

**Saving Lives by Making
Genetic Diagnosis for Cancer
Faster and Accessible**



OACP IE LTD

[13485 CERTIFIED] | [CE-IVDR COMPLIANT] | [MISA KSA LICENSED]

DNA Tests Are Crucial in Cancer Diagnosis — But Plagued by a Dual Crisis

The Time Barrier

**3+ Days
Testing Time**

The Talent Gap

**Critical Shortage Of
Pathologist**

Workflow Inefficiencies have a Massive Impact

Increased Mortality Risk

+10%

Death Risk

Per month of treatment delay

Preventable Deaths

126K

Deaths/Year

Attributable to diagnostic delays (est.)

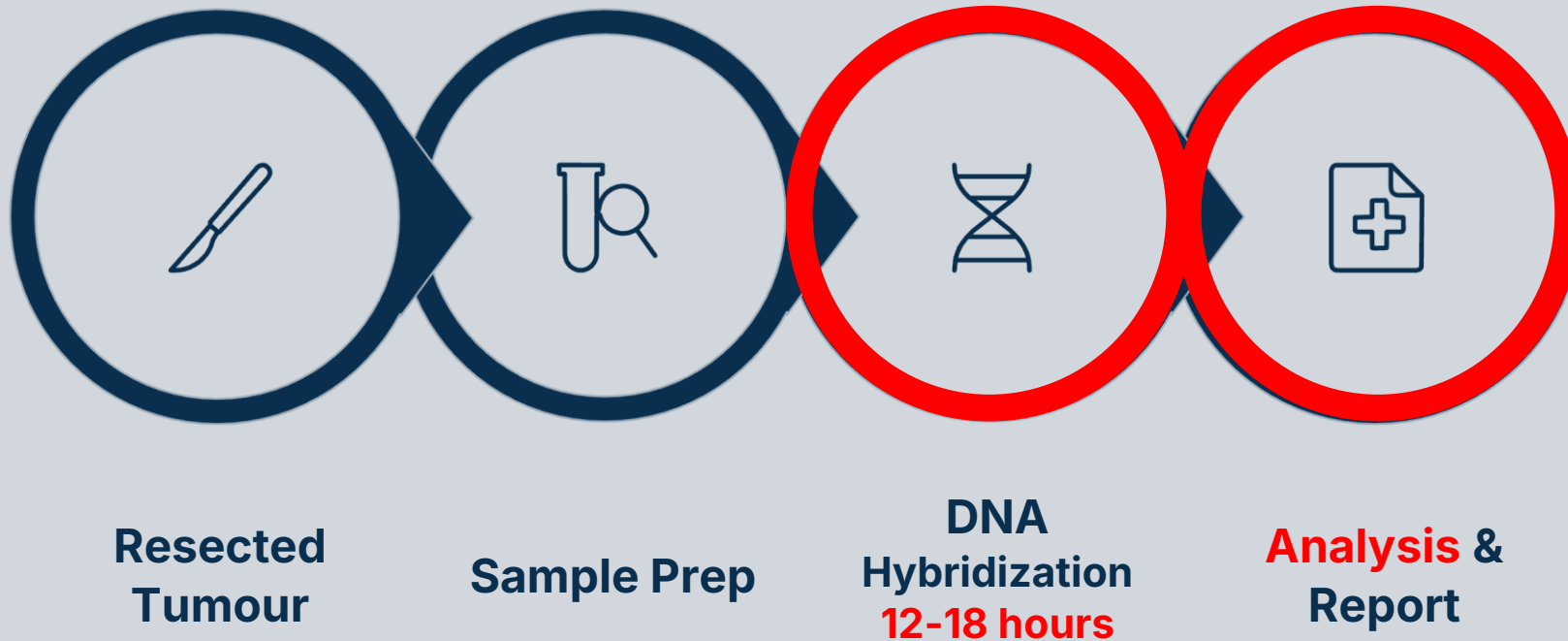
Excess Healthcare Costs

\$20B

Extra Costs

In avoidable annual healthcare spending

Current DNA Testing Workflow Bottleneck



The Solution: Two Engines. One Integrated Platform.



