

Forward looking statement

This presentation contains forward-looking statements and estimates with respect to the anticipated future performance of mdxhealth and the market in which it operates. These statements are often, but are not always, made through the use of words or phrases such as "potential," "expect," "will," "goal," "next," "potential," "aim," "explore," "forward," "future," and "believes" as well as similar expressions. Forward-looking statements contained in this presentation include, but are not limited to, statements regarding the acquisition of Oncotype DX® GPS prostate cancer business from Exact Sciences including statements regarding the anticipated benefits of the acquisition; statements regarding expected future operating results; statements regarding product development efforts; and statements regarding our strategies, positioning, resources, capabilities and expectations for future events or performance. Such statements and estimates are based on assumptions and assessments of known and unknown risks, uncertainties and other factors, which were deemed reasonable but may not prove to be correct. Actual events are difficult to predict, may depend upon factors that are beyond the company's control, and may turn out to be materially different. Examples of forward-looking statements include, among others, statements we make regarding expected future operating results; product development efforts, our strategies, positioning, resources, capabilities and expectations for future events or performance. Important factors that could cause actual results, conditions and events to differ materially from those indicated in the forward-looking statements include, among others, the following: uncertainties associated with global macroeconomic conditions; our ability to successfully and profitably market our products; the acceptance and reimbursement of our products and services by healthcare providers and payors; our ability to obtain and maintain regulatory approvals and comply with applicable regulations; the possibility that the anticipated benefits from our business acquisitions like our acquisition of the Oncotype DX® GPS prostate cancer business will not be realized in full or at all or may take longer to realize than expected; and the amount and nature of competition for our products and services. Other important risks and uncertainties are described in the Risk Factors sections of our most recent Annual Report on Form 20-F and in our other reports filed with the Securities and Exchange Commission. The Company expressly disclaims any obligation to update any such forward-looking statements in this presentation to reflect any change in its expectations with regard thereto or any change in events, conditions or circumstances on which any such statement is based unless required by law or regulation. The Company obtained the industry, statistical and market data, including its general expectations, market position and market opportunity, in this presentation from its own internal estimates and research as well as from industry and general publications and research. surveys and studies conducted by third parties. The market data used in this presentation involves a number of assumptions and limitations. While the Company believes that the information from these industry publications, surveys and studies is reliable, the industry in which it operates is subject to a high degree of uncertainty and risk due to a variety of important factors. These and other factors could cause results to differ materially from those expressed in the estimates made by third parties and by the Company. This presentation does not constitute an offer or invitation for the sale or purchase of securities or assets of mdxhealth in any jurisdiction. No securities of mdxhealth may be offered or sold within the United States without registration under the U.S. Securities Act of 1933, as amended, or in compliance with an exemption therefrom, and in accordance with any applicable U.S. securities laws.

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Analyst Coverage

Any opinions, estimates or forecasts made by analysts are theirs alone and do not represent opinions, forecasts or predictions of mdxhealth or its management. Requests for copies of analyst reports should be directed at the respective analyst and institution.





Ticker: MDXH

mdxhealth provides highly accurate and clinically actionable urologic solutions to inform patient diagnosis and treatment while improving healthcare economics for payers and providers

mdxhealth fundamentals for growth



Fundamentals in place

- Compelling and comprehensive menu in prostate cancer
- · Robust clinical data
- Established reimbursement and guidelines inclusion



Levers for growth

- Expansion of mdxhealth clinical pathway for prostate cancer (acquisition of Genomic Prostate Score)
- Expanding coverage of current menu (Select mdx LCD published)
- · Expanding US commercial footprint



Established focus & execution

- World-class CLIA certified multi-state lab operations
- Experienced and expanded channel into urology
- Urinary Tract Infection opportunity validated



Potential opportunities

- · Opportunistic decentralization of menu as appropriate
- Expanded channel outside of urology
- Menu Expansion: Monitor mdx and business development opportunities



Experienced leadership team

Track record of success



Michael K. McGarrity

Chief Executive Officer

Joined mdxhealth in 2019
Nanosphere
(Luminex/DiaSornin)
Stryker



Ron Kalfus

Chief Financial Officer

Joined mdxhealth in 2019
Rosetta Genomics
Mabcure



John Bellano

Chief Commercial Officer

Joined mdxhealth in 2019
Assurex Health (Myriad Genetics)
Third Wave Technologies (Hologic)
Roche Diagnostics Molecular
Diagnostics



