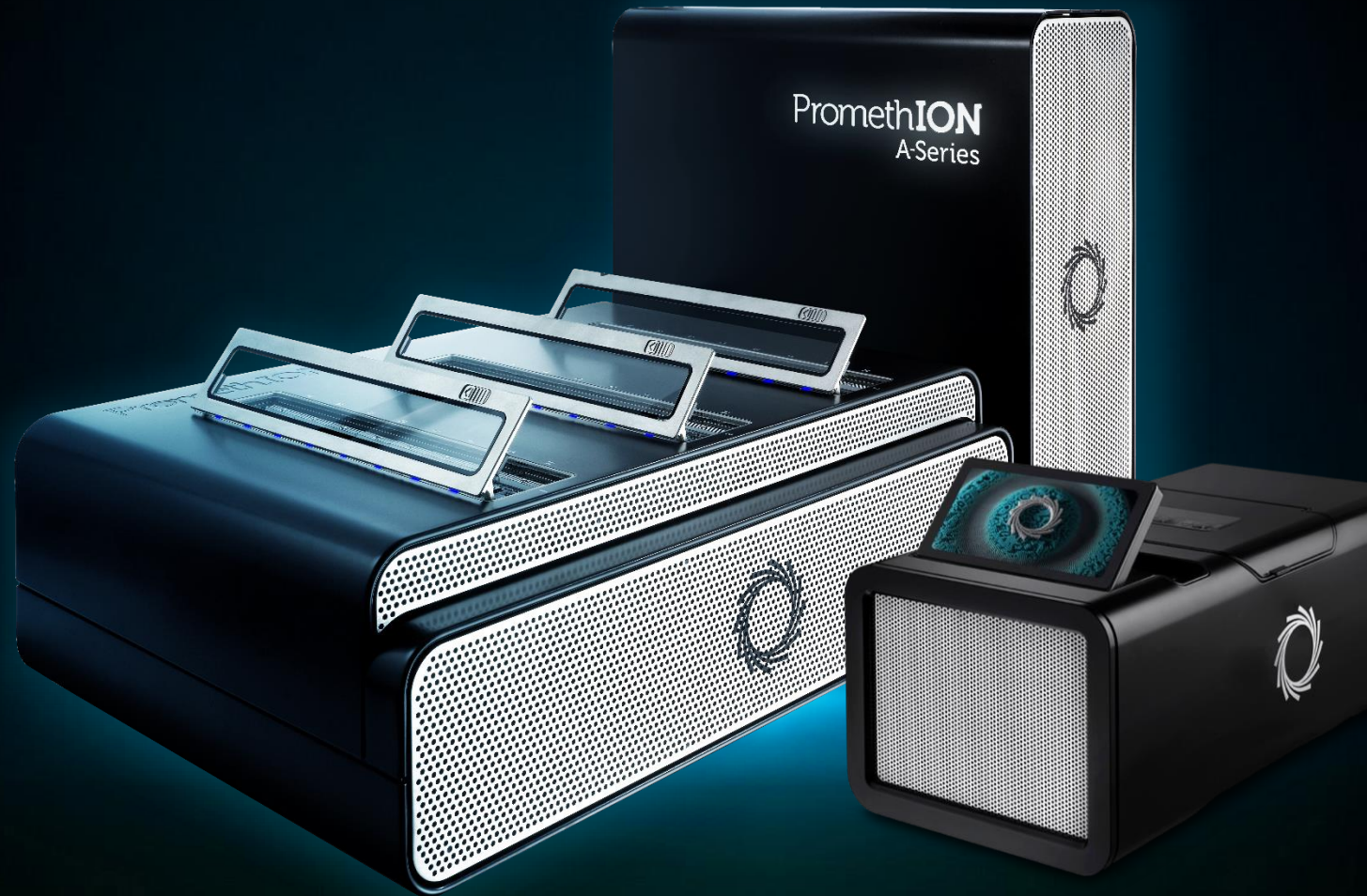


2026 Interim Results Presentation

19 August 2026



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This presentation and the discussion which follows it may contain statements that are forward-looking. For example, statements regarding expected revenue growth and profit margins are forward-looking statements. Phrases such as "aim", "plan", "expect", "intend", "anticipate", "believe", "estimate", "target", and similar expressions of a future or forward-looking nature should also be considered forward-looking statements. Forward-looking statements address our expected future business and financial performance and financial condition, and by definition address matters that are, to different degrees, uncertain.

Our results could be affected by macroeconomic conditions, delays in our receipt of components or our delivery of products to our customers, suspensions of large projects and/or changes in the timing of large projects, or changes in the adoption of pathogen surveillance technologies. These or other uncertainties may cause our actual future results to be materially different from those expressed in our forward-looking statements.

Agenda

01. Opening remarks | Francis Van Parys, CEO

02. H1 performance & FY26 guidance | Nick Keher, CFO

03. Corporate strategy & outlook | Francis Van Parys, CEO

04. Closing remarks | Francis Van Parys, CEO

05. Q&A | Francis Van Parys, CEO & Nick Keher, CFO



Opening Remarks

Francis Van Parys, CEO

Setting the direction for the next phase

First 6 months: Listening, learning and setting priorities

01

Customers

Listening to customers and partners to understand adoption, workflows and unmet needs

02

Operations

Understanding R&D, manufacturing, supply chain and commercial execution

03

Strategic refresh

Comprehensive review of markets, applications, innovation priorities and capital allocation

04

Organisation

Assessing leadership capability, operating processes and how the business scales from here



Francis Van Parys opens London Calling 2026, welcoming more than 3,000+ attendees both in person and online from 105 different countries.

Clear near-term path to profitability and cash generation

FY26 guidance unchanged; on track for adjusted EBITDA breakeven in FY27 and positive free cash flow in FY28

FY26

16–20% CC revenue growth*

~62% gross margin

FY27

Adjusted EBITDA breakeven

FY28

Positive free cash flow

*Excludes \$20m upfront payment from cross-licensing agreement

H1 2026 Performance

Nick Keher, CFO

H1 26 performance

Strong Margin and adjusted EBITDA progression; revenue below our expectations

Revenue

£116.7m

(H1 25: £105.6m)

Gross margin

62.2%

(H1 25: 58.2%)

Adjusted EBITDA

£(22.1)m

(H1 25: £(48.3)m)

Revenue growth

12.3% CC

(10.5% reported)

Adj. OpEX

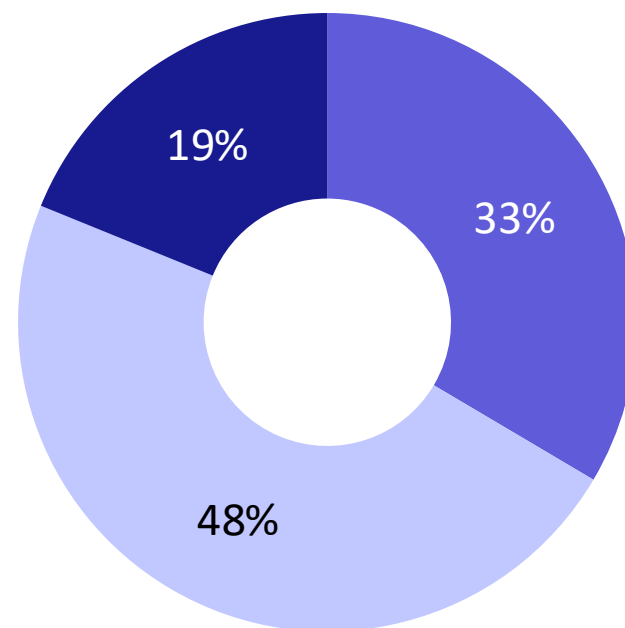
£(123.4)m

(H1 25: £(132.6)m)

Balance sheet*

£234.5m

(31 Dec 25: £302.8m)



