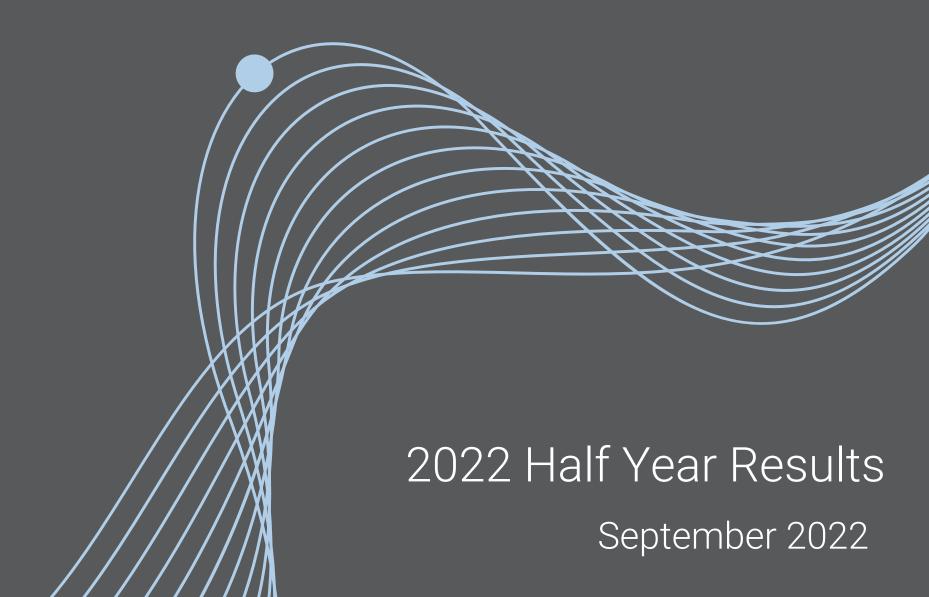
## NOVACYT



## Disclaimer

The document attached hereto and the presentation of which it forms part (together the "Materials") have been prepared by Novacyt" or the "Company"). The Materials are confidential and personal to you and are furnished to you as background information to provide a basis for you, as a potential investor, to consider whether to pursue an acquisition of shares in the Company. The Materials do not constitute an offer or invitation for the sale or purchase of any securities, nor do they, nor do they purport to, set out or refer to all or any of the information a potential investor might require or expect in making a decision as to whether or not to deal in shares in the Company. The Materials do not comprise an admission document, listing particulars or a prospectus relating to the Company and the information contained in, and communicated to you during, this Presentation does not constitute, or form part of, and should not be construed as, an offer or invitation or recommendation to purchase or subscribe for any securities in the Company. Prospective investors should only subscribe for shares in the Company on the basis of information contained in any prospectus which may be published by the Company in connection with the Admission.

The Materials have not been approved by an authorised person within the meaning of the Financial Services and Markets Act 2000. Reliance on the Materials for the purpose of engaging in any investment activity may expose an individual to a significant risk of losing all of the property or other assets invested. The Materials do not constitute and are not a prospectus or listing particulars (under either the Prospectus Regulations 2005 (as amended), the Financial Services and Markets Act 2000 ("FSMA") or the Prospectus Rules of the Financial Conduct Authority) and should not be construed as such. No reliance may be placed for any purpose whatsoever on the information, representations or opinions contained in the Materials or on the Company or any of their respective directors, officers, employees, advisers or any other persons as to the fairness, accuracy or completeness of the information or estimates or opinions or other statements about the future prospects of the Company or any of their respective businesses contained in the Materials or referred to in the presentation given in connection therewith and no responsibility, liability or duty of care whatsoever is accepted by any such person in relation to any such information, representation, projection, forecast, opinion, estimate or statement including in the case of negligence, but excluding any liability for fraud.

SP Angel Corporate Finance Limited (the Company's nominated adviser and joint broker) and Numis Securities Limited (the Company's joint broker) have not approved the Materials as a financial promotion for the purposes of section 21 of FSMA or otherwise. Whilst all reasonable care has been taken to ensure that the facts stated in these presentation materials are accurate and that any forecasts, opinions and expectations contained therein are fair and reasonable, SP Angel Corporate Finance Limited and Numis Securities Limited have not independently verified the contents of these Materials and no reliance whatsoever should be placed on them.