Joseph Sollee

Executive Vice President Corp. Dev. General Counsel

Joined mdxhealth in 2008
Triangle Pharmaceuticals
TherapyEdge

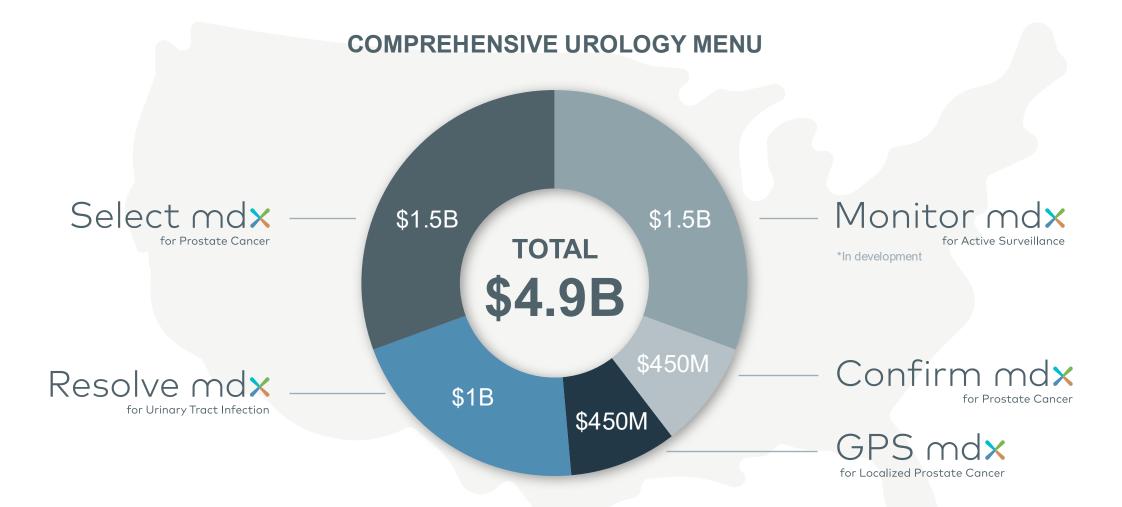


Miriam Reyes

Executive Vice President Laboratory Operations

Joined Mdxhealth in 2011 CombiMatrix Agendia Labcorp

Our menu addresses a \$4.9B U.S. market opportunity(1-5)



Commercial levers to drive growth

One of the most compelling, comprehensive and accurate menus in urology



Standardized laboratory partner for urology group practice

- One rep
- One laboratory
- One patient support program
- ONE PARTNER in the diagnosis and treatment of prostate cancer



Experienced distribution channel and broad KOL network

• Expanded commercial team to >70 people



Acquired Exact Sciences' Genomic Prostate Score (GPS) test

- Established brand with broad customer base
- · Covered by Medicare



Validated advanced Urinary Tract Infection (UTI) opportunity

· Launched in second half of 2021



ACQUIRED AUGUST 2022



UTI TEST LAUNCHED 2021



Current challenges with diagnosing prostate cancer



Prostate cancer screening



elevated PSA results annually⁽¹⁻²⁾



of biopsies DO NOT reveal cancer and may lead to increased complications and hospitalization⁽³⁻⁶⁾



Prostate cancer diagnosis



men undergo biopsies annually⁽²⁾



of cancer-negative biopsies are false negatives, meaning these patients actually have cancer⁽⁷⁾



Prostate cancer risk stratification



prostate cancers diagnosed annually⁽⁸⁾



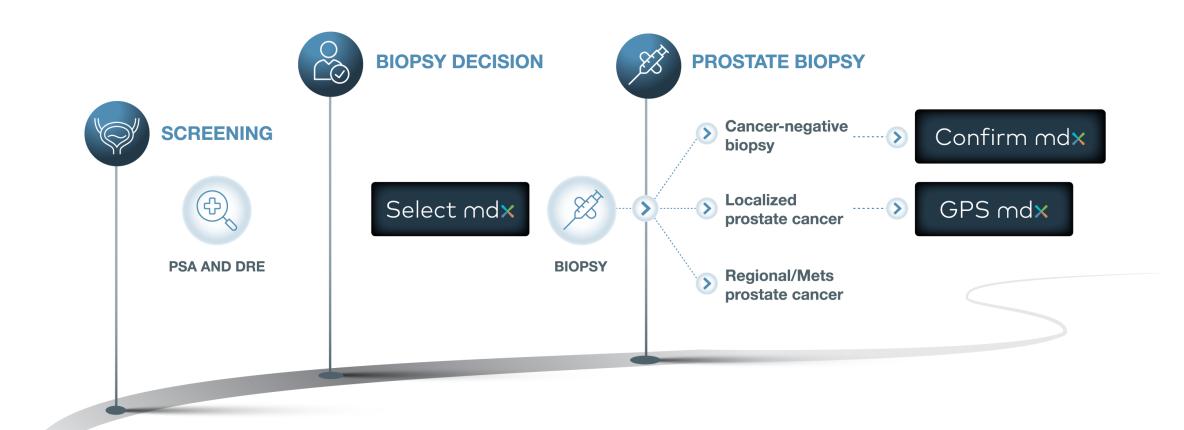
of new prostate cancer diagnosis are localized; Active Surveillance or treatment decision⁽⁸⁾