AMR
£39.1m
+12.5% CC

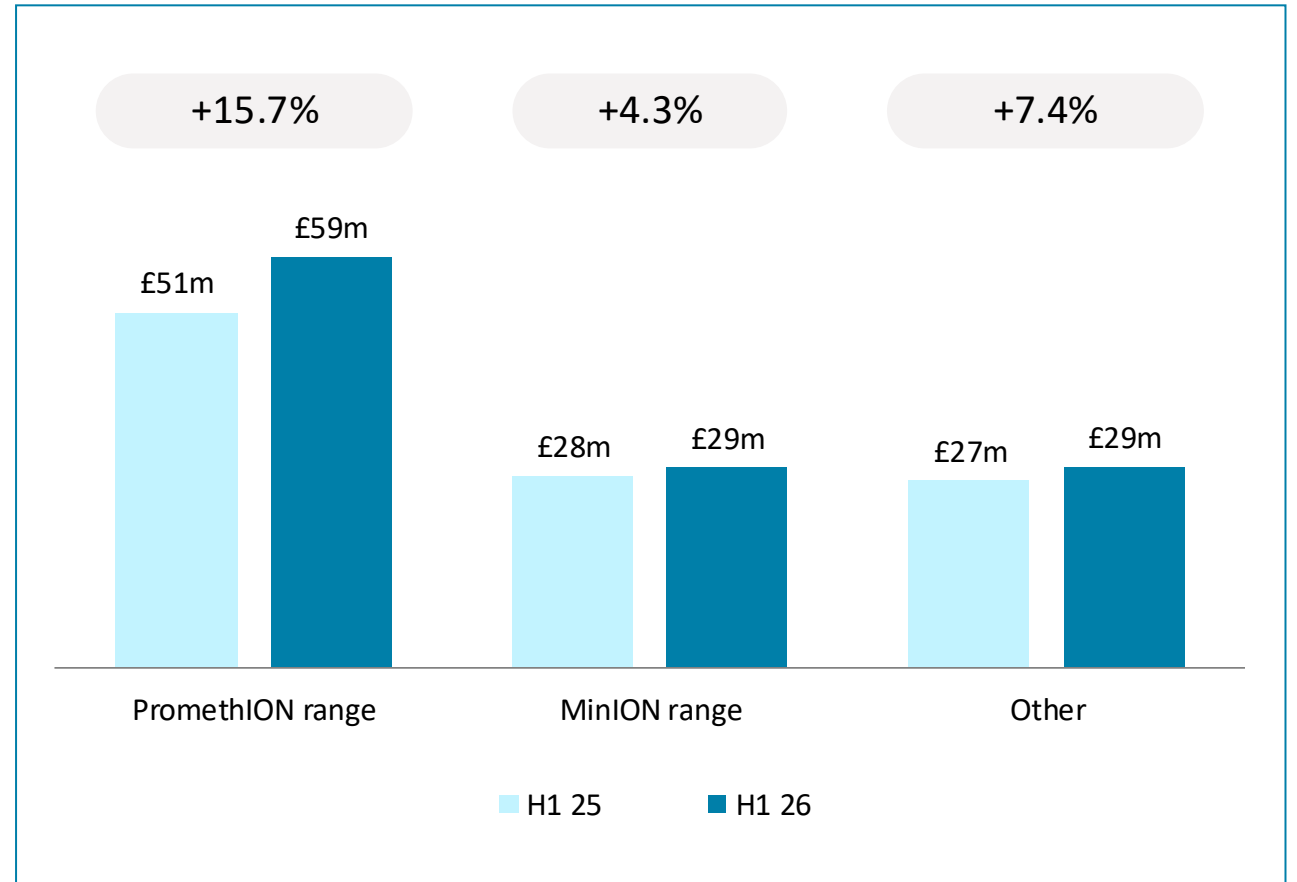
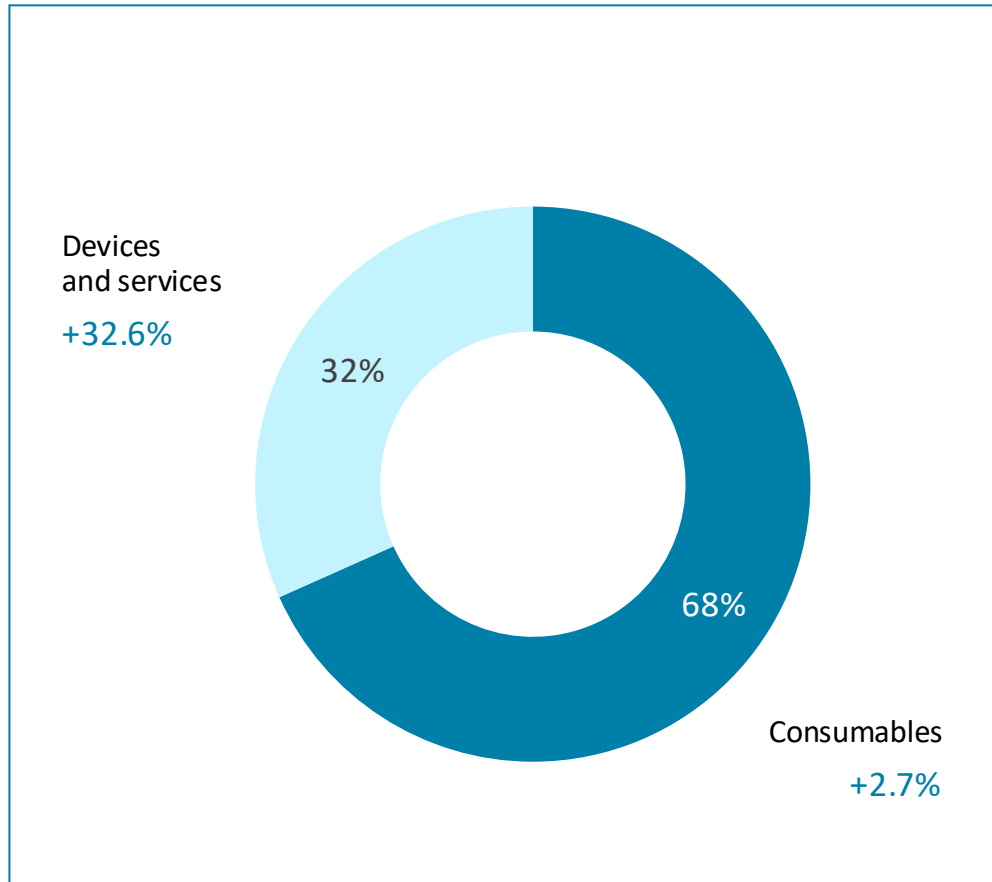
EMEAI
£55.6m
+23.8% CC

APAC
£22.0m
(8.4)% CC

* Cash, cash equivalents and other liquid investments

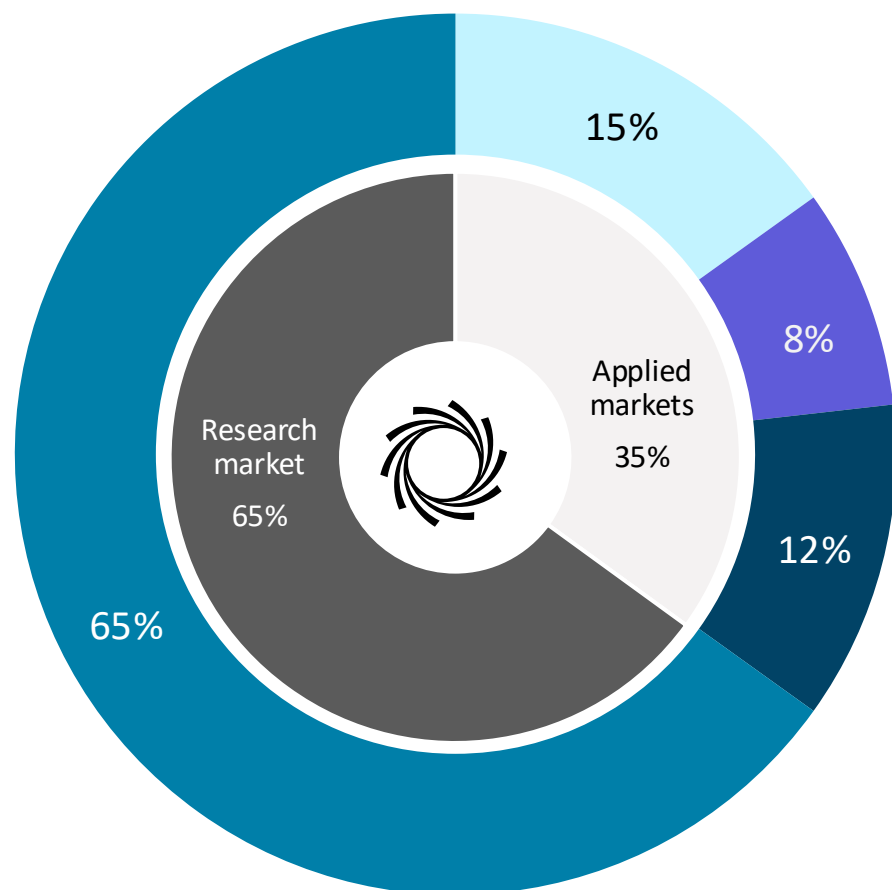
Growth driven by PromethION range

Strong device sales, slower consumables growth expected to be temporary in nature



The MinION product range includes all MinION and GridION devices and MinION Flow Cells, the PromethION range includes all PromethION devices and PromethION Flow Cells. Other revenue includes kits, services and other devices.

