ASX Market Announcement



GTG presenting at HC Wainwright Conference

Melbourne, Australia, 9 March 2021: Genetic Technologies Limited (ASX: GTG; NASDAQ: GENE, "Company", "GTG"), a diversified Genomics and AI driven preventative health business advises that Chief Executive Officer, Simon Morriss will be presenting at HC Wainwright's Global Life Sciences Conference at 9am EST/ 11pm AEDT, Tuesday 9 March 2021.

Additionally, management will be hosting a 'Meet the CEO' event for Australian based investors to have the opportunity to hear from Chief Executive Officer, Simon Morriss on the strategy for the Company and the recently announced Infinity BiologiX agreement.

Date: Thursday 11th March 2021

Time: 1:00pm AEDT

Registration: https://us02web.zoom.us/webinar/register/WN 6K30zFxQSKug6DAgg6hqQg

Attached is the associated presentation.

-END-

Authorised by the Board of Genetic Technologies



Investor Relations (AUS)

Stephanie Ottens Market Eye

M: +61 434 405 400

E: stephanie.ottens@marketeye.com.au

Investor Relations and Media (US)

Dave Gentry 1 800 RED CHIP (733 2447)

Cell: 407 491 4498 E: dave@redchip.com

About Genetic Technologies Limited Genetic Technologies Limited (ASX: GTG; Nasdaq: GENE) is a diversified molecular diagnostics company. GTG offers cancer predictive testing and assessment tools to help physicians proactively manage patient health. The Company's lead products GeneType for Breast Cancer for non-hereditary breast cancer and GeneType for Colorectal Cancer are clinically validated risk assessment tests and are first in class. Genetic Technologies is developing a pipeline of risk assessment products.

For more information, please visit www.gtglabs.com





Genetic based preventative health platform

Genetic Technologies Investor Presentation March 2021

Authorised by the Board of Genetic Technologies

Notice: Forward looking statements



The purpose of the presentation is to provide an update of the business of Genetic Technologies Limited ACN: 009 212328 (ASX:GTG; NASDAQ:GENE). These slides have been prepared as a presentation aid only and the information they contain may require further explanation and/or clarification. Accordingly, these slides and the information they contain should be read in conjunction with past and future announcements made by Genetic Technologies and should not be relied upon as an independent source of information. Please refer to the Company's website and/or the Company's filings to the ASX and SEC for further information.

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*Defined terms are outlined in the Appendices to this presentation

Genetic Technologies

Empowering a healthier life!

A diversified genomics and Al driven preventative healthcare company, partnering to provide individualised risk assessment of serious disease.

Genetic Technologies - Leader in Genomics



Strong focus on R&D

Over a decade of R&D on the development of polygenic risk scores

Multi-year distribution agreement

US License and distribution agreement for COVID-19 Risk Test with IBX for minimum of US\$2.9 million over 3 years

Launched CIT in USA & Aus for other tests

Robust patent portfolio

15 patents granted and 7 patent families pending

\$24 million

Strong cash balance with 18-24 month runway¹

Publications and academic collaborations

Multiple peer-reviewed publications and four collaborations with prestigious academic and medical establishments

Up to 70%

Coverage for all mortalities from tests in development for serious disease risk including major oncological, metabolic and degenerative diseases











We aim to offer the most comprehensive suite of genetic risk assessment tests on the market

Product overview – innovation Pipeline



Test kit

Released 2019/2020





Breast Cancer

geneType

Type 2

diabetes

Type 2 Diabetes

A personalized
Type 2 Diabetes

Exempt human specimen

Distribution partnership confirmed



COVID-19

Germline Products Under Development



BRCA Panel



Lynch Syndrome

TGA Approved



Depression (PREDICTIX by Taliaz)