Prostate cancer is the most common cancer and the 2nd deadliest cancer in U.S. men⁽¹⁾

Expanding menu in the prostate cancer diagnostic pathway

One of the most comprehensive menus in prostate cancer





improves patient selection prior to prostate biopsy

A highly predictive test to identify men at low risk for aggressive prostate cancer



Non-invasive

Urine-based "rule-out" test improves the diagnostic disposition of patients by avoiding unnecessary prostate biopsies

Accurate

95% negative predictive value (1)

Validated

12 published studies on genes and technology

Cost effective

Potential to avoid invasive and unnecessary prostate biopsies and save the U.S. healthcare system >\$500 million (2) each year

National guidelines

Included in EAU and NCCN guidelines (3-4)

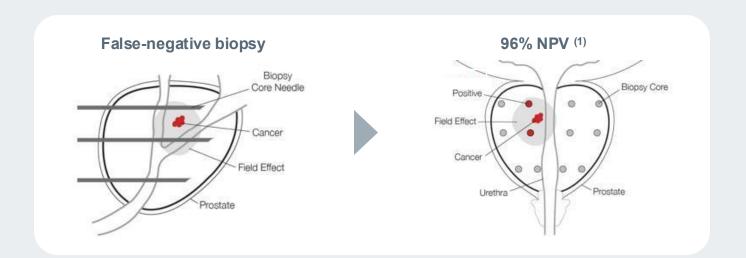


60% of initial biopsies do not reveal cancer (5-8)



improves diagnostic confidence of biopsy result

The only epigenetic test to identify men at risk for aggressive prostate cancer





Non-invasive

"Rule-out" test performed on previous biopsy tissue

Accurate

96% Negative Predictive Value for aggressive prostate cancer⁽¹⁾

Validated

Over 55 published studies on genes and technology

Cost effective

Potential annual U.S. health system savings of \$500K per 1M covered patients(2)

National guidelines

National guidelines: Included in EAU and NCCN guidelines (3-4)



30% of men with a cancer-negative biopsy result actually have cancer (5)



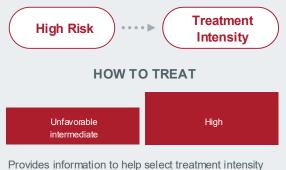
guides treatment decisions for localized prostate cancer

The test analyzes prostate cancer gene activity to predict disease aggressiveness and provide clinically meaningful endpoints⁽¹⁻²³⁾





Provides additional information to help when deciding on whether to pursue active surveillance or more aggressive treatment options



Non-invasive

test performed on previous biopsy tissue

Accurate

Predicts adverse pathology, distant metastasis, prostate cancer mortality and pT3/Extra prostatic extension

Validated

Predicts adverse pathology in AS candidate cohorts in 7 studies >2,000 patients





Hereditary prostate cancer testing

Prostate cancer remains a leading cause of cancer-related death among men.



300,000

estimated new diagnoses



4 35,000

estimated deaths annually



Hereditary prostate cancer accounts for **5–10%** of newly diagnosed patients



Decades of research indicate that prostate cancer has a strong genetic component with implications to ancestry.