H1 26 revenue split by customer type



Research¹ £76.0m (+5.4% YoY)

Customers funded by grants or public funds to conduct research, e.g. academic research institutes with the primary aim of publishing breakthrough science.



Clinical £17.6m (+35.4% YoY)

Customers funded by reimbursement, using either proven assays, or developing new methods for clinical use, e.g. clinical labs for rare disease.



BioPharma £9.5m (+25.0% YoY)

Customers funded to develop, manufacture and sell biopharmaceuticals, e.g. BioPharma making and manufacturing RNA vaccines, cell and gene therapy etc.



Industrial £13.7m (+6.2% YoY)

Customers utilising sequencing for application in industrial or service setting, e.g. manufacturing or outsourced Synthetic biology.



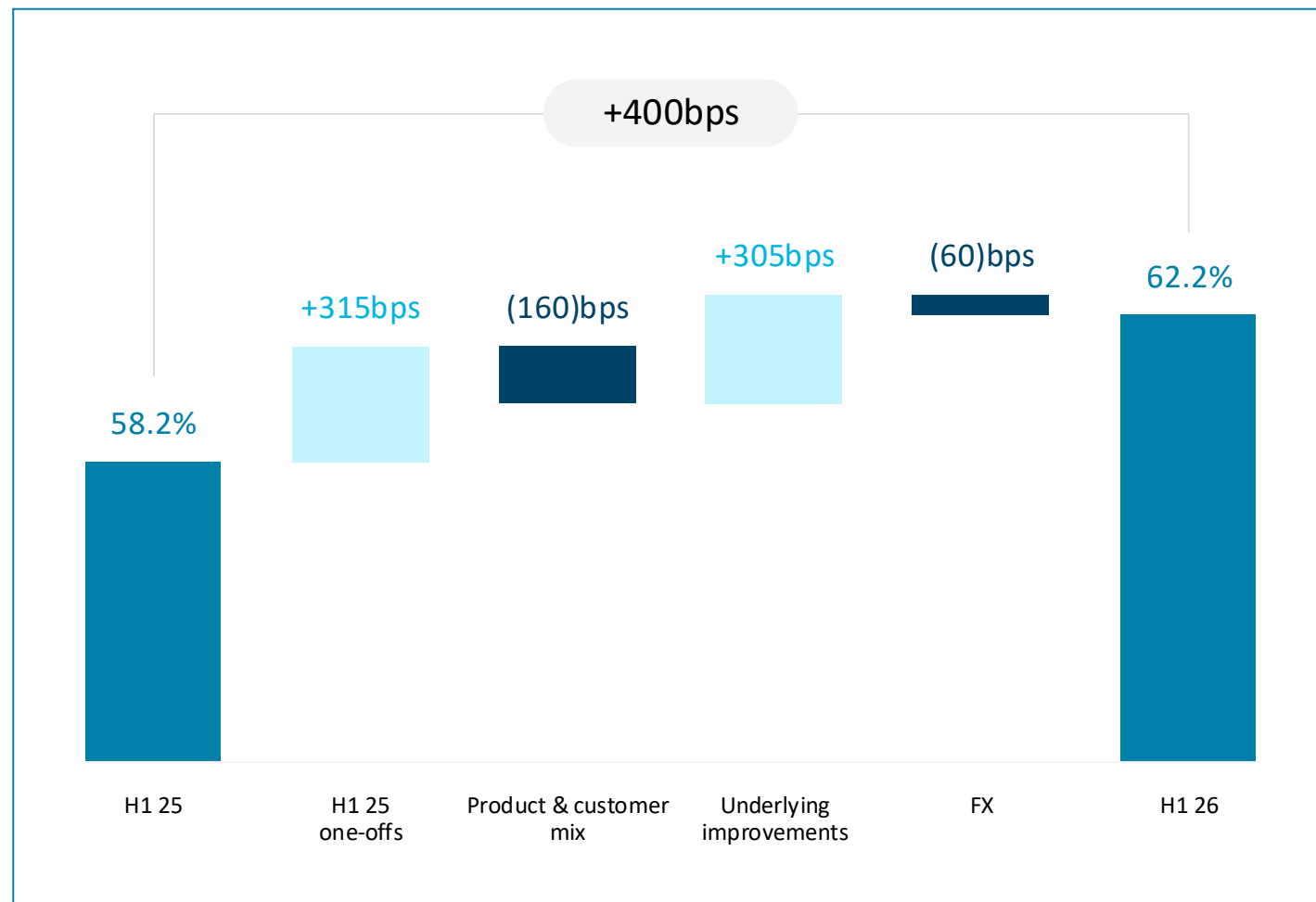
Research includes Government and Distributors

Revenue is split by customer end market categorisation – i.e. the end-market of the company buying Oxford Nanopore Technologies products

Gross margin bridge

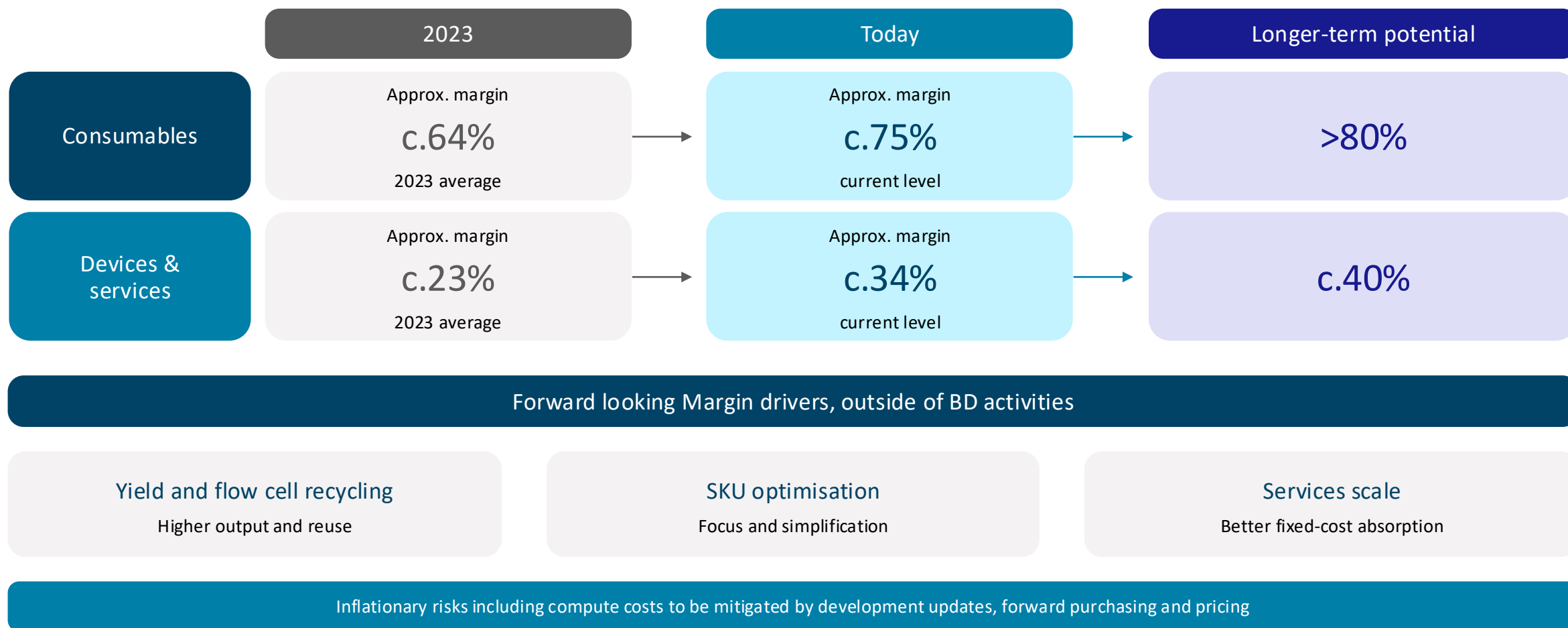
Strong margin expansion in the period

- H1 25 adversely impacted by a one-off non-cash, inventory charge of £3.3 million not repeated in FY26
- Adoption of Capex model alongside Yield improvements and recycling on Consumables drove significant underlying improvement in margin
- Further benefits to PromethION FC margins from recycling and yield improvements to be realised
- Improvements slightly offset by currency (expected to moderate at current rates) and mix (to continue)