ENGINE 1: Rapid Hybridization

Accelerated Chemistry

ENGINE 2: Improved Quality

Digital Pathology AI

How It Works: Zero-Friction Operational Integration

Engine 1: Accelerated Chemistry

Prepare Sample

Add OaCP Buffer

Run Hybridisation

Engine 2: Digital Pathology AI

Capture Slide Image

Upload to AICP

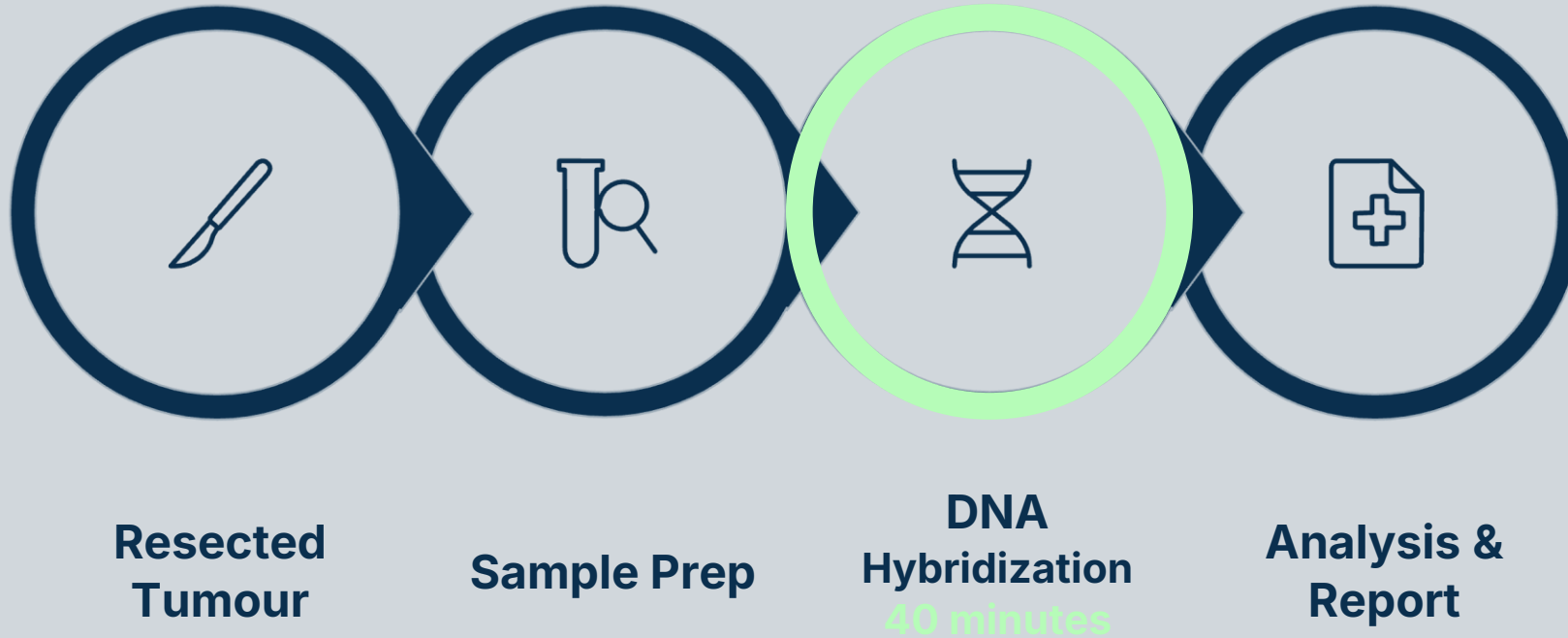
Instant AI Inference



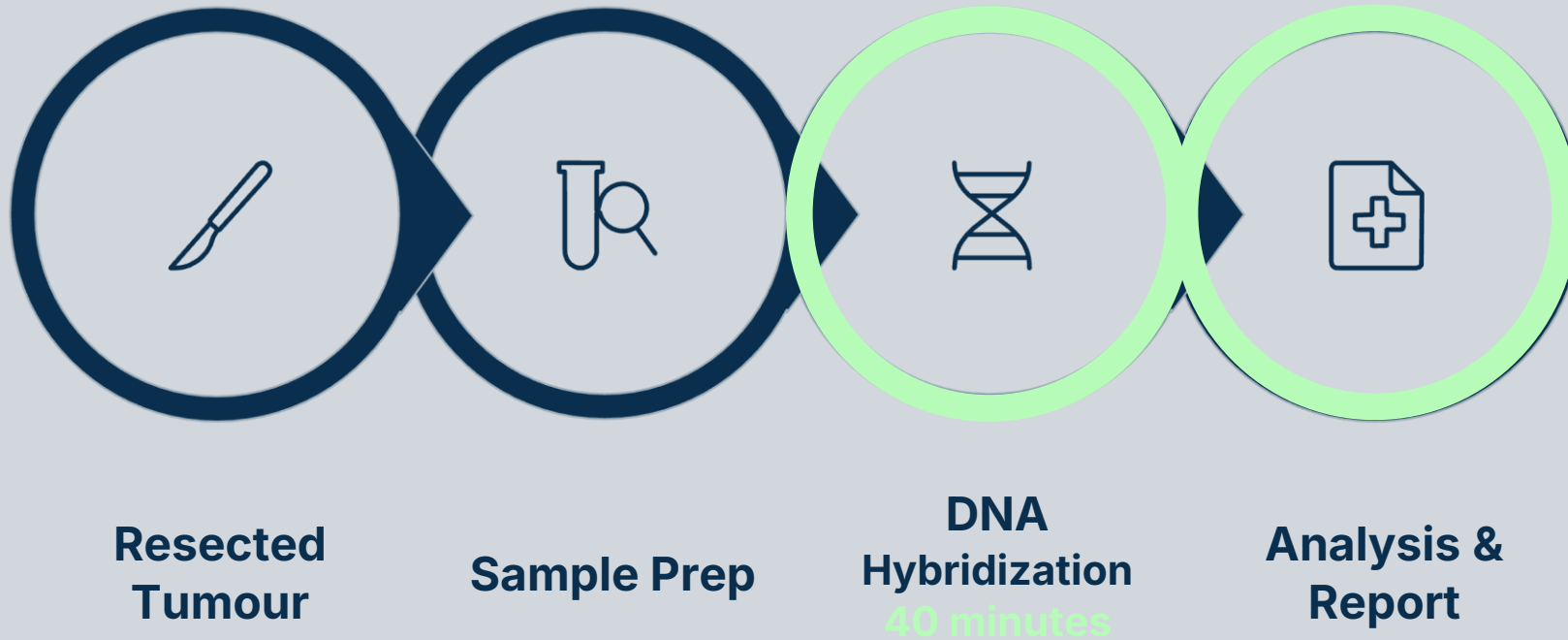
No new equipment,
No protocol redesign,
ZERO switching costs.

▣ Detects hyper/Low-digestion, paraffin artifacts, and confirms neoplastic target cellularity prior to scoring.

Accelerating the Diagnostic Workflow: From 3 Days to 2 Hours With Our Reagents



Improving the Reliability and the Quality of Results With Our AI Tool



Performance Metrics

2hr

Total Test Time

50%

Cost Savings

98%+

Sensitivity

+56.4%

Margin Uplift

ZERO

Hardware Mods

Proprietary and Certified Technology

Reagents Patent Coverage

The EU



The USA



China



India



AI/CP Development Roadmap

1

Q3 2026 — In House model Development

2

Q1 2027 — AI model implementation and Tuning
With the support of the University of Michigan



