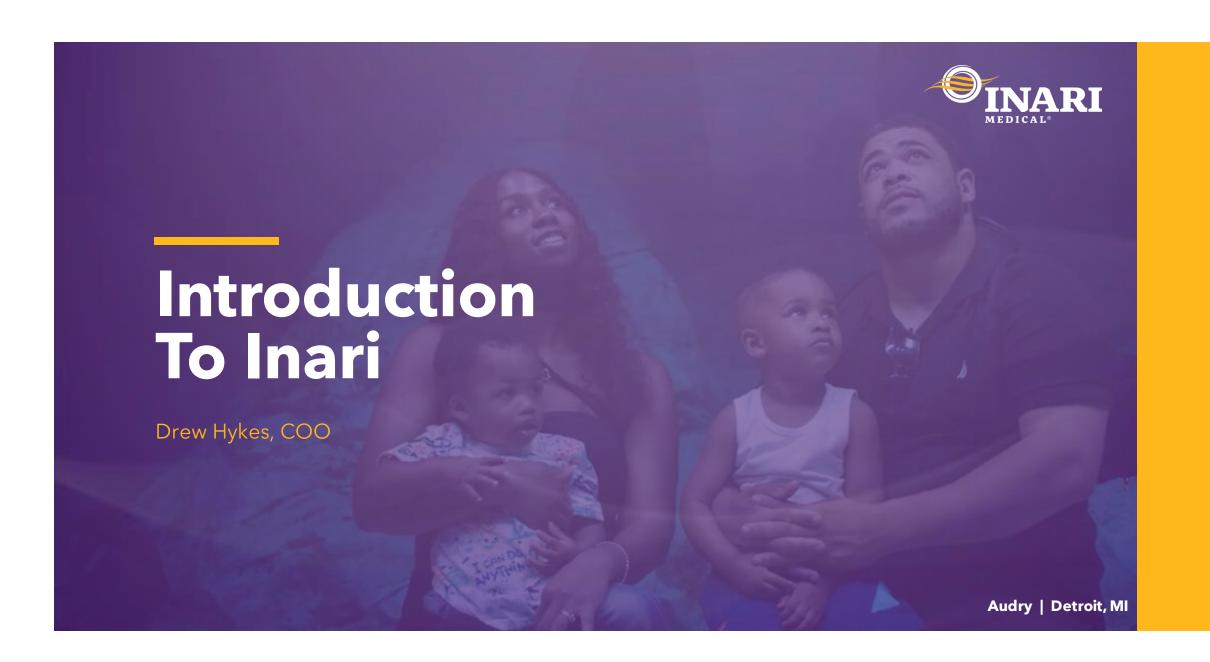
Patients first. **Always.**



Make no small plans. **Ever.**



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





Management Presentation Agenda

Time	Duration	Session
9:00 - 9:20AM	20 min.	Introduction
9:20 - 10:10 AM	50 min.	Inari Five Growth Drivers
10:10 - 10:20 AM	10 min.	Break
10:20 - 10:40 AM	20 min.	Physician Panel
10:40 - 10:50 AM	10 min.	Financials and Closing Remarks
10:50 - 11:30 AM	40 min.	Q&A

Strong leadership team with breadth & depth



Bill HoffmanChief Executive Officer



Mitch HillChief Financial Officer



Drew HykesChief Operating Officer



Dr. Tom TuChief Medical Officer



Angela Ahmad General Counsel



Brian StraussSVP Engineering



Eric Khairy SVP Marketing



Eric LouwVP Manufacturing



Janet BykVP Finance &
Accounting



John Borrell SVP Sales



Justin CrockettVP Inari Solutions
Group



Kevin Strange
VP Strategy &
Business
Development



Kit CariquitanVP Quality
Assurance &
Reg. Affairs



Norman Nie VP Information Technology



Paul KoehnSVP Operations



Randy Hamlin
VP Advanced
Development



Dr. Shon ChakrabartiVP & General Manager
Chronic Venous

Diseases



Tara DunnSVP Clinical Affairs &
Market Development



Dr. Venkat TummalaVP Medical Affairs



Dr. Victor TapsonVP Medical
Affairs



Vitas SipelisVP International

Venous Thromboembolism (VTE)

DVT

Up to

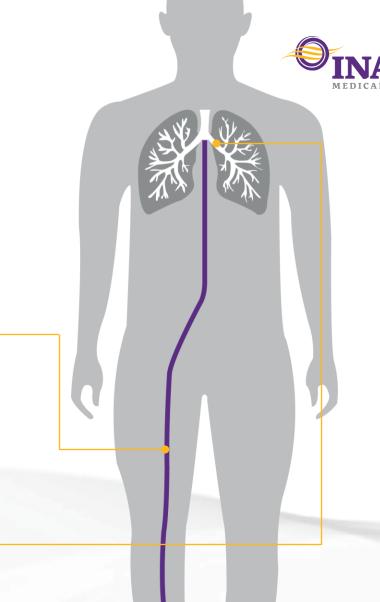
50%

Develop Post-Thrombotic Syndrome (PTS) within 2 years of a proximal DVT

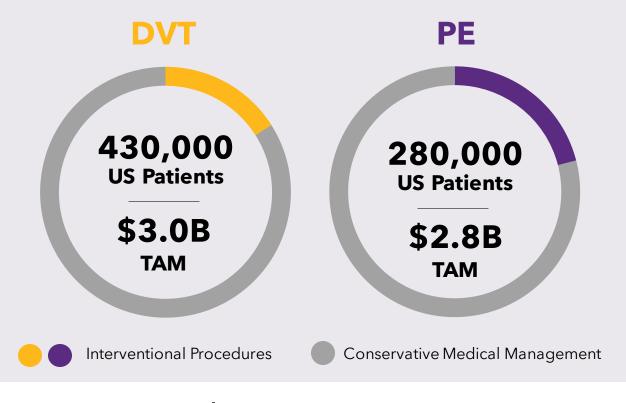
PE

#3

Leading cause of cardiovascular death



VTE is a large and highly underpenetrated opportunity to serve patients in need



\$5.8B

Total US VTE TAM Opportunity

\$15B+

Global VTE TAM Opportunity



Treatment of VTE evolving to definitive mechanical catheter intervention



Myocardial Infarction

Anti-Coagulation Only

Lytics Based

Definitive Catheter Therapy





Stroke

Anti-Coagulation Only

Lytics Based

Definitive Catheter Therapy





VTE (DVT & PE)

Anti-Coagulation Only

Lytics Based

Definitive Catheter Therapy





Highly differentiated, purpose-built solutions

- Simple, intuitive solutions
- Near complete thrombus removal
- **Eliminate need for dangerous lytics**
- Minimal blood loss
- Favorable hospital economics







Taking out all the clot matters



Our five growth drivers remain the roadmap

- Expanding Our U.S. Sales Force
- **Driving Deeper**VTE Penetration
- **3** Building Clinical Evidence
- Innovating
 Purpose-Built
 Solutions
- **5** Expanding Into New Markets











270+

U.S. Sales Territories

<5%

Penetration into U.S. VTE Incidence

250+

Peer Reviewed Publications

5

Distinct Product Toolkits
For 5 Distinct TAMs

>\$20B

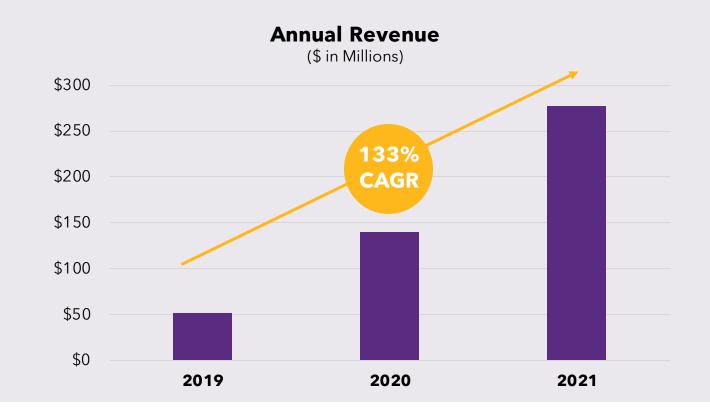
+

~\$10B

US Prevalence Opportunity



Consistent, premium financial performance



\$277M

91%

98%

\$330M

2021 Total Revenue

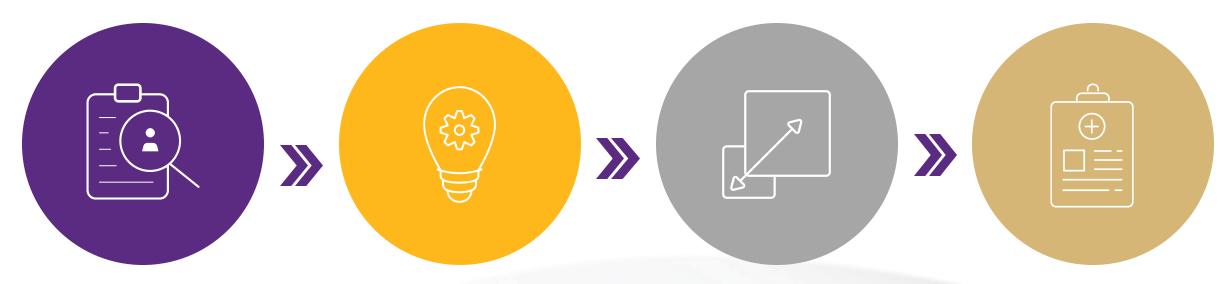
2021 Gross Margin

YOY Growth (From FY20)