This document constitutes a 'financial promotion' for the purposes of section 21 of the FSMA and its distribution in the United Kingdom is restricted. Accordingly, this document will not be offered to the public in the United Kingdom (within the meaning of section 85 of the FSMA) save in circumstances where it is lawful to do so without an approved prospectus (within the meaning of section 85 of the FSMA) being made available to the public before the offer is made. In the United Kingdom, the Materials are only being directed at persons: (a) persons who are outside the United Kingdom; (b) investment professionals falling within Article 19(5) of the Financial Services and Markets Act 2000 (Financial Promotion) Order 2005 (SI 2005/1529) (as amended) (the "Order"); (c) high net worth individuals within Article 48 of the Order who, in this regard, have signed a statement dated within a period of 12 months ending on the date of receipt of this document complying with Part 1 of Schedule 5 of the Order stating that among other things, they have either or both: (i) during the financial year immediately preceding the date on which the statement is signed an annual income of not less than

£100,000; or (ii) held, throughout the financial year immediately preceding the date on which the statement is signed, net assets to the value of not less than £250,000 (excluding the property which is their primary residence or any loan secured on that residence, any of their rights under a qualifying contract of insurance within the meaning of the Financial Services and Markets Act 2000 (Regulated Authorities) Order 2001, or any benefits (in the form of pensions or otherwise) which are payable on termination of their service or death or retirement and to which they are (or their dependants are), or may be entitled; (e) sophisticated investors falling within Article 50 of the Order; (f) self-certified sophisticated investors falling within Article 50A of the Order; and (g) other persons to whom it may lawfully be communicated (all such persons together being "relevant persons"). The investment or investment activity to which the Materials relate are available only to such persons and will be engaged with only with such persons. If you are not such a person: (i) you should not take part in the presentation and nor should you have received the Materials; (ii) please leave the presentation immediately after returning the Materials; and (iv) you may not rely on or act upon the matters communicated by the Materials.

Neither this presentation nor any copy of it, in whole or in part, or any of the Materials may be: (i) taken or transmitted into the United States of America; (ii) distributed, directly or indirectly, in the United States of America or to any US person (within the meaning of regulations made under the Securities Act 1933, as amended); (iii) taken or transmitted into or distributed in Canada, Australia, the Republic of Ireland or Irela

The Materials are being made available on the basis that the recipients keep confidential any information contained therein, whether orally or in writing, in connection with the Company. The Materials are confidential and must not be copied, reproduced, published, distributed, disclosed or passed, directly or indirectly, to any other person or published, in whole or in part, for any purpose at any time without the prior written consent of the Company. By attending the presentation and/or accepting a copy of the Materials you agree to be bound by the foregoing provisions.

The information described in the Materials may contain certain information that is confidential, price-sensitive and which has not been publically disclosed. By your receipt of the Materials may contain certain information in the Materials may be "inside information" as defined in Article 7 of the Market Abuse Regulation EU 596/2014 ("MAR") and constitutes a "market sounding" for the purpose of Article 11 of MAR. You recognise and accept that such information is being provided to you by the Company pursuant to Article 11 of MAR and you confirm, warrant and undertake that you will keep the information confidential and will not: (i) deal, in financial instruments (as defined in MAR) relating to that information, or encourage another person to deal or disclose the information before the inside information is made public; (ii) or cancel or amend an order which has already been placed concerning a financial instrument to which such information relates; (iii) disclose the inside information to another person other than in the proper course of the exercise of your employment, profession or duties; or (iv) engage in behaviour based on any inside information which might amount to market abuse or market manipulation for the purposes of MAR. Recipients should take their own legal advice on the obligation to which they will be subject and the application of MAR and in particular make their own assessment of whether they are in possession of inside information and when such information ceases to be inside information.