3

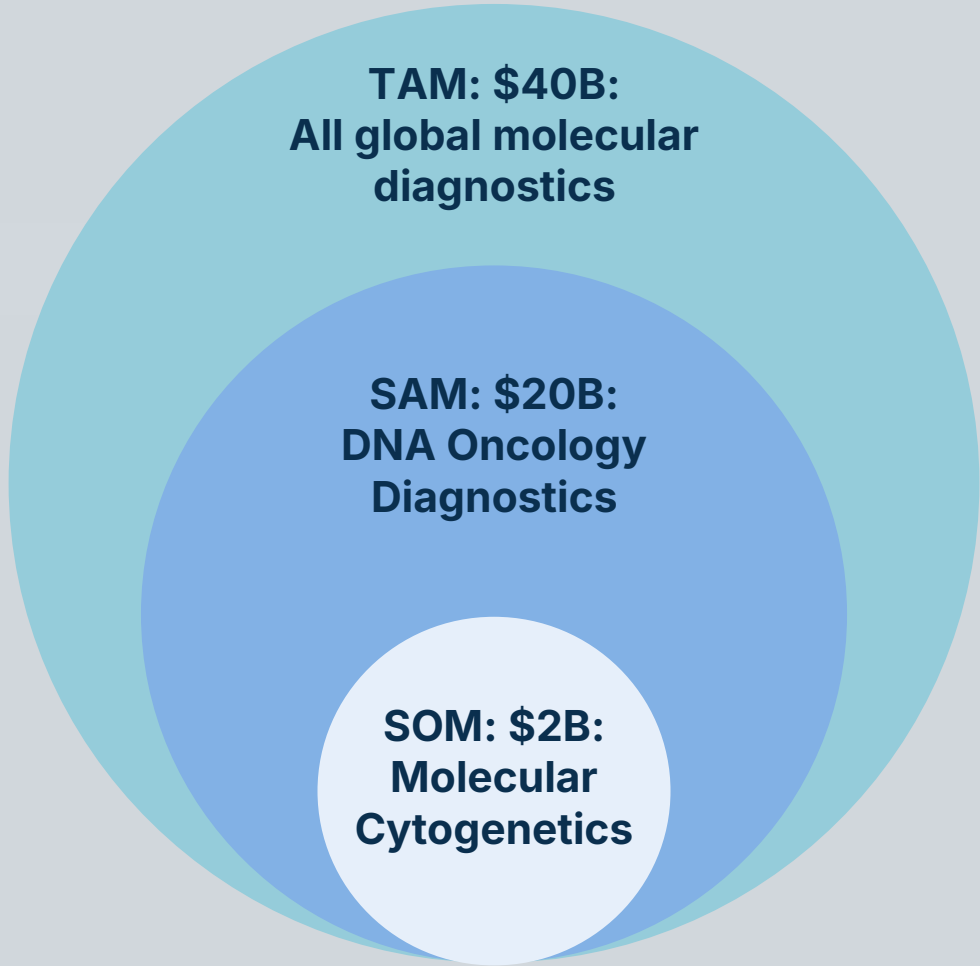
Q4 2027 — AI CP Release

\$40B Market Opportunity

Hospital Market: EU + USA

Country	No. of Hospitals
Germany	1,800
France	1,200
Italy	800
Spain	730
Poland	600
Portugal	200
Denmark	50
Other EU	3,620
The USA	6,000

Market Segmentation



OaCP Competitive Advantages: Faster – Better – Cost-Effective



Metric	OaCP IE Ltd	Abbott	Agilent	Cytocell
Time to Diagnosis	2 hours	1 working day	1+ working day	1+ working day
Overall Test Cost Reduction	Up to 50%	None	None	None
No Switch Costs	✓ Yes	X No	X No	X No

