

Our vision: to enable the analysis of anything, by anyone, anywhere

Gordon Sanghera, CEO

Tim Cowper, CFO

21 March 2023 | Full Year 2022 Results

THE P2: AFFORDABLE, HIGH-OUTPUT SEQUENCING FOR EVERY LAB COMPLETE WITH HIGH END COMPUTE, SCREEN FOR SEQUENCING AND ANALYSIS

P2

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Who we are

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SINGLE MOLECULE SENSING PLATFORM

Building on DNA/RNA to enable the multi-omics world of tomorrow

SUBSTANTIAL MARKET OPPORTUNITY

\$6.2¹ billion, growing existing opportunity for sequencing in 2022 with potential \$10s of billions in future clinical and applied markets

GROWING USER COMMUNITY

Customers in >120 countries doing ground-breaking science



AGILE INNOVATION

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Delivering continuous improvement and intellectual property creation

SCALED OPERATIONS

In-house manufacturing and global distribution

OUR PEOPLE

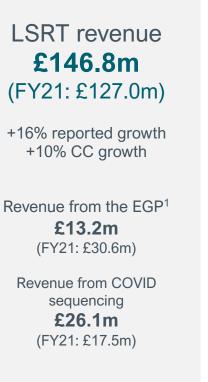
Experienced, driven leadership enabled by a highly ambitious and talented global team expanded to >1,000



3

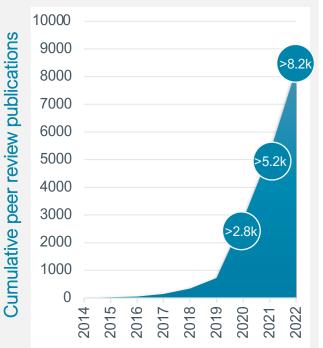
Continued robust underlying growth in core Life Science Research Tools (LSRT) business





Publications

Our accessibility and versatility is reflected in an increasingly large, international scientific evidence base²



¹The Emirati Genome Program

² publications monitoring methodology was updated between 2021-22 resulting in historical numbers increasing. FY22 COVID-19 testing revenue of £51.8m reflects the settlement of the contract with the DHSC



2022 in numbers

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¹Underlying growth – LSRT revenue minus foreign exchange and revenues from the EGP and COVID sequencing ²Following conclusion of the Covid testing contract with the Department of Health and Social Care (DHSC) ³Includes FY22 consumables revenue plus consumables revenue within starter pack sales ⁴Cash, cash equivalents and liquid investments



Delivering across all areas of our business

RESULTS

We are delivering strong revenue growth and margin growth

PLATFORM

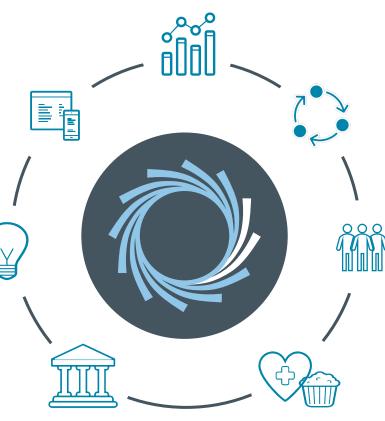
Our platform is highly differentiated from its scalability to information rich

