



Genetic Signatures

ASX: GSS

INVESTOR PRESENTATION

AUGUST 2018

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REVENUE STAGE HEALTHCARE COMPANY LISTED ON THE ASX

Genetic Signatures Limited (ASX: GSS)

A specialist molecular diagnostics company

- Focused on becoming a global leader in the supply of molecular diagnostic solutions
- Developing and commercialisation its proprietary platform technology, **3base™**
- Implementing its commercial strategy through a team of 30+ across Australia, Europe and North America

Financial information

Share price (14-Aug-18)	A\$0.580
Shares on issue	103.9m ¹
Market capitalisation	A\$60.3m
Cash (30-Jun-18)	A\$9.0m
Debt (30-Jun-18)	Nil
Enterprise value	A\$51.3m

Notes:

1: Excludes 1.05m unquoted options (various expiration dates and prices)

Share price (A\$)

Volume (m)



Top shareholders

	%
Asia Union and Christopher Abbott	39.6
Deutsche Bank AG	14.2
Directors, management & advisors	>6.0

GENETIC SIGNATURES DEVELOPS AND MANUFACTURES MOLECULAR DIAGNOSTICS KITS BASED ON ITS PROPRIETARY 3BASE™ TECHNOLOGY



Established revenues

Track record of revenues de-risks development and sales expansion

Focus on profitability

High margin, high volume sales, with focus on production efficiencies

Strong portfolio growth

Substantial product growth with 5 products expected by end of 2018

Geographic expansion

Expansion underway into key US and EU markets, with multiple patents issued (expire 2031+)

Strong sector tail-winds

Surging global MDx market growing from US\$6bn to US\$12bn in 2022

3BASE™ TECHNOLOGY REDUCES COMPLEXITY IN GENETIC TESTING ALLOWS TESTING FOR MORE DISEASES WITH HIGHER ACCURACY MORE RESULTS, MORE ACCURACY, LOWER COST

3base™ technology

- **3base™** platform technology converts original 4-base microbial genome to 3-base
 - Reducing complexity in molecular testing
- Conversion occurs during standard procedures with no additional steps for the technician

3base™ MDx testing is focused solely on the detection of infectious diseases

- ✓ **3base™** MDx can identify a **wider array** of patient infections than current diagnostics
- ✓ Reduces **time to results** (5 vs 24+ hours for conventional methods)
- ✓ Provides greater testing **accuracy by reducing complexity in a single test sample**

3base™ technology in the detection process

Regular cell Lysis and preparation steps

Original genome



3base™
conversion

Converted **3base™**
genome



Standard Real-Time PCR instrument

1,048,576 combinations for a 10 digit number with 4-base

G	C	C	T	C	A	C	T	T	C
↓	↓	↓	↓	↓	↓	↓	↓	↓	↓
G	T	T	T	T	A	T	T	T	T

59,049 combinations for a 10 digit number with 3-base

Powerful evidence of efficacy from clinical trials

Clinical trials demonstrate efficacy

Evaluation study conducted at St Vincent's Hospital, Sydney

221 patient samples tested and compared to traditional culture, microscopy, and antibody based tests

Results highlight the efficacy of 3base™ technology and GSS products

- ✓ **Faster screening:** Generated results in 4 hours, compared to up to 120 hours for traditional testing methods
- ✓ **Greater accuracy:** Identified 44 infections that existing testing missed

St Vincent's Hospital Evaluation Study results

Pathogen	Conventional Methods*	EasyScreen™
Campylobacter	7	9
Salmonella	8	9
Shigella	5	6
C. Difficile	3	7
Yersinia	-	1
Cryptosporidium	-	1
Giardia	9	12
Dientamoeba fragalis	4	20
Blastocystis hominis	16	21
Entamoeba histolytica	1	1
Norovirus group 2	-	7
Adenovirus	-	1
Adenovirus 40/41	-	1
Sapovirus	-	1
Total	53	97

Significantly greater efficacy

GSS is commercialising a range of **MDx tests** based on **3base™** and branded as **EasyScreen™**

EasyScreen™ kits offer faster and more effective detection of infectious disease

- ✓ Four product families available covering **enteric microorganisms, respiratory viruses (including influenza), ESBL & CPO “Superbugs” and STIs**
- ✓ Target customers are high-volume hospitals and pathology laboratories