B2B Sales Model: Global Distribution Network

Direct Customers

Hospitals, Clinical Laboratories, and Independent Distributors



CliniSciences
Group



funakoshi
FRONTIERS IN LIFE SCIENCE

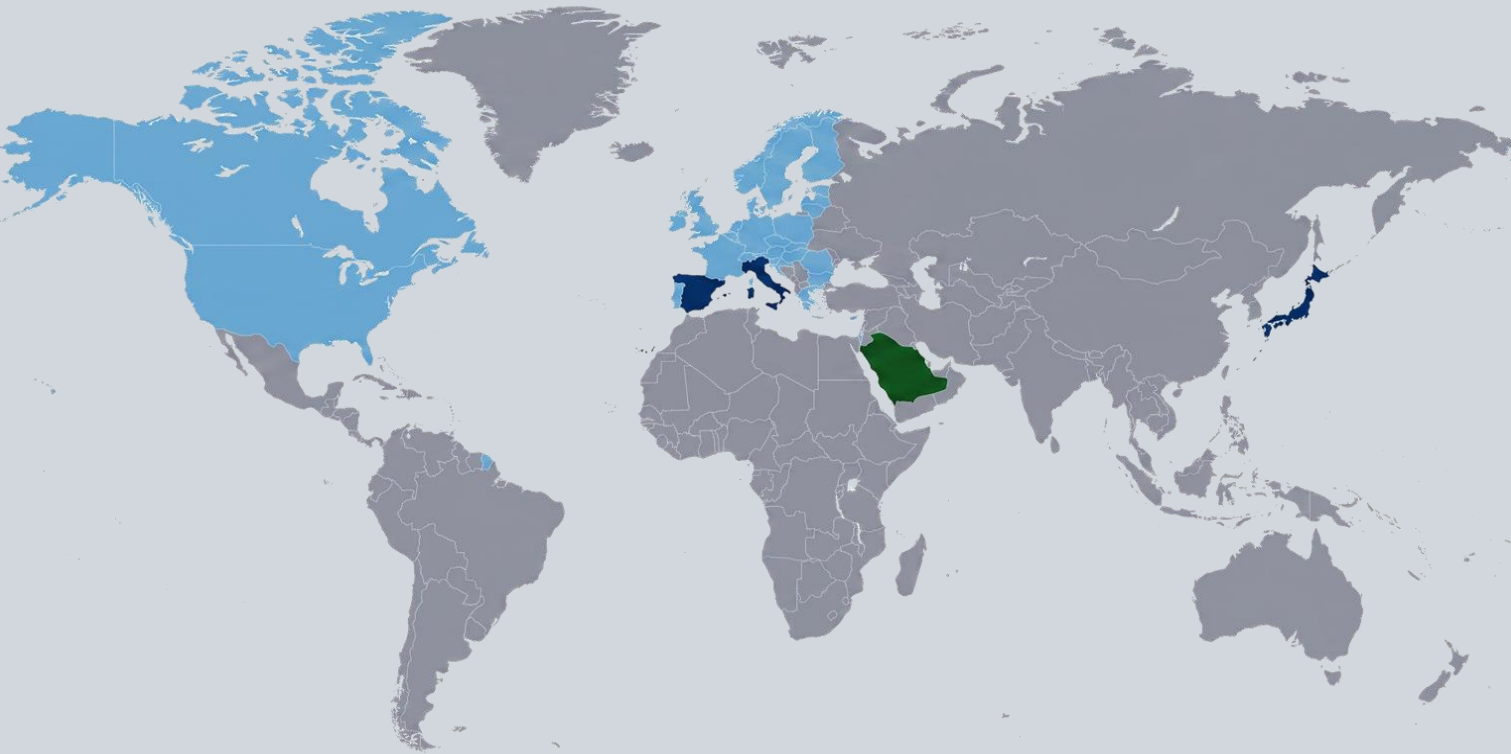
Industry Partners



EVIDENT™
OLYMPUS

ikonisys
FINDING THE CELLS THAT MATTER

Market Footprint: >\$1M Achieved in EU & Japan



- The EU



CliniSciences
Group

- The USA



- Japan



- KSA



Team: 100+ Years of Combined Experience



Enrico Di Oto
CEO

BS, MD, PhD,
18 years clinical experience.