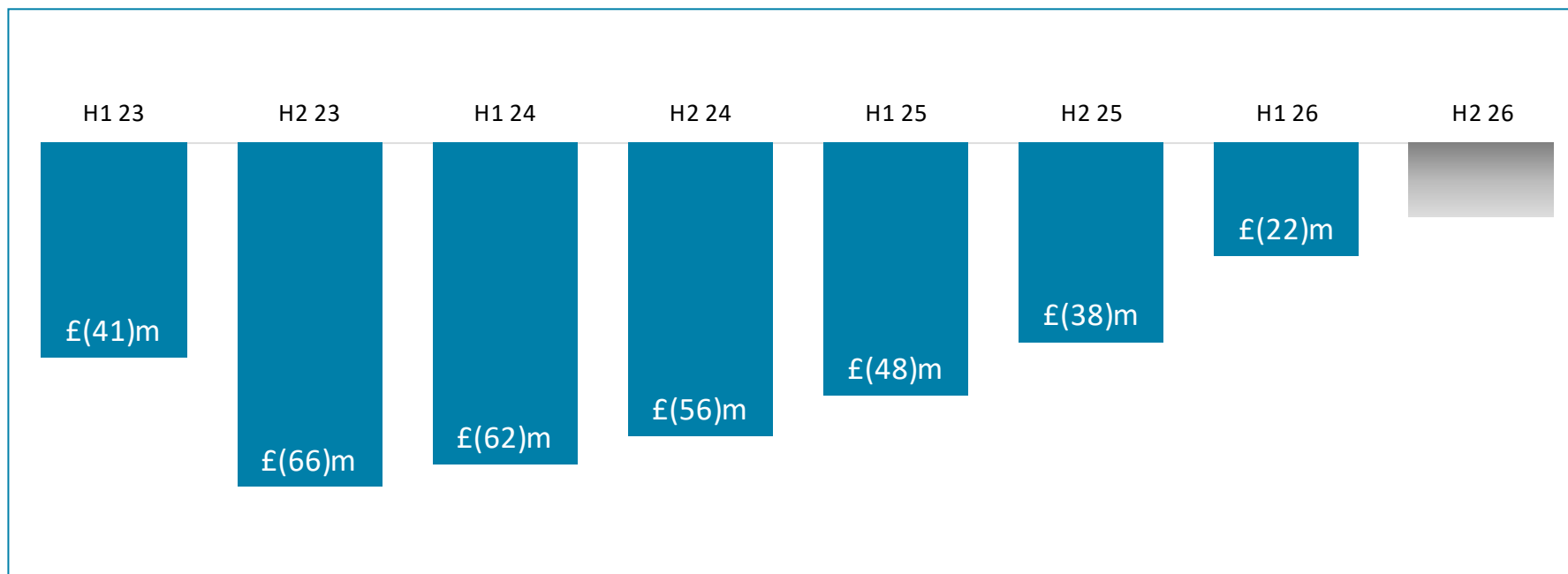
Notable gross margin progression, with further potential over time

Pricing, yield and scale have improved product economics; further gains remain available



Solid progress on pathway to breakeven

Adjusted EBITDA improvement driven by strong revenue growth and disciplined cost control



FY27

Reach breakeven

Key levers

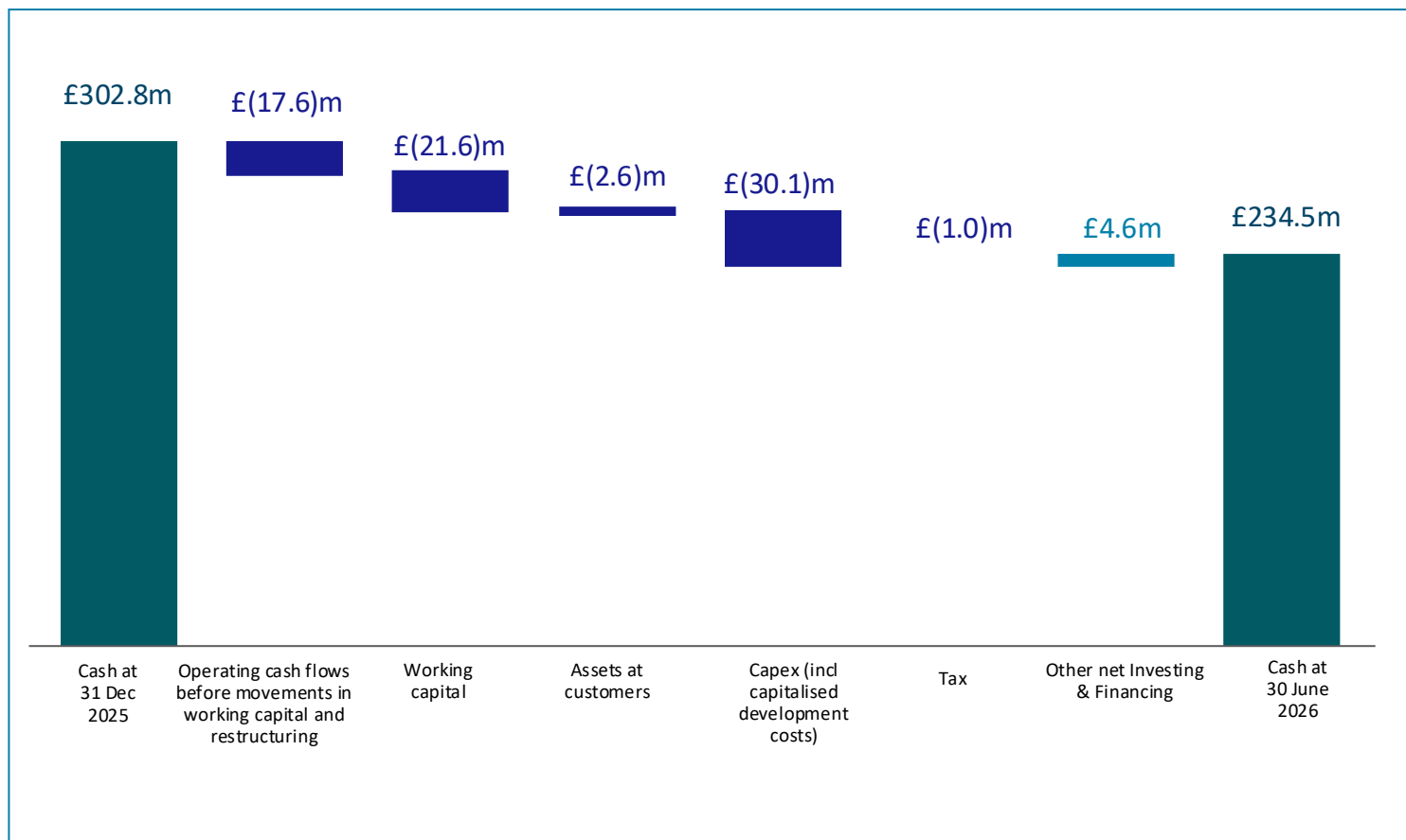
- Continued above market revenue growth; underpinned by expansion in Applied markets
- Gross margin expansion initiatives
- Cost control

+54% YoY improvement to adjusted EBITDA loss driven by:

- Strong gross profit growth
- Operational efficiency and Strategic realignment programmes in FY25
- Good cost control in period; Adjusted Opex down 7% YoY

