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# Our Mission: Treat and Transform the Lives of Patients Suffering from Venous Diseases





## Commercial-Stage Company Focused on Venous Solutions



Commercial-stage company that has developed minimally invasive products designed to remove large clots from veins without the need for thrombolytic drugs.



## **Purpose Built Solutions**

for the

## Venous Anatomy

2 Products
Both Disposable; No
Cap Equip

>16,000
Patients Treated

\$9,100 (1)
Blended Revenue per
Procedure

\$38.7mm 3Q20 Revenue (FY19: \$51.1M YTD20: \$91.1M)

>80% Gross Margin

# Inari Medical: Purpose Built Solutions for Removing Blood Clots from the Venous Anatomy

#### **Venous Focused**



We are pioneering devices specifically designed and purpose-built for the venous anatomy and its unique clot morphology

## 2 FDA-Cleared & Marketed Products



ClotTriever (used in DVT) and FlowTriever (used in PE) safely and effectively remove large volumes of clot while eliminating need for thrombolytic drugs

## Large Market Opportunity



Deep Vein Thrombosis ("DVT") and Pulmonary Embolism ("PE") collectively represent an approximately \$3.6bn annual U.S. market opportunity (1)

## Scaling Commercial Organization



Rapidly growing commercial organization that is designed to harness and leverage unique insights into key business decisions

## **Product Simplicity**



Intuitive, easy to use, single-use devices that do not require capital equipment or the use of thrombolytic drugs and that enable a short learning curve

# **Compelling Economics & Improved Efficiency**



Products allow for short, single sessions and are designed to eliminate need for expensive thrombolytics which require costly ICU stays and carry risks of major bleeding

### **Unique Culture**



Carefully selected team collectively pursuing extraordinary outcomes and improving the quality of life for our patients

## Strong Leadership Team to Capitalize on Our Opportunity



Bill Hoffman
Chief Executive Officer
Visualase, Fox Hollow, RITA
Medical



Chief Financial Officer

Cameron Health, Visiogen,
Buy.com, Walt Disney Imagineering

**Mitch Hill** 



Drew Hykes
Chief Operating Officer
Sequent Medical, Medtronic,
ABN AMRO



Dr. Tom Tu

Chief Medical Officer

Baptist Health Louisville, Massachusetts
General Hospital

| Paul Koehn    | VP Operations                             | Cardiovascular Systems                                   |
|---------------|---|--|
| Tara Dunn     | VP Clinical Affairs & Market Development  | Volcano Corporation, Medtronic, Health Advances          |
| Eric Khairy   | VP Marketing                              | Philips, Volcano Corporation, Corindus Vascular Robotics |
| John Borrell  | VP Sales                                  | Trireme Medical, Fox Hollow, Cardiovascular Systems      |
| Brian Strauss | VP Engineering                            | Reverse Medical, Medtronic, ev3, Micro Therapeutics      |
| Eben Gordon   | VP Quality Assurance & Regulatory Affairs | Sotera Wireless, SenoRx, ReVision Optics                 |



# Poor Outcomes for Venous Thrombectomy Stem from Differences Between Arterial and Venous Clot

| Parameter         | Arterial System  | Venous System   |
|-------------------|--|---|
| Hemodynamics      | High flow, high pressure   | Low flow, low pressure  |
| Vessel morphology | Vessels taper in direction of flow   | Vessels enlarge in direction of flow  |
| Presentation      | Ischemic insult (MI, stroke), sudden, spectacular symptoms, treatment sought quickly | DVT: discoloration, swelling, pain, symptoms emerge over days/weeks, treatment delayed PE: impaired heart & lung functions, shortness of breath, chest pain |
| Clot morphology   | Small amounts of soft clot in small vessels, "floating" in the vessel                | Large amounts of firm/hard clot in large vessels, adhered to vessel wall  |