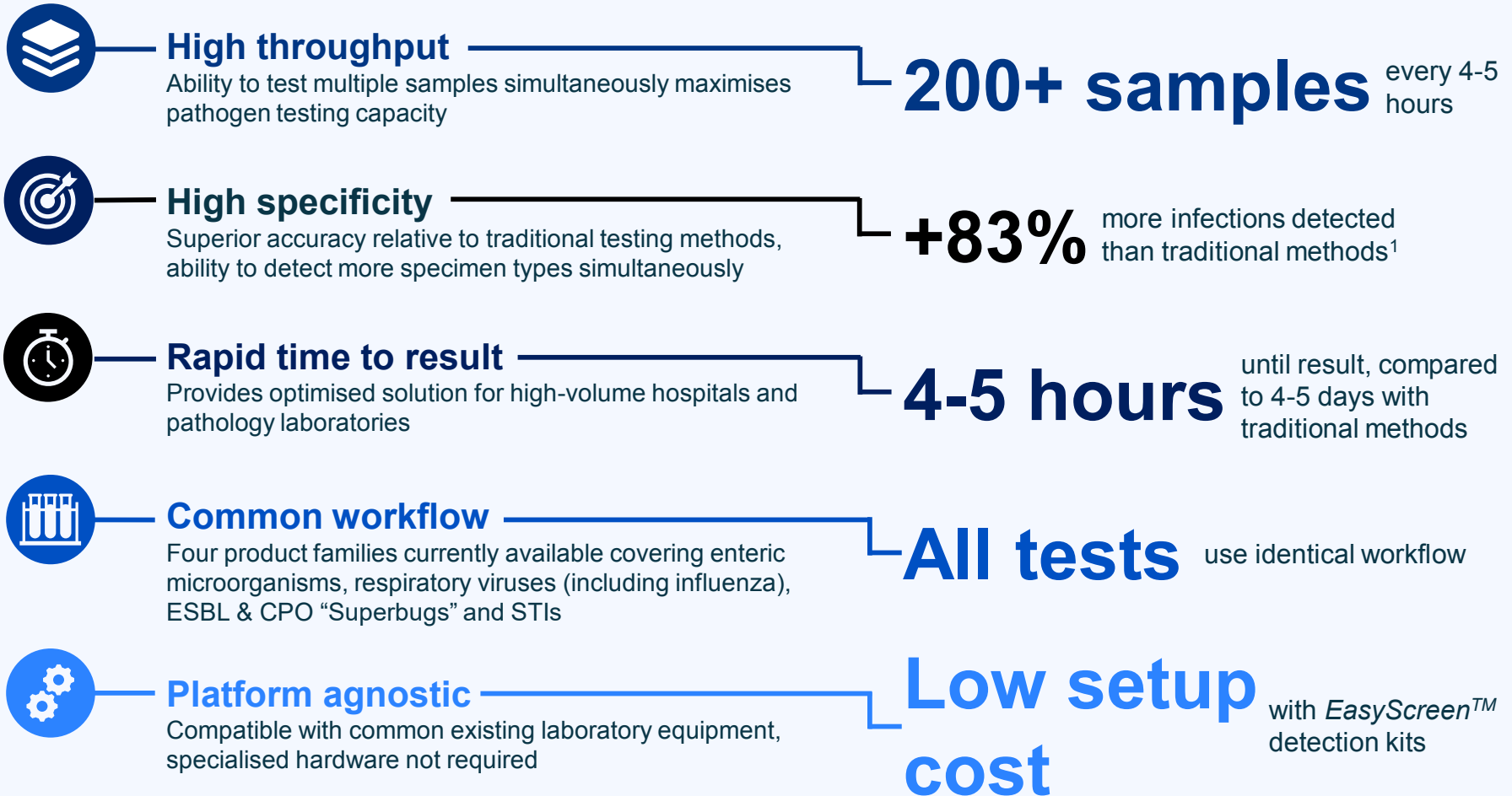


- ✓ **Common Workflow** – *all tests work under identical conditions*
- ✓ **Platform agnostic** – *compatible with common existing laboratory equipment*
- ✓ **High specificity** – *superior accuracy of results*
- ✓ **Rapid time to result (4 - 5 hours)** – *would traditionally take 4 - 5 days*

EASYSscreen™ KITS IDENTIFY MORE PATHOGENS IN LESS TIME

The EasyScreen™ competitive advantage

EASYSCREEN™ KITS OFFER FASTER AND MORE EFFECTIVE DETECTION OF INFECTIOUS DISEASE; OPTIMISED FOR HIGH THROUGHPUT LABS



Notes:

1: St Vincent's Hospital Evaluation Study results, see page 6 for further details on the study

Q4 FY18 trading update

INTERNATIONAL EXPANSION GAINING MOMENTUM

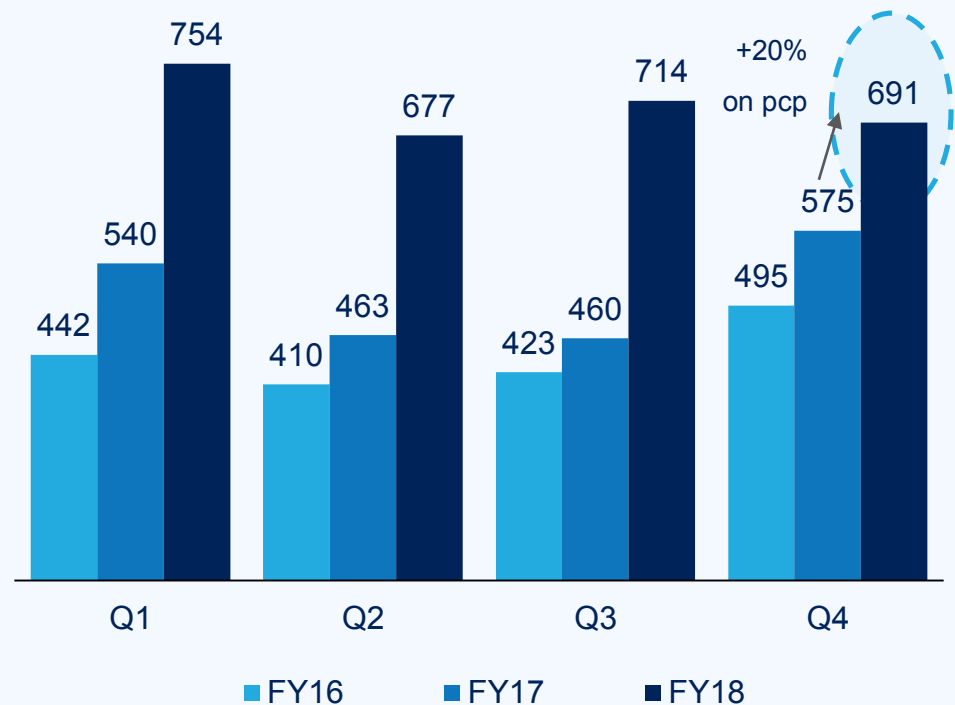
Q4 FY18 Revenue Update

- ✓ **Strong quarterly revenue growth to A\$691k in Q4 FY18, a +20% increase on pcp**
- ✓ **First sales in the US**