#### Forward-looking Statements:

The Materials contain forward-looking statements. These statements relate to the future prospects, developments and business strategies of the Company and its subsidiaries (the "Group"). Forward-looking statements are identified by the use of such terms as "believe", "could", "envisage", "estimate", "potential", "intend", "may", "plan", "will" or the negative of those, variations or comparable expressions, including references to assumptions. The forward-looking statements contained in the Materials are based on current expectations and are subject to risks and uncertainties that could cause actual results to differ materially from those expressed or implied by those statements. If one or more of these risks or uncertainties materialises, or if underlying assumptions prove incorrect, the Group's actual results may vary materially from those expected, estimated or projected. Given these risks and uncertainties, pytential investors should not place any reliance on forward-looking statements and the Company accepts no obligation to disseminate any updates or revisions to such forward-looking statements. These forward-looking statements speak only as at the date

## Company capabilities, vision and purpose



#### **Our vision**

**Committed** to becoming a leading, **global clinical diagnostics** company in infectious diseases

#### Our purpose

We **protect lives** from invisible threats by enabling informed clinical decision-making through quality diagnostics in **the right place**, at the right time

## Presenters



Dave Allmond
Chief Executive Officer

- Appointed CEO of Novacyt in 2021
- Over 25 years of global experience in pharmaceuticals & biopharmaceutical companies
- Internationally experienced, strategic business leader with strong track record in global commercialisation
- Built & led multiple diverse, successful teams in dynamic growth companies including Amgen, Celgene & Amryt Pharma
- · C-level executive with AIM and NASDAQ listed companies



James McCarthy Chief Financial Officer

- Appointed CFO of Novacyt in 2021
- Over 30 years in international manufacturing and industrial businesses in both consumer and B2B
- CFO in both Private Equity and public businesses.
- FCCA qualified over 30 years with broad commercial, supply chain and M&A experience



# H1 2022 and post-period operational highlights



 Developed semi-automated, scalable workflow solution.



 Accelerated menu availability with access to >40 clinical ID assays.



 Launched an LFT digital reader for use in conjunction with 18 non COVID-19 assays



 Launched an RUO monkeypox PCR assay and an RUO assay for adenovirus F41



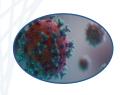
 Launch of defence and counterclaim against DHSC



Completed closure of Microgen
Bioproducts and Lab21 and fully
delivered restructuring programme
announced in July.



 Relaunched RUO portfolio in July with encouraging early contract wins.



 PROmate® COVID-19 1G, 2G and exsig™ COVID-19 Direct Real-Time PCR tests received CTDA approval



Granted a key patent for ORF1a/b, which will lead to a corporation tax credit against future profits on related assays

NOVACYT

## H1 2022 unaudited – Profit & Loss

£'000	H1 2022	H1 2021
Continuing Operations*	Consol	Consol
Revenue	16,508	52,201
Gross profit **	4,010	1,177
Gross profit %	24%	2%
OPEX	(11,148)	(13,301)
EBITDA	(7,138)	(12,124)
EBITDA %	-43%	-23%
Adjusted EBITDA **	(7,138)	23,646
Recurring operating loss ***	(8,179)	(12,958)
Operating loss	(8,712)	(12,958)
Other financial income and expenses	1,628	(1,421)
Income tax credit	2,041	2,295
Loss after tax from continuing operations	(5,043)	(12,084)
Loss from discontinued operations	(3,656)	(591)
Loss after tax attributable to the owners	(8,699)	(12,675)

<sup>\*</sup> Following the 28 April 2022 announcement where Novacyt announced its intention to close Microgen Bioproducts and Lab21 Healthcare the net results of the Lab21 Products segment for 2021 and 2022 has been reported on a separate line 'Loss from discontinued operations' in accordance with IFRS 5, "Non-current Assets Held for Sale and Discontinued Operations".

- Group revenue of £16.5m in H1 2022 compared with £52.2m in H1 2021 predominantly driven by the expected decline in COVID-19 related sales.
- Group gross profit improved to £4.0m (24%) in H1 2022 (H1 2021: £1.2m (2%)). The latter was impacted by the one-off exceptional costs relating to the DHSC dispute.
- The H1 2022 gross profit was reduced as a result of stock provisions and write-offs relating to lower forecast COVID-19 sales. Excluding the impact of these items the Gross Margin would have been in excess of 60%.
- Group adjusted EBITDA loss of £7.1m in H1 2022 before exceptionals (H1 2021: £23.6m profit).
- The discontinued operations losses widen to £3.7m in H1 2022 from £0.6m in H1 2021.
- Loss after tax has decreased to £8.7m in H1 2022 from £12.7m in H1 2021.