The Business School
for the World®



创 造 无 限



Andrea Faviere
CBDO

MBA, 10+ years driving
commercial success.



Simone Di Giacomo,
CTO

BS, PhD 15+ year in molecular
biology.



Peter Mac
Peter MacCallum Cancer Centre
Victoria Australia



Nidhal Louhichi
AI Lead

Biomedical Engineer
Digital Health – AI expert.



Feliziano Balladori

35+ years institutional sales.



Matteo Stefanini

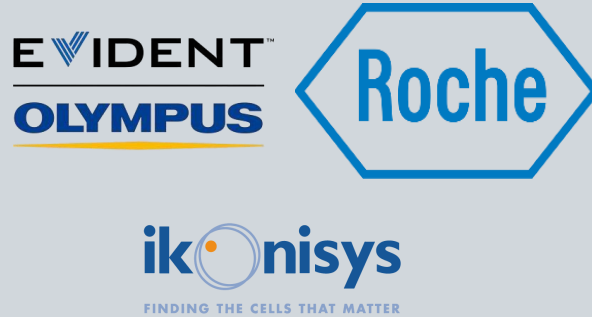
PhD in AI.



De-Risked for Explosive Growth



**>\$1M Revenue Achieved
in the EU and Japan**



**Partnership with Market
Leader**



**IVDR Certified &
Compliant
ISO 13485 Certified**



Patented Reagents



**Partnership with a
Leading University in the
USA**



**Aligned with SDG and
Vision 2030 Program**



**OACP Diagnostics
KSA RN: 24926253927**

Roadmap to Leadership & Financial Growth — Next 16 Months

Global Expansion

Deepen Current partnership
USA Market Entry
FDA 510k

AI Tool Launch

Expected for Q1 2027

Saudi Market Entry (Phase 1)

Al-Borg Labs / KAUST / KFSHRC
partnership

SFDA & Product distribution

SUPPORTED BY

Bio4Dreams



وزارة الاستثمار
Ministry of Investment

منشآت
monsha'at
الهيئة العامة للمنشآت الصغيرة والمتوسطة
Small & Medium Enterprises General Authority