Repurposed arterial thrombectomy systems

Leads to inadequate results

Typically requires use of thrombolytics

Results in inadequate safety, effectiveness and economic outcomes

**Poor Overall Results** 

**INADEQUATE** 

**TREATMENT OF** 

**VENOUS** 

**PATIENTS** 



# Inari Devices Are Specifically Designed for Venous Applications

### Penumbra Indigo System<sup>(1)</sup> Designed For:

- Arterial system
- Small, acute clot
- <3 mm diameter vessel (middle cerebral artery)</li>

#### **Stroke Treated with Indigo**

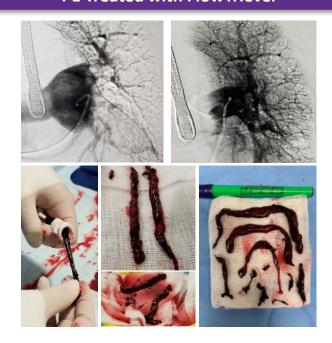


Figure 1. Occluded right MCA (A); revascularization of the MCA (B); removed thrombi (C)

### **Inari Products Designed For:**

- Venous system
- Large, acute/chronic clot
- 6-25 mm diameter vessels (pulmonary arteries)
- 6-16 mm diameter vessels (peripheral vasculature)

#### PE Treated with FlowTriever



# Inadequate Thrombectomy Options Lead to Use of Thrombolytics, An Ineffective Option for Venous Clot

### For Venous Clots, Thrombolytics Are Generally:

1 Ineffective

- Because symptoms from venous clot often appear gradually, the underlying clot can become significant in size and hardened
- Clot morphology changes over time
- The older the clot, the fewer "targets" of thrombolytics remain, which can render thrombolytic treatment ineffective

2 High Risk

- Thrombolytics can carry significant rates of bleeding complications
- Conservative patient selection and lowering dosage do not always eliminate bleeding risks
- Up to 50% of patients with venous thromboembolism ("VTE") are relatively or absolutely contraindicated to thrombolytics

3 Expensive

- Thrombolytic drugs can be highly costly
- Administration of thrombolytics requires multiple procedures and prolonged hospital stays
- Bleeding risks necessitate ICU stay (the most expensive bed in the hospital)
- Reimbursement for thrombolytics is relegated to low-paying, medically-orientated DRGs<sup>(1)</sup>

## Most Venous Clot Does Not Respond to Thrombolytics

Acute

## ClotTriever







## **FlowTriever**









# VTE: the Most Recent Example of Vascular Evolution to Catheter-Based Treatments

Development of new tools and supporting data continue to drive treatment away from thrombolytic drugs to definitive endovascular mechanical interventions

Myocardial Infarction



**Thrombolysis** 

Balloon Angioplasty, Bare Metal Stent, Drug-Eluting Stents Primary Angioplasty in Acute Myocardial Infarction (PAMI), Stent PAMI



Stroke



Systemic Thrombolysis

Stentriever, Aspiration Thrombectomy REVASCAT, MR CLEAN, EXTEND-IA, SWIFT PRIME



Expected Path for Venous Thromboembolism (DVT and PE)





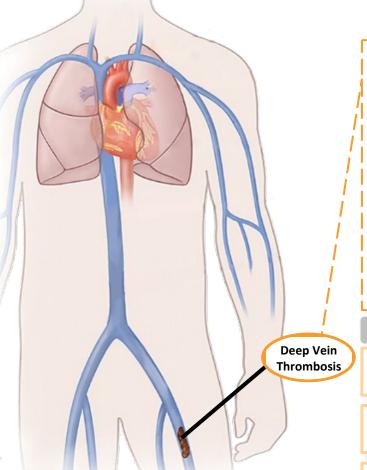
Catheter Directed
Thrombolysis,
Pharmacomechanical
Thrombolysis