Highest Heritability

Prostate cancer has the highest heritability of any major cancer in men

Hereditary Prostate Cancer Testing:



is recommended in the National Comprehensive Cancer Network (NCCN) guidelines



is critical to identify patients eligible for novel therapies (e.g., PARP inhibitors, immunotherapies, and targeted agents for inherited prostate cancers)



is covered by Medicare and many commercial insurance plans, for eligible patients



Prostate Cancer Pipeline

Active surveillance monitoring (Localized prostate cancer)





Current Standard of Care

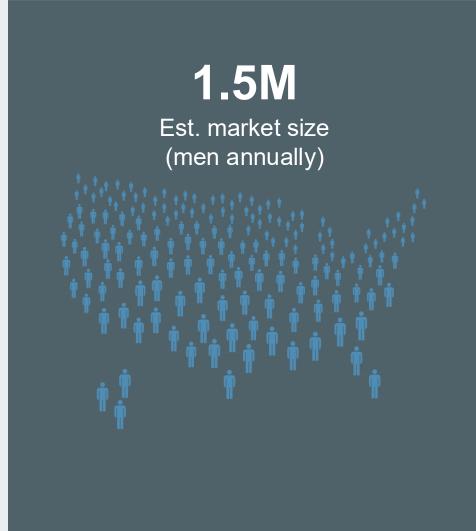
Patients under active surveillance are currently monitored by invasive and costly prostate biopsies



Monitor mdx

Monitor mdx will be a non-invasive alternative that risk-stratifies patients for continued active surveillance vs. intervention, which may also improve patient compliance







Urinary Tract Infection (UTI) annual market opportunity

UTIs are the most common outpatient infection⁽¹⁾



10M

suspected UTI cases present annually⁽²⁾

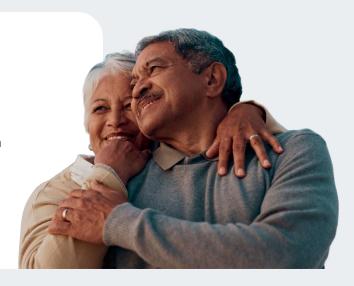


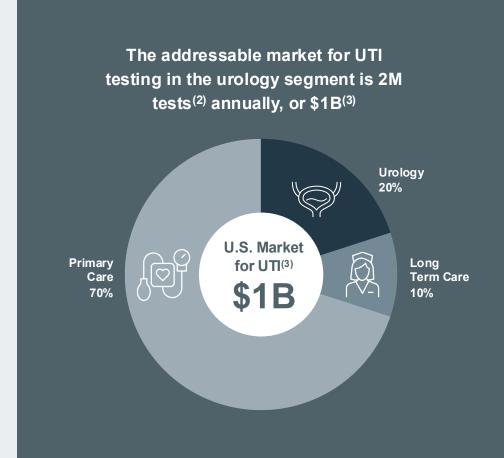
20%

of volume presents to urology⁽³⁾

The current UTI testing market is underserved

- · Current standard is based on dated culture methodologies
- · Complex molecular methods target both organism and susceptibility markers
- · Market conversion comps: Virology and infectious disease
- · Reimbursement well characterized (Medicare/commercial)









Advance molecular urinary tract infection testing



As many as

of urine cultures are polymicrobial, especially in elderly populations

Up to



67%

of recognized uropathogens that traditional urine culture may miss

Resolve mdx

- identifies and quantities uropathogenic bacteria and associated antibiotics susceptibility
- improves antibiotic stewardship

Guide antibiotic Positive selection Rule out Negative infection

Non-invasive

Urine-based test that provides personalized antibiotics options for urinary tract infections.

Accurate

19 pathogens, 6 classes of resistance genes and susceptibility to guide antibiotic selection

Turnaround Time

Results within 24 - 48 hours

Selected Financial Data

Three months ended (unaudited)					Full year ended (unaudited)			
In \$'000 (except EPS)	March 31, 2025	March 31, 2024	% Change	Dec 31,	2024 Dec 31, 2023	% Change		
Total revenue	\$24,292	\$19,834	+22%	\$90,0	49 \$70,193	+28%		
Gross profit	\$15,504	\$12,063	+29%	\$55,14	41 \$43,929	+26%		
Gross profit %	63.8%	60.8%	+3.0pp	61.29	% 62.6%	-1.4pp		
Net loss	(\$9,209)	(\$8,511)	+8%	(\$38,0	69) (\$43,100)	(12%)		
Adjusted EBITDA ¹	(\$1,331)	(\$4,537)	(71%)	(\$14,6	72) (\$19,382)	(24%)		
EPS	\$(0.19)	\$(0.31)	(39%)	\$(1.1	\$(1.66)	(30%)		

¹ A reconciliation of IFRS to non-IFRS financial measures has been provided in the tables included in the appendix. An explanation of these measures is also included under the heading "Non-IFRS Disclosures"

Unaudited 1Q25 results and updated 2025 guidance

2025 REVENUE GUIDANCE OF

\$108-110M ►



representing year-over-year revenue growth of approximately 210 at the midpoint



mdxhealth is well-positioned for sustainable growth and value creation



Revenue growth

- Multi-billion-dollar addressable market opportunity fortified by acquisition of Oncotype DX GPS from Exact Science
- 2025 guidance of \$108-110 million



