

Nasdaq: PAVM, PAVMZ

Q4 2023 Business Update

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- The Companies may not yet have received clearance from the FDA or any other regulatory agency for some of the products described in this presentation.



Recent Highlights

Lucid Diagnostics

- 4Q23 EsoGuard revenue +33% quarterly
- High-volume health fair #CYFT testing events continue to gain traction
- Improving out-of-network reimbursement with stable pricing
- Expansion of clinical validity and clinical utility data to support in-network coverage, including Medicare

Veris Health

- Veris strategic shift to target large academic and regional cancer centers; first such engagement expected in very near-term
- Final, successful FDA pre-sub meeting for implantable monitor; clear path to FDA clearance pending independent financing

PMX Incubator

 Launched wholly-owned incubator, PMX, to complete development and commercialization of existing medtech portfolio technologies starting with PortIO



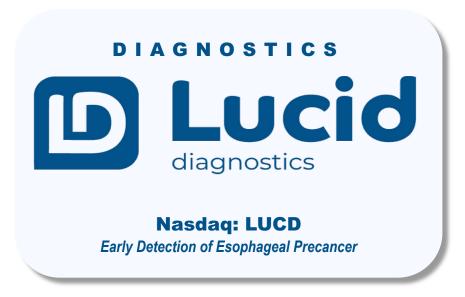
PAVmed Strategy

- Drive shareholder value through holdings in independently financed subsidiaries managed through a shared services structure
- Follow successful Lucid path and seek financing opportunities directly into Veris, subsidiaries based on PMX technologies, and future subsidiaries
- Veris shift to large academic centers driven to enhance financeability
- PMX launching initial effort to independently finance PortIO
- Actively seeking out new groundbreaking, independently financeable technologies with large market opportunities, agnostic of sector, to leverage existing PAVmed infrastructure



PAVmed Corporate Structure









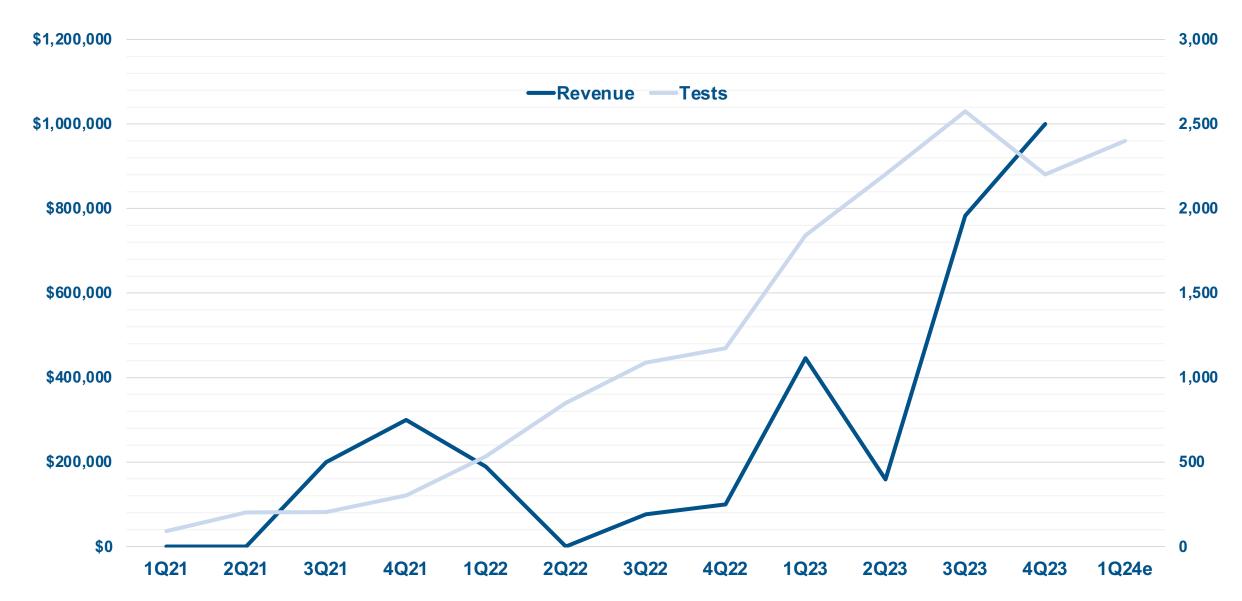








EsoGuard Revenue & Test Volume





Highlights

Commercial Execution

- High-volume health fair CYFT testing events fully booked through July
- Active EsoGuard testing programs over a dozen strategic accounts
- ~50% of claims allowed with stable payments averaging ~\$1,800

Key Strategic Accomplishments

- Strengthened balance sheet with \$18.1 million preferred stock financing
- Significant expansion of clinical validity (CV) and clinical utility (CU) data to support broad EsoGuard medical policy coverage, including MoIDx re-engagement
- Held meetings with major commercial payors to formally request positive medical policy determinations for EsoGuard
- Launched direct contracting program, with EsoGuard offered as covered benefit



Enhancing Cancer Care



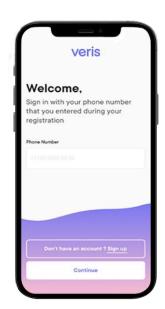




Commercial-Stage Digital Health Company

Enhance Personalized Cancer Care

Veris Cancer Care Platform





Implantable Monitor



MISSION: Utilize modern RPM tools to improve care through early detection of complications, longitudinal trends and risk management



Revised Commercial Strategy

- Target large academic and regional cancer centers
 - Large staffs, large number of patients on infusion therapy
 - Concentrated in metropolitan areas
 - NCI-designated comprehensive cancer centers
 - Venture arms
- Robust Pipeline
 - Over a dozen targets and multiple active

