Bell Potter Healthcare Conference 2025

PERTH, Australia, 20 November 2025: 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 John Konstantopoulos, Co-Founder and CEO will provide a presentation on the Company to investors at the virtual Bell Potter Healthcare Conference on 20 November 2025.

This forum will bring together leading industry thought leaders and the investment community and will be showcasing the best of Australasia's small and mid-cap healthcare offering.

Bell Potter Securities is a leading Australian stockbroking, investment and financial advisory firm that provides a comprehensive offering of financial services to individuals, institutions and corporations.

A copy of the Presentation is attached.

- 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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For more information:

Corporate Enquiries

Investor & Media Enquiries

John Konstantopoulos
Co-Founder and Chief Executive Officer
Tel: +61 8 6478 7816
Email: contact@artrya.com

David Allen or John Granger
Hawkesbury Partners
Tel: +61 499 100 038 or +61 410 577 155
Email: investors@artrya.com

ARTRYA®

(ASX: AYA)

Bell Potter Healthcare Conference

Company Presentation

20 November 2025

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Artrya at a Glance – an attractive market and model

Salix® is the only real-time, point of care platform globally delivering end-to-end CCTA assessment

Attractive Market

$4.4Bn[1]

Market Size

> 6%

YoY Growth

Commercial Momentum

- (ASX: AYA) - ~AU$550m market cap
- Salix® Platform & Plaque FDA cleared
- Targeted U.S. GTM
- Foundation customers going live, generating revenue, growing pipeline
- Strong balance sheet

Differentiated Product

- Unified CCTA Reporting Platform within 10 mins
- Real-time report, Plaque, and FFRCT[3] analysis
- Streamlines workflow and care

Backed by 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 14 November 2025
3. Salix Coronary Flow not yet cleared by FDA

Current methods for diagnosing coronary artery disease are outdated

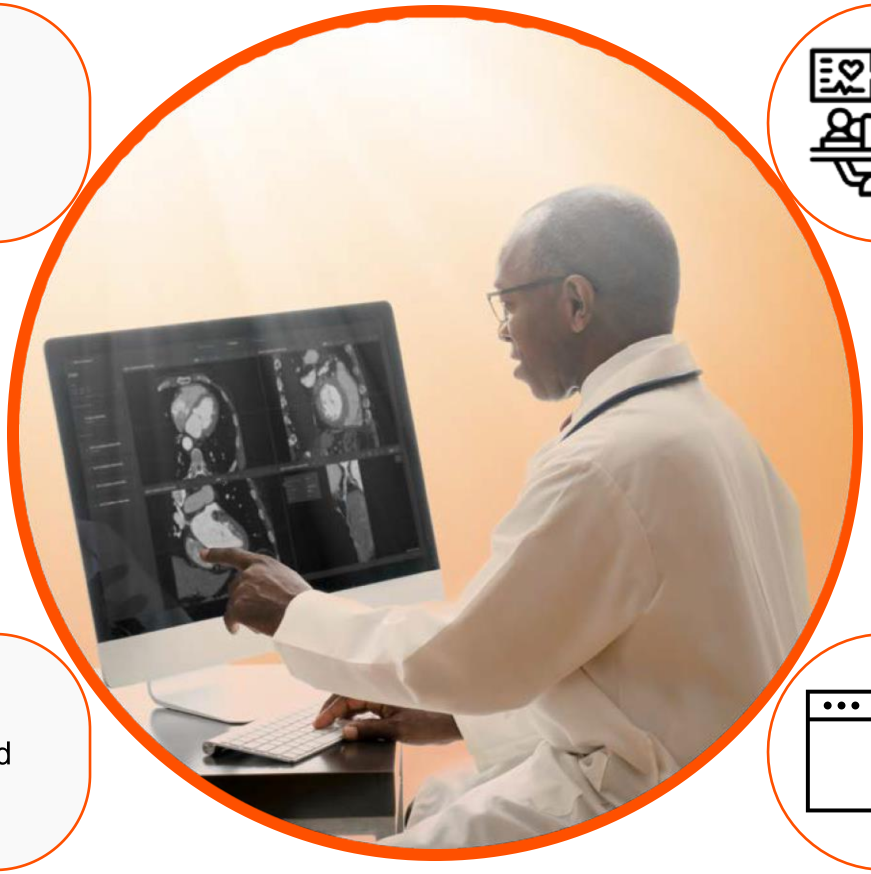
A point of care solution for rapidly diagnosing high risk plaque is a holy grail for clinicians