Q4 performance continues **consistent quarter-on-quarter revenue growth** since listing in 2015

Strong full year revenue of A\$2.8m for FY18 representing a **+39% increase** from FY17

Quarterly revenue (A\$000)



Key corporate and operational milestones achieved

GSS HAS MADE STRONG PROGRESS OVER THE LAST 18 MONTHS

- ✓ Successful clinical trial results for **Flavivirus Detection Kits** at prestigious European conference in April 2017
- ✓ TGA and CE-IVD Mark approvals received for the entire **Enteric product suite** by June 2017
- ✓ Approval received for **3base™** patent in the USA until 2033
- ✓ First sale and delivery of **Flavivirus and Alphavirus Detection Kits** to an offshore customer in June quarter 2017
- ✓ Relocation to larger facilities in Sydney to accelerate next phase of operational growth
- ✓ Successful beta-release of second-generation **Respiratory Detection Kits** in September quarter 2017
- ✓ Record quarterly revenues of A\$754k in September quarter 2017 driven by strong domestic flu season
- ✓ Maiden sales for **ESBL & CPO (“superbug”) Detection Kits** for use in a trial in December quarter 2017
- ✓ Expansion of European team with appointment of field scientist and sales staff, with further recruiting continuing
- ✓ CE-IVD Mark approvals received for **ESBL & CPO Detection Kits** (‘antibiotic resistance’) in April 2018
- ✓ Successful trial results for **ESBL & CPO Detection Kits** presented at major European conference in April 2018
- ✓ Maiden sales in the USA with sales order for **ASR kits** in May 2018
- ✓ TGA registration received for **ESBL & CPO Detection Kits** in May 2018
- ✓ **Major customer contract signed** with large Australian pathology service provider in August 2018

GENETIC SIGNATURES HAS A CLEAR FOCUS ON REVENUE GROWTH IN FY19 DRIVEN BY ITS INTERNATIONAL EXPANSION AND NEW PRODUCT RELEASES

Roll-out of expanded product suite

- Current expectations of 5 product groups to be approved in the near term across target regions
- Continued progress with international registrations

Targeted international sales expansion

- Sales expansion into North America (US\$4.6bn market) and Europe (US\$1.6bn market)
- Multiple trials underway in the US, Europe and Australia

Capitalise on the growing MDx opportunity

- Independent projections of total MDx revenue reaching >US\$12bn by 2022
- Substantial current corporate activity within the MDx sector several acquisitions underway

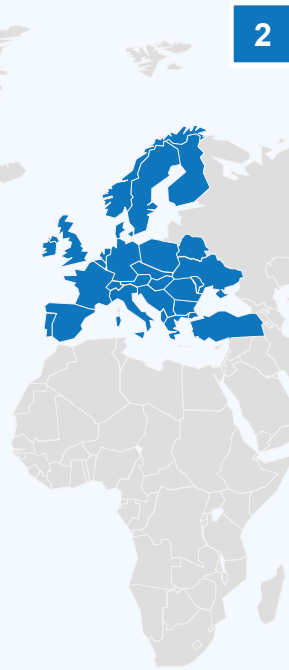
MULTIPLE TRIALS UNDERWAY IN THE US, EUROPE AND AUSTRALIA



3 North America: Accelerate direct sales approach
Trials underway

- ✓ First ASR sales achieved in the US
- ✓ Sales permitted through US FDA listing and ASRs
- ✓ Certified for Health Canada

Immediate focus on securing trials with larger pathology labs
ASR product trials underway



2 Europe: Rapidly increasing footprint to be ready for product approvals
Trials underway

- ✓ Distributors appointed (Italy, Israel, Ireland and Poland)
- ✓ CE-IVD approvals granted for enteric and ESBL & CPO
- ✓ 3 products approaching CE-IVD approval in 2018
- ✓ Appointment of a European based field scientist and other sales staff and still actively recruiting

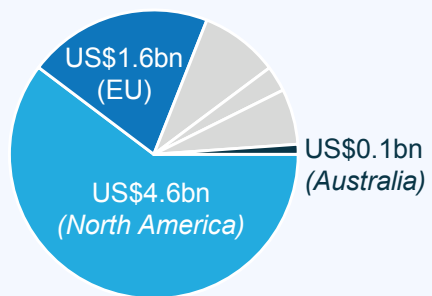
Plan to appoint additional distributors in short term
Product trials underway



1 Australia: Continued rapid growth
Small base of customers with a number of trials underway

- ✓ Major new contract signed for respiratory kit
- ✓ 2 products approaching TGA approval in 2018

