

This presentation has been prepared by Genetic Signatures Limited ACN 095 913 205 (the Company or GSS) and approved by the Board of Directors for release. It comprises written materials/slides for a verbal presentation concerning the Company and should be read in that context. This presentation is proprietary to GSS. It may not be reproduced, disseminated, quoted or referred to, in whole or in part, without express consent of GSS.

No representation or warranty, express or implied, is or will be made in relation to, and no responsibility or liability (whether for negligence, under statute or otherwise) is or will be accepted by the Company or by any of its officers, directors, shareholders, employees or advisers as to or in relation to the accuracy or completeness of the information, statements, opinions or matters (express or implied) arising out of, contained in or derived from this presentation or any omission from this presentation or of any other written or oral information or opinions provided now or in the future to any interested party or its advisers. In particular, no representation or warranty is given as to the achievement or reasonableness of any plans, future projections, management targets, prospects or returns and nothing in this presentation is or should be relied upon as a promise or representation as to the future.

The Company expressly disclaims all liability for any loss or damage of whatsoever kind (whether foreseeable or not) which may arise from any person acting on any information and opinions relating to the Company contained in this presentation or any information which is made available in connection with any further enquiries, notwithstanding any negligence, default or lack of care. In furnishing this presentation, the Company undertakes no obligation to provide any additional information.

Subject to any continuing obligation under applicable law or relevant listing rules of the ASX, the Company disclaims any obligation or undertaking to disseminate any updates or revisions to any forward-looking statements in these materials to reflect any change in expectations in relation to any forward-looking statements or any change in events, conditions or circumstances on which any statement is based. Nothing in these materials shall under any circumstances create an implication that there has been no change in the affairs of the Company since the date of the presentation.

This presentation is for information purposes only and does not constitute or form part of any offer or invitation to acquire, sell or otherwise dispose of, or issue, or any solicitation of any offer to sell or otherwise dispose of, purchase or subscribe for, any securities, nor does it constitute investment advice, nor shall it or any part of it nor the fact of its distribution form the basis of, or be relied on in connection with, any or contract or investment decision. Without limiting the foregoing, this presentation does not constitute an offer to sell, or a solicitation of an offer to buy, any securities in the United States. The securities of Genetic Signatures have not been, and will not be, registered under the U.S. Securities Act of 1933, as amended (Securities Act) or the securities laws of any state or other jurisdiction of the United States, and may not be offered or sold in the United States except in compliance with the registration requirements of the Securities Act and any other applicable securities laws or pursuant to an exemption from, or in a transaction not subject to, the registration requirements of the Securities Act and any other applicable securities laws.

The receipt of this presentation by any person and any information contained herein or subsequently communicated to any person is not to be taken as constituting the giving of investment advice by the Company or any other person to any such person. No such person should expect the Company or any of its officers, directors, shareholders, employees or advisers to owe it any duties or responsibilities and should take its own professional advice. The Recipient must rely solely on its own knowledge, investigation, judgement and assessment of the matters which are the subject of this presentation and to satisfy itself as to the accuracy and completeness

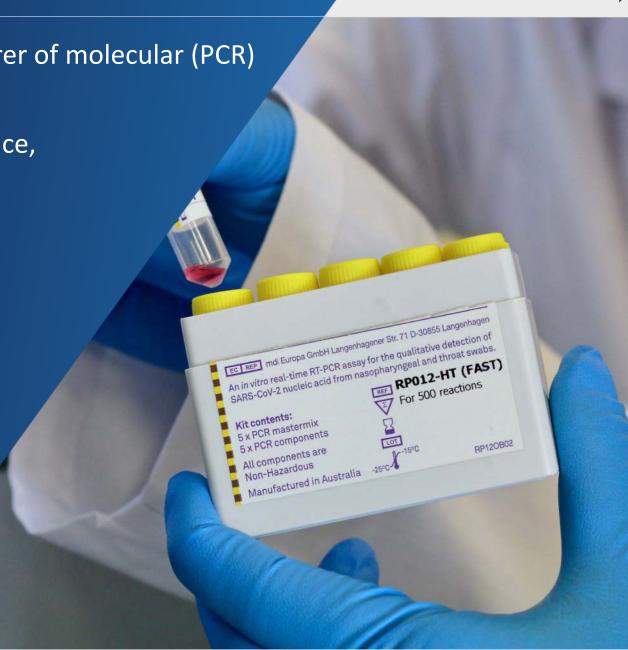
Established ASX listed developer and manufacturer of molecular (PCR) based test kits for infectious diseases