<sup>\*\*</sup> Due to the ongoing commercial dispute with the DHSC, £35.8m exceptional cost of sales were incurred in H1 2021 (H1 2022: £nil) that were one-off in nature. The two largest items were a £26.1m stock provision, as a result of the Group buying stock to fulfil expected future DHSC orders that did not materialise; and the expensing of £6.9m of stock delivered to the DHSC which has not been paid for as it is now part of the ongoing contract dispute.

<sup>\*\*\*</sup> H1 2022 recurring operating loss is stated before £0.5m of non-recurring charges in relation to the ongoing DHSC contract dispute. September 27, 2022 at 12:43 PM

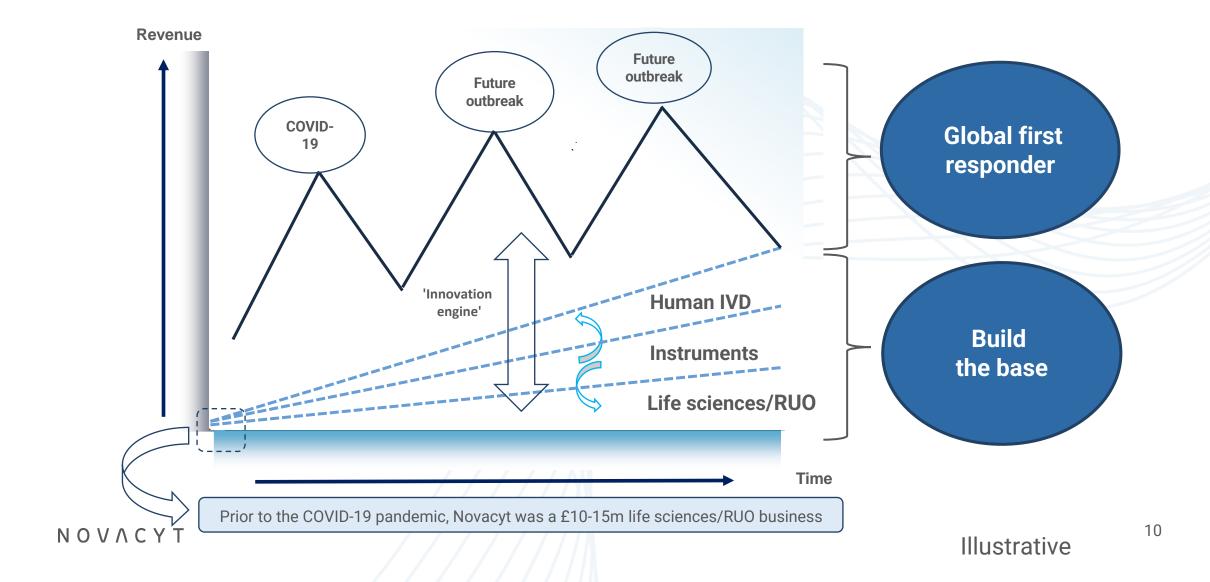
## H1 2022 unaudited balance sheet and cash

	Jun-22	Dec-21		Jun-22	Dec-21
	£'000	£'000		£'000	£'000
Goodwill	11,638	11,471	Share capital and premium	54,632	54,646
Right-of-use assets	552	1,788	Retained earnings and reserves	78,035	87,169
Property, plant and equipment	3,439	4,594	Total equity	132,667	141,815
Deferred tax assets	4,796	3,143			T = T
Other non-current assets	3,625	3,918	Deferred tax liabilities	1,245	1,224
Total non-current assets	24,050	24,914	Lease liabilities long-term	1,324	1,446
			Other provisions and long- term liabilities	425	308
Inventories	4,255	11,461	Total non-current liabilities	2,994	2,978
Trade and other receivables	35,293	38,499			
Tax receivables	1,000	5,034	Lease liabilities short-term	347	424
Other current assets	1,889	2,043	Trade and other liabilities	8,128	17,190
Cash and cash equivalents	99,641	101,746	Other provisions and short- term liabilities	21,992	21,290
Total current assets	142,078	158,783	Total current liabilities	30,467	38,904
					<u>/ X</u>
TOTAL ASSETS	166,128	183,697	TOTAL EQUITY AND LIABILITIES	166,128	183,697