Cash, cash equivalents and other liquid assets

Cash bridge FY 2025 to H1 2026



H1 reflects seasonality:

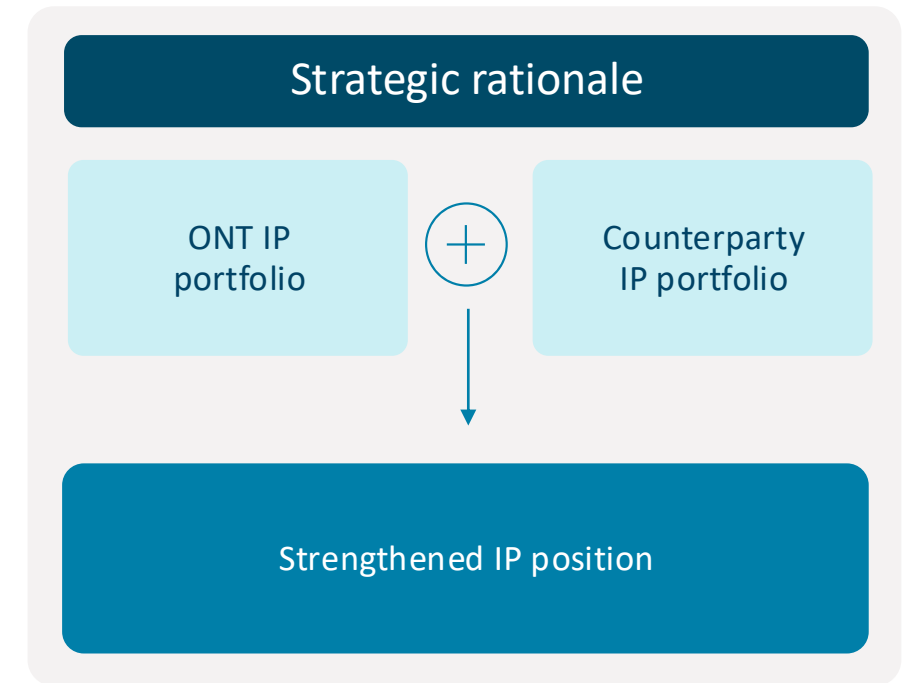
- Operating Cashflows before movements in working capital improve by £27.2m to £(17.6)m (H1 2025: £(44.8)m).
- Working capital outflow of £21.6m driven by payables £18.2m which included bonus payments of £25.7m related to FY25, and £4.0m increase in receivables offset by reduced inventory levels £0.6m.
- Capitalised development costs of £30.1m reflects R&D spend of £24.0m, licence and Patents £4.0m and PPE of £2.1m.

New cross-licensing arrangement with global diagnostics company

Long-term economic benefit with enhanced IP position

FY26	FY27 - FY28	Ongoing
\$20m	\$15m	Low to mid single digit royalty
Upfront revenue 100% gross margin	Total revenue Across the two years related to product purchases	Royalty linked to counterparty revenue For the life of the patents

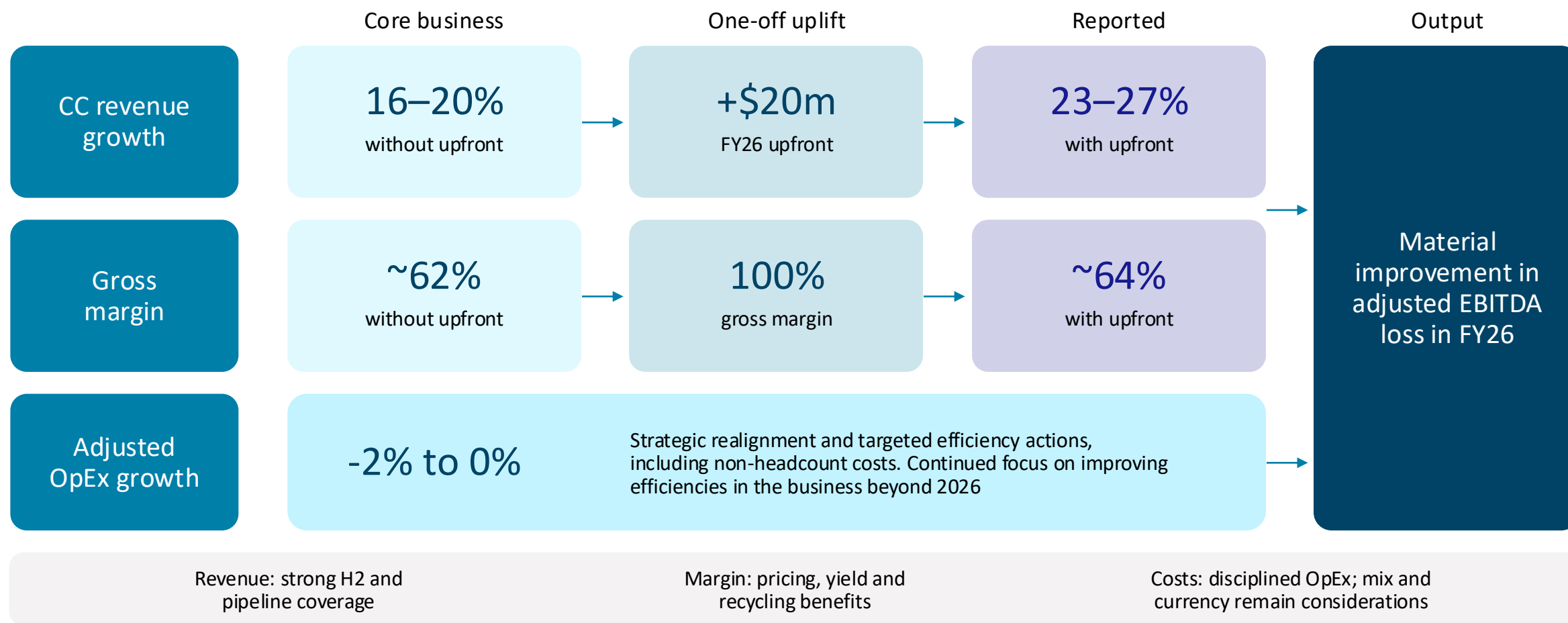
Note: FY26 16-20% CC growth excludes the \$20m upfront payment. Royalties are additive as earned but not included in guidance.



FY26 guidance

16

Strong underlying growth and operating leverage, with a one-off cross-licensing benefit



Corporate strategy

Francis Van Parys, CEO

Why I believe Oxford Nanopore can win

A differentiated platform with a clear path to broader adoption and durable performance



The platform

Differentiated
technology with
global reach



The opportunity

Move from research
into routine use



The potential

Convert
innovation into
customer value



The goal

Build durable
performance
over time

In the long-term, Oxford Nanopore's sensing technology can integrate information about the molecules of life with AI-driven information systems of the future, towards a healthier world, enabled by biological intelligence

Strong foundation, clear areas to build

19

Clear priorities to turn strong foundations into broader adoption and profitable growth

Where we are today

Adoption below potential

Differentiated technology,
not market-led

Broad opportunity,
insufficient discipline

Strong teams, capability
gaps to close

What we will do

Focus on priority applications
and customer experience

Market-led roadmap tied
to customer value

Sharper portfolio and
go-to-market choices

Strengthen leadership, capability
and accountability

What this can deliver

Faster, broader adoption

Technology leadership converted into
product leadership

Predictable, profitable growth

An organisation built to scale

Clear, focused strategy to drive sustainable, profitable growth

20

01



Customer-centric growth

- Prioritise high-value applications where we can win
- Demonstrate clear customer value
- Deepen relationships to accelerate adoption

02



Focused innovation

- Innovate for customer needs
- Deliver robust, easy-to-use products
- Invest where value is greatest

03



Disciplined execution

- Simplify our portfolio, structure and go-to-market
- Standardise processes, ownership and decision-making
- Deliver a consistent customer experience

04



High-performance culture

- Build energised, accountable and collaborative teams
- Build leadership depth and critical capabilities
- Retain agility and innovation as we scale



01 Customer-centric growth

Sharper focus on selected high-value applications and clearer value propositions.