Cash, Cash Equivalents, & Short-Term Investments (Q2 2022)



Our core competencies are scalable and allow us to treat more patients



Identify major unmet patient needs

R&D innovation engine to rapidly design purpose-built devices

High-Touch, scalable commercial org & market development capabilities

Clinical infrastructure generating data to change standard of care

FOUNDATION OF OPERATIONAL AND MANUFACTURING EXCELLENCE



No small plans. And we're just getting started



1	EXPANDING US SALES FORCE	BUILDING THE LARGEST INTERVENTIONAL SALES FORCE
2	DRIVING DEEPER PENETRATION	STANDARDIZING PATIENT PATHWAYS
3	BUILDING CLINICAL EVIDENCE	EXECUTING GUIDELINE-CHANGING CLINICAL TRIALS
4	INNOVATING NEW PRODUCTS	DEVELOPING PURPOSE-BUILT SOLUTIONS
5	EXPANDING INTO NEW MARKETS	LAUNCHING INTO NEW ADJACENCIES & GEOGRAPHIES





Significant growth in sales territories, increasing density of coverage

2020 (Q2)

2022 (YTD)



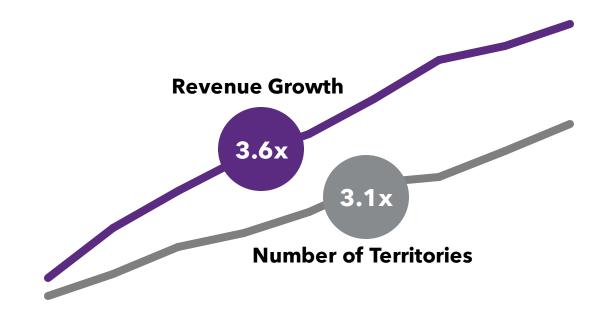


Accounts per
$$\sim 9$$
 $-0.6x \rightarrow \sim 5$



Focused on growth but remaining efficient

PRODUCTIVITY REMAINS HIGH DESPITE GROWTH AND AGGRESSIVE TERRITORY SPLITS

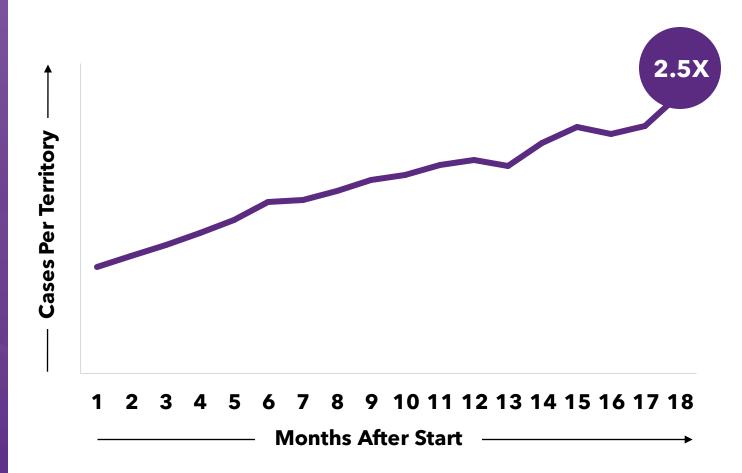


02.2020 03.2020 04.2020 02.2021 02.2021 04.2021 02.2022



Sales rep productivity ramps up quickly after start date

AVERAGE SALES REP PRODUCTIVITY OVER FIRST 18 MONTHS







Continuing growth to fully tackle VTE and address new disease states



Sales team efforts amplified by robust non-sales commercial team

+12

National Accounts

COMMERCIAL / MARKET DEVELOPMENTTEAM

Health Economics & Market Access (HEMA)

Inari Solutions Groups (ISG)

Medical Affairs (including doctors)

Medical Education Group

Upstream/Downstream Marketing & Sales Ops.



High-powered, hightouch commercial system designed to solve patient needs

Intentional fit-based hiring and promotion from within

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

Mining information across all sources, informing every decision we make

Solution-based toolkits, not widgets

Deliberate territory splits & alignment of incentives



Our commercial system enables us to scale in new markets with significant unmet needs



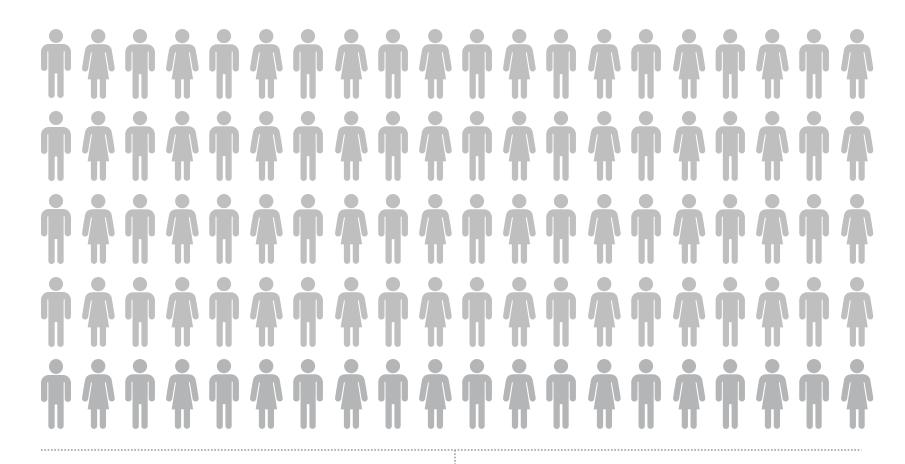






A significant responsibility

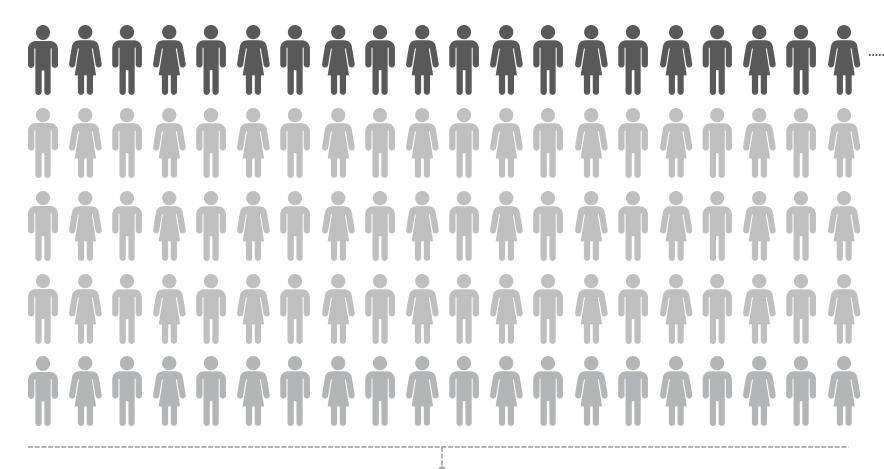




~710,000 Addressable US VTE patients per year

A significant responsibility





15-20%

Receive any intervention

~710,000 Addressable US VTE patients per year

A significant responsibility





patients per year



VTE lacks a systematic approach to identify, screen, and triage patients

MYOCARDIAL INFARCTION

- Consistent screening protocols
- Algorithms for triage and treatment
- Tracked metrics

STROKE

- Consistent screening protocols
- Algorithms for triage and treatment
- Tracked metrics



Interventionalist

Interventionalist

Interventionalist





Annual US VTE Incidence

710K

US Hospital Beds

~740K

Annual US Incidence per Bed

~1



500 Bed Mid-sized hospital



 ~ 500 addressable VTE patients/year

We're helping hospitals build programs that connect VTE patients to VTE experts

Excellent clinical outcomes

- Positive hospital economics
- Systematic patient pathway (i.e., a "VTE program")











ENGAGE

~1,250 Inari accounts

Find champions, build the foundation

EMPOWER

~120 Inari accounts

Create patient pathway and build awareness

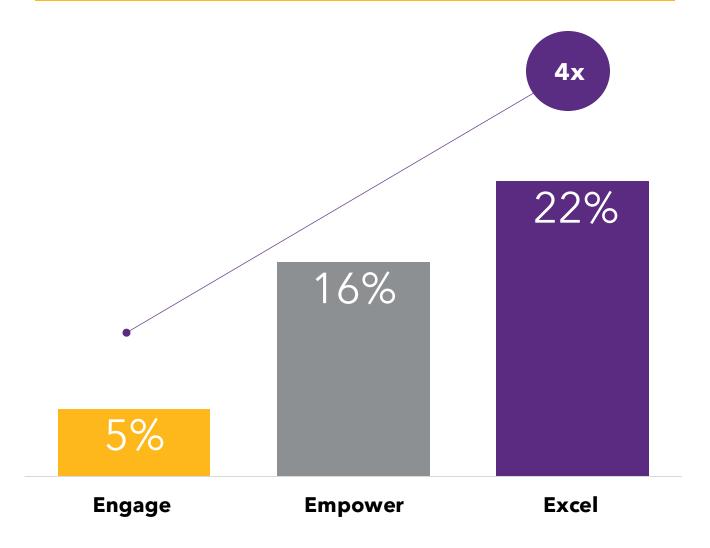
EXCEL

~20 Inari accounts

Solidify consistent patient identification, triage, tracking

VTE Excellence activities are beginning to drive deeper account-level TAM penetration

