

Company Presentation

Australian Non-deal roadshow and Bell Potter Healthcare Conference

PERTH, Australia, 9 March 2026: Artrya Limited (ASX: AYA) (**Artrya** or the **Company**), a medical technology company commercialising its Salix[®] AI-powered cloud platform, for the near real time, point of care assessment and management of coronary artery disease, is pleased to announce that Co-Founder and Chief Executive Officer, John Konstantopoulos, and Executive Chair, Bernie Ridgeway, will be conducting a non-deal roadshow this week, meeting with existing and prospective investors and research analysts in Sydney and Melbourne.

The Company's Presentation for this roadshow is attached.

Additionally, Co-Founder and Chief Executive Officer, John Konstantopoulos, will attend the Bell Potter Healthcare Conference in Sorrento, Victoria on 12 March 2026, where he will participate in an analyst interview and meet with investors.

The Bell Potter Healthcare Conference brings together companies and investors to showcase leading small and mid-cap healthcare companies from across Australasia.

- Ends -

This ASX Announcement is authorised for release by the Board of Artrya Limited.

About Artrya

Artrya Limited (ASX:AYA) is an Australian medical technology company developing AI-powered solutions to improve the detection and management of coronary artery disease. Its proprietary software analyses coronary CT scans to identify key biomarkers of heart disease, supporting clinicians in diagnosing patients more accurately and efficiently. Artrya's mission is to advance cardiac care through innovative technology, with regulatory and commercial activities underway across key international markets.

For more information visit www.artrya.com or follow us on LinkedIn at www.linkedin.com/company/artrya

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ARTRYA[®]

(ASX: AYA)

Company Presentation

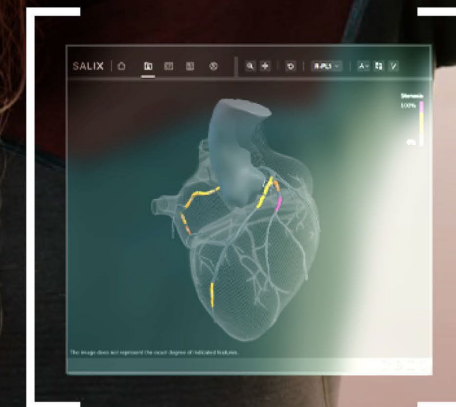
Advancing U.S. Commercialisation

09 March 2026

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Reshaping
the Future of
Heart Care



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BY THE BOARD OF ARTRYA LIMITED THIS PRESENTATION HAS BEEN APPROVED FOR RELEASE

Artrya at a glance – an attractive market and model

Salix® is the only real-time, point of care platform globally improving patient care and streamlining workflow

Attractive Market



Commercial Momentum

- Salix® Platform & Plaque FDA cleared
- Three U.S. commercial customers and growing pipeline
- U.S. Customer Success team in place
- SAPPHIRE Study with major centres
- Strong balance sheet - \$76.5m cash³
- A\$535m market cap (ASX: AYA)²

Differentiated Product

- Single, unified CCTA Reporting Platform within 10 mins
- Real-time report, Plaque, and FFRCT⁴ analysis
- Streamlines clinical workflow and improves patient care

Leading Institutional Investors:

REGAL
FUNDS MANAGEMENT

Wilson
Asset Management

PARADICE
INVESTMENT MANAGEMENT

1. Company Presentation – slide 7 – Released to ASX on 1 September 2025
2. As of 06 March 2026
3. Pro forma cash including cash and term deposits As at 31 December 2025
4. Salix® Coronary Flow not yet cleared by FDA

Recent Achievements

FY2026 marks commercial scale-up following the platform laid in 2025

FY2026 Execution & Growth	
 FDA activity	FDA clearance for Salix® Coronary Plaque Successful FDA Q-Sub meeting for Salix® Coronary Flow Anticipated FDA clearance of Salix® Coronary Flow by 30 June 2026
 Commercial progress	Three U.S. commercial customers launched - Tanner Health, Cone Health, and Northeast Georgia Further commercial pipeline through SAPPHIRE participants
 U.S. Go to market	General Manager U.S. hired U.S. Customer Success team established Additional U.S. Non-Executive Director appointed
 SAPPHIRE Study	Secured Study participants - Mass General Brigham, Piedmont Health, Huntsville Hospital, Dignity Health, HCA Midwest
 Capital management	\$80M raised to fund U.S. expansion strategy

Coronary artery disease is a large and growing health challenge

Cardiovascular Disease Costs will exceed US\$1 trillion by 2035 – American Heart Association⁴