PRS Products Under Development







Melanoma



Cardiovascular Disease



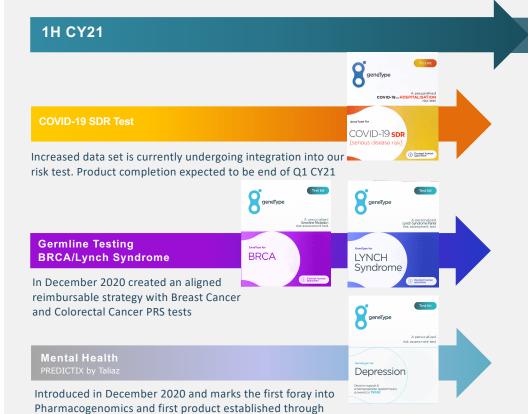
Multitest

Overview of product timeline

Remain on track to deliver products

license and distribution arrangement





2H CY21 and beyond

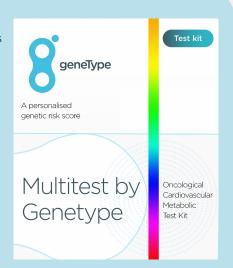
High Density/Multitest

Following the completion of the 1H CY21 tests GTG will progress with the completion of the Multitest products with the intention of releasing a High Density/Multitest following the completion of the included tests.

Market Release to include the following:

- Breast Cancer
- Colorectal Cancer
- Cardiovascular Disease
- Type 2 Diabetes
- Melanoma
- Prostate Cancer
- COVID-19

Covering more than 70% of all mortalities



Responding to unique opportunities and challenges



Genetype for COVID-19 SDR Test - World first serious disease risk (sdr) test that predicts your risk of hospitalisation & life threatening complications

Highlights

- · Rapid response and creation of the test
 - Expected to be market ready Q1CY211
- Allows individual to know their risk of series consequences from contracting COVID-19
- Currently undergoing technical certification²



Product Overview

- · Simple oral swab test
- · Allows for remote screening
- Accurate and low cost³
- Designed to identify who:
 - · may be at risk of serious life-threatening complications
 - should isolate as a precaution
 - should be prioritised for vaccination
- · Combines genetic risk with clinical risk
- Could provide the ability to prioritise medical intervention for high-risk individuals
- Over 100% better at identifying risk than age and gender alone³

Distribution and license Agreement with Infinity BiologiX

- Initial three-year co-exclusive license agreement for sale and distribution of GTG's COVID-19 Risk Test in the US
- Minimum payments US\$2.9 million over three years
- IBX currently has the capacity to process over 100,000
 COVID-19 Risk Tests per day across its two major labs
- GTG to receive US\$10 per test with no additional COGs
- Also can leverage an existing network of SARS-CoV-2 testing partners and associated medical practitioners across the US

1. Increased data set is currently undergoing integration into our risk test. Product launch expected to be end of Q1 CY21

2. GTG anticipates the registration of an LDT COVID-19 severity test should take less than 45 days after completion of validation and technical certification (based on estimates received)

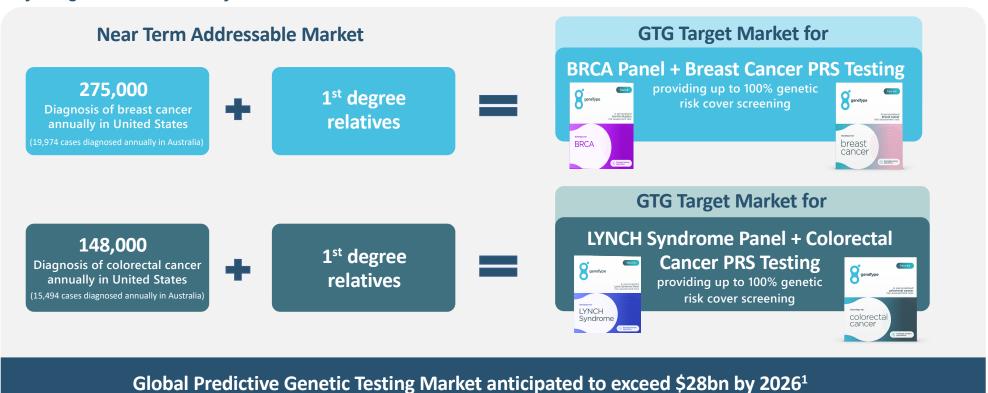
3. "An integrated clinical and genetic model for predicting risk of severe COVID-19" available at: https://www.medrxiv.org/content/10.1101/2020.09.30.20204453v1.full.pdf

