



# **Corporate Presentation**August 2022

# Forward looking statement

This presentation contains forward-looking statements concerning our expectations, anticipations, intentions, beliefs or strategies regarding the future. These forward-looking statements are based on assumptions that we have made as of the date hereof and are subject to known and unknown risks and uncertainties that could cause actual results, conditions and events to differ materially from those anticipated. Therefore, you should not place undue reliance on forward-looking statements. 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; and the anticipated benefits of our acquisitions, including estimated synergies and other financial impacts. 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 the coronavirus (COVID-19) pandemic, including its possible effects on our operations and the demand for our products; our ability to raise additional capital in amounts and on terms satisfactory to us, if at all; our ability to successfully and profitably market our products; the acceptance of our products by patients and healthcare providers; the willingness of health insurance companies and other payers to cover our products and adequately reimburse us for such products; the amount and nature of competition for our products; the effects of any judicial, executive or legislative action affecting us or the healthcare system; recommendations, guidelines and guality metrics issued by various organizations regarding our products; our ability to successfully develop new products and assess potential market opportunities; our ability to effectively enter into and utilize strategic partnerships and acquisitions; our success establishing and maintaining collaborative, licensing and supplier arrangements; our ability to obtain and maintain regulatory approvals and comply with applicable regulations; the possibility that the anticipated benefits from our business acquisitions will not be realized in full or at all or may take longer to realize than expected; and the outcome of any litigation, government investigations, enforcement actions or other legal proceedings. The risks included above are not exhaustive. 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 filings made with the Securities and Exchange Commission. We undertake no obligation to publicly update any forward-looking statement, whether written or oral, that may be made from time to time, whether as a result of new information, future developments or otherwise.

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.

The mdxhealth logo, mdxhealth, Confirm mdx, Select mdx, Oncotype DX, Genomic Prostate Score and Monitor mdx are trademarks or registered trademarks of mdxhealth SA. All other trademarks and service marks are the property of their respective owners.

#### 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.



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



Ticker: MDXH



Ticker: MDXH.BR

# mdxhealth fundamentals for growth



#### **Fundamentals in place**

- The most 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 Oncotype DX)
- Expanding coverage of current menu (Select mdx LCD) published)
- Expanding US commercial footprint



#### **Established focus & execution**

- World-class CLIA certified lab operation
- Best-in-class commercial channel into urology
- Urinary Tract Infection opportunity validated



#### **Potential opportunities**

- Select mdx point of care testing
- Expanded channel outside of urology
- Pipeline: 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