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

Poor Control

Current solutions do not allow for **real-time and point-of-care** management

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

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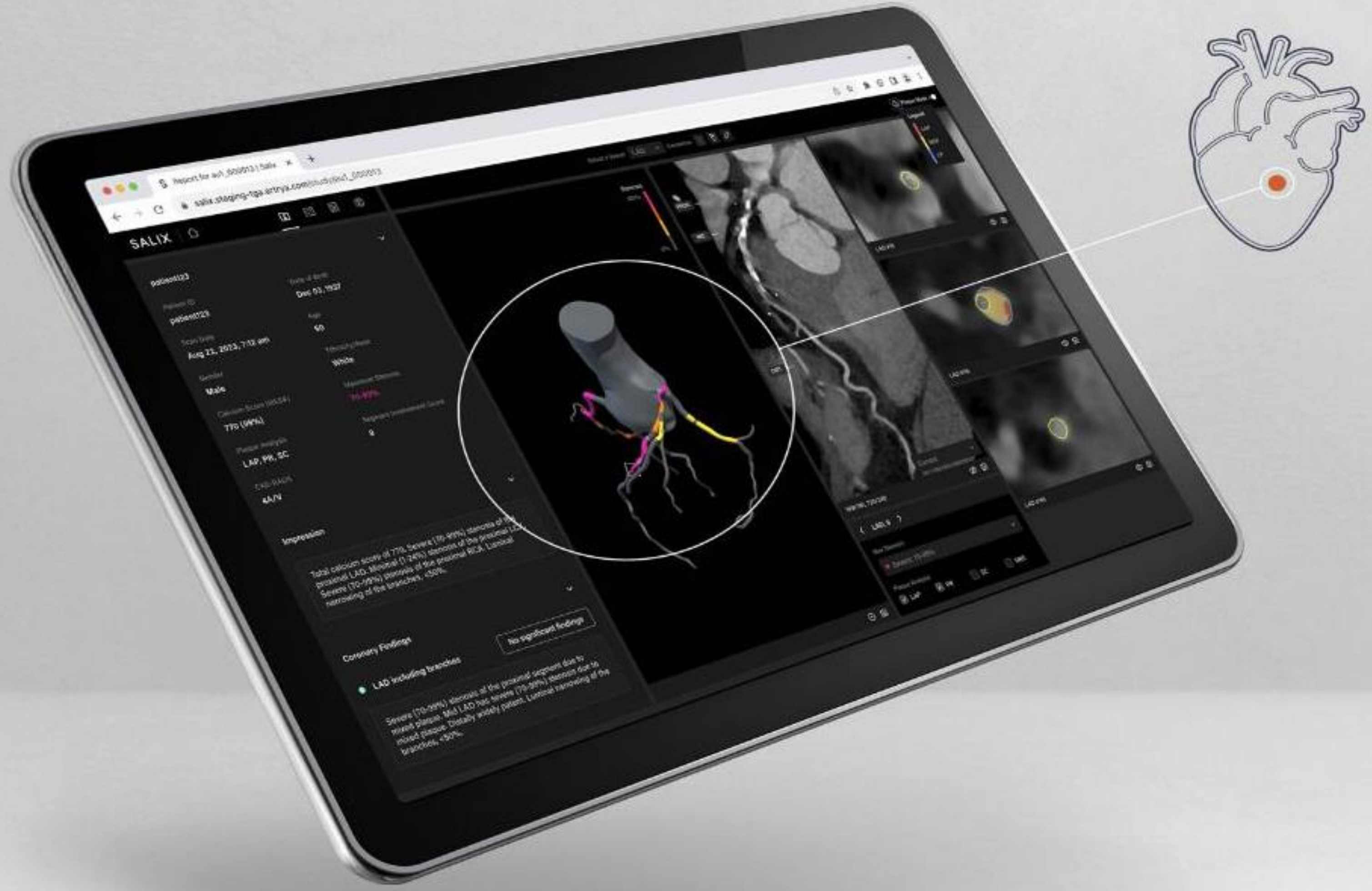
Salix® provides streamlined, real-time patient assessment

Clear clinical insights into actionable decisions in minutes



Point of care enabling real-time analysis – without humans

CCTA analysis in under 10 mins without human intervention.

Detailed Plaque and Flow Analysis

Real-time assessment of plaque and FFRCT* for improved treatment.

Unified Reporting Platform

Consolidates all CCTA reporting into a single platform.