- Cash position at 30 June 2022 was £99.6m, compared with £101.7m at 31 December 2021, and the Company remains debt free.
- WCAP (excluding cash) has fallen by one third from £18.2 at year end to £11.9m at the end of June based on the falling revenues and actions taken in relation to COVID-19 stock.
- to £2.0m in H1 2021, after the Company heavily invested in insourcing manufacturing during 2021.
- Granted the key patent (ORF1a/b), with patent number GB2593010. This means that the effective rate of tax on profits (adjusted for certain rules) derived from the sale of products incorporating this patent is close to 10% rather than the current UK corporation tax rate of 19%.

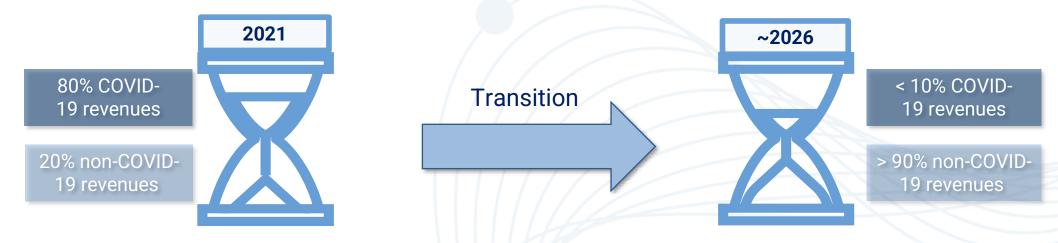


# Corporate strategy – two key imperatives



# Evolving Novacyt beyond the pandemic – building the base

Evolving beyond the pandemic to a sustainable growth business, serving high unmet needs in infectious diseases







## Portfolio development

Relaunch RUO portfolio in Q2 2022, launch e-commerce platform, deliver clinical IVD portfolio



#### **Instrumentation**

Integrated with life sciences/RUO and IVD portfolios



## Geographic expansion

Increasing presence, leveraging adjacent markets and building distributor network



## **Business** development

Innovation for disruptive technologies
Accelerating molecular portfolio
Strategic transactions

# Significant progress in building the base business (1)

#### **Strategy**



## Portfolio development

Relaunch RUO portfolio in Q2 2022, launch e-commerce platform, deliver clinical IVD portfolio

#### **Actions**

- Signed a global distribution agreement with Clonit srl with access to over 40 CE marked clinical assays
- Launched E-commerce platform in Q2 2022
- Relaunched RUO portfolio in July 2022
- Progressing organic R&D for mid-term menu expansion.



#### Instrumentation

Integrated with life sciences/RUO and clinical (IVD) portfolios

- Sourced extraction system to integrate with clinical workflow
- Enhancing Co-prep automation with flexibility for PROmate®, genesig® and dry assays
- Launched a new lateral flow test (LFT) reader for use in conjunction with 18 non COVID-19 Pathflow® assays for patient screening in STI, GI, respiratory and insect-borne infections

#### **Outcomes**

- Broad, approved, clinical menu and enhanced workflow available now in STI and in Q1 2023 in respiratory and GI
- Launch in Europe, our initial target geography, (CE marked) where we estimate a market size of circa £470m growing at 10% pa
- Early RUO wins in Q3 with Salmon testing in Canada and Salmonella testing in Poland
- Advanced design of 2 PCR panels for nearpatient testing in gastro-intestinal viruses and bacteria infections
- Launching decentralised workflow for use in spoke laboratories to enable deployment of expanded clinical (IVD) menu
- Now participating in significant lateral flow tenders across EU

# Significant progress in building the base business (2)

#### **Strategy**



## **Geographic expansion**

Increasing presence, leveraging adjacent markets and building distributor network

#### **Actions**

- Deployed talent in key geographies
- Optimising global distributor network to build coverage in new markets, ensure optimal RUO and clinical portfolio coverage across priority markets