Novel 3base® technology – changes DNA sequence, detects the most common mutations, rapid, low cost and accurate

✓ EasyScreen[™] test kits available for sale in most major markets, expanding into EU and USA

YTD FY22 sales \$30.6m, up 34% pcp

- 3Q FY22 up 108% pcp
- YoY growth since listing
 - 4 year CAGR 93%
- Profitable, \$39.0m cash, no debt
 - \$4.7m profit 1H FY22



Trusted & Proven Technology



A 'Syndromic Screening' approach allows users to test a broad range of clinically relevant pathogens based on patient symptoms, helping clinicians make an accurate diagnosis

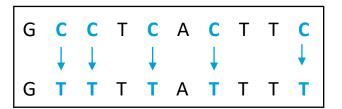
EasyScreen™ Detection Kits

- Simultaneously detect over 20 pathogens from one sample, shortening turnaround from days to hours
- Assays for over 100 organisms, including for SARS-CoV-2, but also other disease states
- Streamlined universal sample processing kits –
 common method across all kits
- 3base® can detect all known COVID variants¹
- Customers high throughput labs, hospital groups and private pathology suppliers
- More than quadrupled GSS branded instruments in the field since start of pandemic



Our proprietary 3base solution...

- 3base® platform technology converts original 4-base microbial genome to 3-base
- Conversion occurs during standard procedures with no additional steps for the technician
- 3base® MDx reduces complexity in a multiplex environment



1,048,576

combinations for a 10 digit number with 4-base



combinations for a 10 digit number with 3-base

...improves workflow and increases throughput



EasyScreen™ kits are compatible with most existing automated nucleic acid extraction and real-time PCR instruments and streamline the preparation process



High throughput labs can achieve further workflow automation with compatible hardware. Creates workflow efficiencies and reduces costs



Currently offering over 150 pathogen targets across enteric, respiratory, anti-microbial resistance, sexual health, tropical diseases and more in development

3base Technology: less sequence variation



	Before	After			
Seq 1	GATG	C G A I A T G G T I G A C A C G A T G G T G A T A T G	G T <u>T</u> G A T A T		
Seq 2	GATG	[GA C ATGGT A GA <u>T</u> A C GATGGTGATATG	G T <u>A</u> G A T A T		
Seq 3	GATG	[GA <u>T</u> ATGGT <u>G</u> GA <u>C</u> A <u>C</u> GATGGTGATATG	G T <u>G</u> G A T A T		
Seq 4	GATG	GA <u>T</u> ATGGT <u>A</u> GA <u>T</u> A <u>T</u> GATGGTGATATG	G T \underline{A} G A T A T		
Seq 5	GATG	GA <u>T</u> ATGGT <u>G</u> GA <u>C</u> A <u>C</u> GATGGTGATATG	GT $\underline{\mathbf{G}}$ GATAT		
Seq 6	GATG	CGA <u>C</u> ATGGT <u>I</u> GA <u>I</u> A <u>I</u> GATGGTGATATG	$G \ T \ \underline{T} \ G \ A \ T \ A \ T$		
Seq 7	GATG	GA <u>T</u> ATGGT <u>G</u> GA <u>C</u> A <u>C</u> GATGGTGATATG	GT $\underline{\textbf{G}}$ GATAT		
Seq 8	GATG	GA <u>C</u> ATGGT <u>A</u> GA <u>T</u> A <u>C</u> GATGGTGATATG	$G \ T \ \underline{A} \ G \ A \ T \ A \ T$		
Seq 9	GATG	GA <u>T</u> ATGGT <u>A</u> GA <u>T</u> A <u>C</u> GATGGTGATATG	$G \ T \ \underline{A} \ G \ A \ T \ A \ T$		
Seq 10	G A T G	GA <u>T</u> ATGGT <u>G</u> GA <u>T</u> A <u>C</u> GATGGTGATATG	G T $\underline{\textbf{G}}$ G A T A T		
Consensus	GATG		G T <u>D</u> G A T A T		
	75% homolo 48 possible	er 20 bases 95% homology over 20 bases r combinations 3 possible primer combinations			