APPROX. ACCOUNT-LEVEL TAM PENETRATION BY STAGE









VTE Today

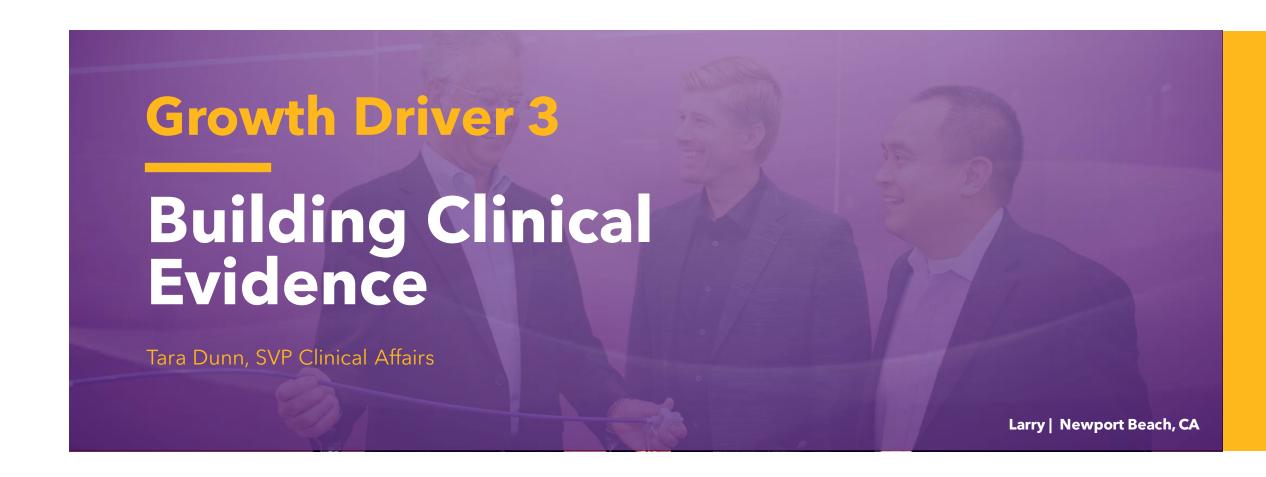
15-20%

Interventional TAM penetration

Myocardial Infarction

~90%

Interventional TAM penetration*





Transforming patient care





for patients





Develop the market



Set the bar high

Clinical by the numbers



2,000+

Patients studied to date

20+

Active or completed IIR engagements

250+

Peer reviewed publications

INCLUDING 2 RCTS

Major prospective Studies

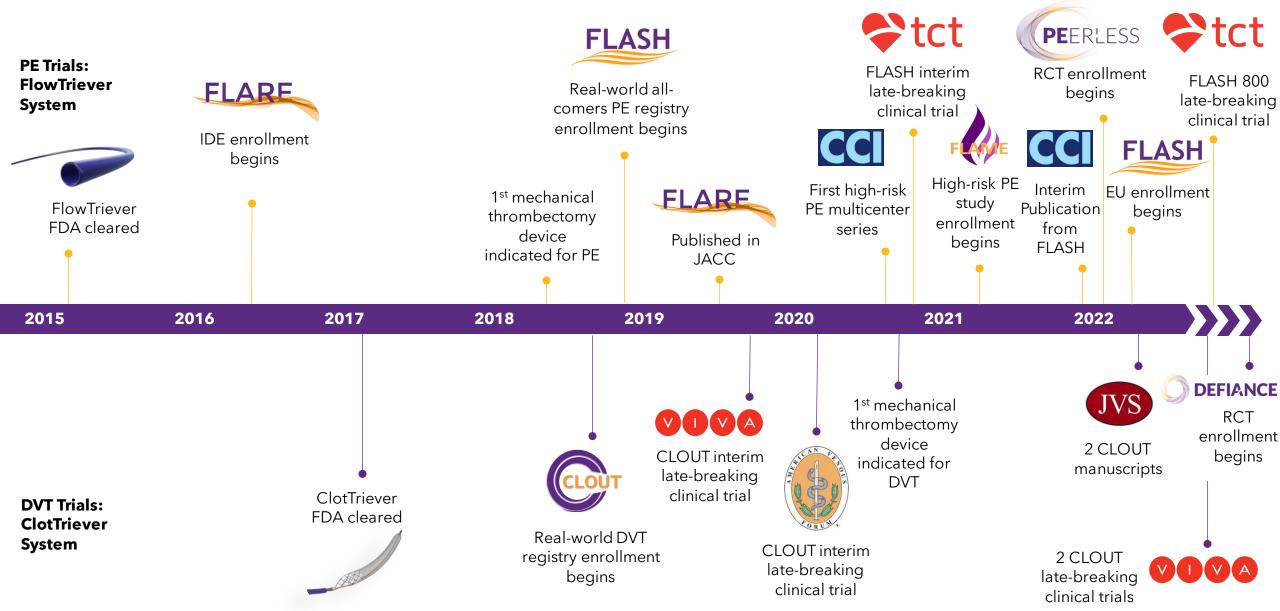


Strong and versatile team driving the quality and pace of best-in-class evidence generation



A tsunami of clinical data





Setting a high bar for VTE evidence





Largest Prospective PE Device Study

~1,000 Patients | 83 Sites

PE STUDIES



Largest Prospective High-risk PE Device Study

100+ Patients | 11 Sites

PEERLESS

First Inari RCT (FlowTriever v. CDT in PE)

550+ Patients | 60 Sites

DVT STUDIES



Largest Prospective DVT
Thrombectomy Study

500 Patients | 47 Sites

DEFIANCE

DVT RCT (ClotTriever v. AC)

300 Patients | 60 Sites

800th & final US patient enrolled. EU enrollment underway

Enrollment near complete

Enrollment commenced in both RCT and registry arms

500th & final patient enrolled

Enrollment expected early 2023

~2,500 patients across 5 studies

Status

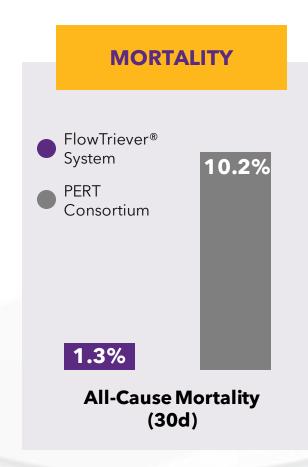


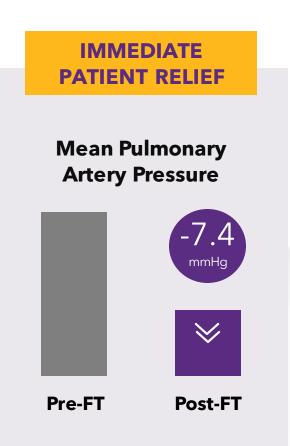
FLASH is the largest prospective registry in PE with exceptional results

EXCELLENT SAFETY

O%

Device related MAEs





LASTING PATIENT
BENEFITS

1.5%
Post-PE Syndrome

FLAME: High-risk PE guidelines are from an 8 patient study. FLAME is 100+ patients

Doing What's Right

Up to 50% of high-risk patients die within 30 days



Changing Guidelines

Designed per AHA recommendations

Developing the Market

Why not treat intermediate-risk?

PEERLESS: Superiority RCT of FlowTriever vs CDT in PE

Intermediate-High Risk PE

550 Patients Randomized 1:1

FlowTriever CDT

Up to 150 Patients in Registry Arm

FlowTriever Only (Contraindication cohort)

Patients Followed for 30 Days



HIGHLIGHTS



Currently >60% of patients intervened on receive Catheter Directed Thrombolysis (CDT)



Primary endpoint via win ratio:

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



Head-to-head definitive treatment trial (US & EU)



Enrollment ahead of schedule

CLOUT demonstrates we can do better for DVT patients

500
Patients Enrolled

47

Sites

2 out of 3

With **Subacute** and/or **Chronic Clot**



EXCELLENT SAFETY

0%

Vessel/Valve Damage

ON-TABLE EFFECTIVENESS

86%

Effectiveness in Core Lab Adjudicated Clot Removal

LASTING PATIENT BENEFITS

91%

Freedom from Moderate to Severe PTS at 6 months

DEFIANCE: Superiority RCT of ClotTriever vs Anticoagulation in DVT

Moderate-Severe Iliofemoral DVT

300 Patients Randomized 1:1

ClotTriever

Conservative Medical Management (AC)