Enhanced profitability for health systems

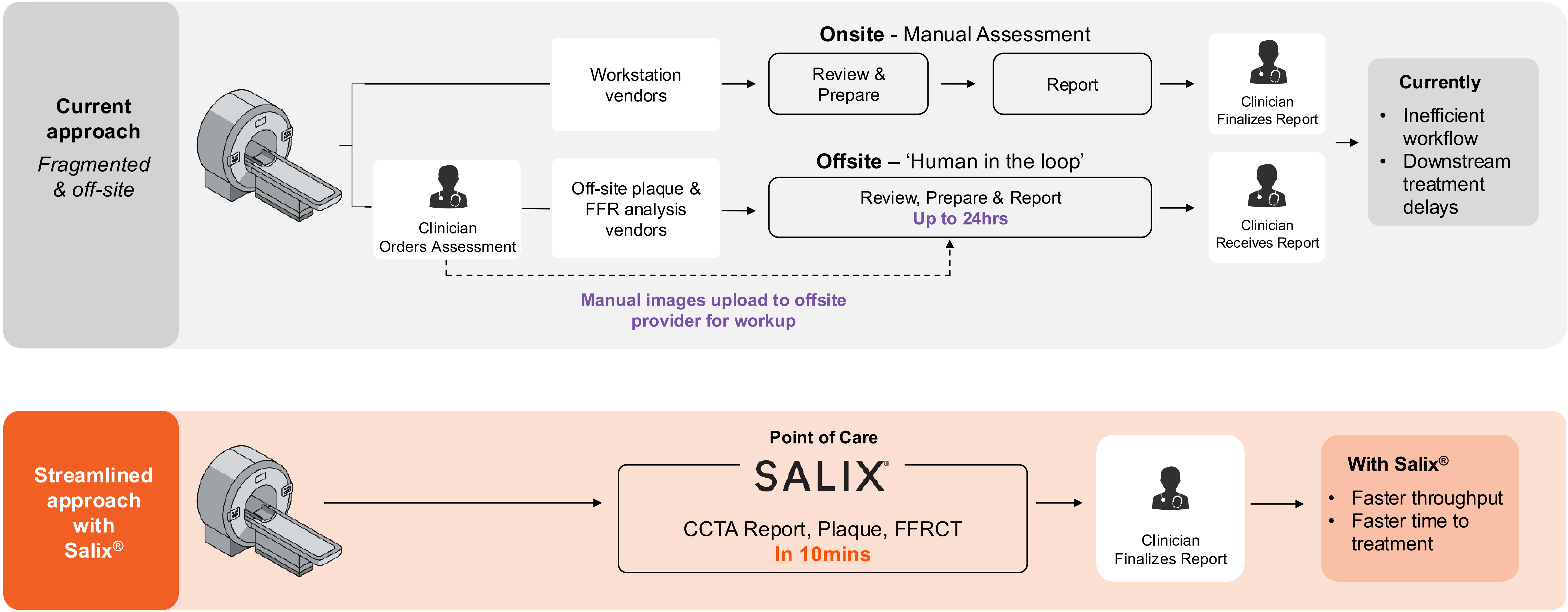
Transforms CCTA into a profitable modality and reduces costs by replacing multiple tools and vendors

* 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



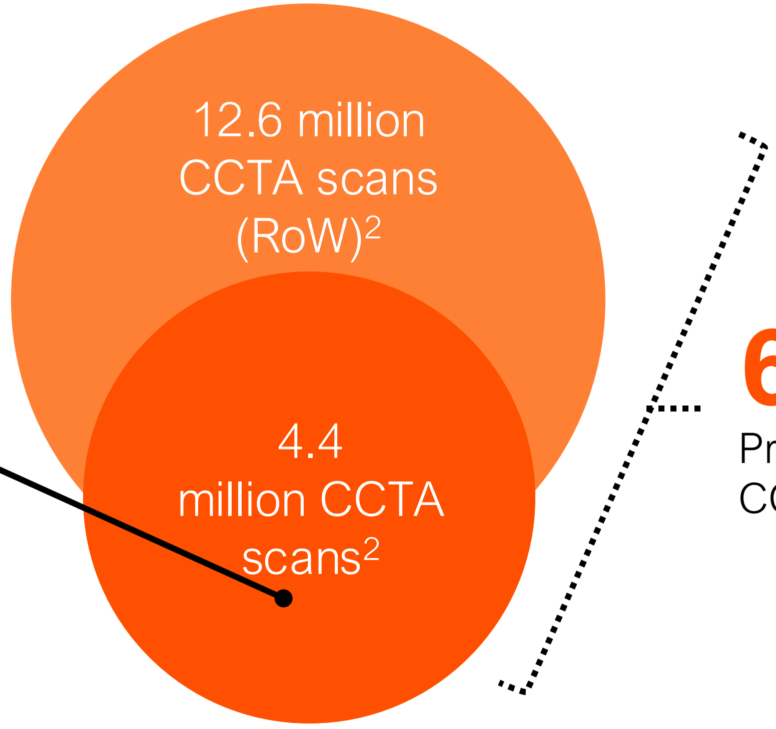
Coronary artery disease is a large and growing health challenge