- discussions
- First engagement expected in the very near-term
- Phased rollout
 - Pilot programs
 - Long-term partnerships
 - Other strategic collaborations



Veris Implantable Monitor

- Implant with vascular access port
- Key features:

Continuous cardiac monitoring Temperature

Activity (steps) Respiratory rate

Patient-triggered event monitor Bluetooth connectivity

- Held final, successful FDA pre-submission meeting
- Clear path to FDA clearance and commercial launch once Veris secures independent financing



Assures 100% patient compliance with RPM billing requirements





PMX Incubator

- Launched to complete development and commercialization of existing portfolio technologies
 - PortIO Implantable Intraosseous Vascular Access Device
 - EsoCure Esophageal Ablation Device
 - CarpX Minimally Invasive Device for Carpal Tunnel Syndrome



- Executed joint venture agreement with Hatch Medical
- Will seek to independently finance a separate subsidiary of PMX to develop and commercialize each technology
- First target: PortIO
 - First implantable Intraosseous Vascular Access Device
 - Offers solution for patients with poor veins and need to preserve veins for dialysis
 - Eliminates need for regular maintenance with flushes, resistant to occlusion and infections
 - Estimated market opportunity >\$500M
 - Completed First-in-Human studies in Colombia in 2022, with excellent device function and no complications
 - Extensive engagement with the FDA has established a clear path to a U.S. IDE clinical study and de novo regulatory clearance







Q4 2023 Financial Update



Balance Sheet

(\$ in thousands)	3Q23	4Q23	Change	
Assets				
Cash	\$26,408 \$19,639		\$(6,769)	
Fixed Assets	1,820	1,783	(37)	
All other Assets	13,792	11,697	(2,095)	
Liabilities				
Accounts Payable	2,165	1,786	(379)	
Senior Secured Convertible Notes	44,990	44,200	(790)	
Other Current Liabilities	7,350	8,191	841	
Long-Term Liabilities	3,343	2,960	(383)	
Shares Outstanding	8.0M	8.6M	0.6M	



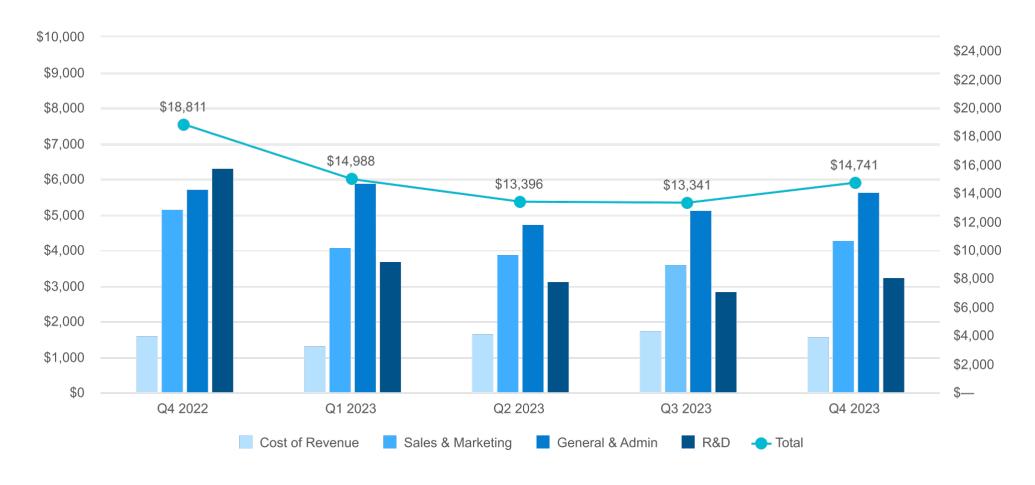
P&L

(\$ in thousands)	4Q22	4Q23	Change	4Q22 - YTD	4Q23 - YTD	Change
Revenue	\$112	\$1,049	\$937	\$377	\$2,452	\$2,075
Operating Expenses	\$24,712	\$17,434	\$(7,278)	\$91,464	\$71,247	\$(20,217)
Net Loss	\$(24,572)	\$(17,408)	\$7,164	\$(103,238)	\$(79,263)	\$23,975
EBITDA	\$(19,746)	\$(13,525)	\$6,221	\$(85,866)	\$(61,767)	\$24,099
Stock-Based Compensation	\$4,949	\$1,968	\$(2,981)	\$19,532	\$11,139	\$(8,393)
Non-GAAP Adjusted Loss	\$(15,638)	\$(10,655)	\$4,983	\$(58,218)	\$(41,781)	\$16,437
Non-GAAP Adjusted Loss per Share	\$(2.52)	\$(1.33)	\$1.19	\$(9.80)	\$(5.78)	\$4.04

To supplement our audited financial results presented in accordance with U.S. generally accepted accounting principles (GAAP), management provides certain non-GAAP financial measures of the Company's financial results. Non-GAAP financial measures are presented with the intent of providing greater transparency to the information used by us in our financial performance analysis and operational decision-making. These non-GAAP financial measures are not intended to be, and should not be, a substitute for, considered superior to, considered separately from, or as an alternative to, the most directly comparable GAAP financial measures. Management believes the non-GAAP financial measures provide useful information to management and investors by isolating certain expenses, gains, and losses that may not be indicative of our core operating results and business outlook and believes non-GAAP financial measures enhance the comparability of results against prior periods. A reconciliation to the most directly comparable GAAP measure of all non-GAAP financial measures included in this presentation and the corresponding press release for the three months and year ended December 31, 2023 are reflected above.



Non-GAAP Operating Expense



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