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



Designed to transform standard of care



Enrollment expected to begin early 2023



Exceptional programproductivity and quality of patient outcomes

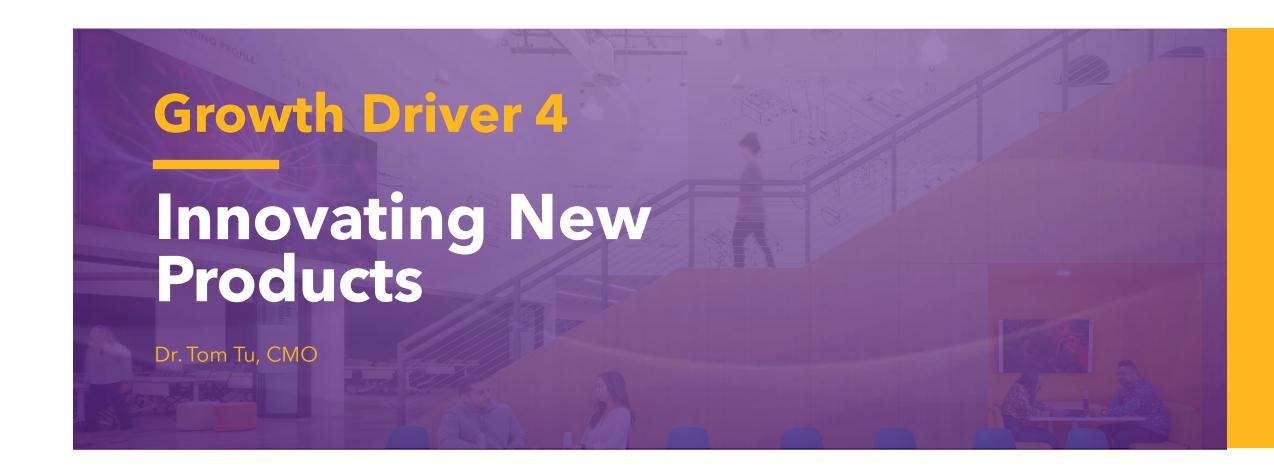
Enrollment began 2022 **PE**ERLESS

Enrollment begins 2023



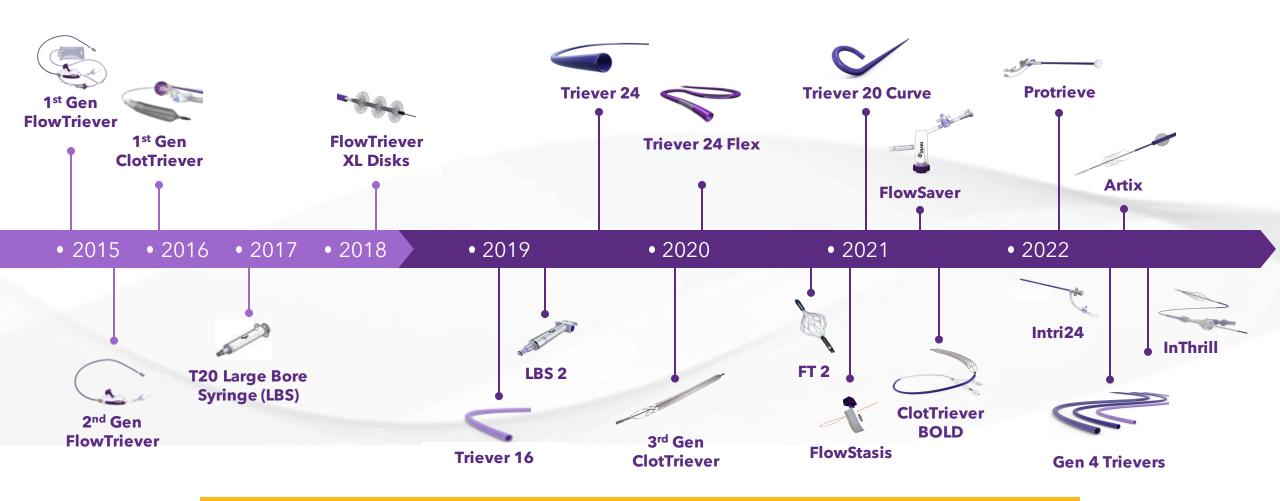


2021





Years of knowledge and commitment. Our mission to address unmet needs

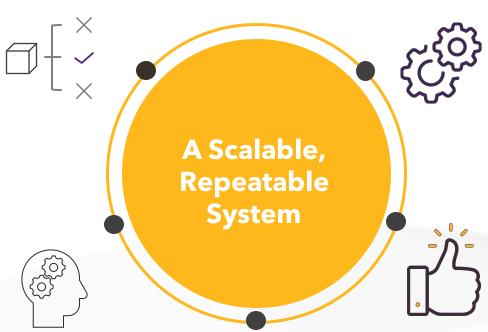




Our Innovation Engine continues to implement purpose-built solutions for unmet patient needs

Rapid innovation

via in house machine shop, 3D-printing, bio-skills lab, laser cutting, etc.



"Engine Room"

R&D infrastructure

Engineering roles defined around solutions,

not products



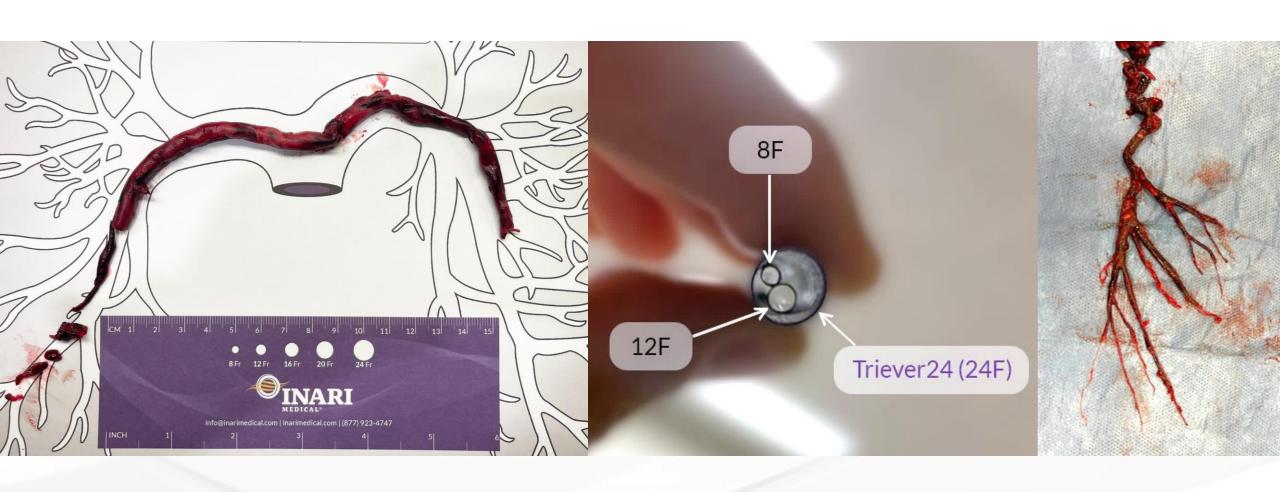
Mining information, informing every project

Market analysis

focused on large and unmet needs

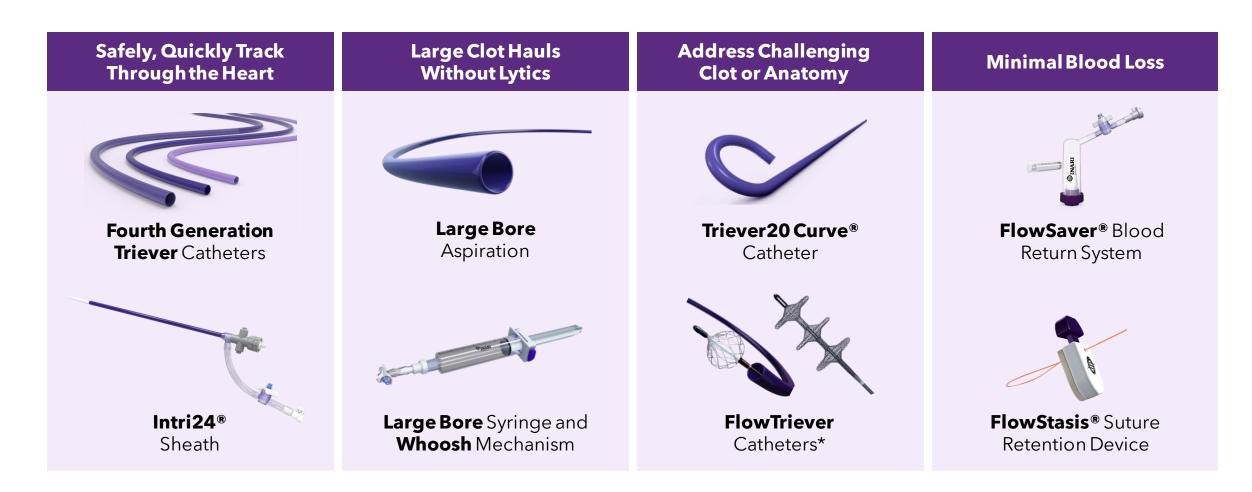
Large bore catheters produce large clot hauls





A comprehensive PE procedure solution years in the making





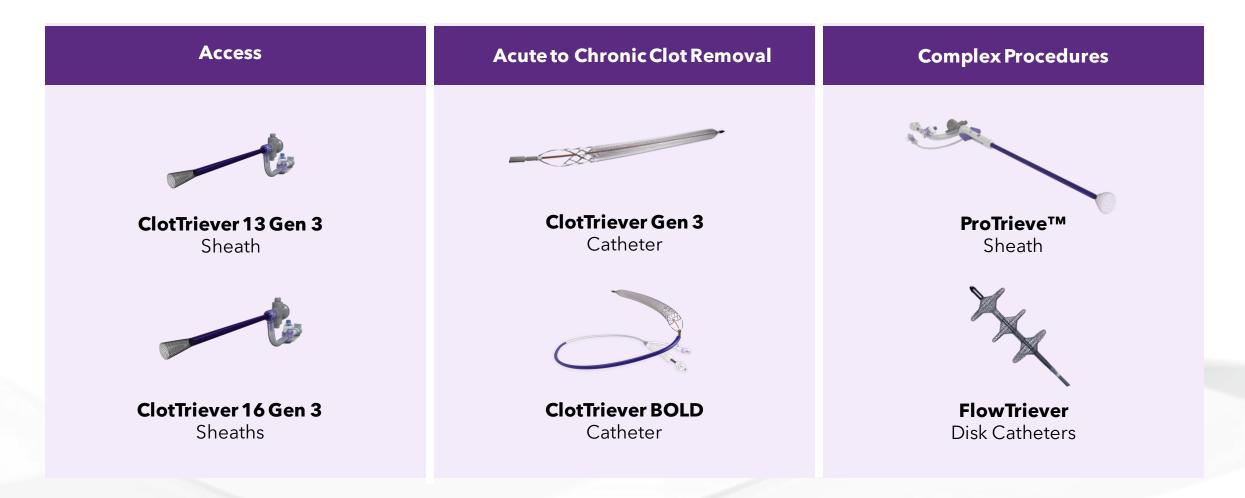
A purpose-built DVT treatment for the full range of clot chronicity