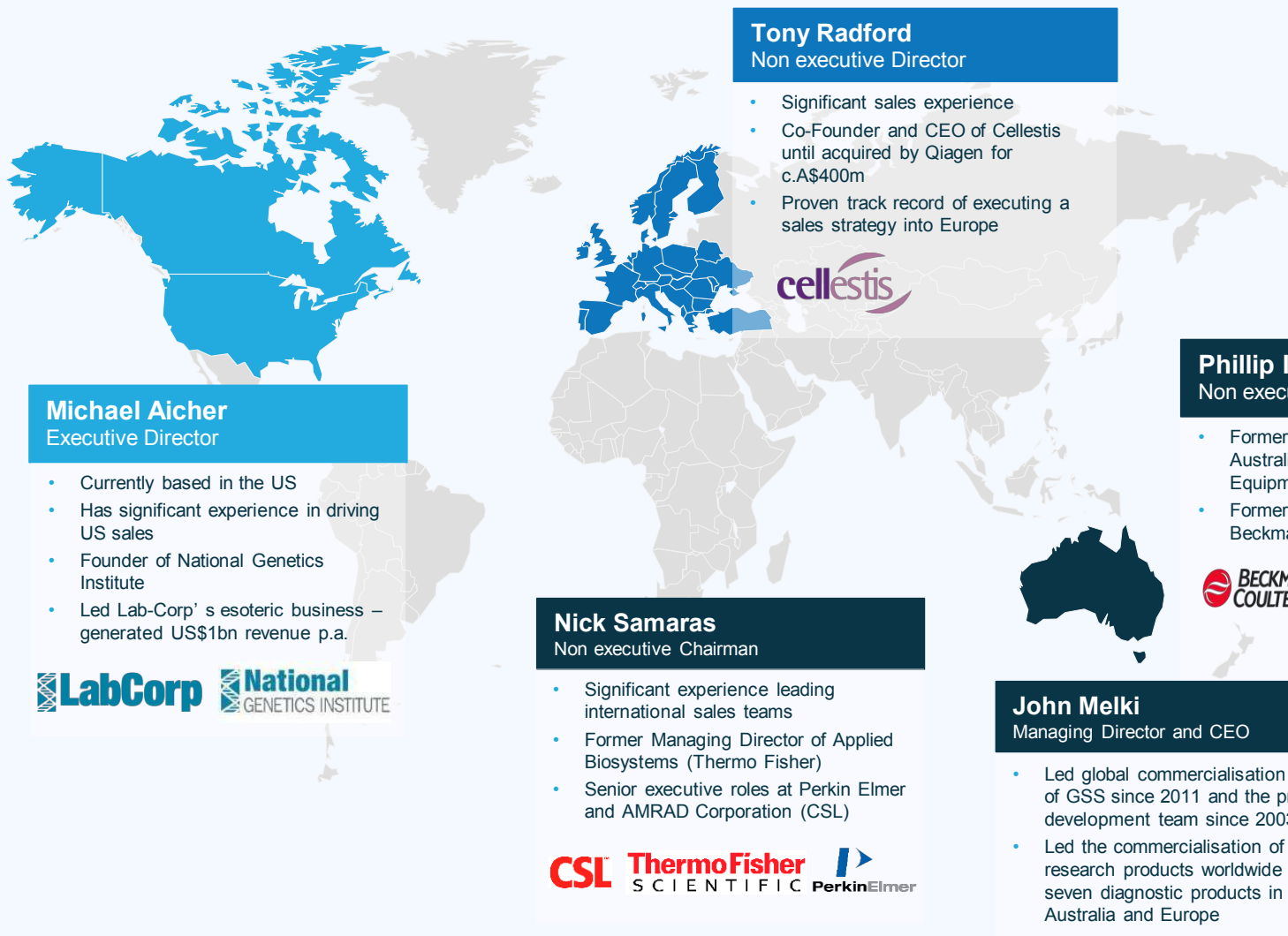
Further customer acquisitions planned in FY19



Estimated MDx market size (2017)

Source: World Market for Molecular Diagnostics, 5th Edition (Infectious Disease, Oncology, Blood Screening, Pre-Natal and Other Areas) Kalorama Information, Published: 1/9/2013, page 94.

Board of Directors have a track record of success



1 Australia: continue rapid growth

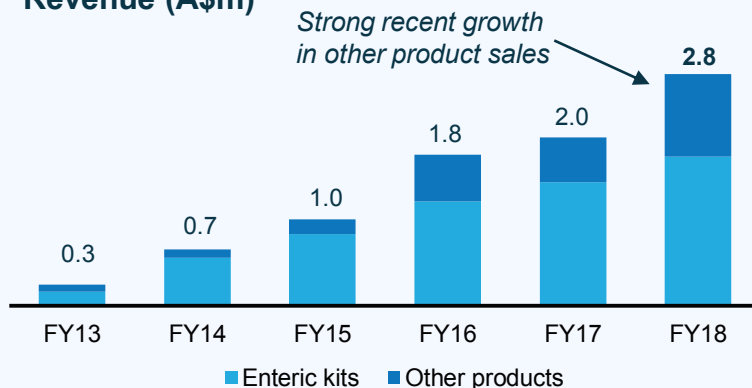
AUSTRALIA IS THE PLATFORM FOR INTERNATIONAL SALES

Commercialisation progress

Established foothold

- Secured relationships with **top tier customers in NSW and Victoria**
- **Major customer contract signed** with large Australian pathology service provider in August 2018
- Established customer relationships **increase ease of new product sales**
- Platform in NSW and Victoria has driven **strong revenue growth over last 4 years**

Revenue (A\$m)



Sources:

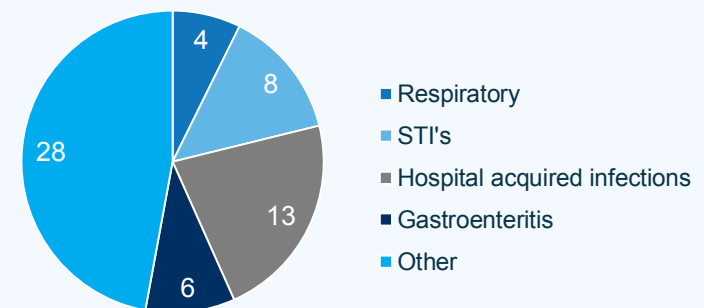
1: World Market for Molecular Diagnostics, 5th. Edition (Infectious Disease, Oncology, Blood Screening, Pre-Natal and Other Areas) Kalorama Information, Published: 1/9/2013, page 94.