ClotTriever FlowTriever FLARE, CLOUT, FLASH, FLAME





## Overview of Deep Vein Thrombosis



- Blood clots that form in the deep venous system of the legs and pelvis
- ~50% expected to develop post-thrombotic syndrome (PTS), a chronic, lifestyle-limiting disease comprising swelling, pressure, chronic pain and ulcers
- Nearly 90% of PTS patients are unable to work 10 years after diagnosis

#### **DVT Symptoms**

Swelling of the leg

Pain that may worsen when standing or walking

Warmth and redness of the leg

Pre-Op



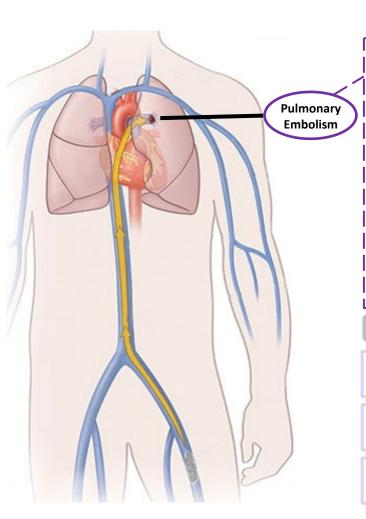
Post-Op



Removing large clot burden quickly improves acute right heart strain and we believe reduced residual clot improves longer-term outcomes



## Overview of Pulmonary Embolism



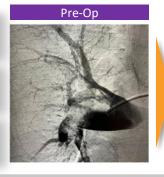
- Blood clots that break loose and travel into the lungs
- 3rd leading cause of cardiovascular death<sup>(1)</sup>; #1 cause of preventable deaths in hospitals<sup>(1)</sup>
- Short-term mortality across Massive and Sub-Massive PE: 12-50%
- Long-term complications are also potentially significant: Residual pulmonary vascular obstruction (RPVO) is common (up to 50%)

#### **PE Symptoms**

Unexplained sudden breathlessness

Sudden sharp chest pain

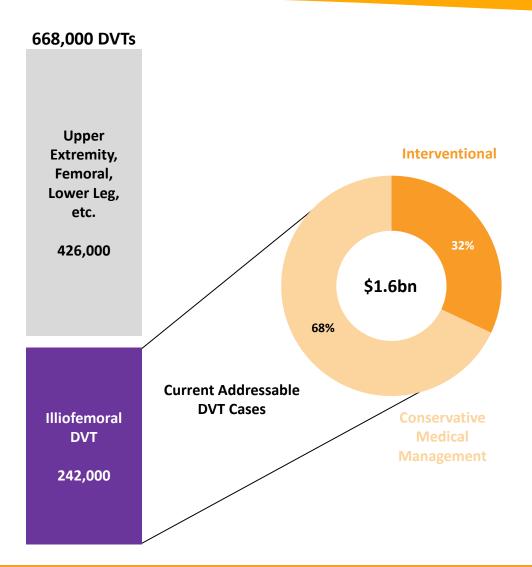
Coughing up blood





Removing large clot burden quickly improves acute right heart strain and we believe reduced residual clot improves longer-term outcomes

## DVT TAM of \$1.6Bn, Out of Combined TAM of \$3.6Bn



#### % of Market Treated Interventionally

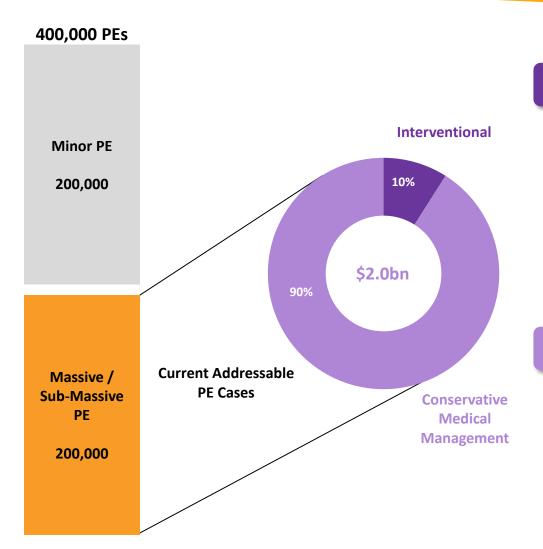
- Interventional treatment: thrombolytics and/or thrombectomy (and anticoagulation)
- ClotTriever, AngioJet (BSX), Indigo (PEN)
- 32% of DVT TAM