INNOVATION

We have transformed our platform and launched new products

RESILIENCE

We have a strong balance sheet and are moving towards breakeven



EASE OF USE

We drive utilisation with simple end-to-end workflows

COMMERCIAL GROWTH

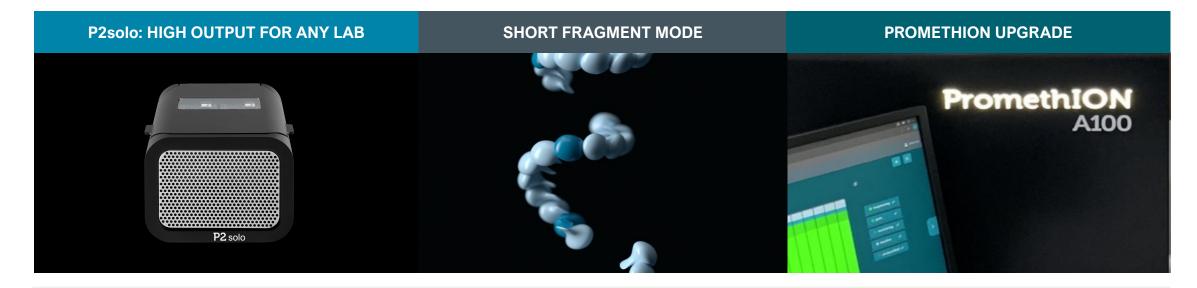
We are expanding our commercial teams to better serve our customers

BEYOND RESEARCH TOOLS

We're executing on opportunities beyond the life science research market



Innovation drives growth: key product releases



In early access Starter packs from:

\$10,455

Human genomes from \$690¹

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Select in software to sequence from

20 bases – Mbases

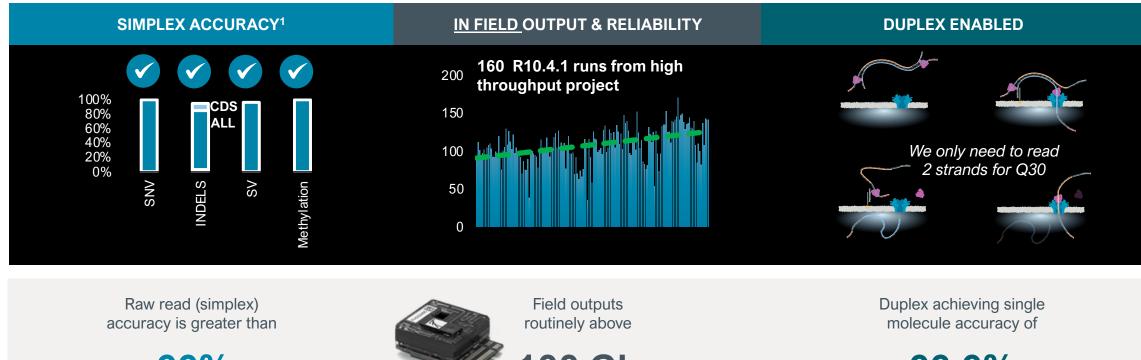
Enables cell free DNA, single cell and more. Oxford Nanopore for any-length reads A100 delivering analysis speed improvements with methylation onboard

Seamless Upgrades

High throughput human genomes from \$345¹



Innovation drives growth: high accuracy sequencing on nanopore (Q20+)



99%

Delivering high precision variant detection

8

🥙 100 Gb

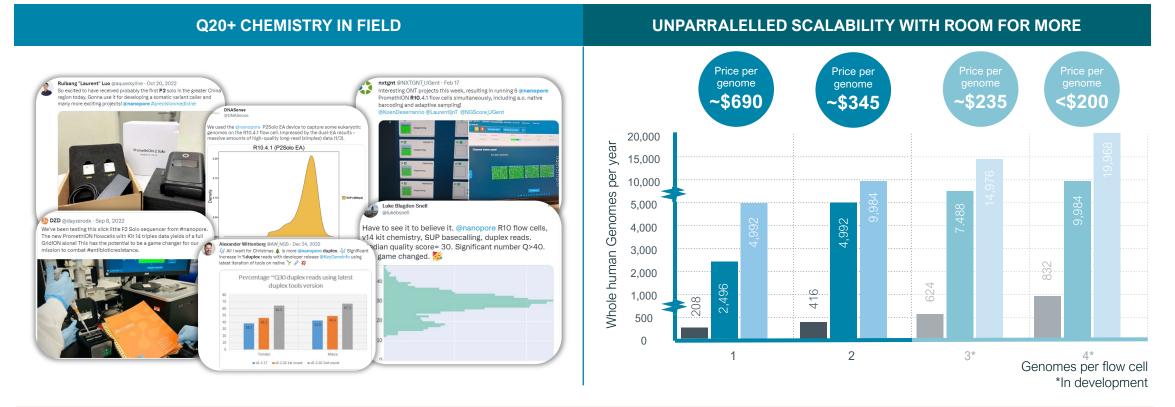
Per PromethION flow cell and increasing steadily

