

Interim report Q2 2022

24 August

Philipp Mathieu – CEO and President



Forward Looking Statements

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Agenda

- Strategic focus & target market
- Q2 Operational highlights
- Q2 Financial highlights
- Commercial update

Fully focused on execution in Pancreatic Cancer



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

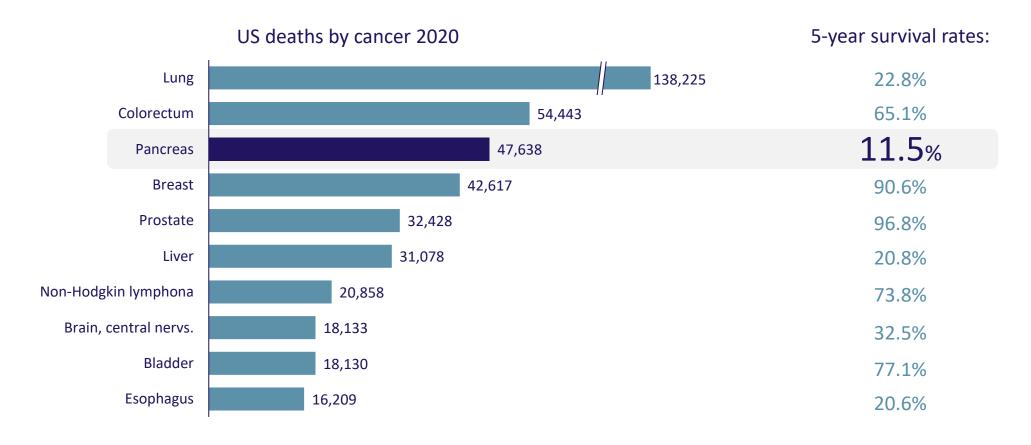
New executive management team

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

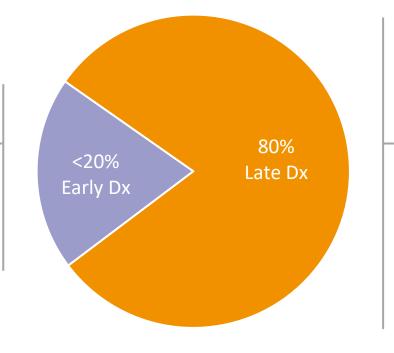




5-year survival rate when diagnosed early (surgical optionality)

Treatment methods:

- Chemotherapy
- Surgery
- Clinical trial therapeutics



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

Treatment methods:

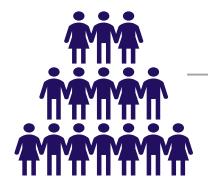
- Chemotherapy
- Clinical trial therapeutics
- Palliative Care

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

U.S. Population estimates from Statistica Research Department based on 2019 survey: https://www.statista.com/statistics/241488/population-of-the-us-by-sex-and-age/.





Q2 – Operational highlights

- Strengthened the US team
- Clinical validation PanFAM-1 results and update on PanDia
- Attained PLA application for IMMray™
 PanCan-d
- IMMRay PanCan-d now available in almost all states in the US
- Physician experience program and collaboration with key opinion leaders





Q2 – Financial highlights

- Net sales of KSEK 103 (38)
- Net earnings amounted to MSEK -33 (-49)
 - Investing in US commercial operations
 - Lower other external expenses Q/Q and Y/Y
 - Currency exchange effects and capitalization of R/D in 2021 impacting net earnings comparison
- Solid cash position of MSEK 197 (382)

^{* (}x) same period as last 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

- 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



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.



Update

Physician experience program almost fully enrolled at select pancreatic cancer highrisk surveillance centers in the US.

The program commenced in April and has enrolled 19 surveillance centers



Physician experience program

- Select pancreatic high-risk surveillance centers
 - Limited number of no-charge tests
 - Providers to become comfortable using IMMray PanCan-d
- Broadly include IMMray™ PanCan-d in standard surveillance routines
 - Develop Key Opinion Leaders to educate other clinicians
 - Cultivate advocates to request payer coverage
- Imaging data generated for comparison

Drive Familiarity, Adoption and Reimbursement

Staged approach to commercializing IMMray PanCan-d



PHASE	LAUNCH (CURRENT)	GROWTH (MEDIUM-TERM)	EXPANSION (LONG-TERM)
	Genetic and familial risk factors	Genetic and familial risk factors	Genetic and familial risk factors
Intended Uses in Pancreatic Cancer Detection		IPMNs (cysts)	IPMNs (cysts)
			Chronic pancreatitis
			New onset diabetes
	High-risk surveillance centers	High-risk surveillance centers	High-risk surveillance centers
Physician Call Points	Interventional GIs & pancreas specialists	Interventional GIs & pancreas specialists	Interventional GIs & pancreas specialists
		Gls	Gls
			Endocrinologists
			Primary care
Geographic Reach	6 territories (18 states)	National	National

Executing reimbursement plan for US insurance coverage





Extensive payer insights obtained; value proposition refined (2021)



Laboratory CAP accreditation received (Mar 2021)



Final publication of peer-reviewed blinded validation study in Clinical and Translational Gastroenterology journal¹ (Feb 2021)



Pioneers in Pancreatic Cancer physician experience program initiated (Apr 2022)



Head of Market Access hired (May 2022)



Pricing recommendation for the Clinical Lab Fee Schedule (CLFS) submitted to Centers for Medicare & Medicaid Services (Jun 2022)



PanFAM-1 study results announced. Specificity further validated. Sensitivity not assessed due to low number of PDACs (Jun 2022)



PLA code approved (Jul 2022)



Discussions with regional & national payers, specifically payer validation studies of IMMray to generate evidence for policy decisions (ongoing)



Engage KOLs and clinicians to advocate with payers (ongoing)



Sign first commercial payer demonstration project (projected Q4 2022-Q1 2023)



PLA code and CLFS pricing active (Jan 2023)



Recognize initial commercial reimbursement (projected Q4 2022 or 2023)

2022 – Executing on strategic priorities



Priorities	Achievements
Additional clinical validation for IMMray™ PanCan-d across risk groups	 ✓ Publication of peer-reviewed validation study ✓ Results for PanFAM-1 & PanDia update
Strengthening US team for successful commercial scale up	✓ Hired an experienced US Leader✓ Hire of Head of Market Access
	✓ CAP and CLIA accreditation of Lab
Execution of US reimbursement plan for pancreatic cancer	 ✓ PLA code approved ✓ Launched physician experience program and KOL collaboration
Roadmap to capture the potential of discovery programs in other indications	✓ Fully focus on Pancreatic Cancer and other programs deprioritized
Prioritization for market access in non-US geographies	

Q&A

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