ابدأ
EBDA

The Ask: \$2.7M

45%

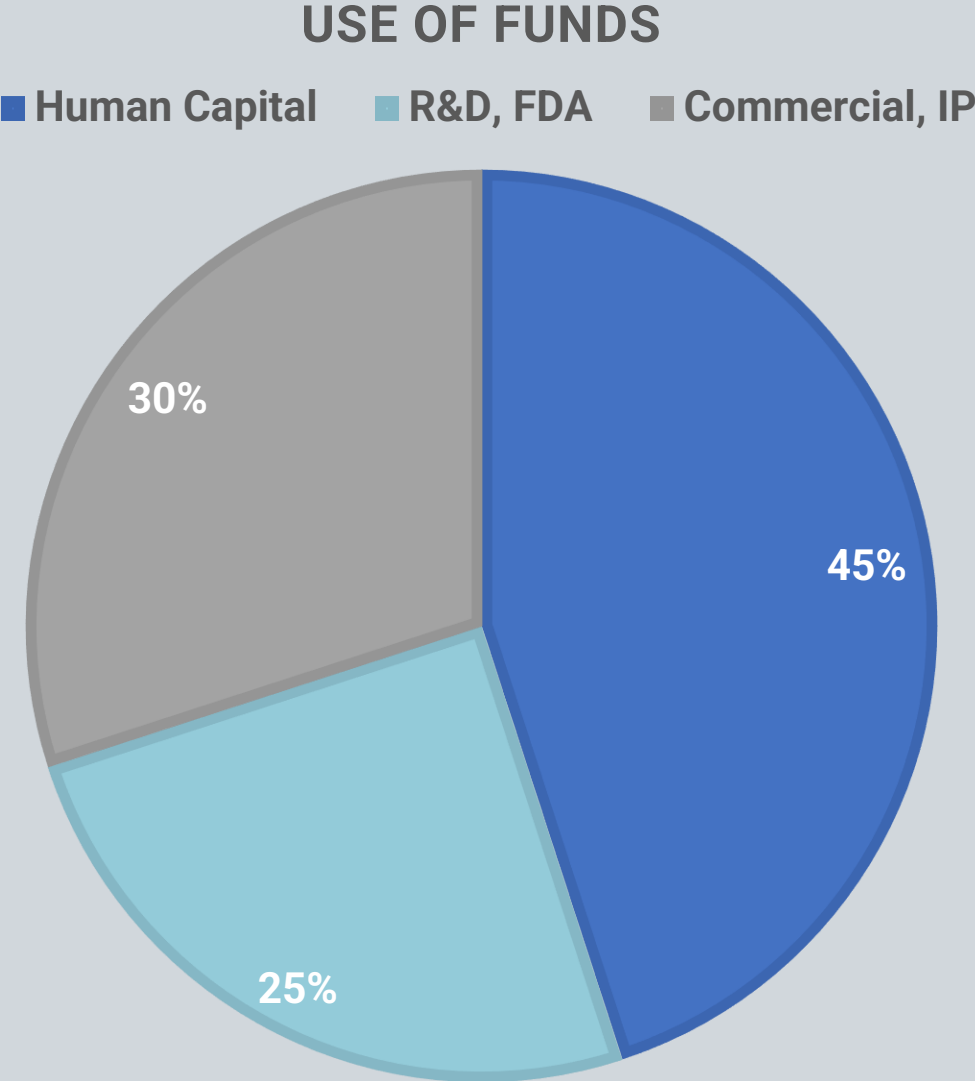
Human Capital

30%

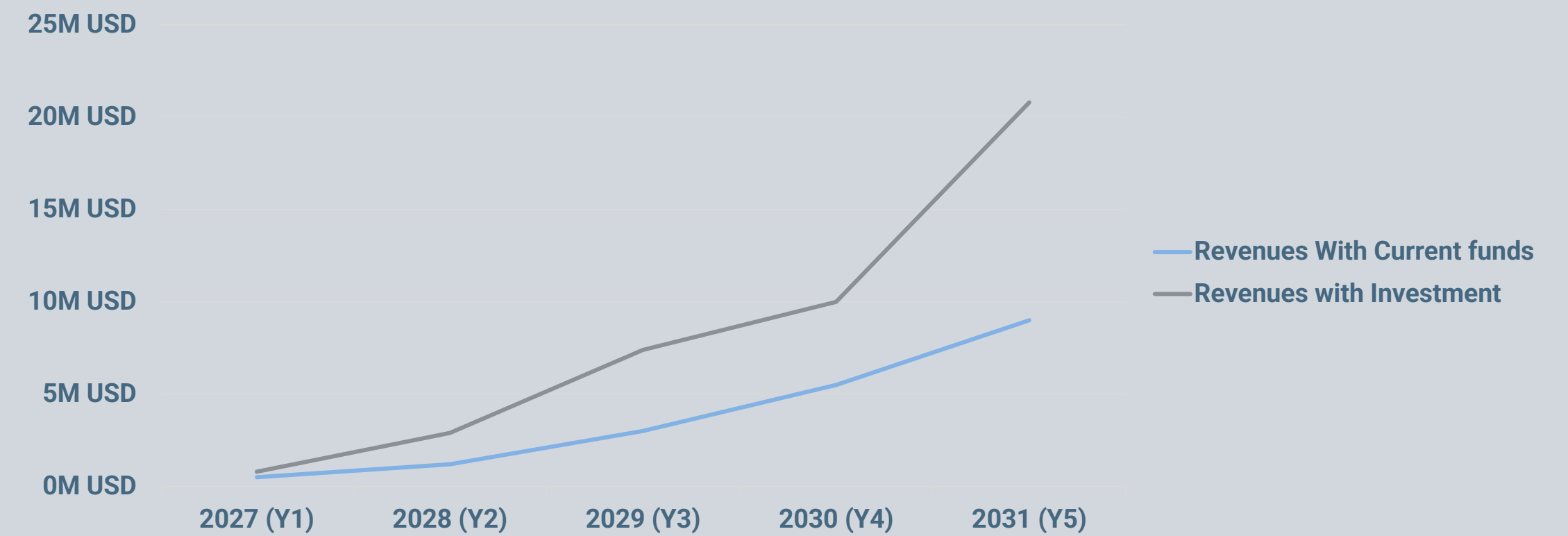
Commercial & IP

25%

R&D, FDA & Roche



OaCP Revenues Gap Analysis



Year	Revenues With Current funds	Revenues with Investment	The Gap
2027 (Y1)	USD 0,5 M	USD 0,8 M	+60% (Early US pilots)
2028 (Y2)	USD 1,2 M	USD 2,9 M	+125% (Network Labs USA)
2029 (Y3)	USD 3,0 M	USD 7,4 M	+130% (Scale-up nazionale USA)
2030 (Y4)	USD 5,5 M	USD 10,0 M	M&A Target Range
2031 (Y5)	USD 9,0 M	USD 20,8 M	Exit Strategy Peak



OACP IE LTD

**Less Time for Diagnosis
More Time for Life**

CEO@OaCP.IT | WWW.OaCPDIAGNOSTICS.COM | @OaCPIELTD

Three Adoption Paths. One Flexible Architecture.

Designed to integrate into any existing lab environment — Roche, Leica, Agilent — without disruption.

OPTION A

Rapid-ISH Chemistry

Reduces test turnaround from **3 days to 2 hours**. 50% cost savings. ISO 13485 certified. Compatible with any manual or automated platform.

Sample Prep → OaCP Rapid Buffer (40-min Hybridisation) → Pathologist Review

OPTION B

AICP Digital Co-Pilot

Automated QC for any pathology lab or OEM microscope provider. Eliminates protocol errors, hyper-digestion, and diagnostic variability.

Slide Imaging → JPG Upload → AICP Analysis → Staining/Protocol Report

OPTION A + B

Full Integrated Stack

Same-day diagnosis with guaranteed protocol standardisation. 2-Hour wet-lab processing + 30-second AI QC. Highest clinical margins at **+56.4%**.

Maximum synergy. Maximum clinical value.

HARDWARE-AGNOSTIC

INDEPENDENT COMMERCIAL STREAM