### % of Market Treated via Conservative Medical Management

- Conservative medical management
- Anticoagulation alone
- 68% of DVT TAM



## PE TAM of \$2.0Bn, Out of Combined TAM of \$3.6Bn



#### % of Market Treated Interventionally

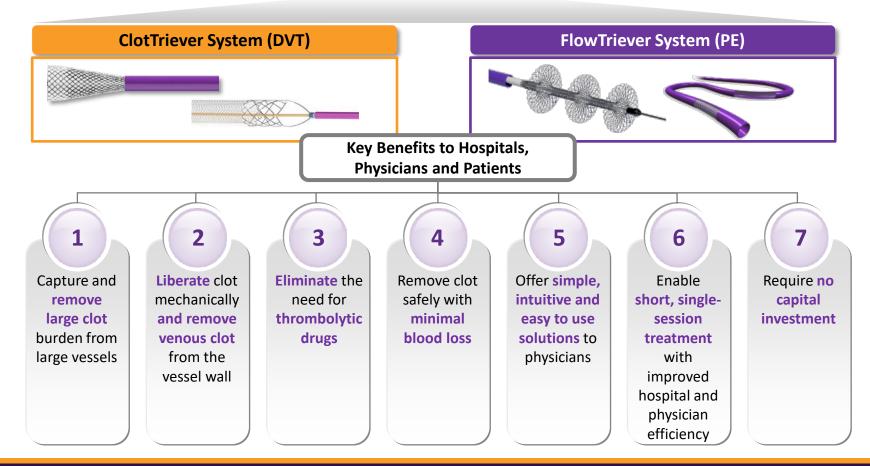
- Interventional treatment: thrombolytics and/or thrombectomy (and anticoagulation)
- FlowTriever, EKOS (BSX), Indigo (PEN)
- 10% of PE TAM

### % of Market Treated via Conservative Medical Management

- Conservative medical management
- Anticoagulation alone
- 90% of PE TAM

# Our Solutions are Designed to Offer Significant Benefits to Hospitals, Physicians and Patients







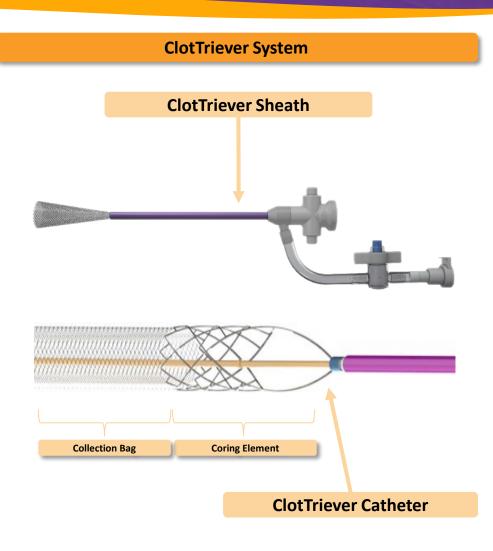
## ClotTriever System Designed Specifically to Treat DVT

#### **Product Overview**

- Designed to core, capture and remove large clots from large vessels and is used to treat DVT
- FDA-cleared for the non-surgical removal of soft thrombi and emboli from the peripheral vasculature in February 2017 and received clearance for the treatment of DVT in September 2020
- Consists of a sheath (15 cm) and catheter (74 cm)

#### **Procedure Details**

- Estimated device time: 30-45 minutes
- Complete or near complete removal of clot in 70% of patients<sup>(1)</sup>
- Estimated blood loss: 40cc (1)



