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## **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 accurate diagnoses

# **EasyScreen™** Detection Kits

Streamlined universal sample processing kits linked to highly multiplexed real-time PCR screening assays

Applicable to bacterial, fungal, protozoan and viral (DNA & RNA) targets

Simultaneously detect over 20 pathogens from one sample, shortening turnaround from days to hours

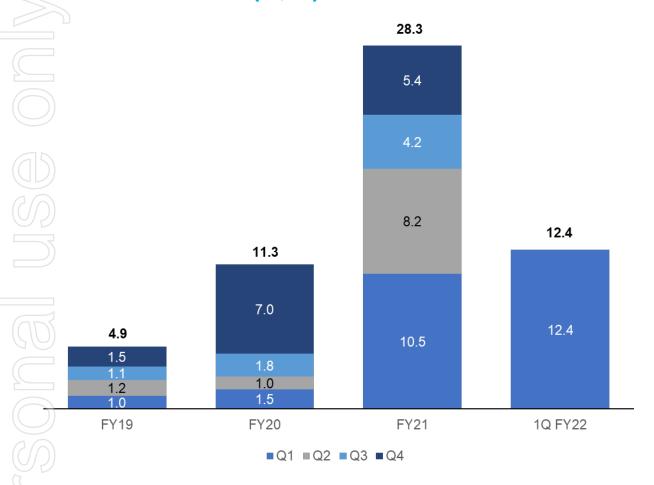
**3base™ can detect all SARS-CoV-2 variants**, including Delta; *EasyScreen™* compatible with existing lab technology



# **Financial Summary - Sales**

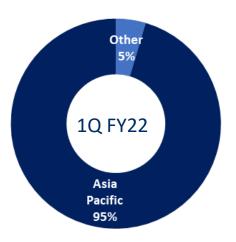


### Revenue from sales (A\$m)



#### **Continued Strong Revenue Growth**

- Record quarterly revenue \$12.4m from sales
- Demand for COVID tests continues due to ongoing outbreaks
- New instrument placements continue to support future demand for tests
- \$33m cash, no debt; drives future growth



# Financial Summary - Cashflow



A\$'000s	1Q FY22
Receipts from customers	10,340
Payments to suppliers and employees	(7,302)
Other	3
Net operating cash	3,041
Payment for plant & equipment	(92)
Net investing cash	(92)
Net proceeds from issue of shares	50
Principal elements of lease payments	(89)
Net financing cash	(39)
Net increase in cash and cash equivalents	2,910
Opening cash and cash equivalents	30,121
Effects of exchange rate changes on cash	7
Closing cash and cash equivalents	33,038

- Positive Q1 cashflow of \$2.9m
- Receipts from customers \$10.3m
  - Trade receivables balance @ 30 Sep \$8.1m,
    up \$2.7m on 30 Jun
- Payments to suppliers & employees 8% lower than pcp
  - Higher R&D and staff costs
  - Offset by reduced inventory purchases

## **Product Portfolio and Development Pipeline**





## **Beyond SARS-CoV-2**

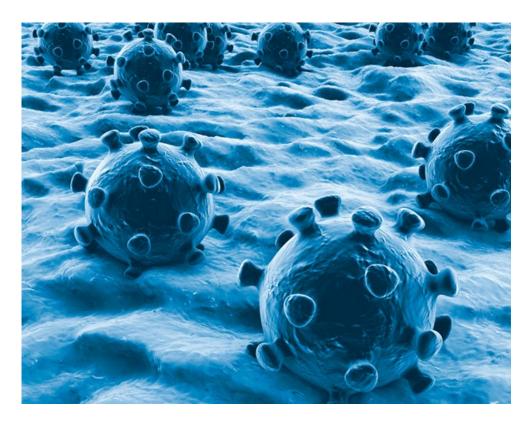


Initially developed SARS-CoV-2 test based on existing expertise in seasonal coronaviruses

**Driving global sales** – new customers in Europe and USA previously difficult to convert. Now interested in other  $EasyScreen^{TM}$  tests

