

Corporate Presentation

July 2022



Forward looking statements

- IMPORTANT: The following applies to this document, the oral presentation of the information in this document by Immunovia AB (publ) (the "Company") or any person on behalf of the Company, and any question-and-answer session that follows the oral presentation (collectively, the "Information").
- The Information has been prepared and issued by the Company solely for use at the presentation held by the Company in relation to the Company's operations and position. The Information has not been independently verified and will not be updated. Unless otherwise stated, and any market data used in the Information is not attributed to a specific source, are estimates of the Company, and have not been independently verified. The Information, including but not limited to forward-looking statements, applies only as of the date of this document and is not intended to give any assurances as to future results.
- THE INFORMATION IS BEING MADE AVAILABLE TO EACH RECIPIENT SOLELY FOR ITS INFORMATION AND BACKGROUND.
- The Information does not constitute or form part of and should not be construed as an offer or the solicitation of an offer to subscribe for or purchase any securities issued by the Company.
- The Information contains forward-looking statements. All statements other than statements of historical fact included in the Information are forward-looking statements. Forward-looking statements give the Company's current expectations and projections relating to its financial condition, results of operations, plans, objectives, future performance and business. These statements may include, without limitation, any statements preceded by, followed by or including words such as "target," "believe," "expect," "aim," "intend," "may," "anticipate," "estimate," "project," "will," "can have," "likely," "should," "would," "could" and other words and terms of similar meaning or the negative thereof. Such forward-looking statements involve known and unknown risks, uncertainties and other important factors beyond the Company's control that could cause the Company's actual results, performance or achievements to be materially different from the expected results, performance or achievements expressed or implied by such forward-looking statements. Such forward-looking statements are based on numerous assumptions regarding the Company's present and future business strategies and the environment in which it will operate in the future. The Company disclaims any obligation to update or revise any forward-looking statements, whether as a result of new information, future events or otherwise.





Revolutionizing blood-based diagnostics to advance early

detection of pancreaticcancer and increase patientsurvival rates

Company overview



Front-runner in non-invasive early detection of pancreatic cancer (PDAC)

Incorporated 2007 in Sweden, 2017 in US

Listed on Nasdaq Stockholm under IMMNOV

CLIA and CAP-accredited laboratory in Marlborough, MA and HQ in Lund, Sweden

Revolutionizing blood-based diagnostics to advance early detection of pancreatic cancer and increase patient survival rates