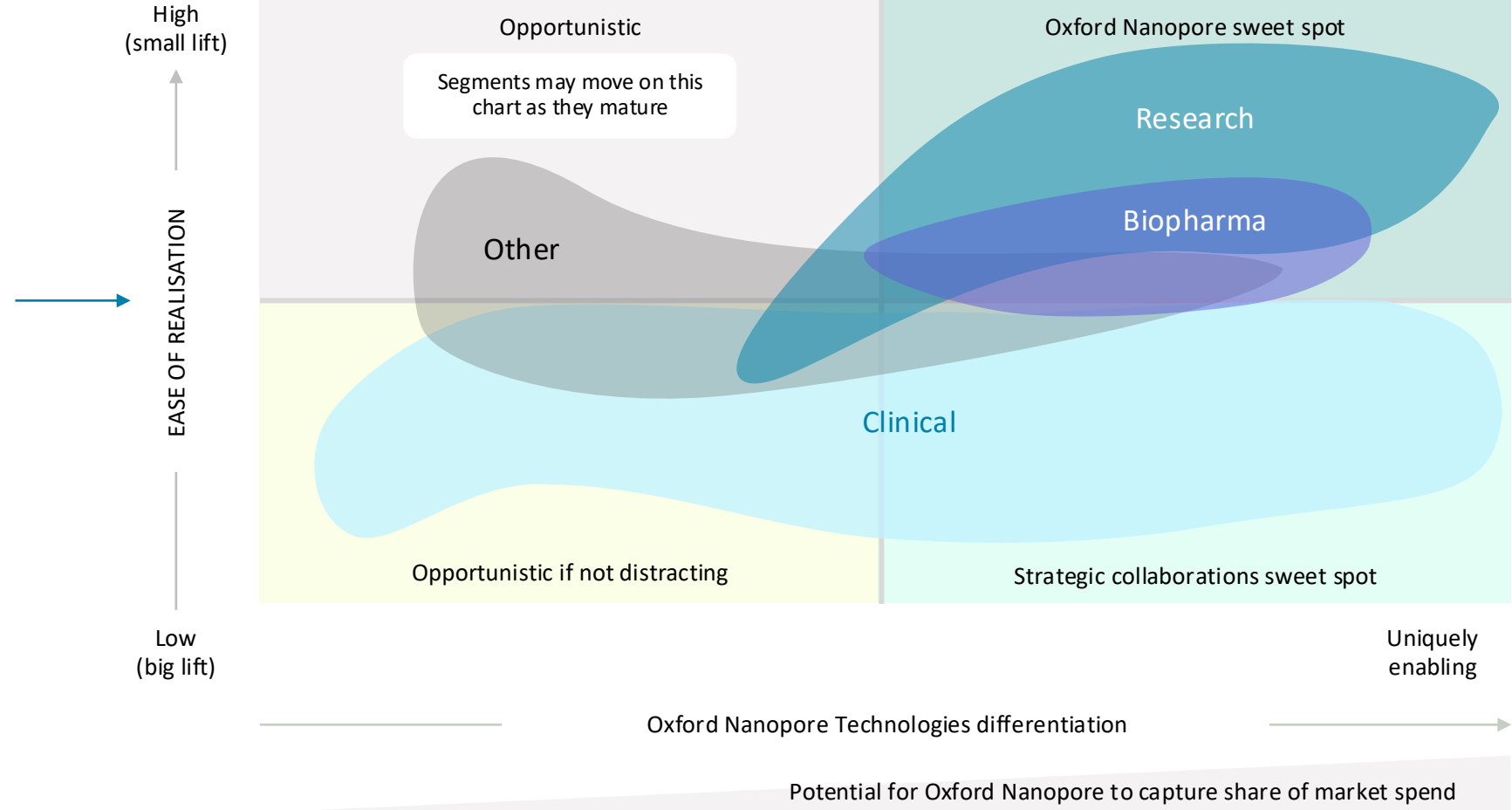
Differentiation and ease of realisation informs our participation model

ILLUSTRATIVE

EVALUATION MATRIX

Factors contributing to ease of realisation:

- Solution performance
- Form factor
- Evidence
- Regulation
- Reimbursement
- Education and market building requirements



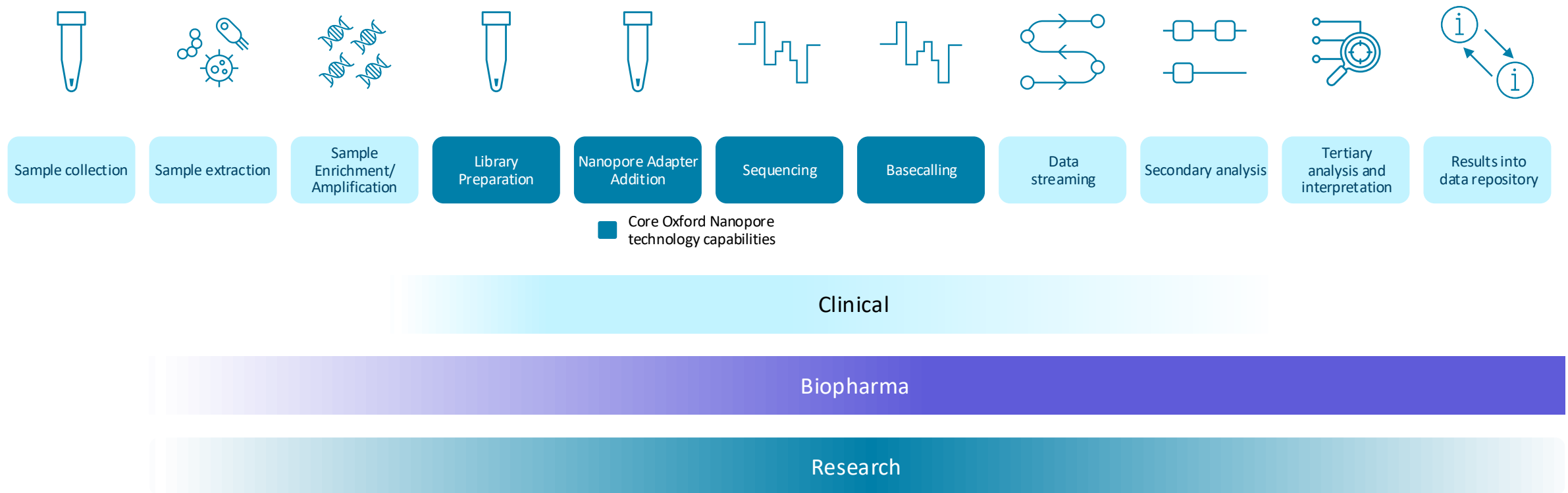
Clear participation model informs how we partner

Route to market dictated by fit and ROI

Category	Model	Our role	Partner role	How we earn	Investment	Best fit
Tech enablement	Out-licence	None	Develops and commercialises	Royalties	\$	Opportunistic
Strategic partner	3rd-party compatible	Off-the-shelf compatibility	Builds on our platform	Product sales	\$	Clinical
	OEM partner	Components made to fit	Full solution using Oxford Nanopore Technologies	Pricing plus royalties	\$\$	Clinical
	Co-development	Shared build	Shared build	Sales plus royalties	\$\$\$	Clinical <i>possible</i>
	Commercialisation partner	Build and own solution	Provides channel	Sales less channel margin	\$\$\$\$	Research, BioPharma R&D
Full Oxford Nanopore Technologies solution	Go-it-alone	Build and own solution	None	Product sales	\$\$\$\$\$	BioPharma R&D and QC

Focused partnering strategy across the value chain by end market

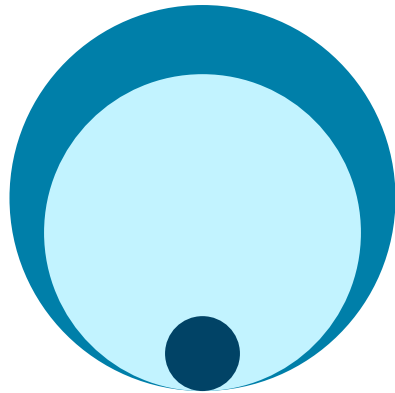
Sample prep, automation and analysis partners extend our utility across markets



A focused path to 2030

Clear focus on high-value applications, across BioPharma, Clinical and Research markets