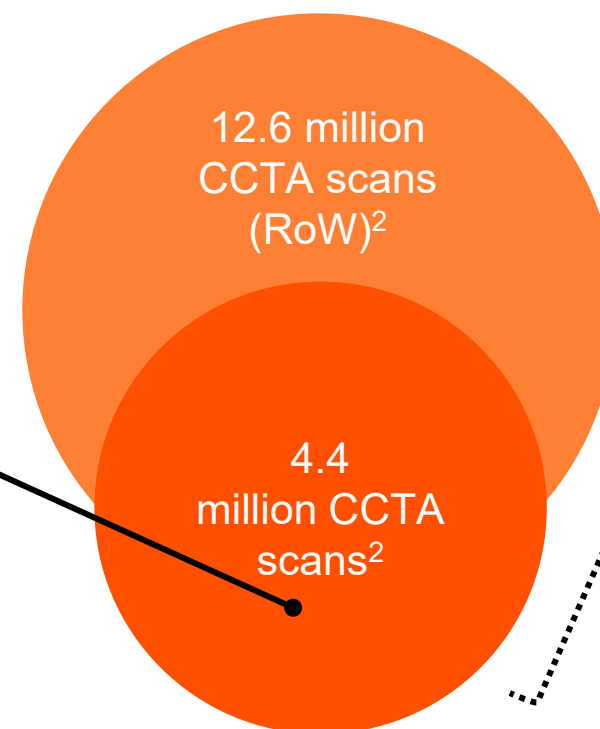
US\$4.4 billion*

Serviceable Addressable Market (US)

U.S. patients undergoing CCTA + Plaque Assessment

Tailwinds impacting TAM growth

- **Guideline Support** – AHA/ACC recommend CCTA as a first-line diagnostic for patients with chest pain⁵.
- **Attractive reimbursement – Category I CPT codes** support CCTA, plaque assessment, and FFRCT³.
- **CT-First Adoption** – Strong shift from invasive angiography and SPECT toward **non-invasive CT-based pathways**.



6.2% YOY

Projected growth in Cardiac CCTA scans to 2028¹

* Assumptions

- National Average Medicare Rate
 - Reimbursement for Automated Plaque – US\$950 (CPT 0625T³)
 - Reimbursement for FFRCT – US\$887 (CPT 75580³)
- TAM estimated as total of all scans receiving Plaque reimbursement only
- FFRCT procedures have not been included in TAM

1 <https://www.dicardiology.com/article/rising-demand-cardiac-ct-positions-market-major-growth>
2 Frost & Sullivan Analysis – Artrya Prospectus – <https://wcsecure.weblink.com.au/pdf/AYA/02456983.pdf>
3 <https://cardiovascularbusiness.com/topics/cardiac-imaging/cms-increases-medicare-payments-cardiac-ct-ccta>
4 Cardiovascular Disease: A Costly Burden for America – Projections Through 2035. American Heart Association.
5 American College of Cardiology – 2021 AHA/ACC Chest Pain Guideline Perspectives

Current methods for diagnosing coronary artery disease are outdated

A point of care solution for rapidly diagnosing high-risk plaque is a holy grail for clinicians