## **Business** development

Innovation for disruptive technologies

Accelerating molecular portfolio
Strategic transactions

- Agreements for extraction, sample handling digital reader and broad clinical assay menu already completed
- Further work on portfolio development in progress across target disease areas to accelerate approved menu to commercialise
- Significantly enhanced BD funnel and deal flow for M&A

#### **Outcomes**

- Reduced active distributor network by over 75% to focus on key partners
- Added coverage in 18 new countries across EMEA. Optimising distributor network in US, APAC and LATAM
- Commenced distributor training on full portfolio including expanded clinical portfolio and workflow
- Accelerated launch of clinical ID menu by ~2 years and developing integrated workflow to be completed by Q2 2023
- Identified potential mini-GI panels approved under IVDD for validation and potential distribution
- Strategically aligned/executable opportunities coming though the funnel

13

# Promoting comprehensive product portfolio to drive near term growth

#### **PROmate**®

Simple, convenient direct to PCR for near patient workflow. Investing in R&D on this platform.



#### **PathFlow**®

Extensive lateral flow portfolio and digital reader for clinical diagnostics/screening



MyGo, Q Series CO-prep™ & VersaLab™

qPCR instruments, liquid handling, extraction and mobile laboratory taking testing to the front line



### genesig<sup>®</sup>

Broad clinical & research use PCR portfolio for pathogen detection in veterinary, human, food, and environment settings



### **SNPsig**<sup>®</sup>

For the rapid detection, identification and monitoring of SARS-CoV-2 variants of concern



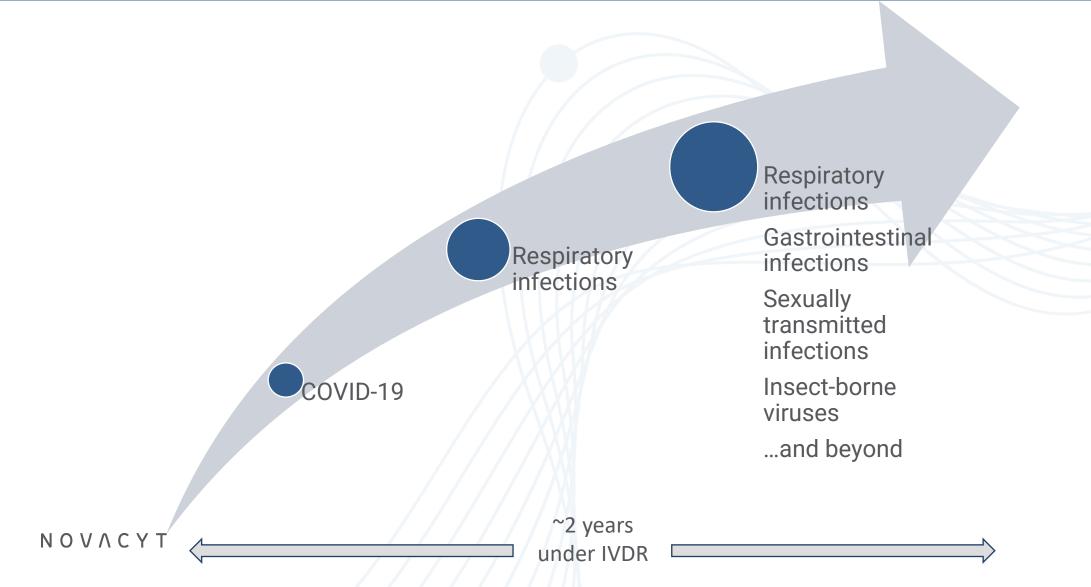
# Relaunching world class, broad range of research use only/life sciences assays and instrumentation

- Veterinary
- Food
- Environment
- Human



MyGo range of 16 and 32 well, open platform qPCR instruments offer an ideal solution for research customers