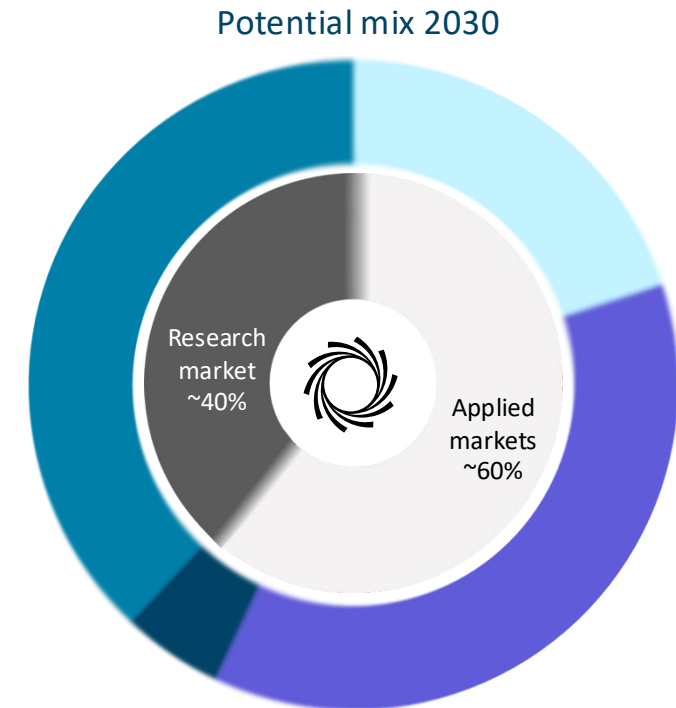
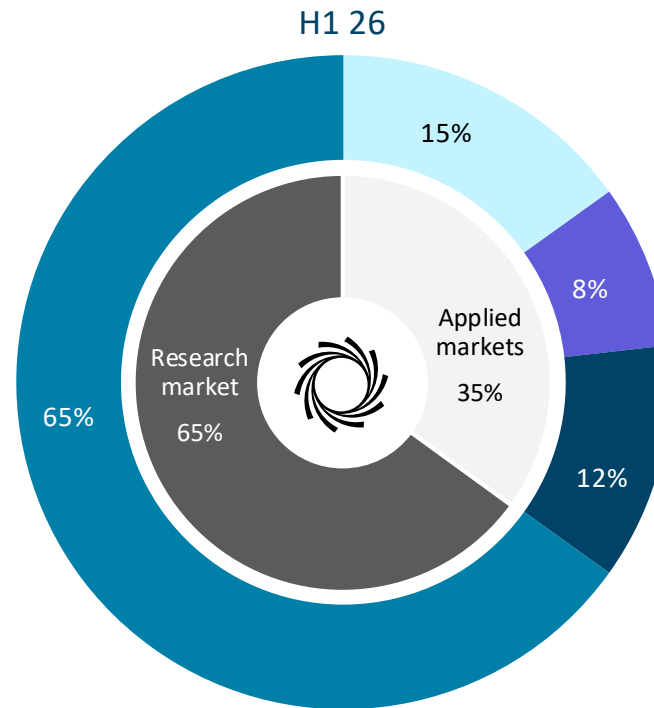
Market opportunity funnel



c.\$20–25B
Total market (TAM)

c.\$14–16B
Serviceable market (SAM)

>c.\$700m
2030 revenue opportunity



Research Clinical BioPharma Industrial

Partnerships could unlock further opportunity,
with the potential to be accretive to top-line growth

Research

Winning where complex genome and methylation insight change the answer

Target applications represent a SAM
of
c.\$610–740m
by 2030

Foundations for adoption

Greater throughput and
lower cost per genome

Productised bioinformatic pipelines

Demonstrate utility of long
read and methylation

Where Oxford Nanopore can win

Genome centres, biobanks, core
labs and disease-research programmes focused
on SVs, phasing, difficult loci and
native methylation.

Right to win

Native reads of any length, with phasing and
methylation in one workflow. Scales from
discovery to large cohorts

Customer requirement

Researchers need deeper biological insight than
current methods provide. Legacy workflows
often miss methylation, a key layer of disease
biology.

Participation model

Production-ready P24 workflow for genome
centres, cores and biobanks. PromethION plus
containerised pipelines, support and roadmap
visibility.

Our advantage: Sequence + methylation in a single run

Richer biological information creates differentiated, scalable products

The biology

DNA sequence

The genetic code

Instructions you are born with



Methylation

Chemical changes on the genetic code

~28m human CpG sites can switch genes on or off



Same DNA, different outcome

Methylation can distinguish and classify healthy from diseased cells.

Oxford Nanopore advantage

Sequence variants
+
methylation
in a single run

Creates differentiated products
and new clinical applications
without an additional assay

Where we are today

Revenue today

4 panels shipping

Methylation, PGx and cancer digital panels sold now; tumour profiling in H2.

Clinical example

Prader-Willi trio

Sequence variants alone gave no answer; phased methylation identified the cause.

Built-in advantage

No added cost and complexity

Same run. Same consumable. Richer information.

Biopharma

Opportunity to carve out a differentiated position within a large and growing market

Target applications represent a SAM
of
c.\$4.1–5.5bn
by 2030

Foundations for adoption

- Kit development
- Productised bioinformatics
- CFR-ready instrument path
- Head-to-head evidence
- Go-to-market plan

Where Oxford Nanopore can win

Biopharma R&D and Biomanufacturing QC
Faster, simpler workflows with richer readouts
reduce time, cost and analysis burden.

Right to win

Richer information: native long reads reveal
biological context and reduce library
complexity.

Speed: Shorter time to result.

Test consolidation: Identity, integrity, purity
and contamination in a single test.

Customer requirement

Customers need more information to advance
pipelines. Current methods rely on multiple
assays, vendors and hand-offs.

Participation model

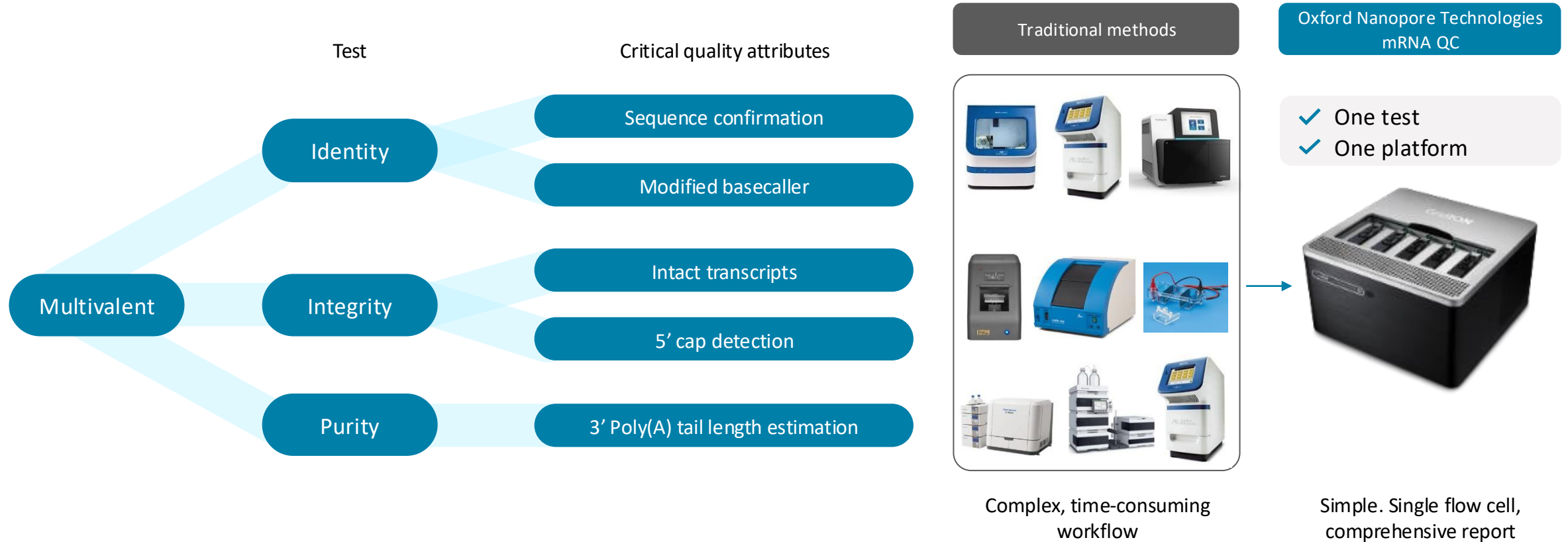
Sell direct through with production-ready
QC workflows, documentation and
support.