- Reduced sequence variation
- Improved subtype similarity
- No cross-reaction with 4base native sequences, reduced laboratory contamination
- 3base® greatly simplifies the multiplexing process

Conventional Sequence			Tm
Primer 1	GTACACAC	CGCCGTCGCTCCTACC	77°C
Primer 2	GAAGGAGA	A G T C G T A A C A A G	56°C
Probe 1	TGAATAAA	GAGGTGAAATTCTAGG	59°C
Probe 2	GAAGGGCC	GCGAGCCCCGCGC	87°C

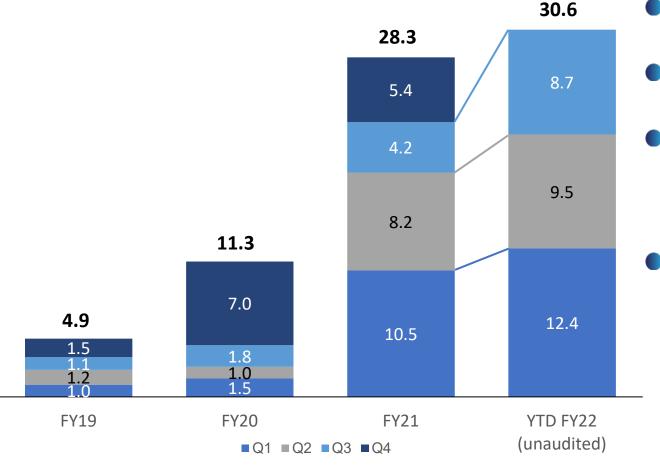
3base™ S	equence		Tm
Primer 1	GTATATA	TTGTTTGTTGTTTTTATT	52°C
Primer 2	GAAGGAG	i A A G T T G T A A T A A G	50°C
Probe 1	TGAATAA	A G A G G T G A A A T T T T A G G	59°C
Probe 2	GAAGGGT	TGTGAGTTTTTGTGT	62°C

- Simple Multiplex Primer/Probe Design
- Improved PCR Efficiency

Financial Summary – Sales



Revenue from sales (A\$m)



Continued Strong Revenue Growth

- 3Q FY22 \$8.7m, up 108% on pcp; increased contribution from overseas
- SARS-CoV-2 test demand tapering, being replaced with other tests
- International sales teams actively engaged with existing customers to promote broader *EasyScreen*™ range European orders received for Enteric range, including 3yr contract with Public Health Wales
- New instrument placements continue to support future demand for tests

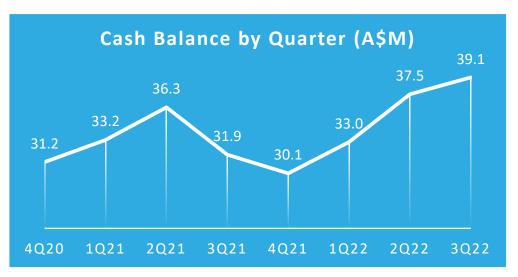


Financial Summary – Cashflow

A'000s	3Q FY22	YTD FY22
Receipts from customers	11,834	34,258
Payments to suppliers and employees	(9,750)	(24,187)
Other	15	95
Net operating cashflow	2,099	9,976
Payment for plant & equipment	(54)	(334)
Payment for intangibles	(390)	(548)
Net investing cashflow	(444)	(882)
Net proceeds from issue of shares	18	132
Principal elements of lease payments	(92)	(272)
Net financing cashflow	(74)	(140)
Net increase in cash and cash equivalents	1,581	8,954
Opening cash and cash equivalents	37,496	30,121
Effects of exchange rate changes on cash	(26)	(24)
Closing cash and cash equivalents	39,051	39,051

Planned use of funds

- Instrument development \$10-12m
- Future US FDA clearances
 - 3+ products up to \$2m per product
- Additional personnel
 - Sales, support, clinical, regulatory
- Marketing and launch costs new products
- Development of new amplification technology, decreasing time to result
- Ongoing R&D and IP costs