infinity biologix

Significant market opportunity



GTG aims to provide up to 100% predictive, pre-symptomatic testing - Germline (5%-10%) and Non Familial (90%) Key insights to inform lifestyle choices and healthcare discussions



^{1.} Genetic Testing Market Size By Test Type (Predictive Testing, Carrier Testing, Prenatal and New-born Testing, Diagnostic Testing, Nutrigenomic Testing), By Application (Cancer, Genetic Disease, Cardiovascular Disease), Industry Analysis Report, Regional Outlook, Application Potential, Competitive Market Share & Forecast, 2020 – 2026; Published Date: Feb 2020; Authors: Sumant Ugalmugle, Rupali Swain

^{2.} PRS = Polygenic Risk Score

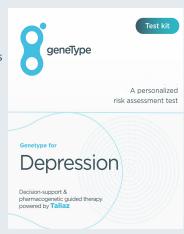
Partnership to provide expanded product offering



Genetype for Depression - Powered by Taliaz's PREDICTIX platform

Highlights

- Key opportunity with strong product alignment to integrate into our platform in line with our mandate to expand our product offering
 - Provides entry into mental health and pharmacogenomics segments
- Regulatory status:
 - CE Marked
 - TGA approved, expected to be market ready by the end of 1H CY21
- Anticipated for market release in 2H CY21
- Distribution Agreement
 - Minimum distribution of 8,000 tests over the initial three-year term
 - Pricing not yet determined but expected to be in line with current test pricing of ~A\$350 per test



Product Overview

- · Utilising a combination of:
 - Genetic, metabolic, clinical and demographic background data; in conjunction with
 - Artificial Intelligence and Machine Learning
- Creates diagnostic and pharmacogenetic solutions that are
 47% better than current best in class practices.
- Better individual outcomes due to superior therapeutic drug selection

1 in 8

Australian prescribed antidepressants annually¹

47% Improvement on accuracy of prescribing antidepressant²

1 in 10
Americans prescribed antidepressants annually³

- 1. Source: Psychwatchaustralia
- 2. Based on a retrospective analysis of STAR*D study medications versus current clinician treatment selection prescribing accuracy (Chekroud et al., 2016). STAR*D is one of the world's largest prospective studies for optimal antidepressant administration.
- 3. Source: https://www.health.harvard.edu/blog/astounding-increase-in-antidepressant-use-by-americans-201110203624

Strategy



GTG have established a clear product pipeline and direction:

- Transitioned from one product two years ago to 10 products in development
- Ability to accelerate the development of new products and tests
- Expanding into reimbursable space

Focused on:

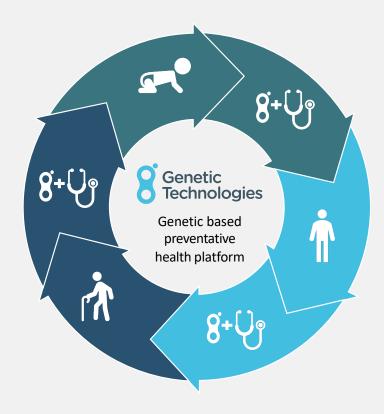
- Establishing leadership through peer reviewed publications, providing reimbursable products and leveraging key opinion leaders
- Forming relationships with large US based labs to provide a clear avenue for increased scale and product distribution through licensing agreements
- Providing individuals with management and lifestyle insights that can be implemented 15-20 years before onset of disease to extend quality of life through:
- Products that are designed to improve medical and lifestyle outcomes mediated through environmental changes, supplements and medications where appropriate