Experienced and expanded channel into urology

- Field sales team of 70 in the US
- Urinary Tract Infection opportunity taking hold
- Additional channel opportunities as they present



Gross margin leverage

- Coverage for Select mdx indication
- Driving additional payer coverage for full menu
- Accretion of (f/k/a Oncotype DX) GPS and UTI test



Leadership team and operating discipline

Focus and execution across all operating discipline

Thank you



INVESTOR RELATIONS

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References

Slide 6 - Our menu addresses a \$4.9B U.S. market opportunity

- 1. MDxHealth management estimates
- Welch. et. Al., Prostate-Specific Antigen Levels in the United States: Implications of Various Definitions for Abnormal. JNCI 2005.
- NIH Cancer Trends Progress Report. July 2021. https://progressreport.cancer.gov/detection/prostate_cancer
- NIH Surveillance, Epidemiology and end Results Program. July 2021. https://seer.cancer.gov/statfacts/html/prost.html.
- Paul et. al. State of the Globe: Rising Antimicrobial Resistance of Pathogens in Urinary Tract Infection. J Glob Infect Dis. 2018.

Slide 8 - Current challenges with diagnosing prostate cancer in U.S.

- 1. NIH 6/10/2024 Website: https://seer.cancer.gov/statfacts/html/prost.html
- Mdxhealth management estimates.
- Moyer VA, U.S. Preventive Services Task Force. Screening for prostate cancer: U.S. Preventive Services Task Force recommendation statement. Ann Intern Med. 2012;157:120–134.
- Bhindi B, Mamdani M, Kulkarni GS, et al. Impact of the U.S. Preventive Services Task Force recommendations against prostate specific antigen screening on prostate biopsy and cancer detection rates. J Urol. 2015;193:1519–1524.
- 5. Loeb et al. European Urology 2013.
- 6. Loeb et al. Journal of Urology 2011.
- 7. Stewart et al. Journal of Urology 2013.
- NIH Cancer Stat Facts: Prostate Cancer. https://seer.cancer.gov/statfacts/html/prost.html

Slide 10 - SelectMDx improves patient selection prior to prostate biopsy

- Haese, A, et al. (2019) Multicenter Optimization and Validation of a 2-Gene mRNA Urine Test for Detection of Clinically Significant Prostate Cancer Prior to Initial Prostate Biopsy. J Uro. doi: 10.1097/JU.00000000000000293.
- Govers TM, et al. (2018) Cost-Effectiveness of Urinary Biomarker Panel in Prostate Cancer Risk Assessment. J Urol. doi: 10.1016/j.juro.2018.07.034A.
- 2022 National Cancer Center Network Guidelines. Early Detection for Prostate Cancer. Version 1.2022 – July 16, 2022.
- 4. 2021 European Association of Urology Prostate Cancer Guidelines.
- Moyer VA, U.S. Preventive Services Task Force. Screening for prostate cancer: U.S. Preventive Services Task Force recommendation statement. Ann Intern Med. 2012;157:120–134.
- Bhindi B, Mamdani M, Kulkarni GS, et al. Impact of the U.S. Preventive Services
 Task Force recommendations against prostate specific antigen screening on
 prostate biopsy and cancer detection rates. J Urol. 2015;193:1519–1524.
- 7. Loeb et al. European Urology 2013.
- 8. Loeb et al. Journal of Urology 2011.

Slide 11 - ConfirmMDx improves diagnostic confidence of biopsy result

- Van Neste, et al. (2016) Risk Score Predicts High-Grade Prostate Cancer in DNA-Methylation Positive, Histopathologically Negative Biopsies. J Urology.
- 2. Aubry. Et al., Budget Impact Model: Epigenetic Assay Can Help Avoid

- Unnecessary Repeated Biopsies and Reduce Healthcare Spending. American Health & Drug Benefits 2013.
- 2022 National Cancer Center Network Guidelines. Early Detection for Prostate Cancer. Version 1,2022 – July 16, 2022.
- 4. 2021 European Association of Urology Prostate Cancer Guidelines.
- Stewart et al., Clinical Utility of an Epigenetic Assay to Detect Occult Prostate Cancer in Histopathologically Negative Biop sies: Results of the MATLOC Study. Journal of Urology

Slide 12 –(f/k/a Oncotype DX) Genomic Prostate Score (GPS) to guide treatment decisions for localized prostate cancer