Next steps

Rapid customer acquisitions

- Targeting trials with large hospitals and clinical labs in Australia to provide **significant domestic revenue growth**
- 2 products approaching TGA approval in 2018 expected **drive new customer acquisitions**
- Target customers include **pathology labs, hospitals** and large **research laboratories**

Australia MDx market potential (US\$m)



2 Europe: strong progress on multiple fronts

ACTIVELY INCREASING PROFILE IN PREPARATION FOR PRODUCT APPROVALS

Commercialisation progress

Distributors hired and evaluations underway

- Currently **adding to European team**
- **Field scientist appointed** based in Europe
- Recent **CE-IVD registration for ESBL & CPO “superbug”** Detection Kits
- **Sales logistics being finalised** with EU-based warehouse in final legal review
- Building traction with customers with **sales order received** across a number of products
- Established VAT deferment program

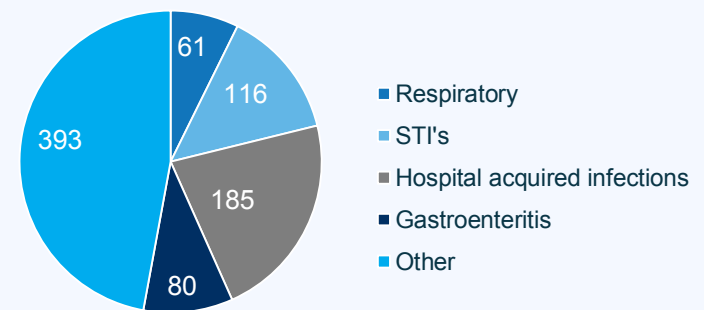
Genetic Signatures is developing a mix of direct sales and partnerships with distributors across Europe to optimise the sales network

Next steps

Strategic market with large opportunity

- Europe has an addressable MDx market of **c.US\$435m** - *c.20% of the global market¹*
- Continue to drive revenue growth and pursue regulatory registration for STI and respiratory products in Europe
- **Actively recruiting sales and support staff** in multiple countries
- Advanced discussions with new distributors in other European countries

European MDx market potential (US\$m)



Sources:

1: World Market for Molecular Diagnostics, 5th. Edition (Infectious Disease, Oncology, Blood Screening, Pre-Natal and Other Areas) Kalorama Information, Published: 1/9/2013, page 94.

3 North America: accelerate direct sales approach

FIRST ASR SALES AND PROGRESS TOWARD FDA CLEARANCE

Commercialisation progress

Increasing traction with ASR approach

- **First ASR sales contract received** with US pathology laboratory
- Several **labs assessing the potential for ASR products** available for sale in the USA
- **Initial trials established** with local clinical labs using GSS supplied products in FY17
 - Trials reaching conclusion
- Also progressing towards **securing FDA clearance** for Enteric Protozoan Detection Kit
- **3base™** technology has patents issued in the US
- Quality Management System **certified for Health Canada**

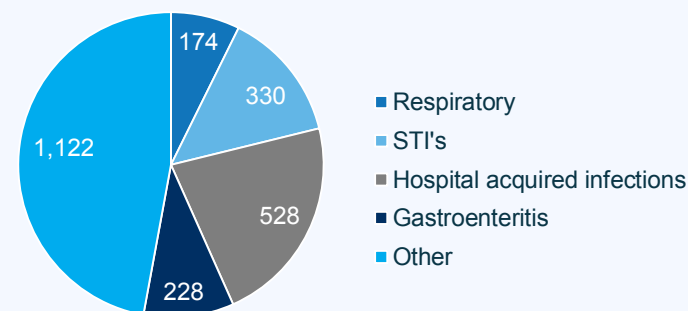


Next steps

Largest market for MDx globally

- Addressable market of approximately **US\$1,265m**, c.55% of the global market
- Over 5,600 registered hospitals in the USA and 11,000 CLA certified laboratories
- FDA listings of Clinical Concentrators in place to allow initial sales