Ineffective

Old methods ineffective in **detecting high-risk plaque**¹



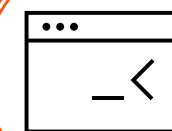
Avoidable

55% of invasive procedures are unnecessary^{2,3,4}



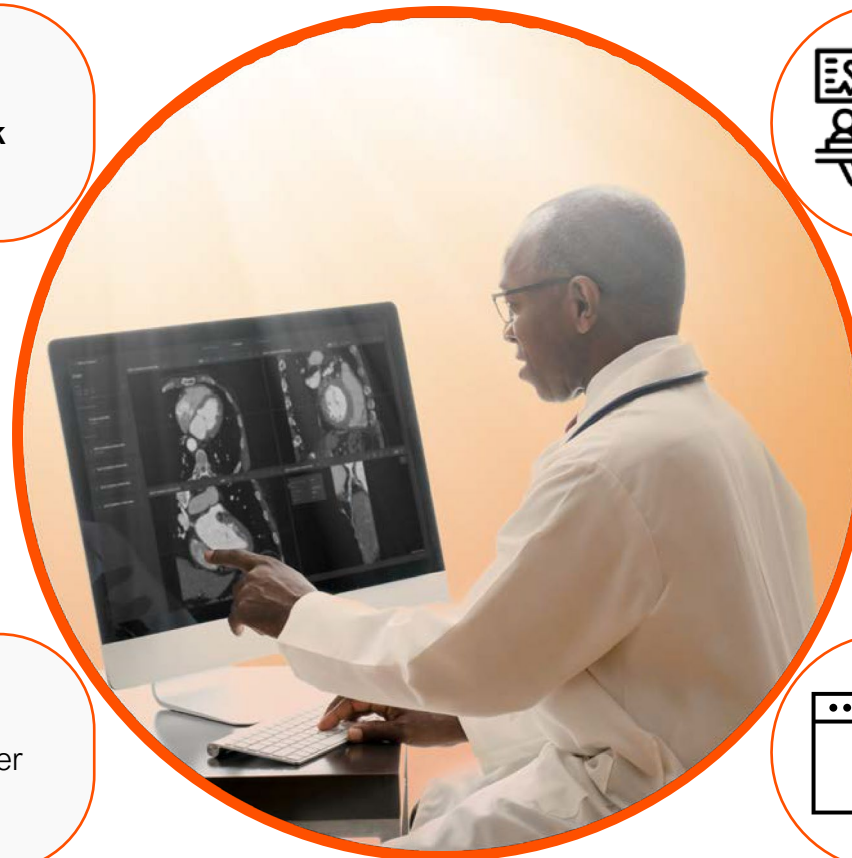
Slow & Costly

Outdated methods are time consuming and higher cost to deliver



Pressure on Hospitals

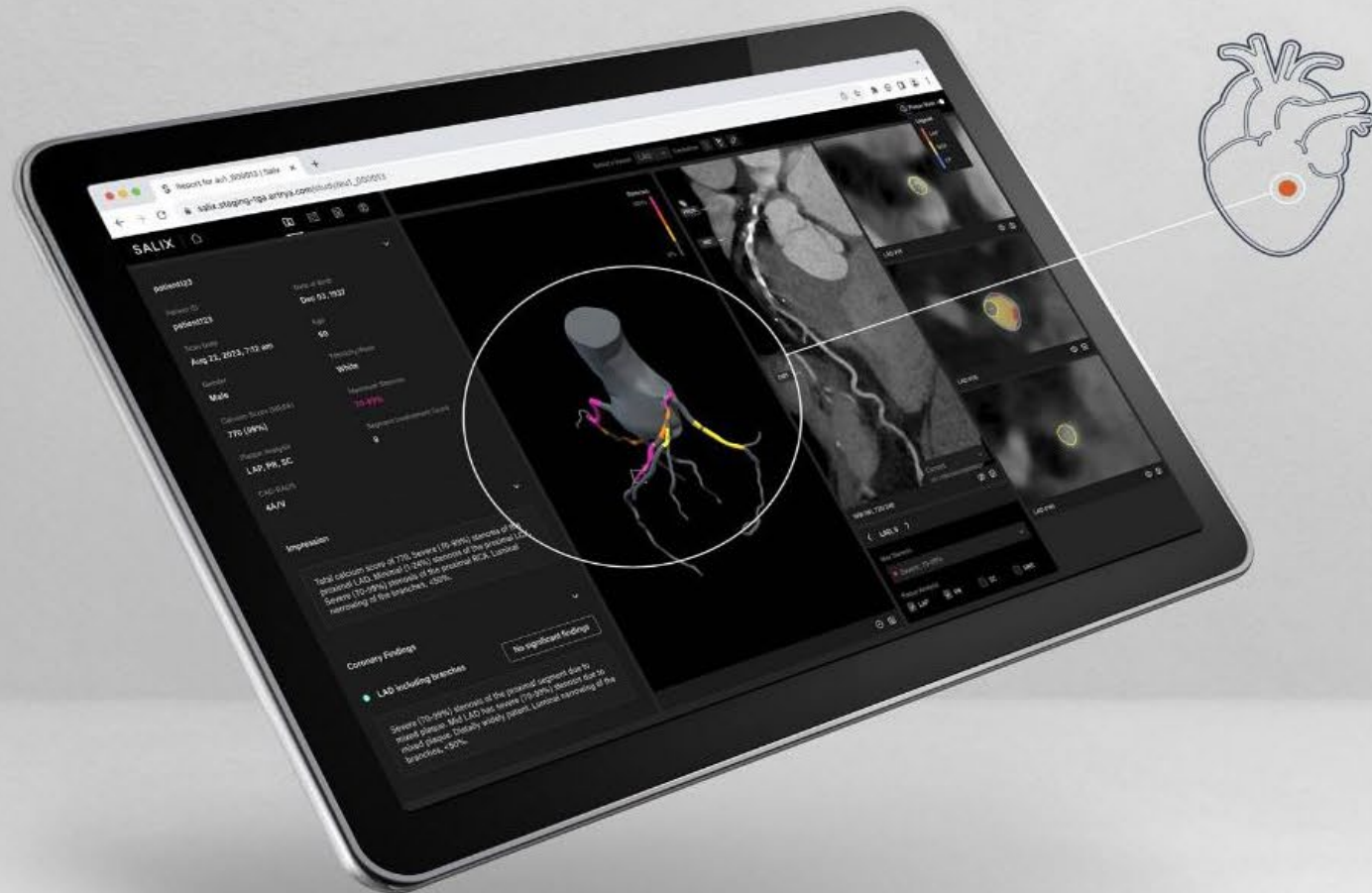
CCTA **demand is outpacing** clinical capacity
Hospitals **need higher throughput**
Current solutions **delay care / increase backlogs**



1. Comprehensive plaque assessment by coronary CT angiography. Nat Rev Cardiol 11, 390–402 (2014)
2. Low Diagnostic Yield of Elective Coronary Angiography. New England Journal of Medicine. March 11, 2010.
3. Temporal Trends in the Frequency of Inducible Myocardial Ischemia During Cardiac Stress Testing. Journal of the American College of Cardiology (JACC). March 12, 2013.
4. Trends in U.S. Cardiovascular Care: 2016 Report from 4 ACC National Cardiovascular Data Registries. JACC. March 2017