ISO 13485 CERTIFIED

Three Stakeholders. Three Transformative Benefits.

24/7 EXPERT COPILOT

For Pathology Labs & Technicians

Instant QC feedback detects staining failures, hyper-digestion, and paraffin residue in seconds. Exact corrective parameters provided in real time — empowering junior technicians to achieve senior pathologist accuracy.

COST & RISK REDUCTION

For Hospitals & Health Systems

Eliminates thousands in wasted reagents from repeat testing. Compresses patient turnaround to enable same-day treatment decisions. Ensures identical QC metrics across all satellite network laboratories.

LIFE-SAVING ACCURACY

For Cancer Patients

Eliminates inconclusive FISH results that delay treatment. Guarantees accurate targeted therapy assignment through standardised diagnostic signals.

📄 Validated on **30,000+ clinical cases**, developed with support of the University of Michigan.

The Ask

A single, structured raise. Four concurrent workstreams. One clear trajectory to Series A.



\$2.7M Pre-Series A

18–20 month runway across all four workstreams executing simultaneously.



18–20 Month Runway

Designed to reach FDA clearance, Roche adapter launch, and KSA contract execution before Series A.



Series A-Ready Milestones

US clearance, recurring hospital contracts, and IP ownership create a compelling, de-risked Series A narrative.

Backed by ISO 13485, MISA KSA Licence, University of Michigan endorsement, and a €200K bridge tranche led by Bio4Dreams SPA — this round deploys into validated, de-risked commercial infrastructure.