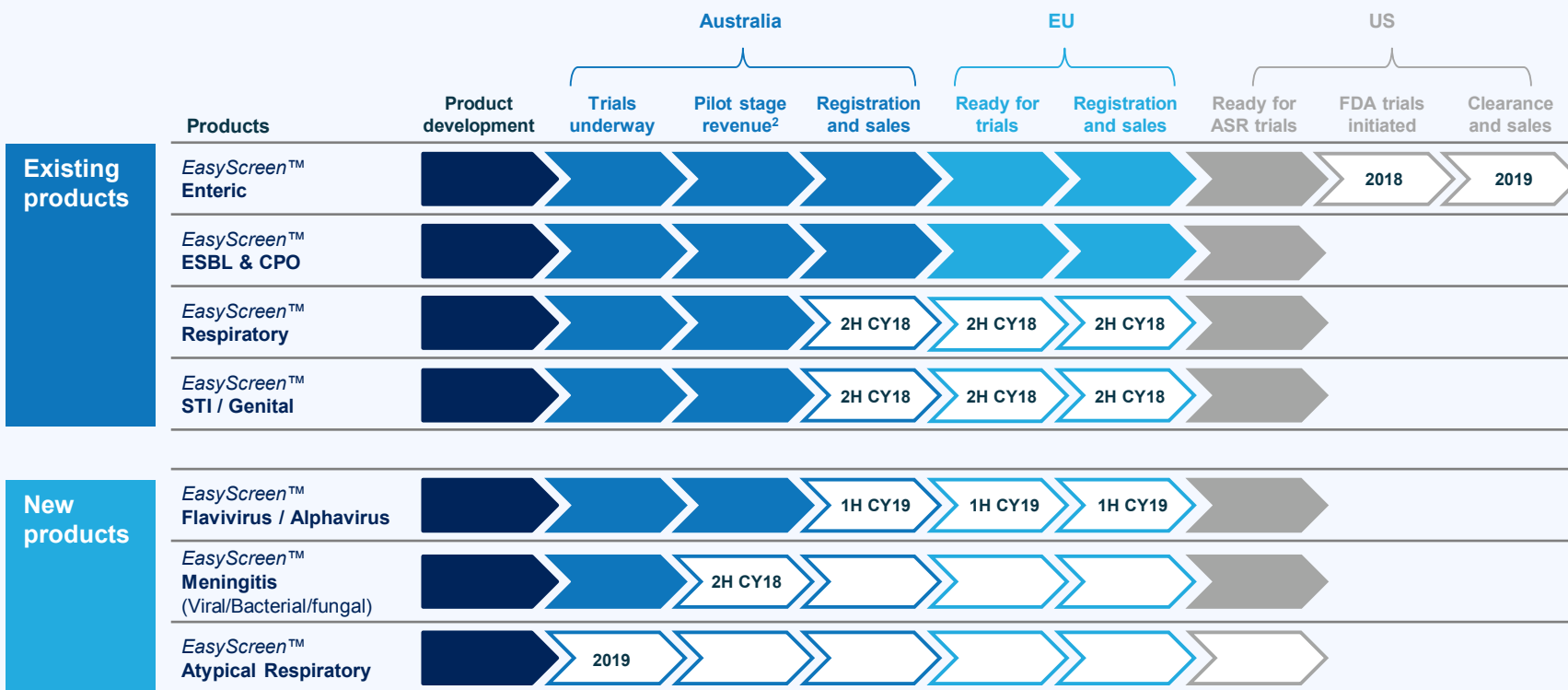
North American MDx potential¹ (US\$m)



Sources:

1: World Market for Molecular Diagnostics, 5th. Edition (Infectious Disease, Oncology, Blood Screening, Pre-Natal and Other Areas) Kalorama Information, Published: 1/9/2013, page 94.