Salix[®] provides streamlined, real-time patient assessment

Clear clinical insights into actionable decisions in minutes



Point of care real-time analysis – no human in the loop

CCTA analysis in under 10 mins increases capacity and throughput

Real-time Plaque and Flow Analysis

Automated assessment of plaque and FFRCT for improved treatment

Unified Reporting Platform

Consolidates CCTA reporting on platform, reduces physician fatigue and reporting costs by consolidating tools

Streamlined Workflow and Enhanced profitability

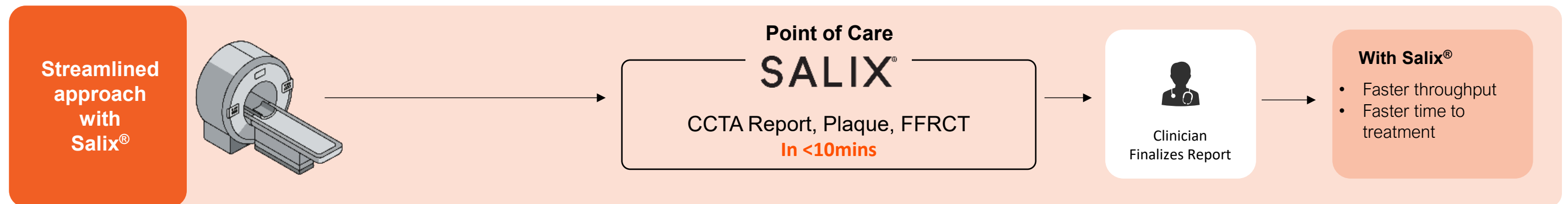
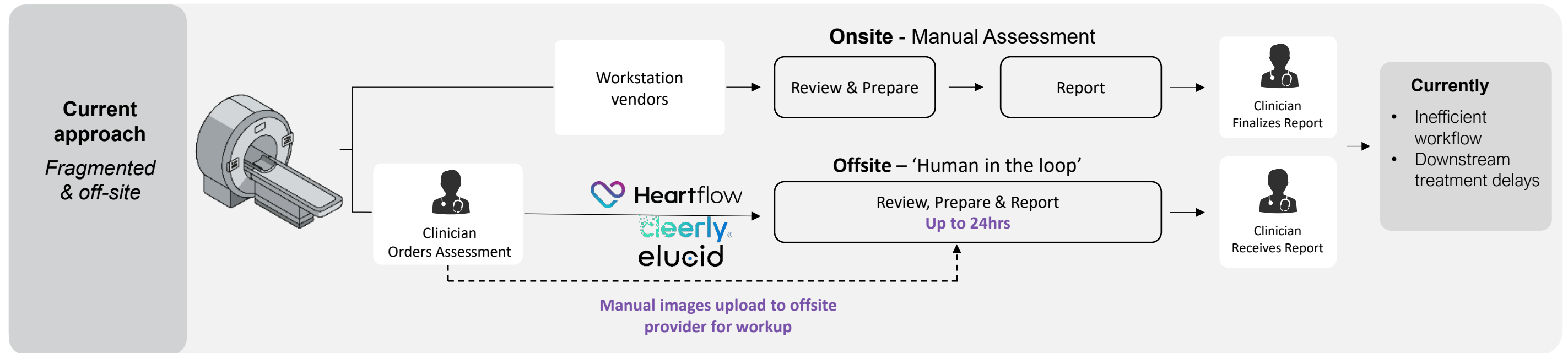
Improved efficiency, transforms CCTA into a scalable, high-margin service line

* Salix Coronary Flow not yet cleared by FDA

ARTRYA[®]

From off-site workflows to 10-minute decision

Salix[®] fits directly into the existing clinical workflow; no uploads, no portals, no off-site vendors