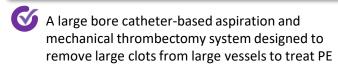
# ClotTriever Actual Case Examples: Designed For Consistent, Safe, Large Volume Clot Removal





## FlowTriever System Designed Specifically to Treat PE

#### **Product Overview**



- FDA-cleared for the non-surgical removal of thrombi and emboli from blood vessels in the peripheral vasculature in February 2015 and received clearance for the treatment of PE in May 2018
- Consists of an aspiration catheter (16, 20, 24 French sizes) and catheter (ranges from 6 to 25 mm)

#### **Procedure Details**

- Estimated device time: 46 minutes(1)
- Estimated removal of target clot: 75%
- Estimated blood loss per procedure: 250cc(1)
- Leverages per procedure pricing strategy to reduce variability and uncertainty

#### FlowTriever System



#### **Triever Aspiration Catheter**

## FlowTriever Catheter



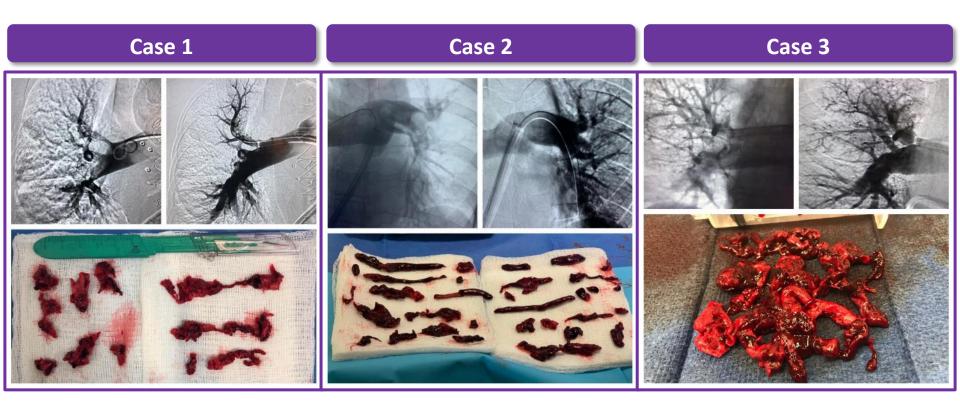
#### Available in 3 sizes

T16: 16 French lumen T20: 20 French lumen T24: 24 French lumen



Available in 4 sizes XL (19-25MM), L (15-18MM), M (11-14MM), S (6-10MM)

# FlowTriever Actual Case Examples: Designed For Consistent, Safe, Large Volume Clot Removal





## Clinical Research Investment – Real World and Broad Evidence Generation to Drive Adoption



- and chronic clot
- Core lab imaging
- **Outcomes: safety, functional** and QoL metrics
- **Utility metrics: single** session, ICU time, tPA use



- All comers, high- and intermediate-risk
- Outcomes: safety, on table hemodynamics, longer-term functional and QoL
- Utility metrics: ICU time, tPA use

## **FLASH AC Substudy:** Intermediate-Risk - PE



- Data collection to mirror FlowTriever arm with the exception of acute hemodynamics
- **Targeting Q1 rollout**

## **FLAME Registry:** High-Risk - PE



- All comer high-risk PE (FT and all standard of care options)
- Primary endpoint: mortality, bailout, clinical deterioration. and major bleeding
- **Targeting Q1 rollout**

#### **Investigator Initiated Research**

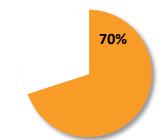
- Several IIR studies in process/under development on scientific topics of interest that do not fit within the evidence construct of our major studies
- Examples: VTE clot morphology and enzymatic activity; Cancer-related DVT, Ventilation-perfusion (RPVO) post FlowTriever, patient risk stratification, etc.