99.9%

Improving rare variant detection and de novo assembly



Innovation drives growth: transforming platform capabilities today and in future



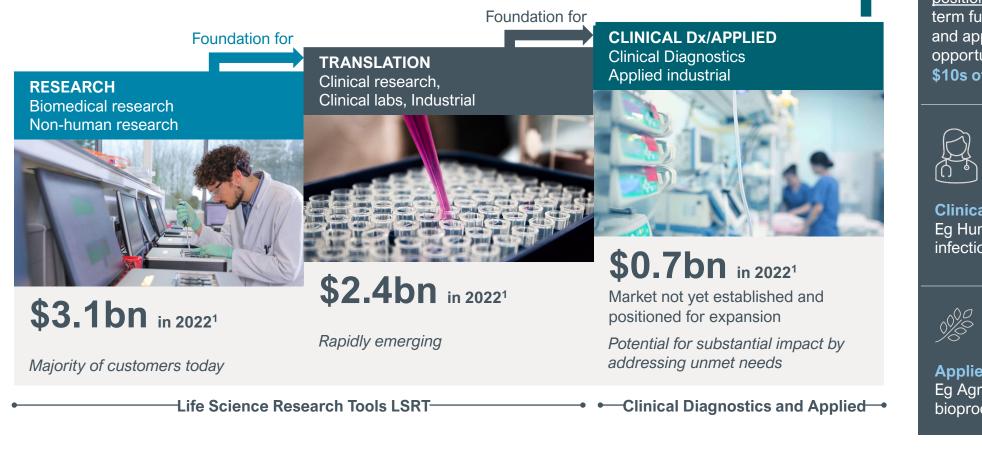
Users achieving high accuracy (99.9% single molecule Duplex), high outputs during a broad early access programme PromethION has room for growth with internal max output >350Gb / flow cell with fast motors in development





Substantial and growing market opportunity: DNA/RNA sequencing

\$6.2 billion¹ in 2022, expected CAGR 2022-25 ~15%¹ Substantial opportunity to penetrate, reshape and expand



Oxford Nanopore is <u>uniquely</u> positioned to unlock long term future potential clinical and applied market opportunities \$10s of billions



Clinical opportunities Eg Human genetics, cancer, infectious disease



Applied Industrial Eg Agriculture, food, bioprocessing, environment





FY22 Financial Review

Tim Cowper, CFO



FY22 key numbers



customer acquisition. Reported growth up 16% manufacturing processes

and improved

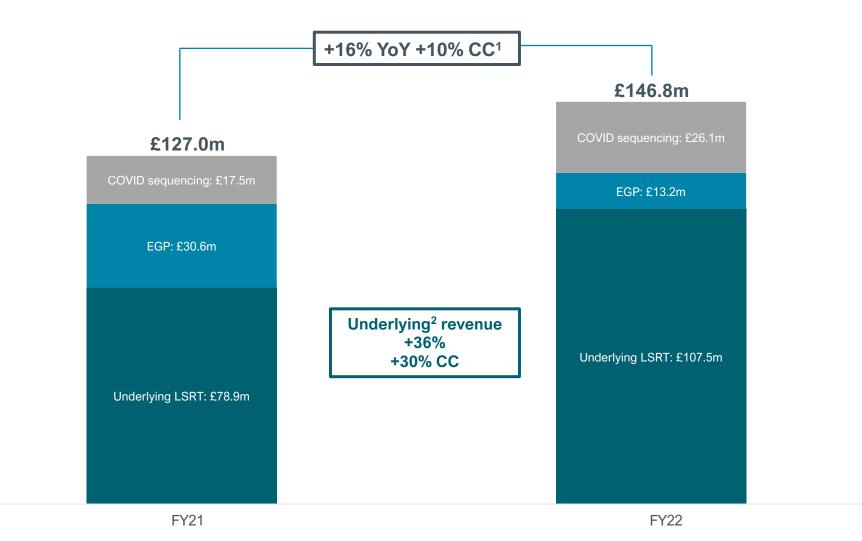
Covid testing revenue from DHSC settlement