Salix[®] demonstration

Go to Market strategy

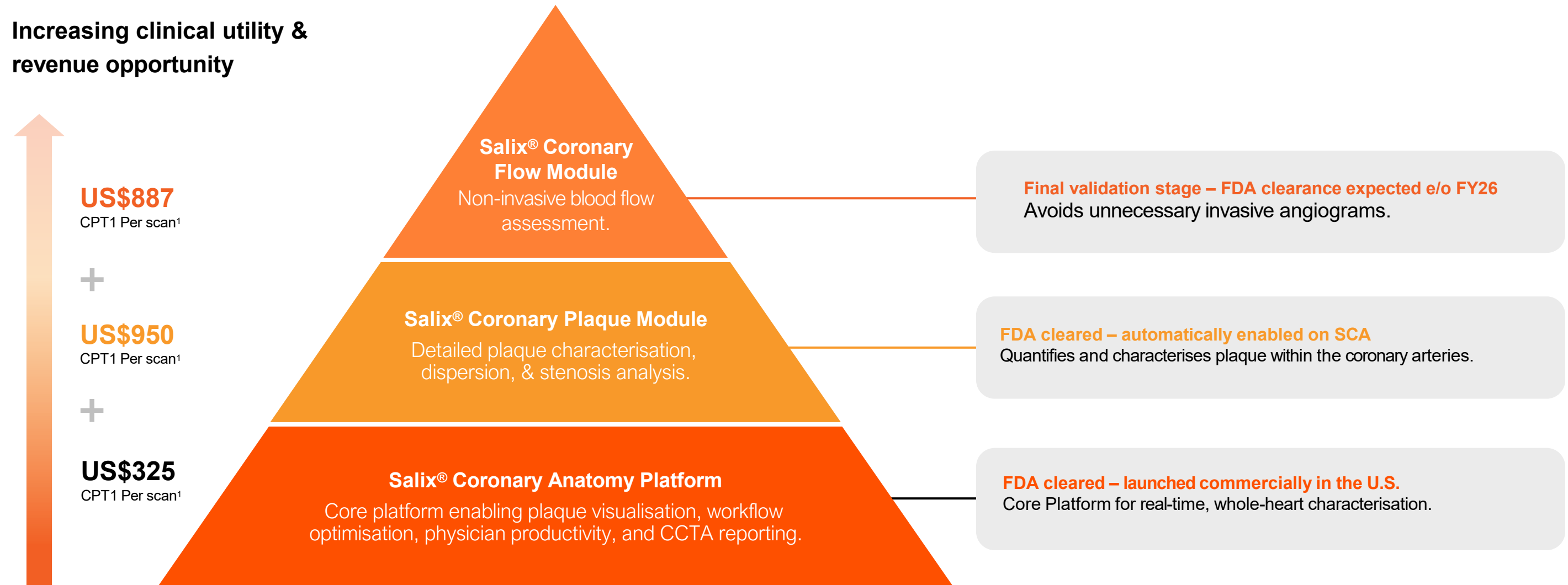
Commercial pathway for 2026 and beyond



Salix[®] is a single platform solution built to scale

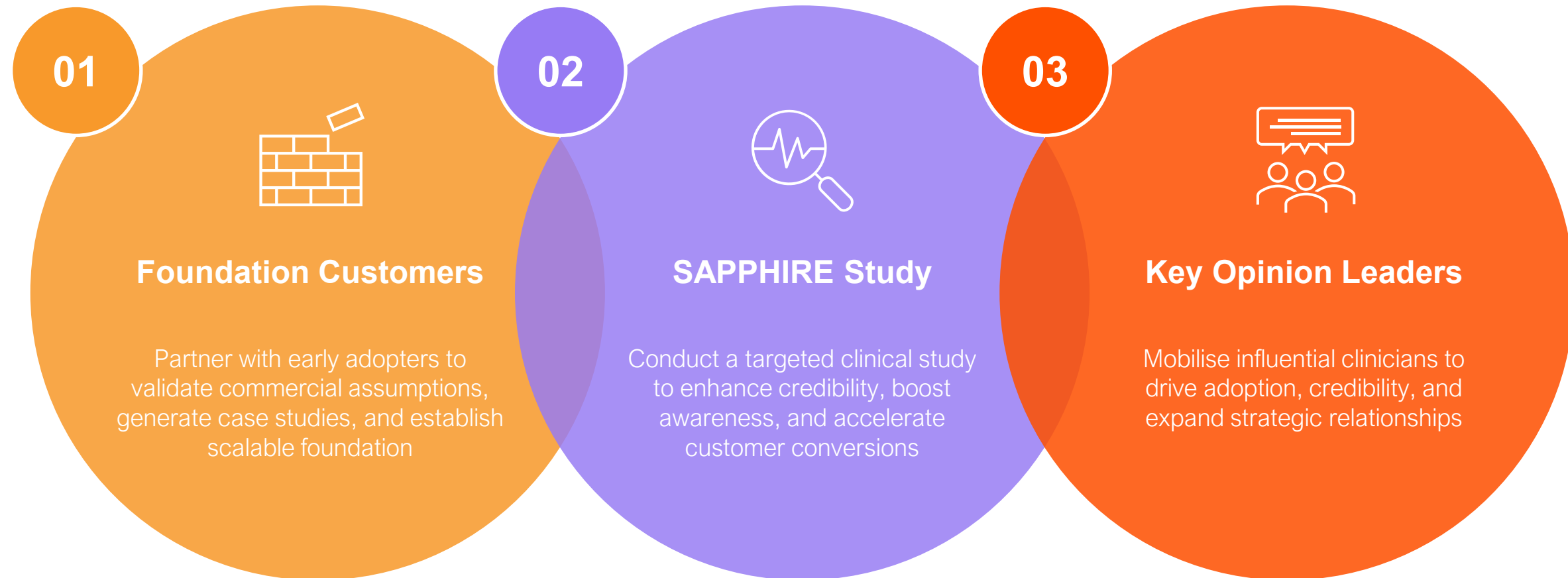
Modular software powering complete, scalable CCTA analysis for coronary artery disease