Initial ~1.7-1.8 million US patient population in pancreatic cancer

Commercial build out in US following launch in August 2021

Execute US reimbursement plan

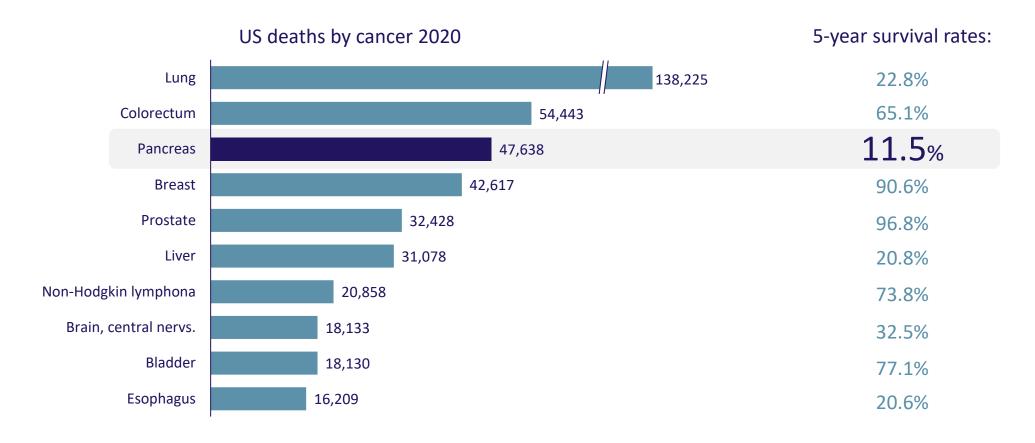
New executive management team

DISCOVERY, DEVELOPMENT AND VALIDATION

TRANSITIONING FOCUS TO COMMERCIAL EXECUTION AND US MARKET PENETRATION

Pancreatic is one of the most lethal cancers with limited diagnostic innovation





Limited industry spending is dedicated to addressing the third deadliest cancer

Patients are often diagnosed too late when surgery is no longer an option

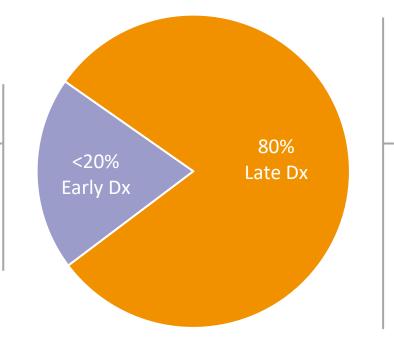




5-year survival rate when diagnosed early (surgical optionality)

Treatment methods:

- Chemotherapy
- Surgery
- Clinical trial therapeutics



5-year survival rate when found late (metastatic, non-resectable)

Treatment methods:

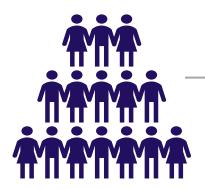
- Chemotherapy
- Clinical trial therapeutics
- Palliative Care

Traditional diagnostic methods for pancreatic cancer have resulted in low patient survival rates

US patient population of approximately 1.7 to 1.8 million



~1.7-1.8 million
Patients



Familial/Hereditary

315,000 – 350,000 individuals that need 1-2 tests per year

First risk group commercialized in US, August 2021

- High-risk individuals typically identified by gastroenterologist or oncologist treating family member with pancreatic cancer
 - Referred for genetic counseling and surveillance
- High-risk Surveillance Programs located at centers throughout U.S.
 - Typically located at academic centers with experience treating pancreatic cancer patients
- Some individuals under surveillance by local GI, rather than formal program at academic center

Symptomatic

596,000 patients/year with concerning gastric symptoms

New Onset Diabetes

856,000 patients/year with 3 years follow up and 2 tests per year

Limitations in current standard of care for pancreatic cancer diagnosis



Too few patients under surveillance

- Only 21% of patients who qualify for high-risk pancreatic cancer surveillance enroll
- Biggest reason cited: lack of awareness
- The nearest center with a surveillance program is too far for many high-risk individuals

Imaging is burdensome for patients

- Both MRCP and endoscopic ultrasound generally require travel to a surveillance center
- Endoscopic ultrasound (EUS) is an invasive procedure that carries the risk of pain, bleeding or acute pancreatitis
- Some patients experience claustrophobia with MRIs

Imaging results can be inconclusive

- Small tumors are difficult to detect with imaging
- Meta-analysis indicates the specificity of MRCP is 89% and EUS is 86%
- Interpretation of imaging results can vary by radiologist

Imaging frequently fails to identify pancreatic cancer early

- Imaging fails to identify some PDACs, especially small tumors
- Diagnosis of pancreatic cancer frequently occurs at stage 3 or 4, when surgery is not an option
- Pancreatic cancer can progress quickly in the year-long interval between imaging

Revolutionary blood-based test: IMMray® PanCan-d



First-to-market advantage

First US blood-based pancreatic cancer monitoring test

Accurate microarray patented technology

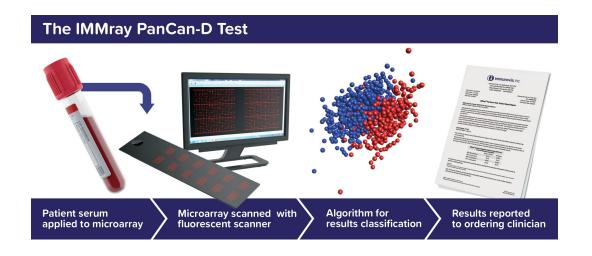
Unique "disease fingerprints" from a blood sample

Significant unmet medical need

US addressable market size of ~1.7-1.8 million patients

Product advantages

Performance and patient experience advantages vs. current surveillance methods



- Test measures 9 biomarkers to detect pancreatic cancer; protected by patents across 7 patent families
- Proprietary algorithm classifies sample into 1 of 3
 actionable results; biomarker weighting is a trade secret
- Results reported 5 -7 days after specimen receipt