- Brooks MA, et al. Validating the associate on of adverse pathology with distant metastasis and prostate cancer mortality 20-years after radical prostatectomy. Urol Oncol. 2022;40(3):104.e1-104.e7.
- Mehrali vand S, et al. A grading system for the assessment of risk of extraprostatic extension of prostate cancer at multiparametric MRI. Radiology. 2019;290(3):709-719.
- Brooks MA et al. GPS assay association with long-term cancer outcomes: twenty-year risk of distant metastasis and prostate cancer-specific mortality. JCO Precis Oncol. 2021;5:PO.20.00325.
- Cullen J, et al.,. The 17-gene genomic prostate score test as a predictor of outcomes in men with unfavorable intermediate risk prostate cancer. Urology. 2020;143:103-111.
- Klein EA, et al. A 17-gene assay to predict prostate cancer aggressiveness in the context of Gleason grade heterogeneity, tumor multifocality, and biopsy under sampling. Eur Urol. 2014;66(3):550-560.
- Cullen J, et al. A biopsy-based 17-gene genomic prostate score predicts recurrence after radical prostatectomy and adverse surgical pathology in a radially diverse population of men with clinically low- and intermediate-risk prostate cancer. Eur Urol. 2015;68(1):123-131.
- Van Den Eeden SK, et al. A biopsy-based 17-gene genomic prostate score as a predictor of metastases and prostate cancer death in surgically treated men with clinically localized disease. Eur Urol. 2018;73(1):129-138.
- Eggener S_n et al. A 17-gene panel for prediction of adverse prostate cancer pathologic features: prospective clinical validation and utility. Urology. 2019:126:76-82.
- Lin DW, et al. 17-gene genomic prostate score test results in the Canary Prostate Active Surveillance Study (PASS) cohort. J Clin Oncol. 2020;38(14):1549-1557.
- Badani KK,, et al. The impact of a biopsy based 17-gene genomic prostate score on treatment recommendations in men with newly diagnosed clinically prostate cancer who are candidates for active surveillance. Urol Pract. 2015;2(4), 181-180
- Dall'Era MA, et al., Utility of the Oncotype DX® prostate cancer assay in clinical practice for treatment selection in men newly diagnosed with prostate cancer: a retrospective chart review analysis. Urol Pract. 2015; 2(6), 343-348.
- Albala D, et al. Health economic impact and prospective clinical utility of Oncotype DX® Genomic Prostate Score. Rev Urol. 2016;18(3):123-132.

- Eure G, et al. Use of a 17-gene prognostic assay in contemporary urologic practice: results of an interim analysis in an observational cohort. Urology. 2017;107:67-75.
- Lynch JA, et al. Improving risk stratification among veterans diagnosed with prostate cancer: impact of the 17-gene prostate score assay. Am J Manag Care. 2018;24(1 Suppl):S4-S10.
- Leapman MS, et al. Association between a 17-gene genomic prostate score and multi- parametric prostate MRI in men with low and intermediate risk prostate cancer (PCa). PLoS One. 2017;12(10):e0185535.
- Kornberg Z, et al. Genomic Prostate Score, PI-RADS version 2 and progression in men with prostate cancer on active surveillance. J Urol. 2019;201(2):300-307.
- Salmasi A, et al. A17-gene genomic prostate score assay provides independent information on adverse pathology in the setting of combined multiparametric magnetic resonance imaging fusion targeted and systematic prostate biopsy. J Urol. 2018;200(3):564-572.
- Magi-Galluzzi C, et al. The 17-gene genomic prostate score assay predicts outcome after radical prostatectomy independent of PTEN status. Urology. 2018;121:132-138.
- Cullen J, et al. Multicenter comparison of 17-gene genomic prostate score as a predictor of outcomes in African American and Caucasian American men with clinically localized prostate cancer. J Urol. 2021;205(4):1047-1054.
- 20. Murphy AB, et al. A 17-gene panel genomic prostate score has similar predictive accuracy for adverse pathology at radical prostate domy in African American and European American men. Urology. 2020;142:166-173.
- Moschovas M, et al. Association between Oncotype DX genomic prostate score and adverse tumor pathology after radical prostatectomy. Eur Urol Focus. 2021;S2405-4569(21)00094-8.
- Aboushwareb, et al. Active surveillance or watchful waiting in clinically low-risk prostate cancer patients in the SEER database with and without an Oncotype Dx genomic prostate score assay. J Urol. 2021;206(3S):e1094 (MP62-06).
- Brand TC, et al. Patient-specific meta-analysis of 2 clinical validation studies to predict pathologic outcomes in prostate cancer using the 17-gene genomic prostate score Urology. 2016;89:69-75.