Financial Summary – 1H FY22 Profit & Loss



A'000s	1H FY22	1H FY21
Sales revenue	21,838	18,693
Cost of materials & freight	(6,890)	(6,086)
Gross profit	14,948	12,607
Employee benefits expense	(5,673)	(4,913)
Scientific consumables	(1,619)	(1,286)
Other expenses	(2,213)	(1,491)
EBITDA	5,443	4,917
Depreciation & amortisation	(814)	(655)
EBIT	4,629	4,262
Other income	72	235
Profit before tax	4,701	4,497
Income tax expense	-	-
Net income	4,701	4,497

Profitable while growing business

- 17% increase in 1H sales over pcp
- Gross margin **improved 1% to 68.4%** including freight costs
- Additional personnel, particularly overseas
- Ongoing R&D activity, clinical trials for FDA clearance
- Includes consulting costs for next generation instrument
- Additional instruments in customer sites & mfg

EasyScreenTM Detection Kit Range





Enteric

Detects 20+ gastroenteritis pathogens including Salmonella, Giardia and Norovirus



Respiratory

Detects 14 common respiratory infections including Influenza types A&B, Rhinovirus and SARS-CoV-2



ESBL & CPO

Detection of antibiotic resistant pathogens also colloquially known as "superbugs"



STI / Genital

Detects the most prevalent pathogen infections (Chlamydia, Gonorrhoeae, Syphilis and Trichomoniasis) plus many others



Flavivirus / Alphavirus

Refers to mosquito born pathogens including Dengue fever, Zika virus, Japanese Encephalitis virus and others



Meningitis

Detects 8 viral meningitis pathogens, a life-threatening infection surrounding the brain and spinal cord



Tick-borne Disease

Detects a range of infectious agents carried by ticks including Lyme disease, typhus and tick-borne encephalitis



Measles, Mumps, Rubella (MMR)

Highly contagious viral diseases that can result in death in severe cases



Dermatophytes

Fungal infections of skin, hair and nails which can become chronic in immunocompromised people

Registrations

Current

In process

TGA (€ IVD

TGA CE IVD

US EUA

TGA CE IVD

CE IVD



Instruments





GS-1000

- Higher throughput platform (188 samples per run)
- Onboard sample & reagent traceability
- Minimal hands-on time, userfriendly wizard driven interface
- Performs both automated sample extraction & PCR set-up



GS-Mini

- Fully automated cartridge design
- Rapid extraction of DNA & RNA
- Reduce wastage/costs; choice of 1 to 12 samples
- Easy to use barcode reader Increased flexibility to perform other in-house laboratory nucleic acid extractions
- Compact footprint (60cmx60cm)



GS1-HT

- Fast sample processing of up to 96 samples
- Sample extraction & both 96 & 384 well PCR set-up on one platform
- Reduced potential for decontamination
- Allows automation of common laboratory tasks



Image is concept only

Sample to Result Project

- Further reduce on-hands time & dramatically reduce time-to-result
- Work commenced & several suitable partners engaged
- Informed by market research & tailored to suit market needs

Enteric Product Range - Protozoan, Viral and Bacterial



Product Summary

- EasyScreen™ enteric testing kits allow for flexible syndromic testing of over 20 clinically relevant pathogenic microorganisms
- Improved diagnostics allows for earlier & more effective treatment, driving significant cost savings across health systems
- Common workflow with other EasyScreen™ diagnostic kits

Contract with Public Health Wales

- National tender won for molecular detection of enteric pathogens, including Protozoan, Viral and Bacterial targets
- Seven hospitals across Wales to screen all patients with 3base® technology
- 3 year term, option to extend for further 2 years
- A\$1.8M per annum equivalent

Enteric Protozoan Detection Kit – US Opportunity

- Est. 5.5m Enteric Protozoan tests pa in the US
- Initial focus on high throughput centralised labs
- Clinical trial recruitment close to complete
- Target to achieve up to 40% market share 5 years after clearance