Immunovia's product aims to increase survival rates for patients with cancer



IMMray® PanCan-d provides specific, actionable results

Immunovia Medical **HIGH-RISK SIGNATURE** Director calls ordering Serum is classified as high-risk for pancreatic cancer physician; prompt **PRESENT** clinical evaluation **NEGATIVE FOR** Continue with regular Serum is classified as low-risk for pancreatic cancer surveillance **HIGH-RISK SIGNATURE** Consider EUS based on Serum could not be classified as high-risk or low-risk pre-test probability; **BORDERLINE** for pancreatic cancer (too close to the cutoff) Retest IMMray PanCand in 3-6 months **TEST NOT PERFORMED** Specimen could not be processed due to poor Re-draw with patient if quality sample or CA19-9 value of 2.5 U/ml or less. appropriate* (TNP)

^{*}If CA19-9 value is 2.5 U/ml, sample will not be re-drawn. Assumption is patient is Lewis-null genotype and retesting is not indicated.

Broad clinical validation of IMMray® PanCan-d



Commercial test model study

Verification study

Blinded validation study (Publication: *Clin Transl Gastroenterol*. 2022)

2020

2020

2022

PDAC stage I-II versus healthy

PDAC stage I-II versus healthy

PDAC stage I-II versus familial/hereditary

1,113 patient samples

89 PDAC stages I & II

182 PDAC stages III & IV

8 EU/US sites

519 patient samples

81 PDAC stages I & II

84 PDAC stages III & IV

9 EU/US sites

586 patient samples

56 PDAC Stages I & II

111 PDAC Stages III & IV

11 EU/US sites

Specificity 98%

Sensitivity 85%

Specificity 99%

Sensitivity 78%

Specificity 99%

Stage I-II Sensitivity 89% Stage I-IV Sensitivity 92%

Commercial Test Model

Candidate commercial signature and fine tune algorithm using fresh samples.

310 healthy controls and 488 symptomatic controls.

All CA19-9 samples included

Verification (Case control study)

Locked signature and algorithms on known clinical samples 212 healthy controls and 112 symptomatic controls.

All CA19-9 samples included

Validation

Blinded clinical samples.

216 healthy and 203 high-risk controls

Lewis Null excluded

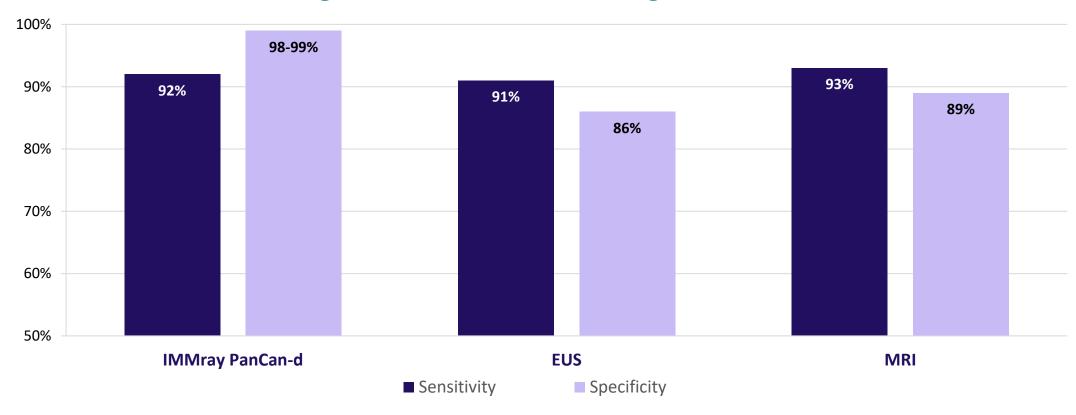
CA19-9 values <2.5 U/ml are Lewis Antigen Null genotype (le/le), patients don't express CA19-9

PanFAM-1 study showed 98% specificity but not enough PDAC's to evaluate sensitivity

Outstanding performance compared to standard methods



Diagnostic Performance for Detecting PDAC I – IV



IMMray® PanCan-d shows comparable sensitivity and superior specificity to imaging and is less burdensome to the patient

Toft J, Hadden WJ, Laurence JM et al. Imaging modalities in the diagnosis of pancreatic adenocarcinoma: A systematic review and meta-analysis of sensitivity, specificity and diagnostic accuracy. Eur J Radiol. 2017;92:17-23.