Increasing clinical utility & revenue opportunity



From strategy to scale: executing with precision

Delivering focused execution through three clear go-to-market strategies



Salix® platform successfully launched in three U.S. systems

All three U.S. integration partners have been converted to commercial customers



Live from July 2025

- ▶ 5-year Commercial Agreement
- ▶ Minimum US\$0.6m *
- ▶ 5 Hospitals, cardiovascular centres and 30 practices

"...we believe Artrya's Salix® platform with its AI-powered, user friendly and near real-time outcomes, provides our clinicians with another key tool to help detect heart disease earlier and save lives."

- Loy Howard, President and CEO of Tanner Health



Contract signed December 2025

- ▶ 3-year Commercial Agreement
- ▶ Minimum US\$0.3m *
- ▶ 5 Hospitals, outpatient centres, and physician practices

"This latest A.I.-enabled technology reflects our commitment to advancing cardiovascular medicine and ensures patients across Georgia Heart Institute can benefit from faster, more accurate diagnoses and improved outcomes."

- Zaid Said, MD, FACC, Medical Director of Cardiac CT and MRI for Georgia Heart Institute



Contract signed December 2025

- ▶ 5-year Commercial Agreement
- ▶ Minimum US\$0.45m *
- ▶ 5 hospitals, ambulatory care centres, outpatient surgery centres, urgent care centres, >120 physician practices

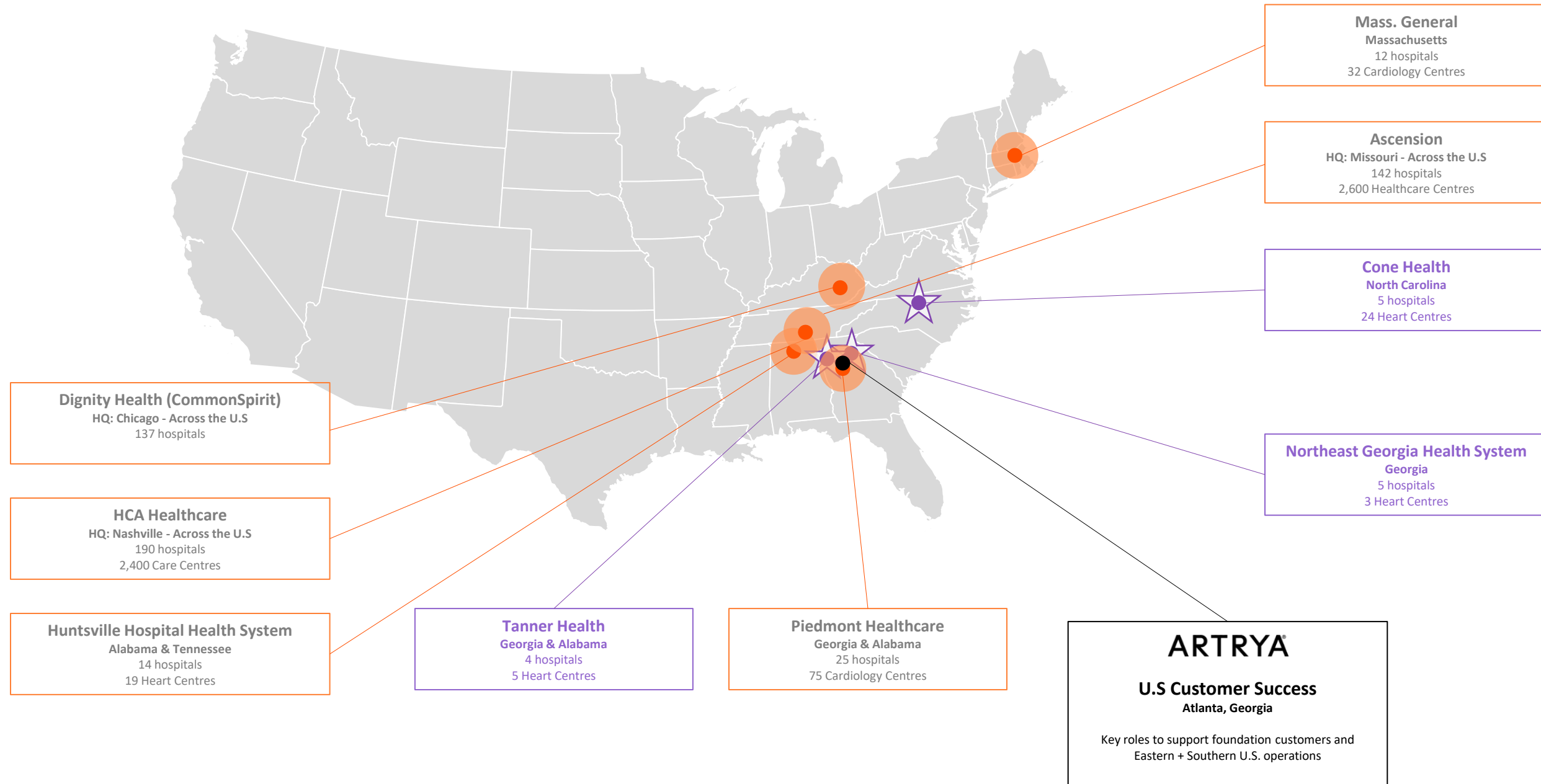
We are delighted to build on our successful collaboration with Artrya and bring this transformative Salix® technology into the clinical workflow across our network. Through this process we can see major benefits in the way that Salix® can provide accurate, point-of-care interpretation of CCTA scans within minutes.