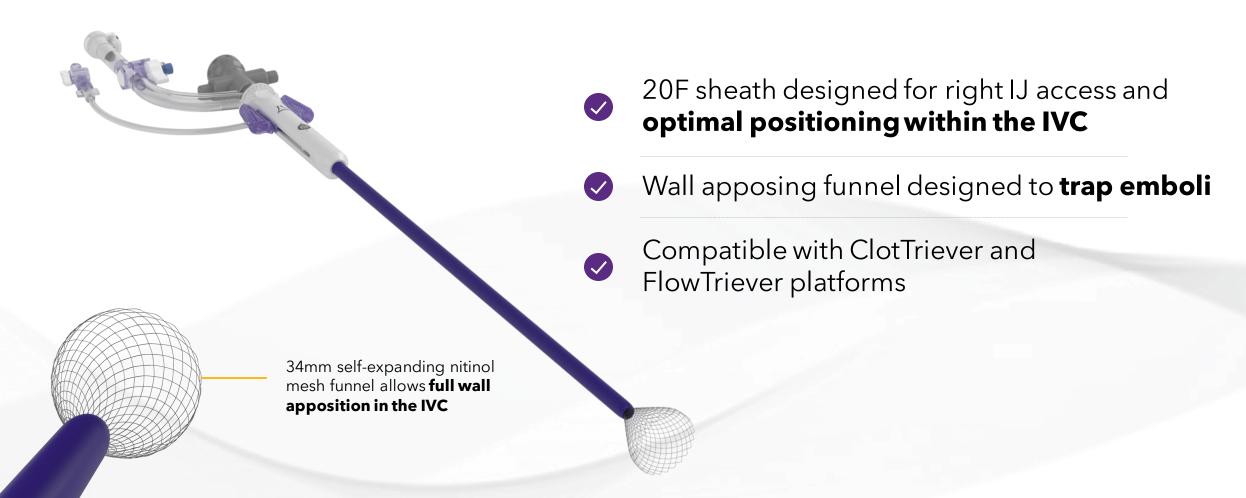


A comprehensive DVT procedure solution years in the making



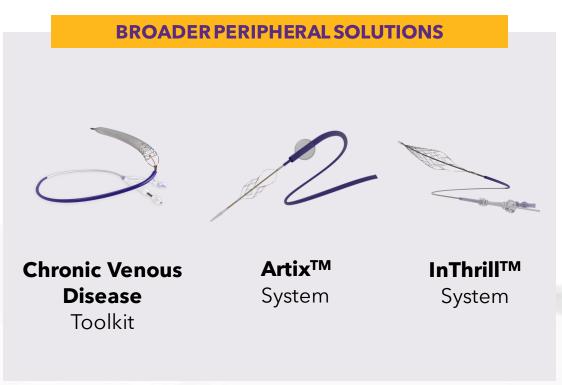


ProtrieveTM provides confidence during complex DVT and IVC procedures



Expanding beyond VTE to develop purpose-built solutions for new diseases





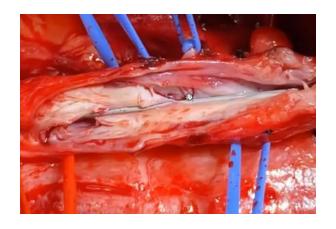
Chronic Venous Disease (CVD)

Chronic clot

Post-Thrombotic occlusions

PTS and venous leg ulcers (VLU)

CVD often progresses from DVT and includes scarred vein walls & wall-adherent **obstructions**



If obstructions are left unaddressed, patients can develop **painful**, **debilitating ulcers**





Conservative treatments for Chronic Venous Disease are inadequate and only address symptoms



Compression Therapy



Anticoagulation

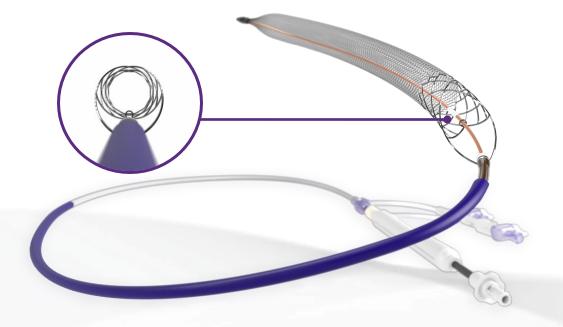




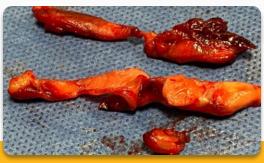
ClotTriever BOLD was designed to extract the full range of clot chronicity



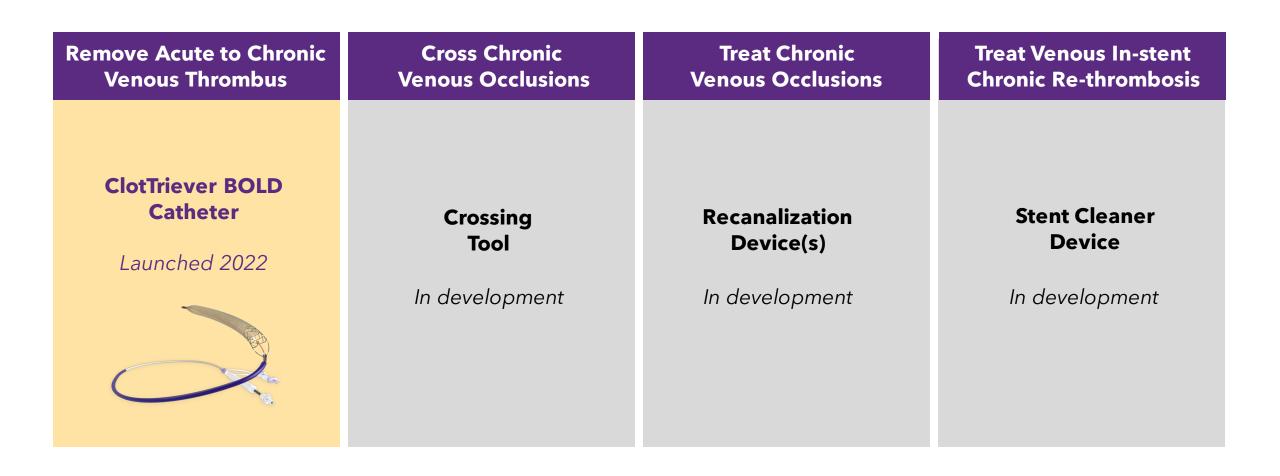
- Clot is often older than symptoms suggest
- ~30% greater radial force for **improved** wall apposition
- Improved thrombus engagement to treat the full range of acute to chronic DVT







We're building a solution-based toolkit to address Chronic Venous Disease

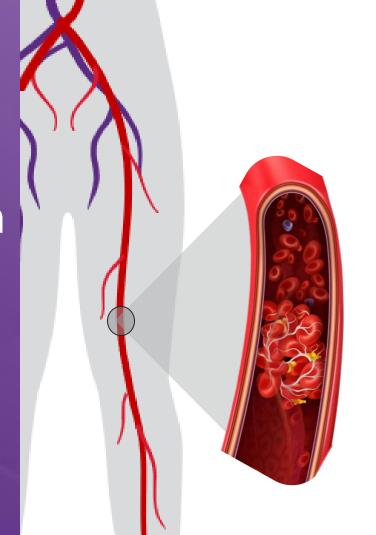


Arterial Thromboembolism

Acute limb ischemia (ALI)

Acute visceral ischemia

Chronic limb ischemia (CLI)



Acute embolization event - **extensive damage** can happen if not treated quickly



Current treatments for arterial thromboembolism have significant drawbacks

- Often requires open surgical procedures
- Distal embolization and vessel trauma
- High rates of lytic use
- Significant blood loss
- Need for a better, purpose-built solution



The Artix System: purpose-built toolkit for peripheral artery thromboembolism

- Combines both aspiration and mechanical thrombectomy
- Sheath has 4X flow rate vs. existing arterial catheters
- Proximal flow arrest to prevent distal embolization



Artix MT

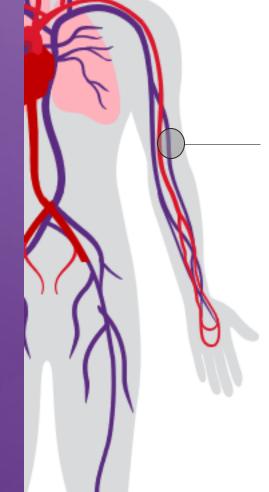


Small Vessel Thrombosis

Upper Extremity

Below-the-knee

AV thrombosis



AV access thrombosis can result in complications and loss of access to life-saving dialysis





Limitations in current treatments for small vessel thrombosis

- AV "declotting" sends clot to the lungs, exacerbating pulmonary hypertension
- High recurrence rates
- Ineffective for chronic clot
- Ineffective for large clot burdens
- Need for a better, purpose-built solution