**Miriam** Reyes

**Executive Vice President Laboratory Operations** 

Joined Mdxhealth in 2011 CombiMatrix Agendia LabCorp



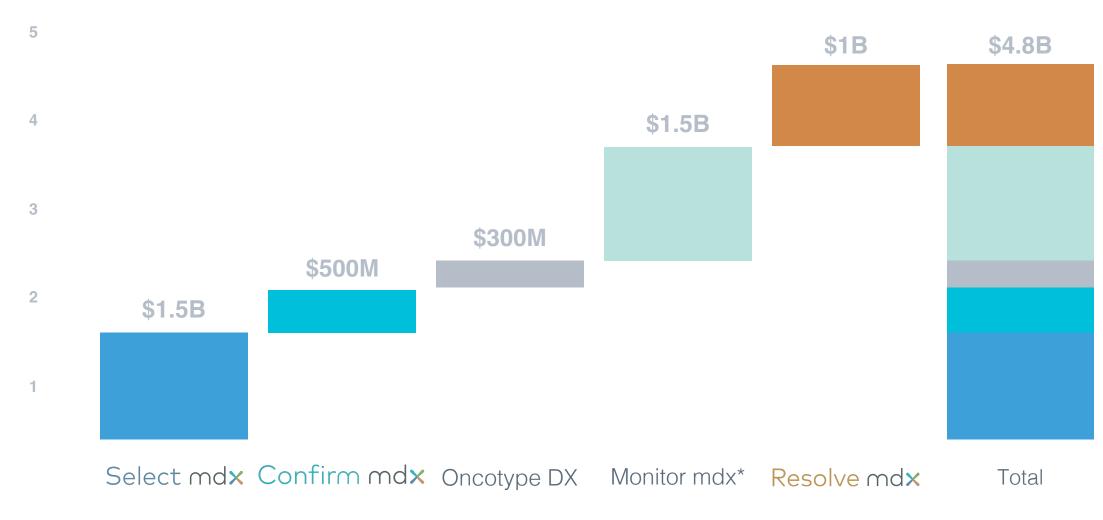
Joseph Sollee

**Executive Vice President** Corp. Dev. General Counsel

Joined mdxhealth in 2008 Triangle Pharmaceuticals TherapyEdge

# Our menu addresses a \$4.8B U.S. market opportunity

Comprehensive Urology Menu



# **Commercial levers to drive growth**

Best-in-class offering into urology with menu of the most accurate clinically-actionable tests for patients and urologists

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 and other urologic diseases

Acquired Exact Sciences' Oncotype DX Genomic Prostate Score (GPS) test

- Established brand with broad customer base
- Covered by Medicare and included in NCCN guidelines

Validated advanced Urinary Tract Infection (UTI) opportunity

Launched in second half of 2021

Experienced distribution channel and broad KOL network

• Expanded commercial team to >70 people

## oncotype bx°

**Genomic Prostate Score** 

Acquired August 2022

#### Resolve mdx

for Urinary Tract Infectio

UTI test launched 2021



# **Current challenges with diagnosing prostate cancer**



#### **Prostate cancer screening**

3 million elevated PSA results annually (1-2)

60% of biopsies DO NOT reveal cancer and may lead to increased complications and hospitalization (3-6)



#### **Prostate cancer diagnosis**

500,000 men undergo biopsies annually (2)

30% of cancer-negative biopsies are false negatives, meaning these patients actually have cancer (7)



#### Prostate cancer risk stratification

268,000 prostate cancers diagnosed annually (8)

74% of new prostate cancer diagnosis are localized Active Surveillance or treatment decision (8)

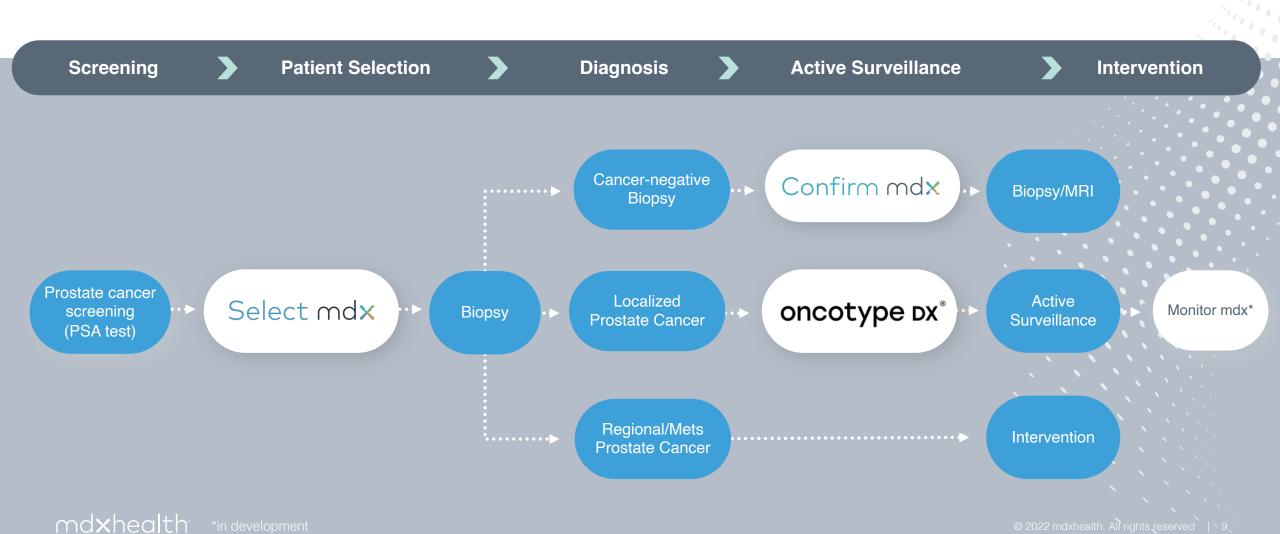
Prostate cancer is the most common cancer and the 2nd deadliest cancer in U.S. men (1)