- Dr. Wesley O'Neal, MD, Medical Director of Cardiac CT and Nuclear Cardiology at Cone Health

* Minimum contract values relate solely to the monthly subscription of Salix® Coronary Anatomy, with additional fee-per-scan revenues from the Salix® Plaque and Flow (once approved) modules

Building a U.S. footprint to support leading health systems

Foundation customers, SAPPHIRE sites, and our Atlanta support pod working together to drive U.S. growth



Expanding clinical utility with Salix[®] Coronary Flow module

Expands use and utility of Salix[®] platform with access to U.S. reimbursement once FDA cleared

01

Robust FDA submission developed

Q-Sub meeting confirms approach and requirements

02

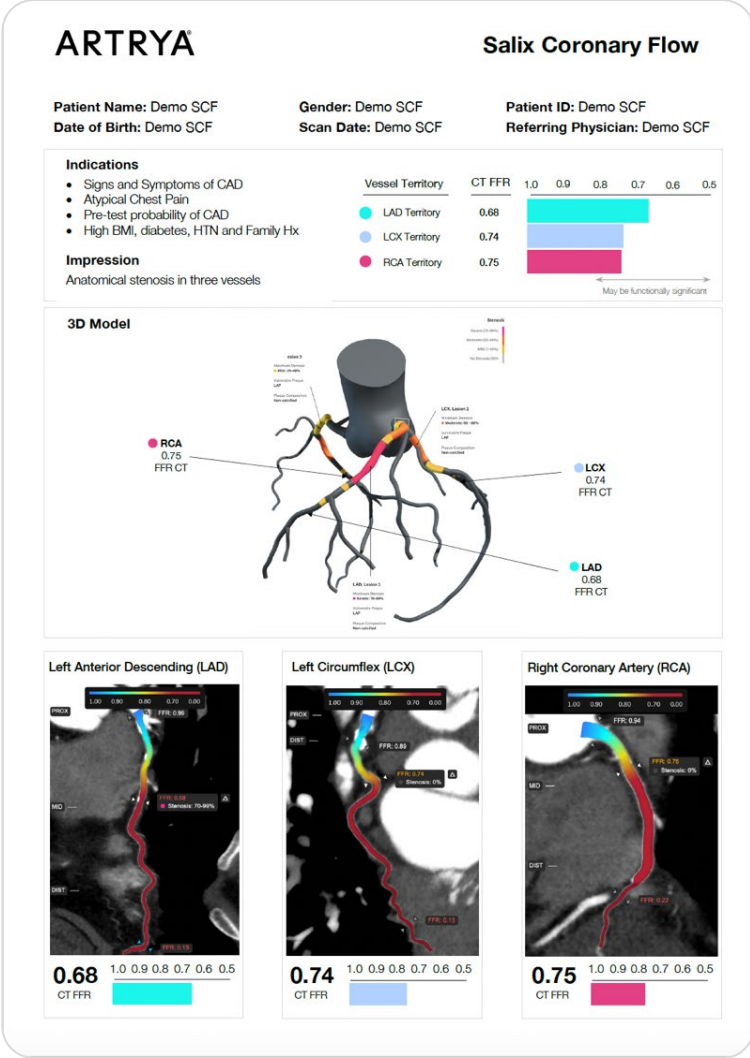
Final module calibration completed early 2026

Sourced high-quality imaging for additional module analysis – sensitivity and specificity meet and exceed FDA targets

03

Planned FDA clearance by end of FY2026

Timing is estimated and subject to FDA process



* Salix Coronary Flow not yet cleared by FDA

Flagship Novel Plaque Study - SAPPHIRE

The SAPPHIRE Study will validate the novel Salix® Plaque Dispersion Score to identify patients at risk of heart attack

Overview

- Evaluate Salix® PDS assessing cardiovascular risk
- Specific focus on CAD in women.
- 6 high volume U.S. Centres
- Multicentre, retrospective study
- **HCA, Dignity Health, Mass General, Ascension, Piedmont Healthcare, and Huntsville** as participants

Key Phases

Phase 1: Identify patients at risk using Salix® Plaque analysis compared to current standard of care. ~ 12 months

Phase 2: Leverage Salix® Plaque Dispersion Score to further risk stratify patients and change treatment. ~ 12 months

Key Benefits

01

Predict future adverse events and improve treatment

02

Build clinical awareness and credibility in US

03

Accelerate commercial adoption in US hospitals

Summary and Outlook for 2026: Scaling Salix® in the U.S.