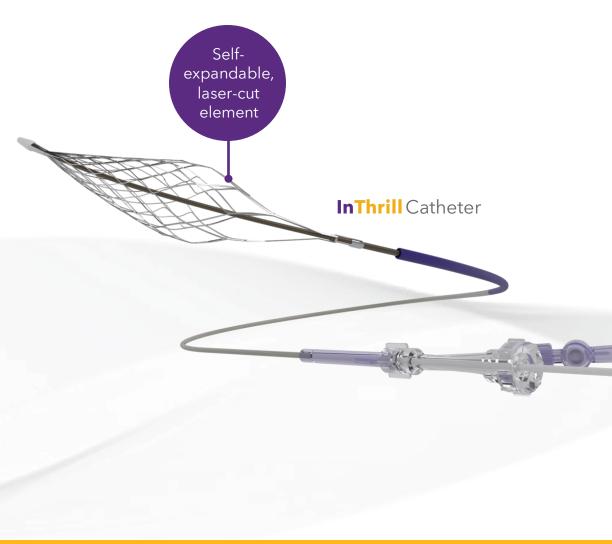


The InThrill System: a solution for smaller vessels



- Effectively extracts clots
- Addresses acute to chronic thrombus
- **▼** Tailor made for 4-10mm vessels

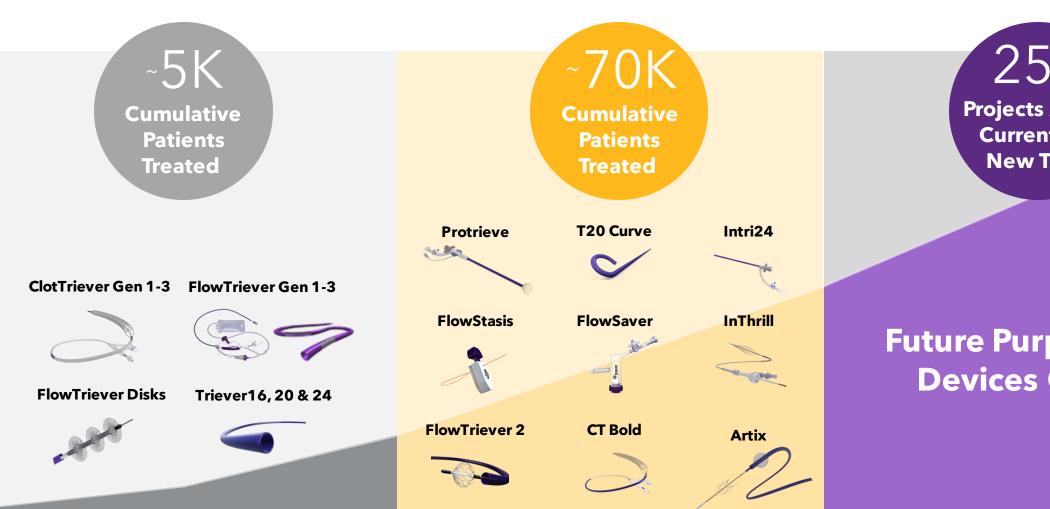




Note: the InThrill device is indicated for use in the peripheral vasculature

We're just getting started!





Projects Across Current and New TAMs

Future Purpose-Built Devices Coming

< 2020

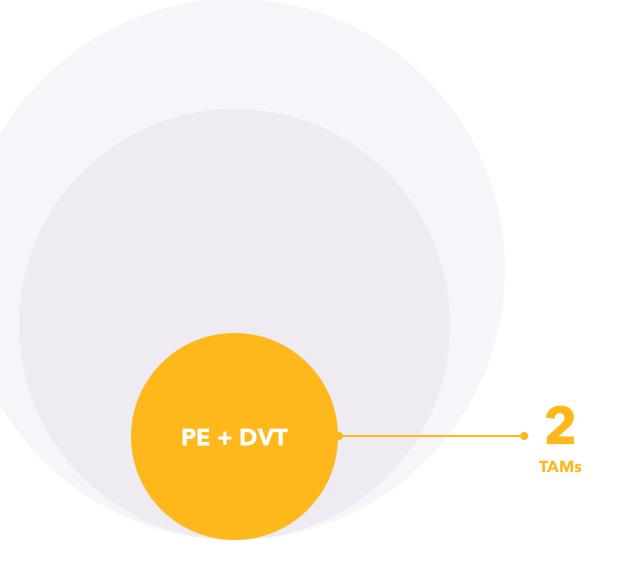
2020 - 2022

2022 - 2024+



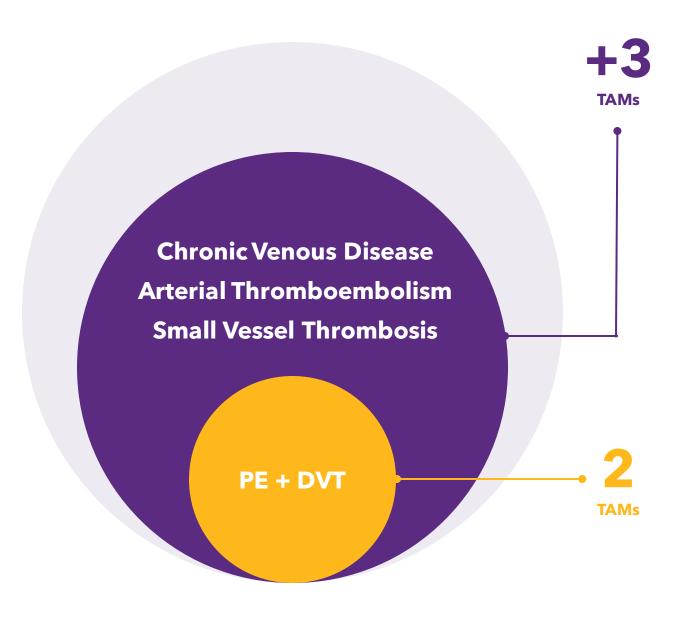


Historically, we have treated patients in two TAMs with two toolkits



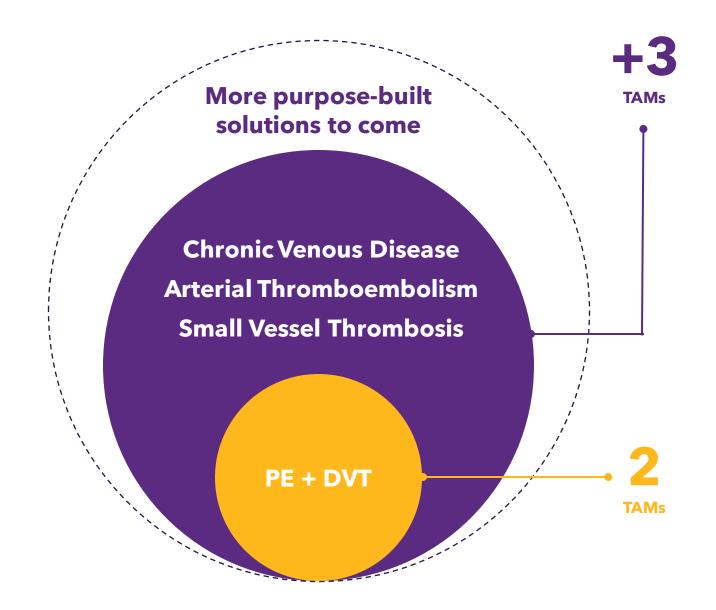


Currently expanding into three new TAMs, continuing to address large unmet needs





We have no small plans. More purposebuilt solutions to come in incremental **TAMs**





Protrieve is a purpose-built solution for complex IVC cases



LMR launched in August

Enables complex IVC cases

New tool for interventionalists treating **DVT** and **PE**



Chronic Venous Disease prevalence opportunity larger than core TAM







INCIDENCE

~100K ~\$1B

Patients

TAM Opportunity

PREVALENCE

 $\sim 1 M$

Patients

~\$10B

TAM Opportunity



Chronic Venous Disease: addressing the underlying cause via purpose-built devices

~3,000 CT-Bold cases completed

Evaluating **go-to-market options**

Common interventional call point

Premium ASP of ~\$10K



Crossing Device

In development

*To treat acute to chronic clot

Recanalization Device(s)

In development

Stent Cleaner Device

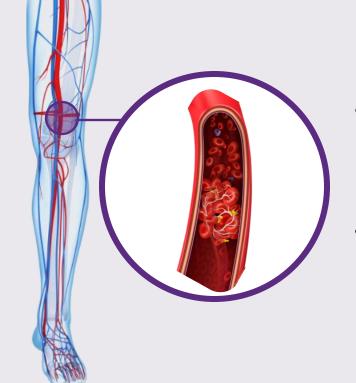
In development

"Bold is amazing for treating more-chronic patients... but I CANNOT WAIT for what is coming with the rest of this dedicated toolkit!"