# Expanding menu in the prostate cancer diagnostic pathway

The most comprehensive menu in prostate cancer



# Select mdx improves patient selection prior to prostate biopsy

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

Abnormal PSA/DRE

At risk for aggressive cancer?

95% NPV



Binary actionable results for patient and HCP



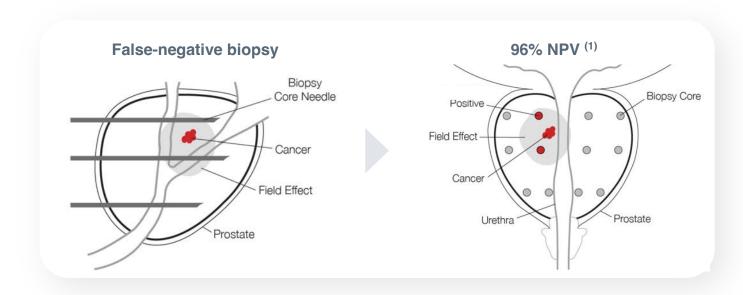
- Non-invasive: Urine-based "rule-out" test improves the diagnostic disposition of patients by avoiding unnecessary prostate biopsies
- **Accurate:** 95% negative predictive value (1)
- Validated: 12 published studies on genes and
- Cost effective: Potential to avoid invasive and unnecessary prostate 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)

# Confirm mdx improves diagnostic confidence of biopsy result

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



 Positive
 Biopsy/MRI

 Negative
 .... ►

 Avoid Biopsy/MRI

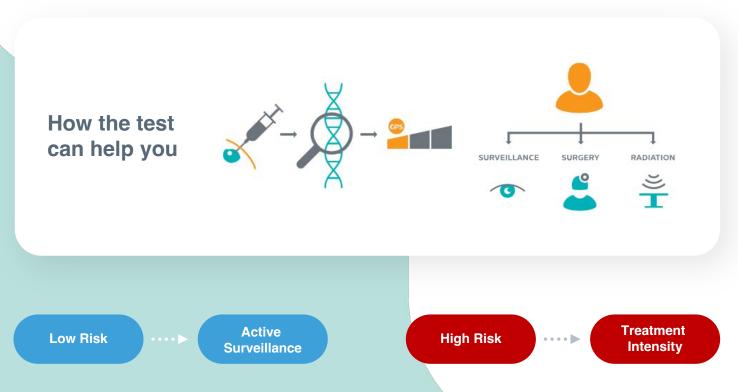
- Non-invasive: "Rule-out" test performed on previous biopsy tissue
- Accurate: 96% Negative Predictive Value for aggressive prostate cancer (1)
- Validated: Over 55 published studies on genes and technology
- Cost effective: Potential annual U.S. health system savings of \$500K per 1M covered patients (2)
- National guidelines: Included in EAU and NCCN guidelines (3-4)



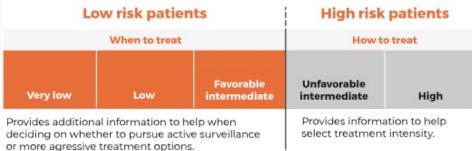
30% of men with a cancer-negative biopsy result actually have cancer (5)

# Oncotype DX Genomic Prostate Score (GPS) 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)



- 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
- National guidelines: Included in NCCN guidelines (24)



# Prostate cancer pipeline

Active surveillance monitoring (Localized prostate cancer)

#### **Monitor mdx**

Patients under active surveillance are currently monitored by invasive and costly prostate biopsies Monitor mdx will be a non-invasive alternative that risk-stratifies patients for continued active surveillance vs. intervention, which may also improve patient compliance