# Investing in organic R&D and business development to expand our clinical portfolio in 4 key therapeutic areas



# Enhancing scalable, semi-automated workflow taking clinical testing to the front line











NOVACYT











"Q Series" PCR, 16 & 32 well Instruments

Rapid results High sensitivity & specificity. Up to 1000 tests per day

Co-Prep™ extraction and automated liquid handling assay set-up

Convenience with walkaway time

Reduced contamination & human error risk

### Combined with a VersaLab to provide ultimate flexibility





# Launching a broad menu of approved assays across respiratory, STI, GI and insect borne infections

### Phase 1

Chemistry supply only

STI

Respiratory

GI

Transplant

Insect-borne

September 2022

NOVACYT

### Phase 2

q16/32 instrument supported



STI

Respiratory

GI

Q1 2023

### Phase 3

Full workflow supported (CO-Prep extraction, automation and PCR instrument validation



GI

Respiratory

Insect-borne\*

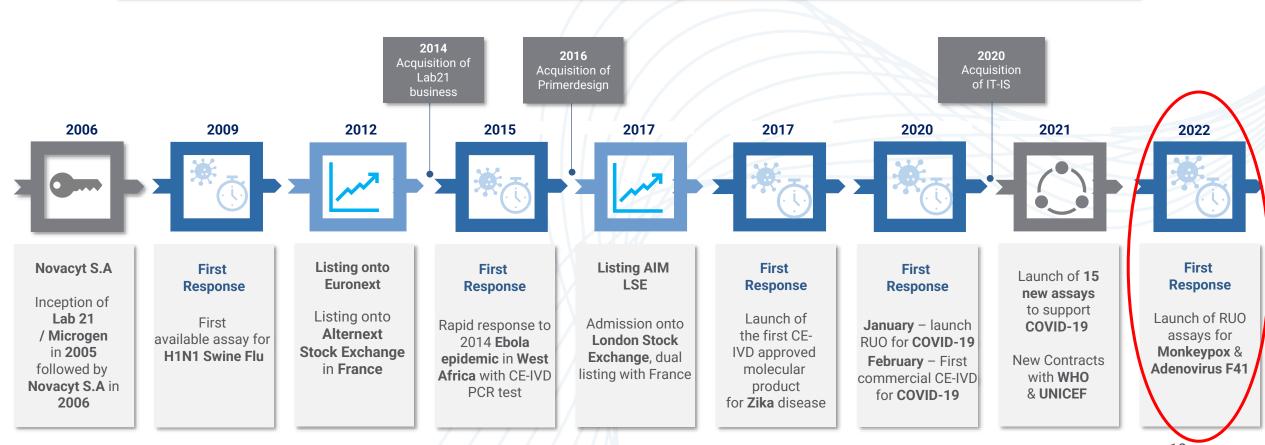
Q2 2023

18

# Continuing to be a global first responder in infectious diseases

Strong track record of being a "first responder" to disease outbreaks.

Rapid response to Monkeypox and Adenovirus F41 in 2022



ΝΟΥΛΟΥΤ

# Three key areas of focus

# Delivering on inorganic growth Objectives through business development



1. Innovation

Innovation to in license & develop disruptive clinical diagnostic technologies and workflow

Point of care Biosensor in development √



2. Acceleration

Accelerating the molecular portfolio (AMP)

- Molecular assay menu expansion √
- LFT digital reader √
- Expanding liquid handler to broader chemistry for automation √
- Extraction capability by Q4



3. Strategy

Strategic transactions/ M&A to diversify and build for scale

 Built significant funnel to deliver strategically aligned, transactable deals √

We plan to deploy capital in strategic transactions to accelerate growth

## Summary and outlook

- Launch of new clinical IVD workflow, in partnership with Clonit, with access to a broad menu
- Launched LFT digital reader for use in conjunction with extensive LFT menu
- Re-launch of RUO portfolio and refreshing route to market with more focussed and streamlined distributors and pilot launch of e-commerce
- Right-sizing cost base in line with falling COVID-19 sales, fully delivered on announced restructuring
- Redoubling business development activities to find the right investment options for cash resources
- M&A funnel delivering opportunities for strategic transactions
- Predicted Q3 2022 revenue of circa £2.0m, with similar levels expected in Q4 2022, resulting in an anticipated EBITDA loss for the full year of circa £13.5m
- The Board believes that Novacyt remains well positioned to leverage its core capabilities to become a leading, global clinical diagnostics company

21

## NOVACYT