by increased investment in R&D and commercial team innovation execution and commercial expansion





Continued robust underlying growth in LSRT business

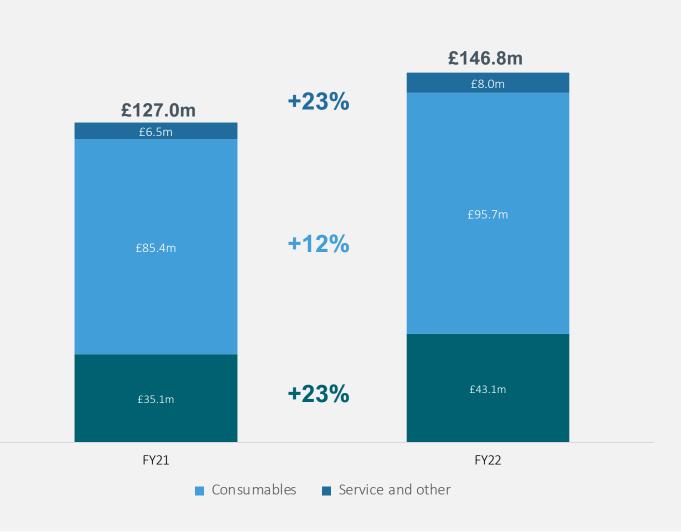


¹CC= constant currency, which applies the same rate to FY22 and FY22 non-GBP results based on FY21 rates ²LSRT revenue excluding EGP and COVID sequencing revenues

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FY22 LSRT revenue driven by consumables revenue



23% growth in 'service and other' revenue driven by service and license fees reflecting increased installed base

43% growth in consumables excluding decline in EGP consumables spend

Starter pack revenue growth of 23% driven by new customer acquisition



FY22 LSRT revenue driven by strong growth across all customer groups





FY22 LSRT revenue driven by customer acquisition

	FY22	FY21	Change
S1 (<\$25k)			
Number of active customer accounts	7,210	5,501	+31%
Avg. revenue per customer account (\$000s)	5.2	5.8	(10)%
S2 (\$25k-\$250k)			
Number of customer accounts	989	782	+26%
Avg. revenue per customer account (\$000s)	66.0	67.7	(2)%
S3 (>\$250k)			
Number of active customer accounts	73	56	+30%
Avg. revenue per customer account (\$000s) ¹	581.0	629.9	(8)%
Indirect			
Number of indirect accounts	11	6	+83%
Avg. revenue per indirect account (\$000s)	2,055	2,238	(8)%