Monitor mdx\*

Low Risk

Intervention

Continued Active Surveillance





# **Urinary Tract Infection (UTI)** annual market opportunity

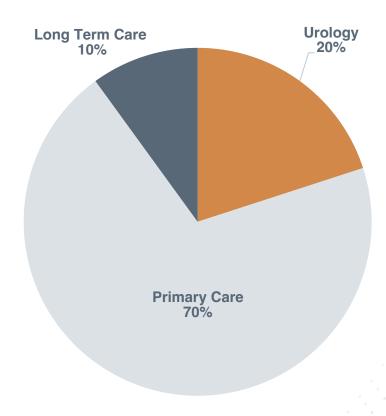
#### UTIs are the most common outpatient infection<sup>(1)</sup>

- 10 million suspected UTI cases present annually<sup>(2)</sup>
- 20% of volume presents to urology\*

#### 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)

The addressable market for UTI testing in the urology segment is 2M tests<sup>(2)</sup> annually, or \$1B\*

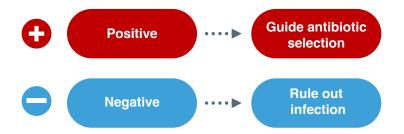


**U.S. Market for UTI\*** 



# Resolve mdx: Advance molecular urinary tract infection testing

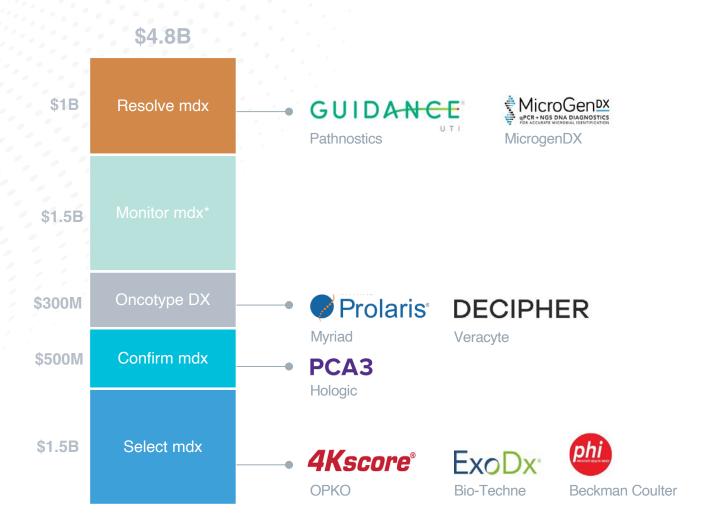
- As many as 33% of urine cultures are polymicrobial, especially in elderly populations, and traditional urine culture may miss up to 67% of recognized uropathogens
- Resolve mdx identifies and quantities uropathogenic bacteria and associated antibiotics susceptibility
- Resolve mdx improves antibiotic stewardship



- 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



# Broad urology-focused menu provides opportunity for growth







# mdxhealth is well-positioned for sustainable growth and value creation



#### Large total addressable market

Selling clinically-actionable diagnostic tests to urologists represents a multi-billion-dollar addressable market opportunity



#### Broad menu provides sustainable growth potential

Generating revenue from clinically-proven commercial products; growth to occur via commercial execution, expanded menu and improved channel access



# Proprietary position into urology call point enables additional growth via menu expansion

Continuing to pursue growth opportunities through R&D and business development



#### Leadership team with commercial focus

Implementing proven strategies to support growth while maintaining operating cost discipline

# Thank you

#### **Investor relations contact**

LifeSci Advisors, LLC US +1 949 271 9223 ir@Mdxhealth.com

#### **Global operations**

US headquarters & laboratory 15279 Alton Parkway, Ste 100 Irvine, CA 92618 United States

**TX laboratory**7000 Preston Road, Ste 1500
Plano, TX 75024
United States

**EU headquarters**CAP Business Center
Rue d'Abhooz, 31
4040 Herstal, Belgium

R&D & laboratory
Novio Tech Campus
Transistorweg 5
6534 AT Nijmegen
The Netherlands