Building the foundation for sustainable commercial growth through revenue, clinical validation, and operational readiness.



Drive Commercial Growth

- 1 Generating annual revenue** from SCA, SCP and SCF from U.S. operations (SAPPHIRE conversions & foundation customers).
- 2 3 foundation customers live** and operational



Clinical & Operational Excellence

- 3 6 hospital systems active** in the SAPPHIRE Study; Initial Phase 1 data analysed.
- 4 SCF cleared through FDA.**
- 5 CX & Support framework implemented**



Build for Scale & Credibility

- 6 Full U.S. organisation & governance operational** (core recruits, executives, Board/CAB).
- 7 Clinical credibility** through papers published and conferences presented.

Note: 2026 Calendar Year

Building a Sustainable Competitive Advantage

Competitive advantage built on product strength, pricing flexibility, and customer experience

Our strategy is built on three reinforcing pillars:

► **Product Experience: Real-time, platform infrastructure**

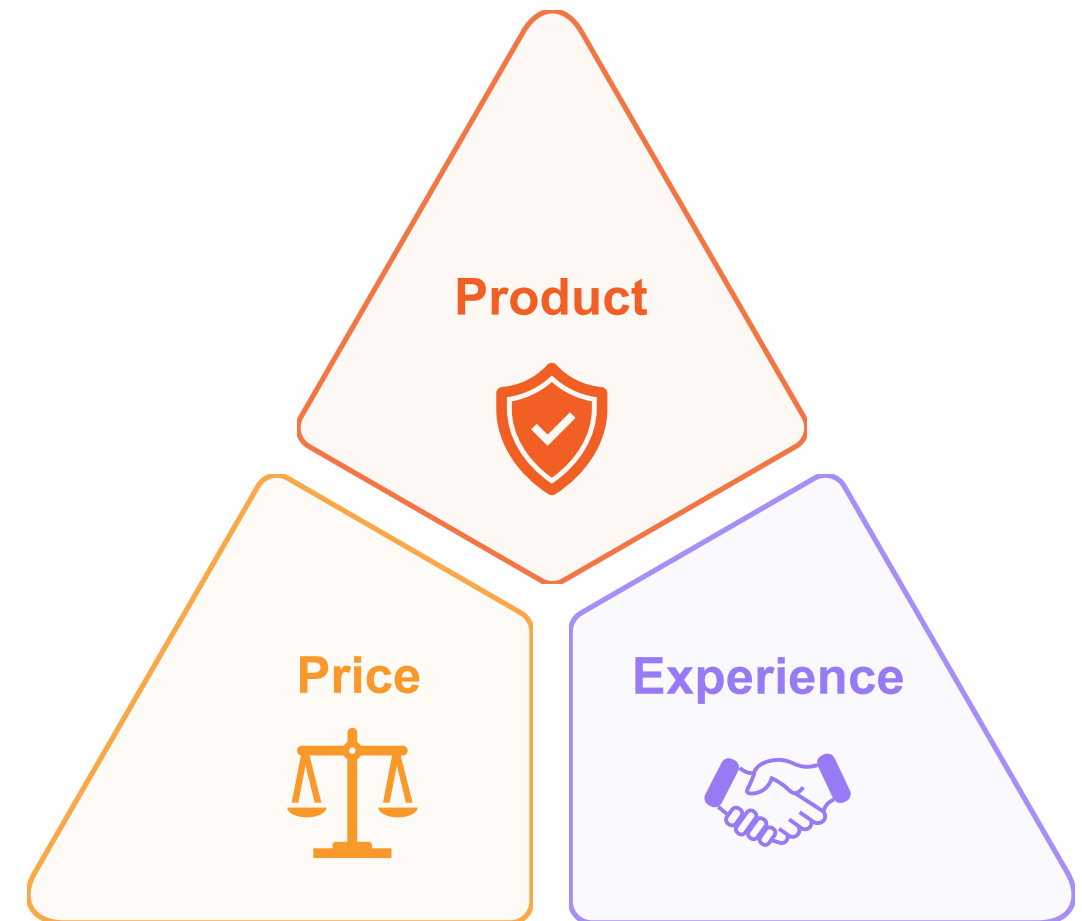
We deliver the strongest product in market, combining real-time diagnosis with integrated workflows and a user experience designed for effortless adoption.

► **Price: Adoption-aligned economics**

Our pricing is competitive and differentiated through a revenue share model that reduces customer risk and directly aligns with our mutual success.

► **Customer Experience: Growth & retention engine**

We provide an easy experience at every stage of the journey; from onboarding to support to reducing friction and making every interaction simple and reliable.



Superior product. Differentiated value. Exceptional customer experience.
Saving Lives

www.artrya.com

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