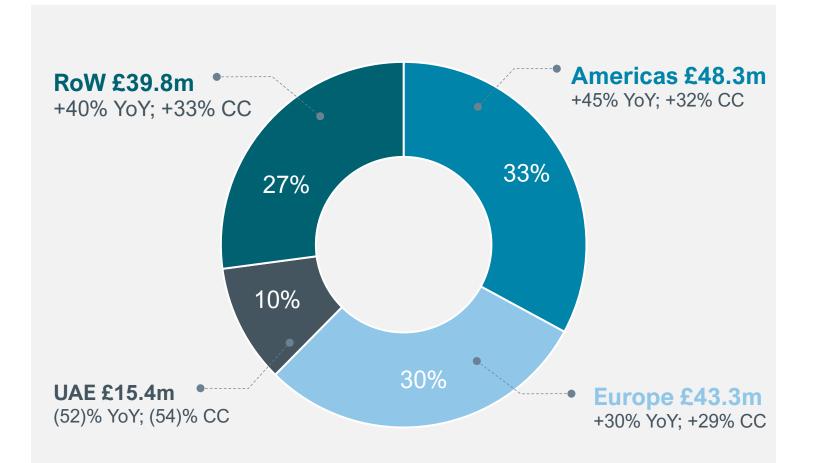
>8,200 active customers²

>1,900

net increase in active customers in FY22



Balanced and global customer base



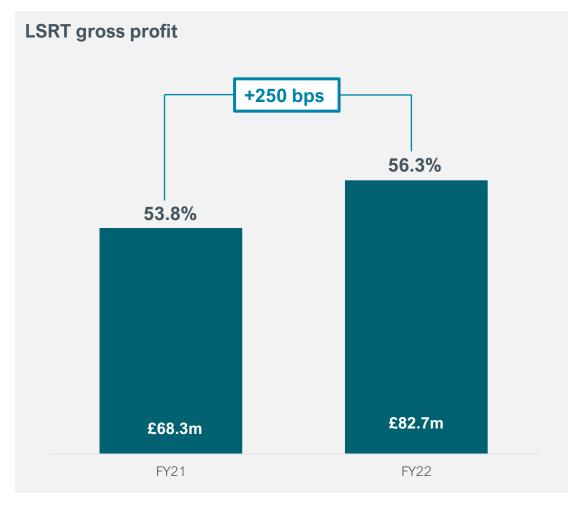
Strong growth in Americas and Europe reflects increased commercial resource

RoW growth driven by strong growth in China (+£8m or 66% CC)

UAE impacted by 57% decline in EGP revenue



Strong FY22 LSRT gross margin expansion despite supply chain headwinds



- Delivered +250 bps increase despite significant global supply chain challenges
- Margin expansion predominantly driven by operational improvements, including: automation, recycling of electronic components and improvements in manufacturing techniques
- H2 margin +300 bps higher than H1
 - H1: 54.8%
 - H2: 57.8%



Continued investment in innovation and commercial infrastructure in FY22

£93.9m¹ total adjusted R&D expenses

£116.0m¹ Adjusted S,G&A expenses

+39%

Principally due to a 31% increase in R&D headcount and an increase in capitalised development costs

+18%

Principally due to a 48% increase in the commercial team and a 30% increase in corporate team



Adjusted EBITDA

	FY22	FY21
	£m	£m
Loss before tax	(83.4)	(166.0)
Depreciation & Amortisation	31.9	24.7
Other	(4.3)	0.7
EBITDA	(55.9)	(140.6)
Founder LTIP	53.2	37.6
Employers Social Security on pre-IPO Share Awards	(21.6)	39.3
Gain on sale of property	(18.6)	-
Settlement of DHSC Covid Testing contract	(37.9)	-
IPO costs	-	4.8
Other	2.2	1.2
Adjusted EBITDA	(78.6)	(57.7)



FY23 and medium-term financial guidance

	FY23Medium-termtargetstargets	
LSRT revenue	>30% underlying revenue growth 16-30% constant currency	>30% underlying revenue growth
LSRT gross margin	>60%	>65%
Adjusted EBITDA		Targeting breakeven by FY26 ¹

Underlying revenue = LSRT revenue minus foreign exchange and revenues from COVID sequencing and the EGP ¹ FY26 = 31 December 2026



Summary

1 Core business is doing well. 2022 was a year of **customer acquisition** which will drive consumables growth in 2023.

2 Delivered **strong gross margin progress** in 2022 despite significant global supply chain challenges. Operational improvements made in 2022 will drive margin expansion in 2023 and beyond.

3 Strong balance sheet of £558.0m in cash, cash equivalents and liquid investments.

Progress made in **new market applications will be a key driver of long-term growth**.