- Dr. Nicolas Mouawad



Arterial Thromboembolism large TAM with significant unmet needs



- Acute event extensive damage can happen if not treated quickly
- Includes Acute Limb Ischemia, Acute Visceral Ischemia, and CLI procedures where distal embolization occurs

~80K

US Patient Incidence

~\$600M

US TAM Opportunity



Artix: purpose-built toolkit addressing unmet needs in arterial thromboembolism

Artix MT LMR in April 2022; Balloon Sheath LMR in July 2022

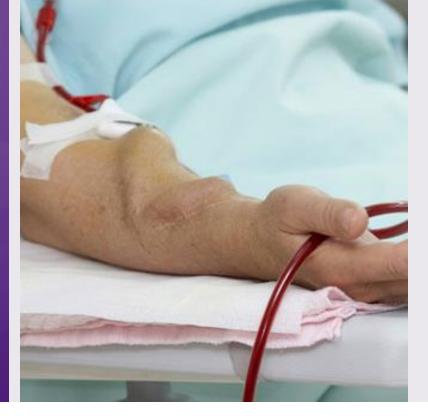
Significant site-of-service and physician overlap

Targeting an ASP of ~\$7.5K

Additional tools in development



Small Vessel Thrombosis: No purpose-built solutions exist for this large patient population



AV access thrombosis can result in complications and loss of access to lifesaving dialysis

~150-200K

US AVF thrombotic events / year

~80K

Addressable US BTK + UE thrombotic events / year

~\$1B

Total US Market
Opportunity



InThrill: a purpose-built, novel solution designed to treat small vessels

LMR launched in **August**

Targeting in-hospital procedures; significant treating physician overlap with DVT and PE

Targeting an ASP of ~\$4K

Additional tools in development





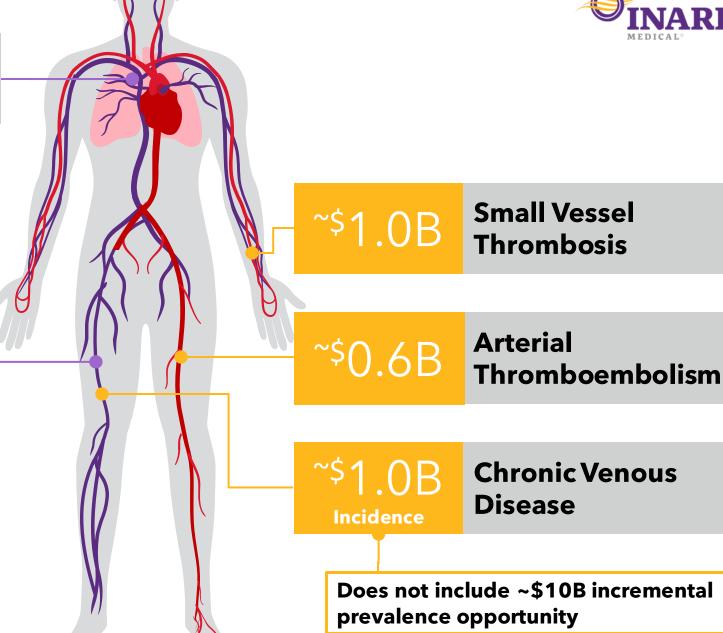
~\$2.8B

Pulmonary Embolism

~\$3.0B

Deep Vein Thrombosis

Large US total addressable market totaling ~\$8B across 5 disease states



Substantial global opportunity exists across VTE and three new disease states



Global Incidence TAM across 5 Disease States

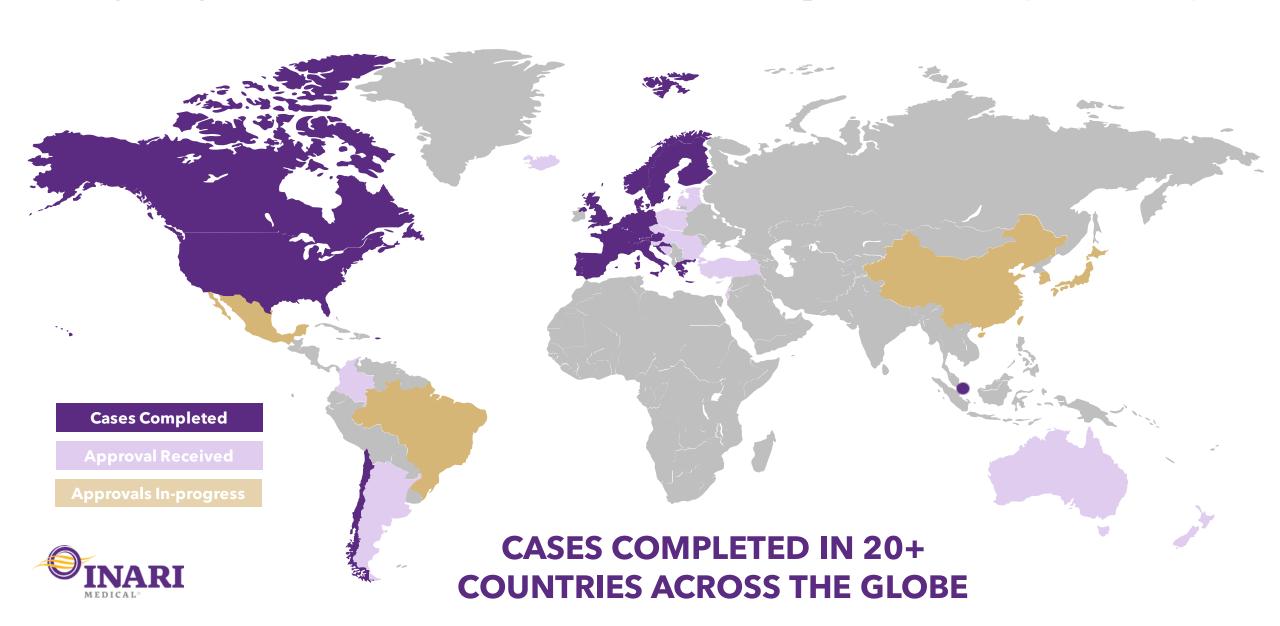


~\$10B

CVD Prevalence Opportunity (US only)



Laying the foundation to treat patients globally



No small plans. And we're just getting started



1	EXPANDING US SALES FORCE	BUILDING THE LARGEST INTERVENTIONAL SALES FORCE
2	DRIVING DEEPER PENETRATION	STANDARDIZING PATIENT PATHWAYS
3	BUILDING CLINICAL EVIDENCE	EXECUTING GUIDELINE-CHANGING CLINICAL TRIALS
4	INNOVATING NEW PRODUCTS	DEVELOPING PURPOSE-BUILT SOLUTIONS
5	EXPANDING INTO NEW MARKETS	LAUNCHING INTO NEW ADJACENCIES & GEOGRAPHIES

10 Minute Break







Inari Investor Day

Physician Panel Discussion



Thomas Tu, MDChief Medical Officer
Inari Medical
Moderator



Christopher M. Huff, MD
Interventional Cardiology
OhioHealth Riverside Methodist Hospital