Clear path to global commercialisation for all products

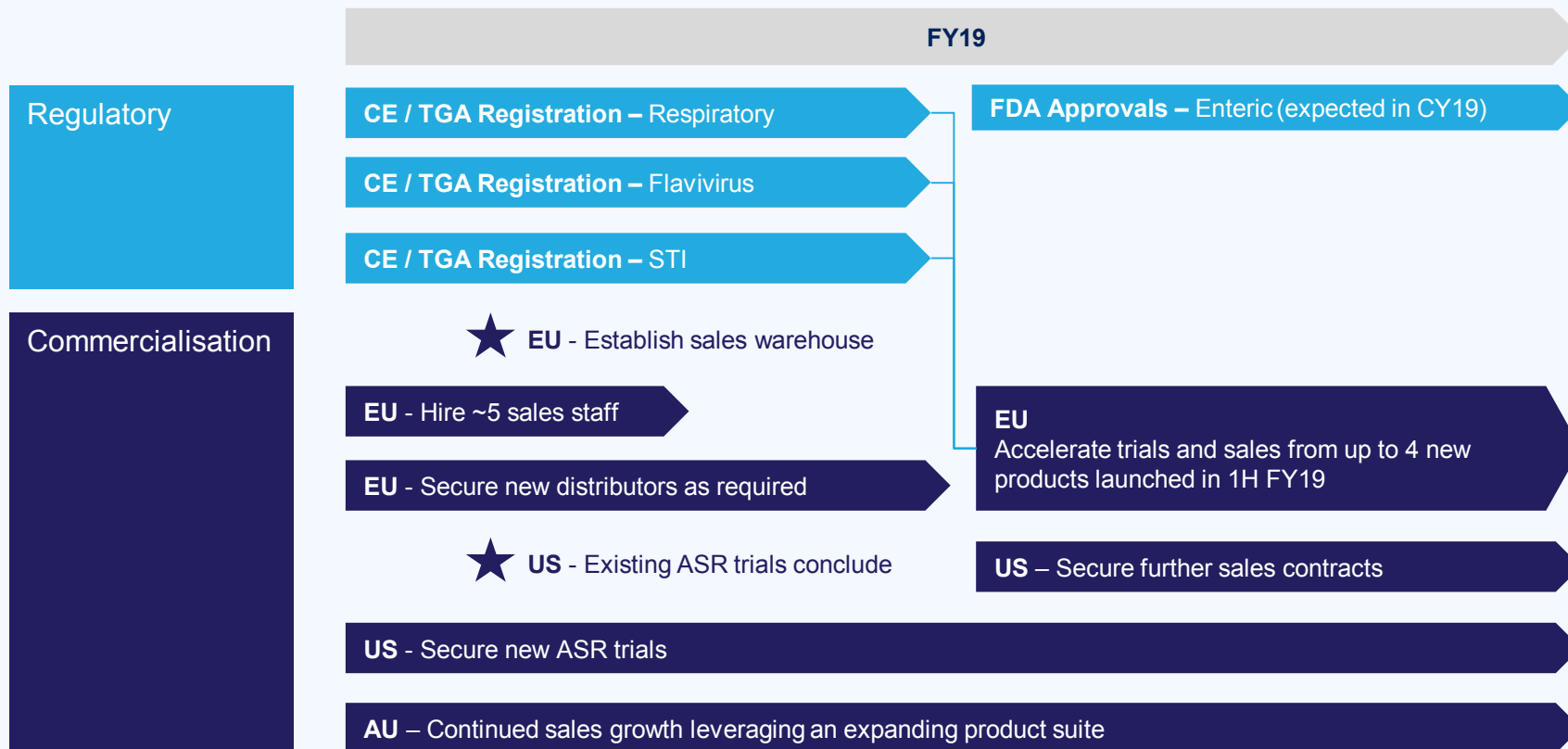


Notes:

- 1: Each of the product suites may contain multiple different individual products, requiring multiple regulatory submissions and registrations for a suite of products
- 2: Pilot stage revenue generated from sales for research use only (medical professionals may use products for diagnosis only where they validate results with independent testing)
- 3: Sold as individual Analyte Specific Reagents (ASR) for a number of pathogens in the USA

GSS HAS MADE SUBSTANTIAL PROGRESS ON ALL PRODUCTS SINCE IPO

GSS IS WELL POSITIONED TO DELIVER ON ITS GLOBAL COMMERCIALISATION STRATEGY IN FY19



Significant opportunity to accelerate global sales and revenue

US\$1.2bn market opportunity for 3 existing products



ENTERIC

Commercialisation underway with **full regulatory registration in Europe received**

3base™ advantages:

- Tests for over 20 potentially deadly enteric pathogens including *Salmonella*, *C. difficile* Bacteria and Protozoan species
- Successful trials and validation with higher accuracy, sensitivity and detection rates

Market opportunity:

US\$380m

Global addressable market (2017)



RESPIRATORY

Significant growth opportunity with **registration in Australia and Europe in progress**

3base™ advantages:

- Simultaneously identifies 15 of the most common respiratory infections
- Rapid screening across high volume sample sets
- Detects significantly more respiratory infections compared to existing methods

Market opportunity:

US\$290m

Global addressable market (2017)



STI

Innovative STI detection kit testing pathogens **contracted by 1 million people daily**

3base™ advantages:

- Proven market (sales underway)
- Highly successful clinical trials and validation
- Rapid testing for 12 pathogens
- Uses existing lab equipment
- Higher accuracy, sensitivity and detection rates compared to existing techniques

Market opportunity:

US\$550m













Global addressable market (2017)

Sources:

1: World Market for Molecular Diagnostics, 5th. Edition (Infectious Disease, Oncology, Blood Screening, Pre-Natal and Other Areas) Kalorama Information, Published: 1/9/2013, page 168

Significant strategic interest in MDx technologies

- Major MDx players are pursuing aggressive acquisition strategies to secure emerging technologies
 - ...and they are prepared to pay high multiples to secure opportunities

Date ↓	Target	Acquirer	Deal value (US\$m)	EV/Revenue multiple (x)	Target Description
Mar-18	 STAT Dx Closer to Care	 QIAGEN	191	27	<ul style="list-style-type: none"> • Targets enteric and respiratory infections and is focused on lower throughput
Feb-18	 RHS Unique Genetic Solutions	 PerkinElmer	20	N/A	<ul style="list-style-type: none"> • Provides platform for MDx processes based on single cell genomic technologies
Dec-17	 ignyta®	 Roche	1,700	N/A	<ul style="list-style-type: none"> • Focuses on MDx for cancer, and treatment through precision medicines
Dec-16	 Multiplicom	 Agilent Technologies	72	6.5	<ul style="list-style-type: none"> • Provides MDx kits for cancer, genetic disorders, pre-natal screening
Nov-16	 Cepheid. A better way.	 DANAHER	3,860	5.5	<ul style="list-style-type: none"> • Specializes in MDx for infectious diseases and cancer
Aug-15	 GeneWEAVE	 Roche	425	N/A	<ul style="list-style-type: none"> • Owns Smarticles, a type of MDx identifying multidrug-resistant organisms

Source: Company filings

A detailed electron micrograph of a spherical virus particle, likely a coronavirus, showing its characteristic surface covered in spike-like proteins. The particle is centered in the frame against a dark blue background. A semi-transparent dark blue horizontal band is overlaid across the middle of the image, containing the text.