Use channel partners for broader pre-QC
R&D applications.

Multi-attribute mRNA QC test

34

Reducing complexity, accelerating results – one test, one platform



Clinical enablement

Building evidence where richer information and faster insights may enable future clinical applications

Target applications represent a SAM
of
c.\$6bn
by 2030

Foundations for adoption

- Locked workflows and Q-Line releases
- Clinical-grade reporting
- LIS / LIMS integration
- Evidence and health-economic utility
- Anchor-account conversion

Where Oxford Nanopore can win

Rare disease / rapid WGS, tumour profiling and selected acute ID workflows

Right to win

Speed to result

Richer information: Long reads resolve genomic variations

Richer information: Native methylation supports tumour classification.

Unique ONT features: Adaptive sampling and device format

Customer problem

Current technologies require batching or time to result outside of ideal therapeutic window

Unresolved genetic diseases require more complete answers

Adoption a multi step, gated process

Participation model

Oxford Nanopore Technologies led enablement in specialist centres, AMCs and reference labs

Partner-led IVD and channel routes for distributed acute ID workflows

Acute Leukaemia: opportunities for workflow consolidation

Conventional pathology assays consolidated into one adaptive sampling run

Conventional pathology

Flow cytometry

Karyotype

Lesion-specific PCR or FISH

Array

Targeted NGS or PCR panel

Lesion-specific PCR

Transcriptome (RNA)

Lesion-specific PCR

Eight separate assays

\$1,650–\$2,050

Consolidates into

Next generation sequencing

Oxford Nanopore DNA adaptive sampling

\$350–\$450

per sample, one run



One assay, comprehensive insights

- Lineage
- Aneuploidy
- Fusions
- Copy number variants
- Single nucleotide variants
- Tandem repeats
- Expression patterns
- Methylation
- Pharmacogenomics

Source: Alexander et al., medRxiv, 2026. Preprint, not peer reviewed. Costs reflect reagent and consumable estimates; nanopore estimate assumes multiplexing.



04 High-performance culture

Successful execution relies on ambition, accountability and collaboration



Strengthening the leadership team for scaleup

Adding scaleup leadership depth across People, Data, Tech, Medical and Marketing



Francis Van Parys

Chief Executive Officer

Joined March 2026. Previously President & CEO of Radiometer (Danaher).



Nick Keher

Chief Financial Officer

Joined January 2024. Previously CFO at both Clinigen Group and Benevolent AI.



Dr. Lakmal Jayasinghe

Chief Scientific Officer

Joined at foundation. Previously member of the Hagan Bayley group.



Tina St Leger

Chief People Officer

Joined June 2026 from Immunocore. Previously Chief Human Resources Officer at GW Pharmaceuticals.



Andrew Watson

Chief Information Officer

Joined June 2026 from Mimecast. Previously senior roles at Healx and Dyson.



Davide Manisero

Chief Medical Officer

Joined August 2026 from Radiometer (Danaher). Previously CMO at QIAGEN; roles at ECDC and the WHO.



Conor McKechnie

Chief Marketing and Communications Officer

Joining October 2026. Currently VP of Marketing at Cytiva (Danaher).



Employee feedback reinforced strategic priorities

Strong alignment across the organisation

Culture review highlighted the same themes as the strategy process:

01

Focus on fewer priorities

02

Simplify ways of working

03

Clarify accountability and speed up decisions

04

Improve collaboration across teams

05

Bringing customers closer to decision-making and product development

Employee surveys carried out in January and July 2026.

67%

Participation

75%

Positive engagement

73%

Would recommend Oxford Nanopore Technologies

Outlook

Francis Van Parys, CEO

2030 medium-term guidance

41

2030 revenue target

Organic revenue growth mid-teens+ from FY26

Accelerating towards 2030

BD / licensing / royalties

additional upside

Industrial

Clinical

BioPharma

Research

c.£260-269m
(c.\$345-355m)

16-20%

CC growth

Excludes \$20m upfront*

2026

Headwinds

China / regional volatility
Customer and contract timing
Evidence and regulatory timelines
Current mix weighted to slower-growth markets

>\$700m
Revenue by 2030

2030

2030 profitability & cash targets

>15%

Adjusted
EBITDA margin

Positive
and growing
Free cash flow
from FY28

*Excludes \$20m non-recurring revenue from the cross-licensing agreement

Capital allocation priorities

Financial discipline and framework for success

01



Organic core investment

Capitalised development
spend and manufacturing capacity
expansion

02



Partnership enablement

Assessed on SAM expansion,
workflow enablement
and ROIC >15%

03



Selective M&A

Drive adoption and
solidify position in key
target applications

Strong balance sheet

The foundation underpinning all three priorities

Ensure flexibility and reduce risk of volatility through maintaining strong balance sheet

Closing remarks

Francis Van Parys, CEO

Key takeaways

A focused strategy to deliver 2030 targets and build towards a \$1bn+ revenue business

01



Differentiated platform
and clear right to win

Focused on high value applications
where richer information, speed
and flexible deployment create
distinctive customer value

02



Clear strategy, now
being operationalised

Priority markets, market-led
innovation, disciplined execution and a
stronger organisation

03



2030 targets are a milestone
in a larger ambition

>\$700 million of revenue, an
adjusted EBITDA margin above 15%
and positive, growing FCF; building
towards \$1 billion+ revenue over
time

Q&A

Appendix

Investing for growth with improved cost control

47

£m	H1 26	H2 25	H1 25	H2 24	H1 24
Revenue	116.7	118.3	105.6	99.1	84.1
Adjusted gross profit	72.6	71.7	61.4	55.9	49.5
Adjusted R&D expenses	(36.9)	(45.4)	(44.0)	(49.9)	(49.4)
Adjusted SG&A expenses	(86.5)	(91.2)	(88.6)	(85.7)	(81.5)
Total adjusted costs	(123.4)	(136.6)	(132.6)	(135.6)	(131.0)
Depreciation & amortisation	28.7	26.5	22.9	23.5	19.8
Adjusted EBITDA	(22.1)	(38.4)	(48.3)	(56.2)	(61.7)



- Adjusted EBITDA loss improved in H1 26 due to improved margins and disciplined cost control
- Adoption of Capex model alongside yield improvements and recycling on Consumables drove underlying margin improvement. H1 25 adversely impacted by a one-off non-cash inventory charge of £3.3m not repeated in H1 26
- Adjusted R&D spend declining as focus increases on later stage development and cost discipline

Revenue breakdown by regions

48

	H1 2026 £m	H1 2025 £m	Reported growth %	Constant currency growth %	Currency impact %
AMR	39.1	36.0	8.6%	12.5%	(3.9)%
APAC	22.0	24.9	(11.6)%	(8.4)%	(3.2)%
EMEA	55.6	44.6	24.7%	23.8%	0.9%
Total	116.7	105.6	10.5%	12.3%	(1.8)%

Adjusted R&D expenses reconciliation

49

£m	H1 26	H1 25
Research and development expenses	36.9	44.1
Adjusting items:		
Employer's social security taxes on pre-IPO share awards	-	(0.1)
Adjusted research and development expenses	36.9	44.0
Amortisation of capitalised development costs	(17.9)	(12.9)
Capitalised development costs	24.0	20.1
Total R&D and capitalised development costs	43.0	51.2

Adjusted SG&A expenses reconciliation

50

£m	H1 26	H1 25
Selling, general and administrative expenses	86.4	95.1
Adjusting items:		
Share-based payment expense on Founder Long Term Incentive Plan (LTIP)	-	(2.0)
Employer's social security taxes on Founder LTIP and pre-IPO share awards	0.1	(0.3)
Restructuring costs	-	(4.2)
Adjusted S,G&A expenses	86.5	88.6

Strategy recap

51

2030 Targets

>\$700m
Revenue

>15%
EBITDA margin

01



Customer-centric growth

Sharper focus on selected high-value applications and clearer value propositions.

02



Focused innovation

Translate technology leadership into product leadership

03



Disciplined execution

Create a focused, scalable model for disciplined execution

04



High-performance culture

Successful execution relies on ambition, accountability and collaboration