Development of **new "fast" PCR test** that reduces batch processing times by 1.5 - 2 hours; now incorporated into *EasyScreen™* SARS-CoV-2 Detection Kit and in use in customer labs with very positive feedback

Conversion of other *EasyScreen*<sup>™</sup> tests to fast methodology underway – **significant benefit to laboratories** 



## **Regional Summary**



## **Europe**

- ~35% of molecular diagnostics market
- Direct sales in Germany & UK, distributors elsewhere
- Currently selling SARS-CoV-2 kit
  - First order for enteric test from UK customer others conducting in-house assessments with view to adopting
  - CE-IVD registration for:
  - Enteric

- SARS-CoV-2
- Respiratory
- ESBL/CPO
- STI

## **North America**

- Largest market, ~40% of PCR testing revenue
- Direct US sales, sales & support teams in place
- Distributor in Canada
- Selling SARS-CoV-2 kits to CLIA laboratories
- Enteric protozoan test in clinical trials for FDA

# Australia/NZ

- Head office, R&D, manufacturing
- Direct sales est. 10% of all testing volume
- First site in Queensland secured
- TGA registration for:
  - Enteric
  - SARS-CoV-2
  - Respiratory
  - ESBL/CPO
- STI registration lodged

# FDA 510(k) Clearance Program





Revenue per test	20% Market Share	30% Market Share	40% Market Share
US\$20	\$22.0m	\$33.0m	\$44.0m
US\$30	\$33.0m	\$49.5m	\$66.0m
US\$40	\$44.0m	\$66.0m	\$88.0m

#### **Market Dynamics**

- Est. 5.5m Enteric Protozoan tests per annum in the US
- Current methodology is microscopy slow and inaccurate
- Initial focus on largest 30 "high throughput" centralised labs
- Smaller decentralised labs more accessible with development of new testing hardware
- CPT code 87506 Infectious agent detection by nucleic acid (DNA or RNA); gastrointestinal pathogen; 6-11 targets (\$262.99)
- Review underway of next products for FDA clearance

#### **Status**

- 3 sites running FDA trials for GSS. Minimum 500 samples per site required
- Goal to complete sample collection by end CY2021, dependent on patient recruitment rate
- Aiming to win 40% market share within 5 years post FDA clearance



### **Financial information**

Share price (22-Oct-21)	A\$1.495
Shares on issue	143m
Market capitalisation	A\$214m
Ave monthly turnover (shares)	2.2m
Cash (30-Sep-21)	A\$33m
Debt (30-Sep-21)	Nil
<b>Enterprise value</b>	A\$181m

## **Top shareholders %**

Asia Union	26.2%
(Chris Abbott private investment)	20.270
Perennial Value Management	14.9%
Fidelity International	7.7%
Directors & management	3.1%









#### Leverage COVID-19 – new customers, new tests

- Continue building interest in EasyScreen<sup>™</sup> kits in US & EU markets using new sales teams and SARS-CoV-2 experience as leverage
- Targeting high throughput pathology groups, hospitals & govt programs
- Build long-term reliable customer contracts/relationships
- Embed EasyScreen<sup>TM</sup> workflows & demonstrate favourable unit economics
- Promote & place GSS branded instruments

#### **Product Development**

- Progress product registrations
  - FDA submission: Enteric Protozoan Detection Kit
  - TGA registration for STI/Genital Pathogen Detection Kits
- Next generation 3base<sup>TM</sup> 'sample to result' instrument
- Develop new test kits including flavivirus, measles, mumps & rubella, tick-borne diseases and dermatophytes





#### Revenue growth

- Maiden full year profit in FY21
- Q1 FY22 cashflow positive and record quarterly revenue



# Significant market opportunities

- Products sold in AU, EU& US
- Demand continues in FY22



# Continued product expansion

- 5 product groups in development
- Next generation 'sample to result' instrument



# Attractive investment proposition

- Business model with favourable unit economics
- Increasing international recognition via EasyScreen<sup>TM</sup> SARS-CoV-2
- Unique technology –
  3base<sup>™</sup> with patents issued with expiry to 2031+