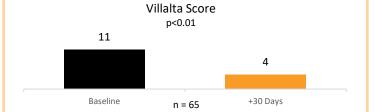
## **CLOUT Interim Results Summary**



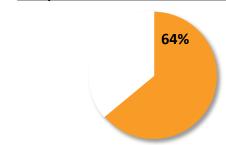
#### **Complete or Near Complete Clot Removal**

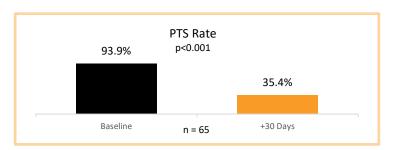






#### **Complete Reversal of PTS Within 30 Days**





### **Key Procedural Information (2)**

66%

Presented with Clot Older than 2 Weeks

27%

Previously Treated for DVT(3)

99%

Treated in a Single Session

31 Mins

Median ClotTriever Device Time

**40cc** 

Median Estimated Blood Loss

2 Days

Median Hospital Stay



## **FLASH Interim Results Summary**

## 230 Patients Enrolled at 19 US Sites (1)

93% Intermediate-risk7% High-risk

**1.6 ± 0.5** RV/LV Ratio

**96.3%** Positive RVD Biomarkers

**69.7%** Concomitant DVT

**38.3%**Contraindicated for Lytics

**Procedure Outcomes** 



**0** days ICU stay post procedure



**46** min thrombectomy time



<5% adjunctive therapy



**0.4%** Access Site Complications

On-Table Improvements



**7 mmHg** average drop in mean PA pressure



**11.8%** average improvement in cardiac index



22.7 bpm (20%) average drop in heart rate

Acute Safety (48-hrs)

0%

Mortality

0

Device-related pulmonary/cardiac injuries or procedural clinical deteriorations

1.3%

Major Adverse Events

**30-day Outcomes** 

0.4%

Mortality

6.7%

**Readmission Rate** 

## Significant Improvements in:

- Dyspnea scores
- RV/LV ratio
- RV systolic pressure
- RV systolic function
- RV dilatation

# Our Products Offer Benefits and Value to Our Hospital and Physician Customers

## **Established Coding & Payment for Mechanical Thrombectomy**

**DVT** 

PE

DRG: 270 - 272

\$17,281 - \$33,302

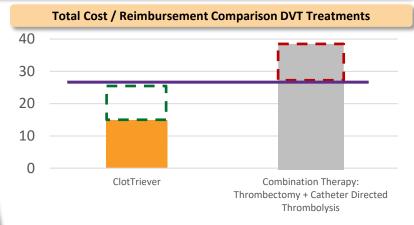
DRG: 163 – 165

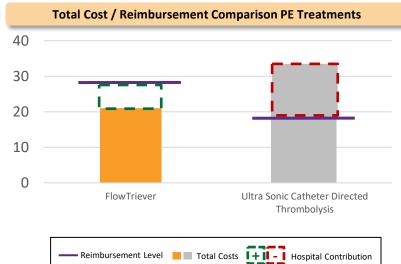
\$12,267 - \$31,875

#### Inari's Products Offer the Potential for:

- Shorter, single-session treatments
- Elimination of thrombolytic drugs
- Reduction of ICU stays
- Shortening total hospital stay
- More efficiency in hospital and physician workflows

### Illustrative Procedural Hospital Contributions<sup>(1)</sup>

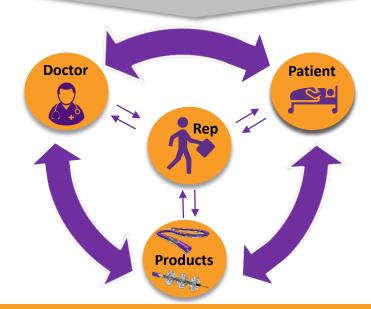




## Meaningful Investment in Our Commercial Organization

- Wide and deep
- Systems and processes to support rapid expansion
- High touch, effective interventional call points
- Refined and established hiring and training process designed to enable rapid sales rep productivity ramp and increased profitability