Steven Abramowitz, MDVascular Surgery
MedStar Washington Hospital Center

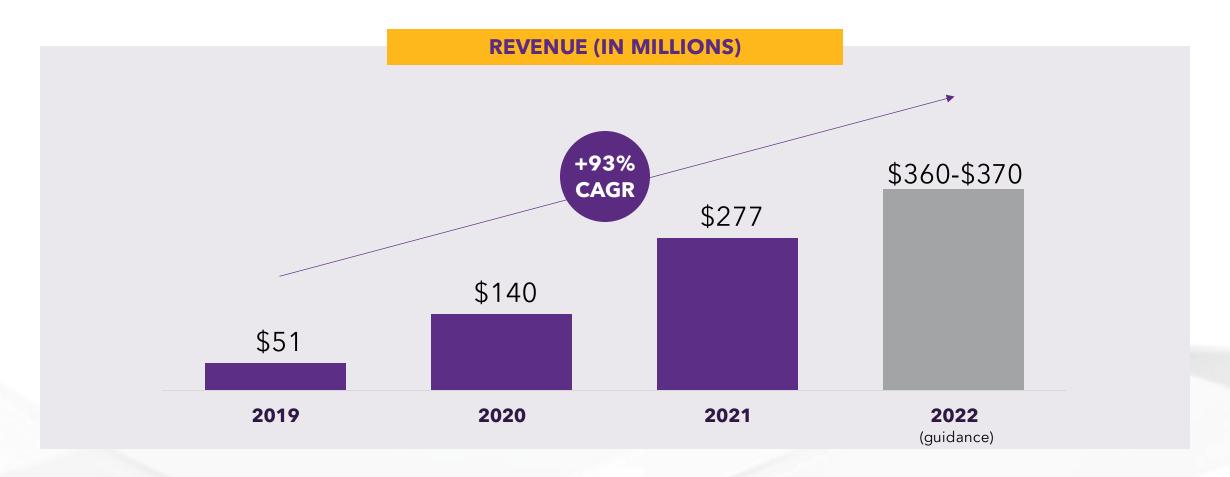






We have a proven track record of industry leading revenue growth



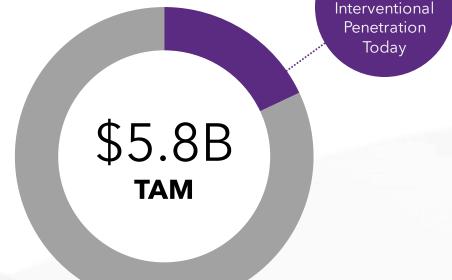


MedTech comps to approximate future VTE market size

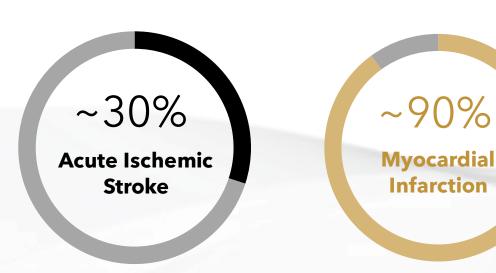


VTE PENETRATION TODAY

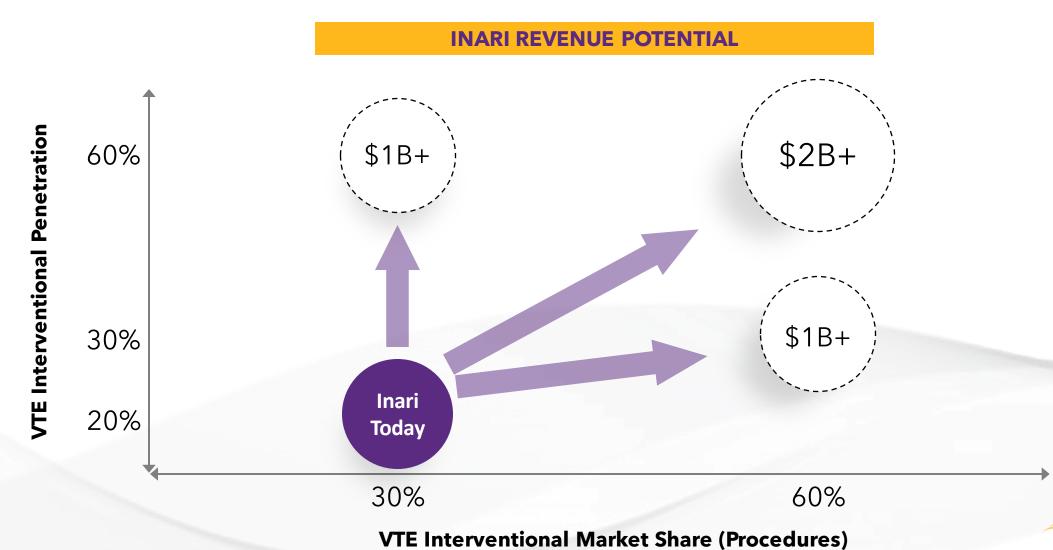




INTERVENTIONAL PENETRATION PROXIES



Substantial revenue opportunity exists for the leader in \$5.8B VTE TAM



Premium growth combined with exceptional margin profile





Well-positioned for sustained operating profitability

2022

Significant investment in our growth drivers

- Commercial team
- Clinical research
- Product development pipeline
- International





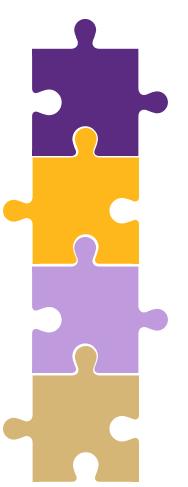
Sustained operating profitability by 1H 2024

- Large, attractive market
- ~85% target gross margin
- Commercial productivity ramp
- Disciplined investment approach



All the components of a premium financial profile



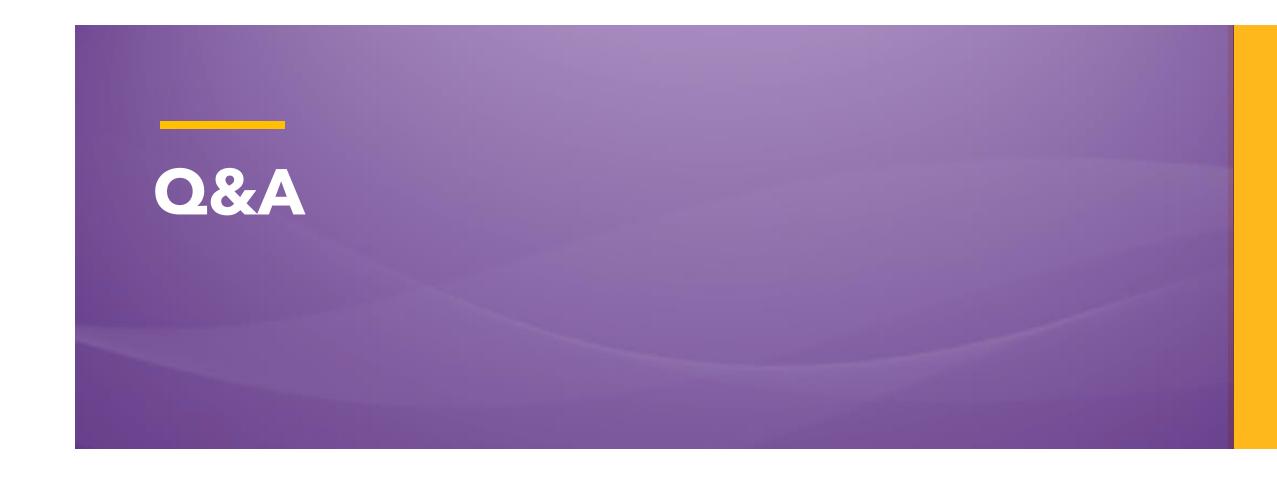


Market leader in \$5.8B underpenetrated US VTE market;
 expanding into global ~\$20B+ TAM

Exceptional gross margin profile

 Disciplined investments driving growth, operating leverage and consistent profitability

Strong balance sheet and ~\$330M cash position, allowing financial flexibility









Appendix



Citations



Slide #:	Source(s):		
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46	• Kucher N, Rossi E, De Rosa M, Goldhaber SZ. Massive pulmonary embolism. Circulation. 2006;113(4):577-82		
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61	 Fife, C. E., Publicly Reported Wound Healing Rates: The Fantasy and the Reality. Advances in wound care 2018 Fife, C.E., From the Editor: The Need for Real Venous Ulcer Data. Today's wound clinic., 2018 		
76	 Donadini, Marco, et. al., "Prognostic Significance of Residual Venous Obstruction in Patients with Treated Unprovoked Deep Vein Thrombosis." Thrombosis and Haemostasis, vol. 111, no. 01, 2014, pp. 172–179 Dronkers, C.E.A. et. al., "Deep vein thrombosis: diagnostic and prognostic challenges" Thromb Haemost. 2018 Aug; 118(8):1428-1438 Hospital claims data and internal analysis 		
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78	 Fleck D, Albadawi H, Wallace A, Knuttinen G, Naidu S, Oklu R. Below-knee deep vein thrombosis (DVT): diagnostic and treatment patterns. Cardiovasc Diagn Ther. 2017 Dec;7(Suppl 3):S134-S139 Elna M. Masuda, Robert L. Kistner, The Case for Managing Calf Vein Thrombi With Duplex Surveillance and Selective Anticoagulation, Disease-a-Month, Volume 56, Issue 10, 2010, Pages 601-613, ISSN 0011-5029 Franco, L, Giustozzi, M, Agnelli, G, Becattini, C. Anticoagulation in patients with isolated distal deep vein thrombosis: a meta-analysis. J Thromb Haemost 2017; 15: 1142–54 Hospital claims data and internal analysis 		
80	 Creager MA, Kaufman JA, Conte MS. Clinical practice. Acute limb ischemia. N Engl J Med. 2012 Jun 7;366(23):2198-206 Howard et. al., Population-Based Study of Incidence, Risk Factors, Outcome, and Prognosis of Ischemic Peripheral Arterial Events. Circulation Vol 132, Issue 19:1805–1815 Conte MS, et. al., GVG Writing Group. Global vascular guidelines on the management of chronic limb-threatening ischemia. J Vasc Surg. 2019 Jun;69(6S):3S-125S.e40 Agarwal S, et al. Burden of Readmissions Among Patients With Critical Limb Ischemia. J Am Coll Cardiol. 2017 Apr, 69 (15) 1897–1908 Internal analysis 		

Device Indications For Use



The **FlowTriever System®** is indicated for (1) the non-surgical removal of emboli and thrombi from blood vessels (2) injection, infusion and/or aspiration of contrast media and other fluids into or from a blood vessel. The FlowTriever System is intended for use in the peripheral vasculature and for the treatment of pulmonary embolism.

Triever Catheters are intended for use in peripheral vasculature and for the treatment of pulmonary embolism. The Triever Catheters are also intended for use in treating clot in transit in the right atrium.

The **FlowTriever2® 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 FlowTriever2 Catheter is intended for use in the peripheral vasculature.

The FlowSaver® Blood Return System is used with Triever Catheters for autologous blood transfusion.

The **Intri24 introducer sheath** is indicated to provide a conduit for the insertion of endovascular devices into the vasculature while minimizing blood loss associated with such insertions.

Device Indications For Use (cont.)



The ClotTrieverTM thrombectomy system consists of the ClotTriever catheter and ClotTriever sheath. The ClotTriever Thrombectomy System is indicated for: (1) the non-surgical removal of thrombi and emboli from blood vessels. (2) Injection, infusion, and/or aspiration of contrast media and other fluids into or from a blood vessel. The ClotTriever Thrombectomy System is intended for use in the peripheral vasculature including deep vein thrombosis (DVT).

The **FlowStasis® Suture Retention Device** is indicated for temporary suture retention following a percutaneous venous procedure.

The **Artix MT thrombectomy device** is indicated for: (1) The non-surgical removal of emboli and thrombi from blood vessels. (2) Injection, infusion, and/or aspiration of contrast media and other fluids into or from a blood vessel. The Artix MT thrombectomy device is intended for use in the peripheral vasculature.

The InThrill Thrombectomy System consists of the InThrill Thrombectomy Catheter and InThrill Sheath. The InThrill Thrombectomy System is indicated for: (1)The non-surgical removal of thrombi and emboli from blood vessels. (2) Injection, infusion, and/or aspiration of contrast media and other fluids into or from a blood vessel. The InThrill Thrombectomy System is intended for use in the peripheral vasculature. The InThrill Thrombectomy System is not intended for use in deep vein thrombosis (DVT) treatment.

Device Indications For Use (cont.)



Caution: Federal (USA) law restricts this device to sale by or on the order of a physician.

Refer to Instructions for Use for complete indications for use, contraindications, warnings, and precautions.

All trademarks are property of their respective owners.