Slide 15 – U.S. Urinary Tract Infection (UTI) annual market opportunity

- Medina M, Castillo-Pino E. An introduction to the epidemiology and burden of urinary tract infections. Ther Adv Urol. 2019;11:1756287219832172. Published 2019 May 2. doi:10.1177/1756287219832172.
- Flores-Mireles AL, Walker JN, Caparon M, Hultgren SJ. Urinary tract infections: epidemiology, mechanisms of infection and treatment options. Nat Rev Microbiol. 2015;13(5):269-284. doi:10.1038/nrmicro3432.
- Mdxhealth management estimates are informed by the Company's knowledge of the industry.





Select mdx robust clinical evidence

12 published studies on genes and technology











PIVOTAL CLINICAL STUDIES

Analytical validation	Hessels et al., Translational Medicine Communications 2017			
Clinically validated for a 95% NPV	Haese et al., Journal of Urology 2019			
Significantly impacts prostate biopsy decision making	Shore et al., Urology Practice 2019			
>\$500M in savings to health care system	Govers et al., Journal of Urology 2018			









Confirm mdx robust clinical evidence

Over 55 published studies on genes and technology











PIVOTAL CLINICAL STUDIES

Analytical validation	Van Neste et al., BMC Urology 2013			
Validation of high NPV	Partin et al., Journal of Urology 2014.			
Meta analysis validating high NPV	Partin et al., Trans. of the Am. Clin. and Clim. Assoc 2016			
Risk score development NPV 96% CS PCa	Van Neste et al. The Prostate 2016			
Validated in African American men	Waterhouse et al., Urology 2016			
Validation of clinical utility/actionability	Wojno., et al 2014			
Savings to health care system	Aubry et al., American Health Drug and Benefits 2013			





Transactions of the American Climatological Association

The Prostate





GPS robust clinical evidence

Over 20 published clinical validation and utility studies











PIVOTAL CLINICAL STUDIES

Analytical validation	Knezevic et al., 2013			
Clinically validated as an independent predictor of adverse pathology	Klein et al., 2014, Cullen et al., 2015, Eeden et al., 2017, Eggner et al., 2019			
Clinical validated in African American men	Cullen et al., 2015, Murphy et al., 2021			
Validation of clinical utility	Badani et al., 2015, D			
Validation of clinical utility/actionability	Badani et al., 2015, Dall'Era et al., 2015, Eure et al., 2017, Lynch et al., 2017, Murphy et al., 2021, Moschovas et al., 2021			
Cost savings by decreasing unnecessary immediate treatment	Albala et al., 2016			











Select mdx, Confirm mdx and GPS technology

The most comprehensive menu in prostate cancer

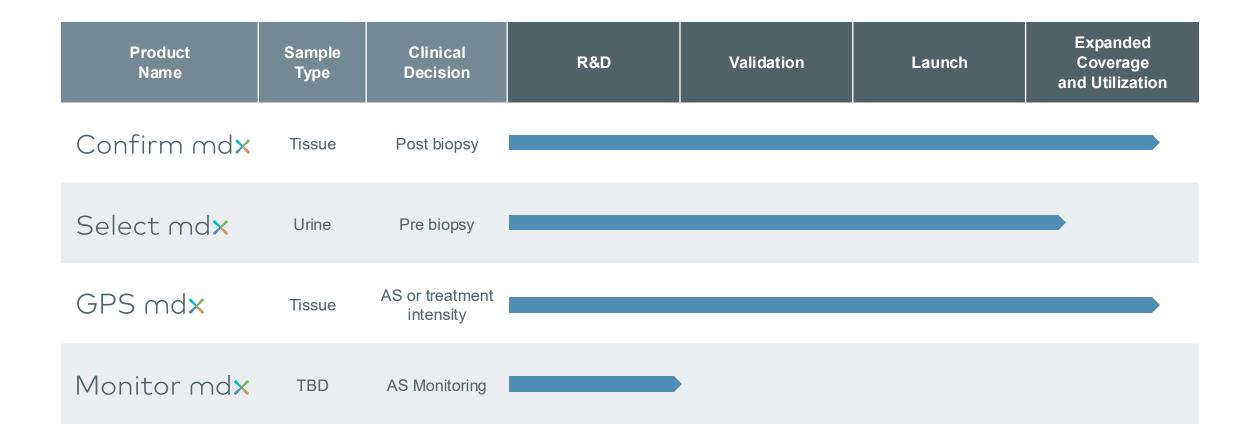
	Select mdx ⁽¹⁾	Confirm mdx ⁽²⁾	GPS ⁽³⁾
Specimen	Urine	Prostate tissue	Localized PCa tissue
Science	mRNA RT-PCR assay	DNA Methylation Specific PCR assay	Multi gene expression RT-PCR Assay
Biomarkers	DLX1, HOXC6	GSTP1, APC RASSF1	17 genes (AZGP1, FAM13C, KLK2, SRD5A2, FLNC GSN, GSTM2, TPM2, BGN, COL1A1, SFRP4, TPX2, ARF1, ATP5E, CLTC, GPS1, PGK1)
Clinical Model	Clinical model combines mRNA with established clinical risk factors	Clinical model combines DNA Methylation markers with established clinical risk factors	Clinical algorithm aggregates expression of 5 reference genes to normalize the expression of the 12 cancer-related genes
Performance	95% NPV for clinically significant prostate cancer	96% NPV for clinically significant prostate cancer	Predicts adverse pathology, distant metastases, PCa mortality