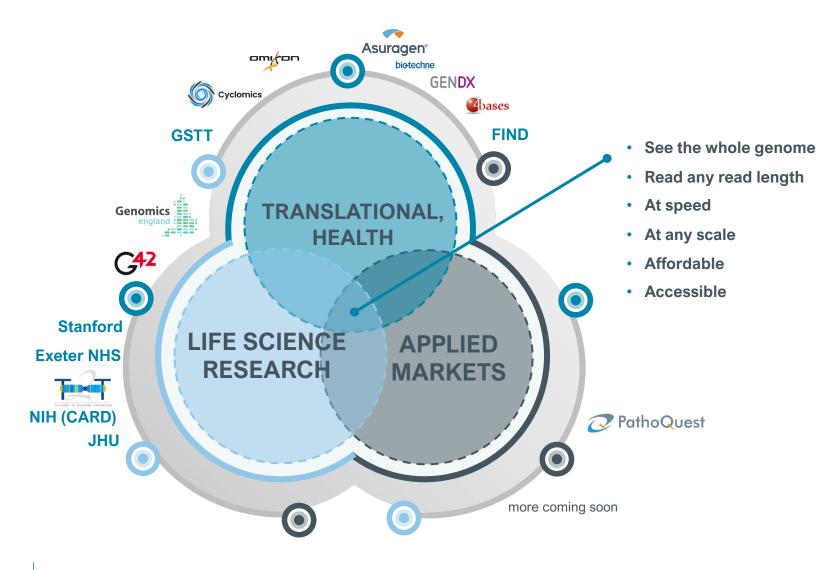


Emerging uses of Oxford Nanopore sequencing

Gordon Sanghera, CEO



Executing on DNA/RNA sequencing collaborations



With strong future innovation pipeline





Scientific discoveries are the foundation for real-world use

Large scale research in neurodegenerative conditions



NIH Centre for Alzheimer's and Related Dementias (CARD)

Thousands of brain samples being sequenced to understand Alzheimer's and related conditions

Cancer 2.0 data-rich cancer genomes, at speed



Transitioning to pilot programme deploying PromethION into NHS environment for near patient cancer sequencing

England

Resolving hard-todiagnose disease, faster than ever before



DNA to 60x genome in <2.5 hrs

Higher diagnostic yield than traditional short read sequencing

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Rapid Whole Genome Sequencing (WGS) in translational setting



Exeter, UK **Royal Devon NHS**

Evaluating benefits of rapid, complete human genomes in NHS settings, in patient care pathways

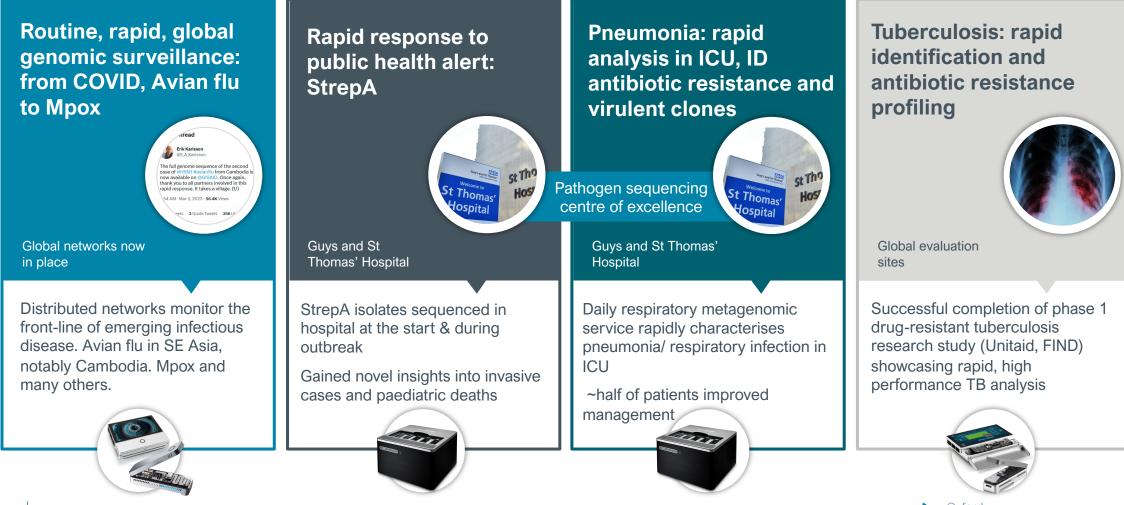


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Developing a unified approach to infectious disease and public health