### References

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

- 1. NIH 8/20/2019 Website: https://seer.cancer.gov/statfacts/html/common.html.
- 2. Mdxhealth management estimates.
- 3. 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.
- 4. 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.
- 8. NIH Cancer Stat Facts: Prostate Cancer. https://seer.cancer.gov/statfacts/html/prost.html

#### Slide 10 – SelectMDx improves patient selection prior to prostate biopsy

- 1. 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.
- 2. 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.
- 3. 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.
- 5. Moyer VA, U.S. Preventive Services Task Force recommendation statement. Ann Intern Med. 2012;157:120–134.
- 6. 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

- 1. 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.
- 3. 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.
- 5. Stewart et al., Clinical Utility of an Epigenetic Assay to Detect Occult Prostate Cancer in Histopathologically Negative Biopsies: Results of the MATLOC Study. Journal of Urology



## References

#### Slide 12 - Oncotype DX Genomic Prostate Score (GPS) to guide treatment decisions for localized prostate cancer

- 1. 022 National Cancer Center Network Guidelines. Prostate Cancer. Version 4.2022 May 10, 2022.
- 2. 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.
- 3. Mehralivand 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.
- 4. 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.
- 5. 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.
- 6. Klein EA, et al. A 17-gene assay to predict prostate cancer aggressiveness in the context of Gleason grade heterogeneity, tumor multifocality, and biopsy undersampling. Eur Urol. 2014;66(3):550-560.
- 7. Cullen J, et al. A biopsy-based 17-gene genomic prostate score predicts recurrence after radical prostatectomy and adverse surgical pathology in a racially diverse population of men with clinically low- and intermediate-risk prostate cancer. Eur Urol. 2015;68(1):123-131.
- 8. 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.
- 9. Eggener S., et al. A 17-gene panel for prediction of adverse prostate cancer pathologic features: prospective clinical validation and utility. Urology. 2019;126:76-82.
- 10. 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.
- 11. 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), 181-189.
- 12.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.
- 13. Albala D, et al. Health economic impact and prospective clinical utility of Oncotype DX® Genomic Prostate Score. Rev Urol. 2016;18(3):123-132.
- 14. 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.
- 15. 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.
- 16. Leapman MS, et al. Association between a 17-gene genomic prostate score and multiparametric prostate MRI in men with low and intermediate risk prostate cancer (PCa). PLoS One. 2017;12(10):e0185535.
- 17.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.
- 18. Salmasi A, et al. A 17-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.
- 19. 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.
- 20. 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.
- 21. Murphy AB, et al. A 17-gene panel genomic prostate score has similar predictive accuracy for adverse pathology at radical prostatectomy in African American and European American men. Urology. 2020;142:166-173.
- 22. 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.
- 23. 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).
- 24. 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.



## References

#### Slide 16 – U.S. Urinary Tract Infection (UTI) annual market opportunity

- 1. 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.
- 2. 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.



# **Appendix**

## Select mdx robust clinical evidence

12 published studies on genes and technology







Clinical utility



Health economics

#### **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.,











## Confirm mdx robust clinical evidence

Over 55 published studies on genes and technology





Clinical



Clinical



Health economics

#### **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 Clinical and Climatological Association

The Prostate





# Oncotype DX robust clinical evidence

Over 20 published clinical validation and utility studies





Clinical validity



Clinical utility



Health economics

#### **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 Oncotype DX technology

The most comprehensive menu in prostate cancer

	Select mdx <sup>(1)</sup>	Confirm mdx <sup>(2)</sup>	Oncotype DX <sup>(3)</sup>	
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	

<sup>3.</sup> 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



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

# Prostate cancer precision diagnostics: menu and pipeline

Product name	Sample type	Clinical decision	R&D	Validation	Launch	Expanded coverage and utilization
Confirm mdx	Tissue	Post biopsy				•
Select mdx	Urine	Pre biopsy				•
Oncotype DX	Tissue	AS or treatment intensity				•
Monitor mdx	TBD	AS Monitoring	•			