- Inari sales representatives are typically present in >80% of all cases<sup>(1)</sup>
- Rich information is generated when patient, physician, and product come together
- Field based information is the primary input into product development and clinical and commercial strategies
- No plans for a bifurcated sales model e.g. clinical specialists
- Our goal is to be a market-driven company

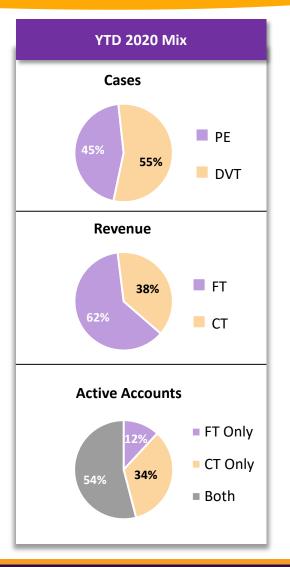


## **Financial Results**



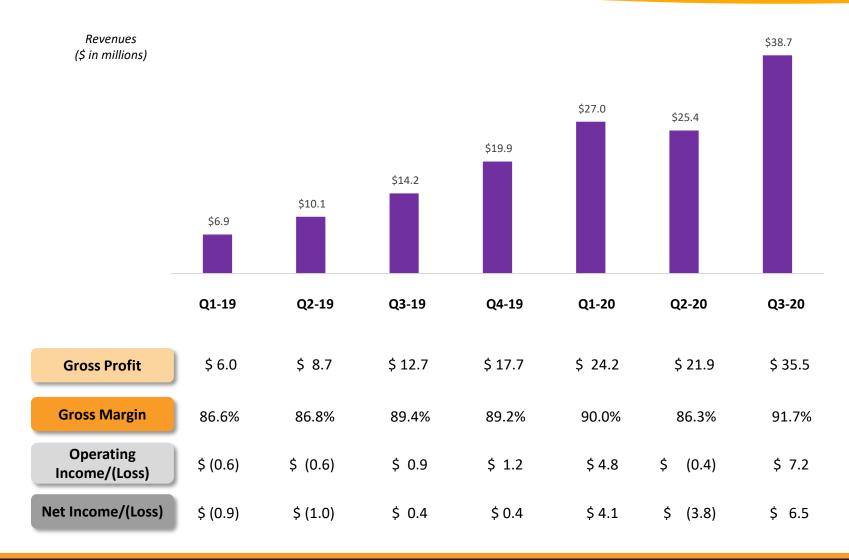
## Despite COVID, Q3 Cases Rebound and Regain Much of Pre-COVID Growth







## Financial Performance Through Q3 2020





## Our Customers and Team Are Better Prepared to Manage C19 Impacts Going Forward

### Clinical "Supply"



We have seen and continue to expect hospitals will prioritize procedures based upon:

- Acuity: Inari procedures can warrant clinical priority
- Safety and efficiency of care pathway: VTE thrombectomy has modest interventional "footprint" (no intubation, elimination of nearly all ICU stays, short LoS)
- Economics: Favorable procedural economics can help hospitals recover financially

### Clinical "Demand"



- As acute phase passed, patient fears have subsided, and we believe patients will be more likely to seek care for high acuity conditions
- Potential "backlog" of deferred VTE patients can be treated: anticoagulation only often defers intervention
- COVID is risk factor for VTE

#### Commercial



- Further developed our leading position in VTE
- Adapted, expanded and improved sales training and customer engagement
- Enhanced our physician outreach and training

## Summary



## Inari's Growth Drivers

- ✓ Continuing to expand our U.S. sales force
- Driving increased awareness and adoption of our products in existing and future hospital customers
- ✓ ) Building upon our base of clinical evidence
- ✓ Continuing to expand our portfolio of venous products
- Pursuing strategically adjacent markets and international opportunities