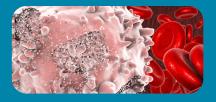
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Liquid biopsy: cancer recurrence monitoring

Translating research discoveries to real world benefits

The challenge:

Recurrent disease after initial remission is a leading cause of cancer death



Today, there are limitations associated with existing (imaging-based) methods

Monitoring Residual Disease requires exceptional specificity and sensitivity



The vision:

Non-invasive, highly accurate method of detecting molecules associated with cancer, with potential to reduce delays, optimise treatment decisions



Prep Sequence

Our unique advantage:

CyclomicsSeg is minimallyinvasive, highly sensitive, fast, affordable and deployable

- MinION based assay in developer testing
- Research shows near 100% accurate sequencing of TP53 gene, able to facilitate detection of mutations at frequencies down to **0.02%**
- Faster turnaround than short-read methods •
- Decentralised model for deployment in centres of excellence





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Human genetics; breast/ovarian cancer

Translating research discoveries to real world benefits

The challenge:

Tests for BRCA1 & BRCA2¹ are well defined but slow



The vision:

Reducing delay and improving accuracy in human genetics and cancer analyses, in future diagnostics, to inform treatment pathways and improve outcomes



- can take weeks to return a result
- Set up labs costs can be prohibitive resulting in lack of equity in access



Our unique advantage:

Our collaborator has developed rapid, cost effective, sample to answer assays



Human genetics kits, first target BRCA targeted panel

- MinION based assay
- Sample to answer in a day
- Decentralised model for deployment in centres of excellence: near sample
- Distribution agreement for Italy and Switzerland

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¹genes associated with breast and ovarian cancer



Carrier screening; uncovering the dark genome

The vision:

Translating research discoveries to real world benefits

Challenge:

Today's carrier screening methods cannot see all potential risks

6/10 genes with most common pathogenic variants in carriers are hard to sequence with short reads



More confident carrier screening,

nanopore native long reads

driven by comprehensive insights using

Our unique advantage:

Our collaborator has developed rapid, cost effective, sample to answer assays

Asuragen[®] biotechne[®]

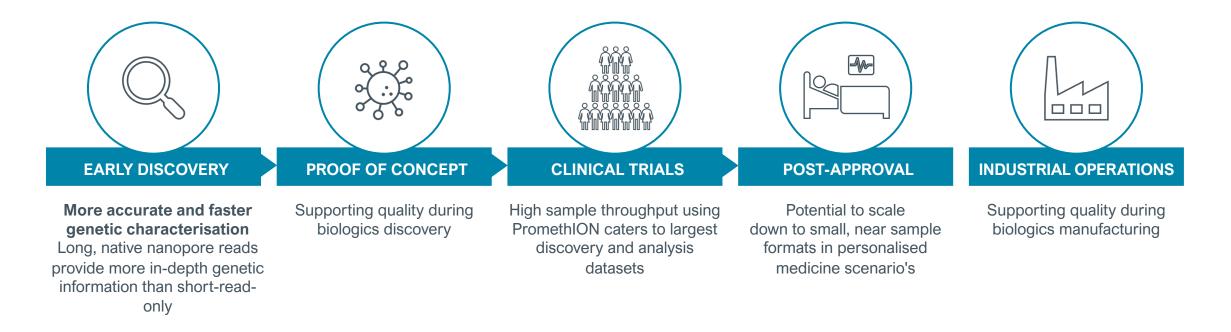
- GridION based assay
- Developing kit for genotyping 11 genes that cover ~70% of pathogenic variants associated with severe genetic disorder
- Including eight notoriously hard-toanalyse genes (e.g. fragileX)