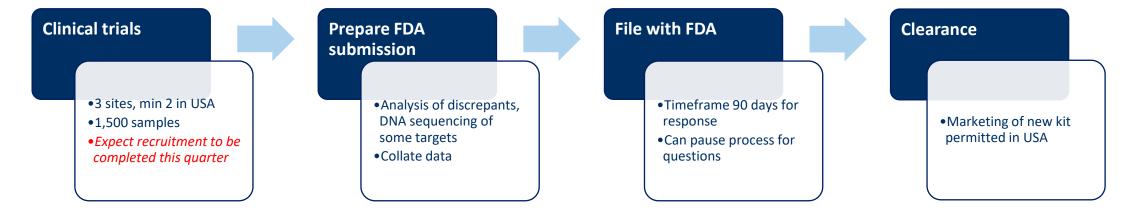


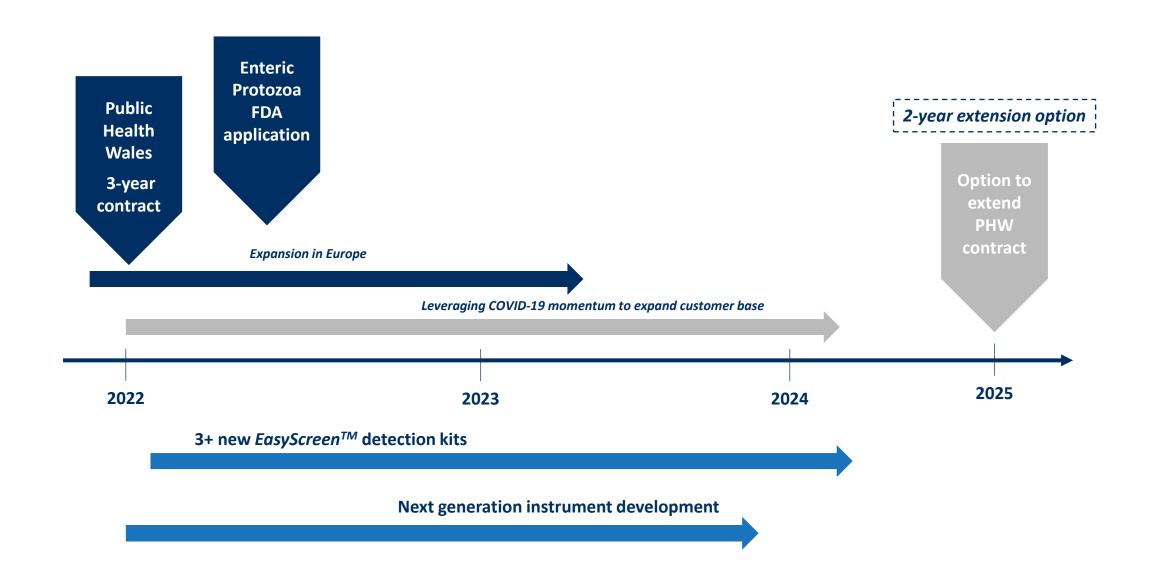


Enteric Protozoan Product – Road to FDA 510(k)



Process





Next Steps – Beyond COVID





Leverage COVID-19 – new customers, new tests

- Continue building interest in EasyScreen[™] kits in US & EU markets using expanded sales teams and SARS-CoV-2 experience as leverage
- SARS-CoV-2 only customers have five other kits with regulatory clearances available
- Promote & place GSS branded instruments



Product Development

- Next generation **3base**® 'sample to result' instrument
- Development of new amplification technology, markedly decreasing time to result, provisional patent filed
- Progress product registrations
 - FDA submission: Enteric Protozoan Detection Kit
 - TGA application for STI/Genital Pathogen Detection Kits
- Develop new test kits

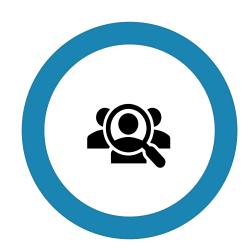






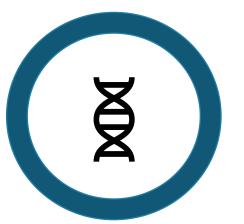
Revenue growth

- Maiden full year profit in FY21, profitable 1H22
- Cashflow positive 1H FY22, and continued revenue growth



Significant market opportunities

- Products sold in AU, EU & US
- Sales teams in key regions to engage with potential & existing customers



Continued product expansion

- 5 product groups in development
- Next generation 'sample to result' instrument
- Development of new amplification technology, reducing time to result



Attractive investment proposition

- Business model with favourable unit economics
- Increasing international recognition via EasyScreenTM SARS-CoV-2
- Unique technology 3base® –
 with patents issued with
 expiry to 2031+



Contact us

Dr John Melki
Genetic Signatures Ltd
Chief Executive Officer
E: john.melki@geneticsignatures.com
P: +61 (0)2 9870 7580

Visit us

www.geneticsignatures.com

Follow us