Cardiovascular Disease Costs will Exceed $1 Trillion by 2035 - American Heart Association[4]

US$4.4 billion*

Serviceable Addressable Market (US)

U.S. patients undergoing CCTA + Plaque Assessment



6.2% YOY

Projected growth in Cardiac CCTA scans to 2028[1]

Tailwinds impacting TAM growth

- **Guideline Support** – AHA/ACC recommend CCTA as a first-line diagnostic for patients with chest pain[5].
- **Attractive reimbursement** – **Category I CPT codes** support CCTA, plaque assessment, and FFRCT[3].
- **CT-First Adoption** – Strong shift from invasive angiography and SPECT toward **non-invasive CT-based pathways**.

*** Assumptions**

- National Average Medicare Rate
  - Reimbursement for Automated Plaque – US$950 (CPT 0625T[3])
  - Reimbursement for FFRCT – US$1,017 (CPT 75580[3])
- 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

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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

US$1,017
CPT1 Per scan[1]

+

US$950
CPT1 Per scan[1]

+

US$325
CPT1 Per scan[1]

Salix® Coronary Flow Module
Non-invasive blood flow assessment.

Final validation stage – Expected FDA clearance 1Q 2026
Avoids unnecessary invasive angiograms.

Salix® Coronary Plaque Module
Detailed plaque characterisation, dispersion, & stenosis analysis.

FDA cleared – automatically enabled on SCA
Quantifies and characterises plaque within the coronary arteries.

Salix® Coronary Anatomy Platform
Core platform enabling plaque visualisation, workflow optimisation, physician productivity, and CCTA reporting.

FDA cleared – launched commercially in the U.S.
Core Platform for real-time, whole-heart characterisation.

1 https://cardiovascularbusiness.com/topics/cardiac-imaging/cms-increases-medicare-payments-cardiac-ct-ccta

Salix® platform successfully launched in the U.S.

Tanner Health collaboration partnership to validate and integrate

TANNER HEALTH SYSTEM

CONE HEALTH

Northeast Georgia Health System

Improving the health of our community in all we do

Integrated and Live from July 2025

- 5-year Commercial Agreement – US$0.6M
- Monthly subscription and future fee per scan revenues
- 5 Hospitals, cardiovascular centres, 30 practices
- Live in clinical workflow – PACS + EMR

…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

Next U.S. launch partners

- Cone Health and Northeast Georgia Health System have been key U.S. validation partners since 2024
- Work underway to integrate and go live shortly
- Each will also execute a Commercial Agreement
- U.S. launch partners – medium sized hospitals all moving to CCTA scans as first line test for patients with symptoms of coronary artery disease

Flagship Novel Plaque Study – SAPPHIRE

Salix-based Analysis of Plaque to identify Patients at Higher Risk of Events

The SAPPHIRE study will validate the novel Salix plaque analysis to identify patients at risk of heart attack and improve treatment.

Overview of Study

- 6 - 8 high volume U.S. Centres
- **Mass General, Ascension, Piedmont Healthcare** and **Huntsville Heart Centre** confirmed
- 3-phased retrospective study
- **Phase 1:** identify patients at risk using Salix Plaque analysis compared to current standard of care. Anticipated timeline – 12 months
- **Phase 2:** leverage Salix Plaque Dispersion Score to further risk stratify patients and change treatment. Anticipated timeline – 12 months

Benefits

1. Predict future adverse events and improve treatment.
2. Build clinical awareness and credibility in US
3. Accelerate commercial adoption in US hospitals

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Key Takeaways

Salix® has moved from R&D to commercial use, and a structured pathway for U.S. expansion.

1 | Commercialisation activities and platform now scaling.

2 | Large growing market in the U.S. with established reimbursement codes

3 | Salix® platform live in the U.S. – generating revenues with a clear pathway to grow.

4 | FDA clearance of Salix® Coronary Plaque module driving increased revenue opportunity.
Salix® Coronary Flow strengthens future market opportunity once FDA cleared

5 | Structured go-to-market approach into the U.S. market with 3 partners and SAPPHIRE Study

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Corporate Enquiries

John Konstantopoulos
Co-Founder and Chief Executive Officer
Tel: +61 8 6478 7816
Email: contact@artrya.com

Investor & Media Enquiries

David Allen or John Granger
Hawkesbury Partners
Tel: +61 499 100 038 or +61 410 577 155
Email: investors@artrya.com