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Pharma/biotech: an emerging customer segment

Meeting unmet needs across the value chain





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Improving biotherapeutic discovery and manufacturing

Moving into the applied markets

The challenge:

Biotherapeutic discovery and manufacturing needs better, faster tools

Discovering new biotherapeutics (e.g. antibodies) is a long and complex process involving screening of many potential candidates using sub-optimal methods.

Manufacturing: Once discovered, regulatory testing to allow the drug product to be released to the market is equally time consuming and complex as it relies on the use of the same testime rethodology.

The vision:

Rapid screening for discovery and GMP-grade testing for manufacturing quality control

To bring to the market the first rapid, accurate, regulatory approved quality control test for genetic characterisation of biotherapeutics at all stages in the drug development pipeline



Our unique advantage:

Rapid, comprehensive assays for discovery and GMP manufacturing needs



- PathoQuest, is a leader in QC testing of biologicals (e.g., ATMP, vaccines, antibodies), enabling rapid and robust analysis at a GMP level.
- Collaboration to optimise and commercialise a testing service for non-GMP and GMP validated Integration Site Analysis (ISA) tests for biomanufacturing quality control



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2023 outlook



CONTINUOUS INNOVATION

Execute on our new product introductions as they continue to drive platform adoption

PLATFORM PERFORMANCE

Drive exceptional accuracy, output and reliability with Q20+ full roll out

CUSTOMER FOCUS

Continue to make products our customers love and expand sales and support teams to ensure their success

OPERATIONAL EXCELLENCE

Drive high performance into our customer experience through key partnerships and digital excellence

COMMERCIAL EXECUTION

Deliver on our commitments from LSRT to the applied markets as we build a true technology company

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Strong balance sheet

	FY22 £m	FY21 £m
Non-current assets	185.9	91.3
Current assets		
Inventory	87.7	63.1
Trade and other receivables	62.9	54.8
R&D tax credit recoverable	9.1	14.3
Cash and cash equivalents	356.8	487.8
Derivative financial assets	2.1	0.0
Other financial assets	119.4	130.6
Total assets	823.9	841.9
Non-current liabilities	(27.8)	(32.8)
Current liabilities	(102.5)	(105.0)
Net assets	693.6	704.0
Total Equity	693.6	704.0

Increase in inventory reflects higher stock levels of core components to manage global supply chain risk

£558.0m

cash, cash equivalents and liquid investments¹



Summary income statement

	FY22 £m	FY21 £m
Revenue	198.6	133.7
Cost of Sales	(74.8)	(60.5)
Gross Profit	123.8	73.2
Gross margin %	62.3%	54.8%
Operating expenses		
Research and development expenses	(64.8)	(76.0)
Selling, general & administrative expenses	(157.4)	(161.8)
Loss from operations	(98.5)	(164.5)
Other	15.1	(104.3)
Loss before tax	(83.4)	(166.0)



Continued investment in innovation and commercial infrastructure

	FY22 £m	FY21 £m
Research and Development expenses	64.8	76.0
Adjusting Items:		
Employer's social security taxes on pre-IPO share awards	9.9	(17.7)
Adjusted Research and Development expenses	74.7	58.3
Capitalised development costs	19.2	9.3
Total R&D and capitalised development costs	93.9	67.6
Selling, general and administrative expenses	157.4	161.8
Share-based payment expense on Founder Long Term Incentive Plan	(50.0)	
(LTIP)	(53.2)	(37.6)
Employer's social security taxes on Founder LTIP and pre-IPO share awards	11.7	(21.5)
IPO costs expensed in Income Statement	0.0	(4.8)
Adjusted Selling, general and administrative expenses	116.0	97.9





% of LSRT sales









LSRT revenue by currency