APPENDIX

FURTHER INFORMATION ON 3BASE™ TECHNOLOGY

7 product groups underpinned by 3base™ technology

	Product group	Target pathogens	Key highlights	
Four existing products groups	<i>EasyScreen</i> ™ Enteric	20+ enteric pathogens Includes Salmonella, <i>C. difficile</i> Bacteria, Protozoan and Viral Detection Kits and <i>C. difficile</i> reflex kits	All tests have full regulatory registration in Europe (10-20x the size of Aus market)	
	<i>EasyScreen</i> ™ ESBL & CPO	Extended Spectrum Beta-Lactamase and Carbapenemase Producing Organisms	Full regulatory registration in Europe (CE-IVD) and Australia (TGA)	
	<i>EasyScreen</i> ™ Respiratory	15 respiratory pathogens (including influenza A & B, Rhinovirus and M. pneumonia)	Registrations in Australian and European markets currently in progress	
	<i>EasyScreen</i> ™ STI / Genital	12 most commonly encountered STIs (including chlamydia, gonorrhoea and syphilis)	Offers superior test efficacy for pathogens contracted by 1m people daily ¹ (market worth ~US\$550M ²)	
Commercial sales underway				
	Three new product groups	<i>EasyScreen</i> ™ Flavivirus / Alphavirus	Multiple Flavivirus/Alphavirus viral families (including Zika and West Nile virus)	Offers testing for complex viruses causing widespread morbidity and mortality but historically labour-intensive to detect
		<i>EasyScreen</i> ™ Meningitis (Viral and Bacterial)	Multiple strains of Meningitis (viral and bacterial infections)	In final development stages – offers rapid diagnosis for life-threatening bacterial meningitis
<i>EasyScreen</i> ™ Respiratory Atypical Pathogen		Simultaneous detection of leading causes of bacterial respiratory infection	In development	
Trials underway				

Sources:

1: <http://www.who.int/mediacentre/factsheets/fs110/en/>

2: World Market for Molecular Diagnostics, 5th. Edition (Infectious Disease, Oncology, Blood Screening, Pre-Natal and Other Areas) Kalorama Information, Published: 1/9/2013, page 168

Detection of infectious diseases

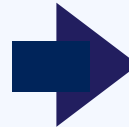
THERE IS POTENTIAL TO TRANSFORM THE MODERN MICROBIOLOGY LABORATORY AND THE CARE OF PATIENTS WITH SUSPECTED INFECTIONS

Current process

Traditional methods for diagnosing infection have relied largely on identifying pathogens via culture

Conventional culture-based microbiology

1. Sample collected from patient
2. Preparation of medium (i.e. agar plate)
3. Incubation
4. Analysis (i.e. microscopy)



Industry problems

Inefficient: window between specimen collection and organism identification is large and can:

- × Enhance spread of infection to other patients
- × Delay onset of appropriate course of treatment

Limited scalability: targets often require separate tests

- × Analysis can be time consuming and require significant technician time

Reduced patient outcomes

- × Broad empirical antimicrobials are given to excessive numbers of patients to avoid the potential grave consequences

Genetic Signatures' 3base™ technology can alleviate many of the pressures evident in the detection of infectious diseases and deliver improvements in time, quality and cost to meet surging demand

3base™ reduces complexity by 94%

SIGNIFICANT REDUCTION IN COMPLEXITY WHILE MAINTAINING OR INCREASING ACCURACY

20% improvement in homology

94% reduction in complexity

- ✓ Sufficient information retained for genotyping equivalent to native (4 base) genomic assays
- ✓ No loss of clinical specificity is observed by this base conversion¹
- ✓ **3base™** delivers greater Sensitivity and Specificity, in a rapid assay

3base™ showed superior performance in HPV clinical trial

	4 base sequences	3base™ base sequences
Seq 1	GATGG C GAT I ATGGT I G A C A C	GATGGTGATATGGT I GATAT
Seq 2	GATGG I G A C ATGGT A G A I A C	GATGGTGATATGGT A GATAT
Seq 3	GATGG I G A I ATGGT G G A C A C	GATGGTGATATGGT G GATAT
Seq 4	GATGG I G A I ATGGT A G A I A I	GATGGTGATATGGT A GATAT
Seq 5	GATGG I G A I ATGGT G G A C A C	GATGGTGATATGGT G GATAT
Seq 6	GATGG C G A C ATGGT I G A I A I	GATGGTGATATGGT I GATAT
Seq 7	GATGG I G A I ATGGT G G A C A C	GATGGTGATATGGT G GATAT
Seq 8	GATGG I G A C ATGGT A G A I A C	GATGGTGATATGGT A GATAT
Seq 9	GATGG I G A I ATGGT A G A I A C	GATGGTGATATGGT A GATAT
Seq 10	GATGG I G A I ATGGT G G A I A C	GATGGTGATATGGT G GATAT
Consensus	GATGG Y G A Y ATGGT Y G A Y A Y	GATGGTGATATGGT D GATAT

75% homology over 20 bases

48 possible primer combinations



95% homology over 20 bases

3 possible primer combinations

Sources:

1: HPV clinical trial for example showed superior performance vs. Digene assay in reducing false positives (J. Clin. Virol. 42:22-6. 2008)

EasyScreen™ customers are strong advocates

*“We use EasyScreen™ at ACL for its **robust user friendly automation, higher levels of specificity** for clinical samples and our confidence in delivering **clinically significant results** to our clinicians.*

Tony Field

Operations Manager

Australian Clinical Labs

*“Same day results and triage options for urgent requests help potentially **save lives**. And we get a high level of service and support from the GSS team!”*

*“I find that the **fast turnaround time** and the number of targets tested in the EasyScreen™ assays allow me to **more rapidly identify highly infectious agents**, potentially stopping the spread to other healthy individuals and thereby **saving the health system money**.”*

Dr Damien Stark

KOL and Senior Microbiologist

St Vincent's Hospital Sydney