^{2.} Van Neste, et al. (2016) Risk Score Predicts High-Grade Prostate Cancer in DNA-Methylation Positive, Histopathologically Negative Biopsies. J Urology.

^{3.} Knezevic et al., (2013) Analytical validation of the Oncotype DX prostate cancer assay – a clinical RT-PCR assay optimized for prostate needle biopsies. BMC Genomics

Prostate cancer precision diagnostics: menu and pipeline





Unaudited reconciliation of ifrs to non-ifrs financial measures

	Three Months Ended March 31,			Year Ended December 31		
In thousands of \$		2025	2024		2024	2023
IFRS net loss Amortization of intangible assets	\$	(9,209) \$ 1,322	(8,511) 1,125	\$	(38,069) \$ 4,905	(43,100) 4,494
Depreciation expense		925	775		3,134	2,365
Share-based compensation expense		391	168		1,725	665
Interest expense, net Debt extinguishment costs		1,859 -	1,218		6,551 3,130	4,494 -
Fair value adjustments ⁽¹⁾		2,547	519		2,961	9,960
Exact Sciences earnout amendment ⁽²⁾		-	-		-	1,128
Other adjustments ⁽³⁾		697	169		609	611
Income tax		137			382	1
Adjusted EBITDA	\$	(1,331) \$	(4,537)	\$	(14,672) \$	(19,382)

¹⁾ Primarily related to GPS contingent consideration, Exact Sciences 5-year warrants, Innovatus derivative instrument, option to pay Exact Sciences earnout in shares, and Noviogendix contingent consideration

Non-IFRS disclosure

In addition to the Company's financial results determined in accordance with IFRS, the Company provides adjusted EBITDA, a non-IFRS measure that it determines to be useful in evaluating its operating performance. The Company defines adjusted EBITDA as net loss less interest expense, depreciation and amortization of intangible assets, share-based compensation, fair-value adjustments, debt extinguishment costs, amendments related to the Exact Sciences earnout, income tax benefit, and other financial and non-cash expenses. Management believes that presentation of non-IFRS financial measures provides useful supplemental information to investors and facilitates the analysis of the Company's core operating results and comparison of operating results across reporting periods. The Company uses this non-IFRS financial information to establish budgets, manage the Company's business, and set incentive and compensation arrangements. However, non-IFRS financial information is presented for supplemental information purposes only, has limitations as an analytical tool and should not be considered in isolation or as a substitute for financial information presented in accordance with IFRS. For example, non- IFRS adjusted EBITDA excludes a number of expense items that are included in net loss. As a result, positive adjusted EBITDA may be achieved while a significant net loss persists. The Company's presentation of expected non- IFRS adjusted EBITDA is a forward-looking statement about the Company's future financial performance. This non-IFRS measure includes adjustments like share-based compensation, debt extinguishment costs, fair-value adjustments related to contingent considerations that are difficult to predict for future periods because the nature of the adjustments pertain to events that have not yet occurred. Additionally, management does not forecast many of the excluded items for internal use. Information reconciling forward-looking non-IFRS measures to IFRS measures is therefore not available without unreasonable effort and is not provided. The occurrence, timing, and amount of any of the items excluded from IFRS to calculate non- IFRS could significantly impact the Company's IFRS results.



²⁾ Amendment fee and issuance of shares as part of amended GPS asset purchase agreement

³⁾ Bank fees and other non-cash expenses