Extensive key opinion leader & advocacy network



Advocacy Partner Organizations







Key Research and Clinical Collaborators





















Staged approach to commercializing IMMray PanCan-d



PHASE	LAUNCH (CURRENT)	GROWTH (MEDIUM-TERM)	EXPANSION (LONG-TERM)
Intended Uses in Pancreatic Cancer Detection	Genetic and familial risk factors	Genetic and familial risk factors Cysts (potential)	Genetic and familial risk factors Cysts Chronic pancreatitis New onset diabetes
Physician Call Points	High-risk surveillance centers Interventional GIs & pancreas specialists	High-risk surveillance centers Interventional GIs & pancreas specialists GIs	High-risk surveillance centers Interventional GIs & pancreas specialists GIs Endocrinologists Primary care
Geographic Reach	6 territories (18 states)	National	National
Sales Staffing	6 field sales reps, 2 inside sales reps	24 field sales reps, 8 inside sales reps	~80 field sales reps, 27 inside sales reps Commercialization partner for PC
Key Additional Go- to-Market Tactics	Pioneers in Early Detection physician experience program Advocacy assoc. partnerships (pilots)	Advocacy assoc. partnerships Digital marketing to at-risk individuals	Advocacy assoc. partnerships Digital marketing to at-risk individuals

Financial highlights & planned investments



Key Investments next 24 months

- Build out field and inside sales teams
- Invest in digital marketing and advocacy partnerships to drive patient demand
- Scale operations to support volume growth
- Bolster clinical utility evidence and payer dossier

Target growth capital

Investment	24-Month Use of Proceeds	
R&D	\$8M (27%)	
Commercial	\$15M (50%)	
Operations	\$7M (23%)	
TARGET GROWTH CAPITAL	\$30M	

Cash position is \$20M as of June 30, 2022, current burn rate is \$1.5-2.0M per month

Establishing the leader in the early detection of pancreatic cancer



AREA	ACCOMPLISHMENTS –	MILESTONES - NEXT 18 MONTHS
OPERATIONAL	 ✓ First to market dedicated liquid biopsy test for the early detection of pancreatic cancer in the US ✓ Secured CLIA and CAP accreditation 	☐ Build out sales team to enable nationwide coverage
		■ Explore potential commercial partnerships
		☐ Build out digital marketing capabilities
	✓ Hired experienced commercial leader as US CEO	☐ Scale lab to accommodate greater test volumes
	✓ Hired sales team	
	✓ Ran over 6,000 tests (research + commercial samples)	
ADOPTION /	✓ Deepened strategic partnerships with PanCAN, National	☐ Payer demonstration pilots with commercial payers
REIMBURSMENT	Pancreas Foundation & Lustgarten	☐ Secure licenses for California and New York
	 ✓ Launched the Pioneers in Early Detection physician experience program 	☐ Explore out-of-network billing, prior authorizations and appeals to assess opportunities for payment
	✓ Hired Head of US Market Access	■ Expand patient advocacy association partnerships
	✓ Obtained PLA code & submitted recommended pricing on the Clinical Lab Fee Schedule	☐ Leverage KOLs and Pioneers to advocate for coverage with regional commercial payers
CLINICAL	✓ Published peer-reviewed, blinded validation study in Clinical & Translational Gastroenterology ¹	☐ Additional clinical studies for PDAC early detection
		☐ Analyze PanDIA samples to develop and assess test
	✓ Announced results from the PanFAM-1 study	accuracy in new onset diabetics over age 50
	✓ Obtained 6,000 samples from patients with new onset diabetes through the PanDIA clinical collaboration	☐ Prioritize additional intended uses and develop an R&D and clinical roadmap

Experienced global management team







Karin Almqvist Liwendahl Group CFO



Jeff Borcherding US CEO, US Commercial Lead



Linda Mellby VP R&D



Collective Experience – Broader Management Team

Myriad genetics









DIGNITANA

LAZARD











Appointed several experienced key leaders for growth phase: CEO & President, CFO and U.S. CEO

Q&A

helloir@immunovia.com www.immunovia.com