Controlled operational costs

• Only ~10% increase associated with expanded product base to date

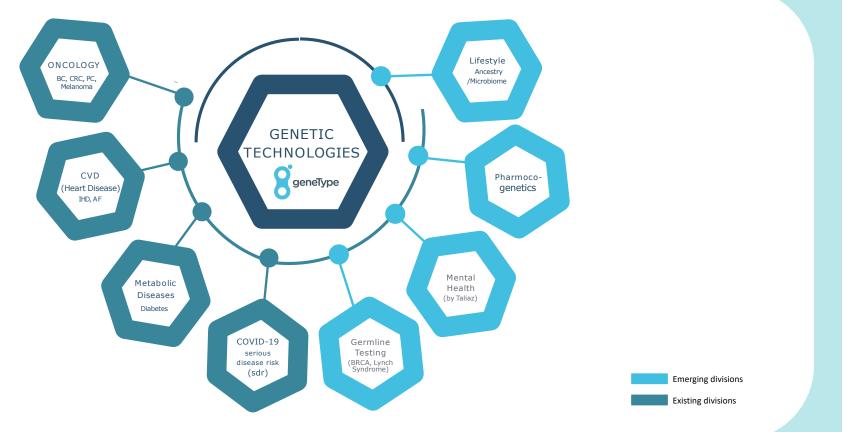
Continually evaluating further opportunities

- Continue to focus on research and development alongside product innovation
- Leveraging in market products to enhance product offering (ie. Taliaz)



Expanding divisions and product offerings





Medical/Pharmacogenetics pathway



Third Party Licensing

- Provides enhanced distribution and product offering via:
 - Licensing of own products for enhanced distribution opportunities (ie. Infinity BiologiX COVID-19 Risk Test)
 - Licensing of novel products for enhanced product offering (ie. PREDICTIX by Taliaz)



Business to business sales via the medical profession

- Adversely impacted by COVID-19 restrictions but remains a key avenue for education and sales
- Combined with an educational program to target mediated healthcare professional industry education content providers

Certifying reimbursible Germline testing platform anticipated to be completed by end of 1H CY2021

- BRCA test: Medicare Benefits Schedule:
 - Item 73296 Fee: \$1,200.00 Benefit: 75% = \$900.00 85% = \$1,115.301
 - Item 73297 Fee: \$400.00 Benefit: 75% = \$300.00 85% = \$340.00²
- LYNCH Syndrome test: Medicare Benefits Schedule
 - Item 73354 Fee: \$1,200.00 Benefit: 75% = \$900.00 85% = \$1,115.30³

^{2.} http://www9.health.gov.au/mbs/fullDisplay.cfm?type=item&q=73297&qt=ItemID

^{3.} http://www9.health.gov.au/mbs/fullDisplay.cfm?type=item&q=73354&qt=item&criteria=lynch%20syndrome

Pathways to market



Medical – Business to Business (B2B)

Third Party Licensing

Business to business sales via the medical profession



Certifying reimbursable Germline testing platform

BRCA test

LYNCH Syndrome test

(More to follow)



Consumer initiated testing (CIT) with medical supervision

Launched US and Australia CIT platforms in 2020 with medical supervision with:

InTeleLabs in the US

Phenix Health in Australia



Current products include:

GeneType for Breast Cancer

GeneType for Colorectal Cancer

priced at AUD\$349 per test



Direct to consumer testing (DTC) with no medical supervision

Will be leveraged for ancestry and gut microbiome testing Scheduled for development following the establishment of the regulated disease and reimbursable segments





Regional Distribution





United States

- Certification required by US regulators **CLIA** to sell into the USA
- One product1 currently certified with further products expected to be submitted in next 12 months
- Colorectal launching March 2021



Australia

- Certification required by Australian regulators NATA, to sell into the Australian market
- Two products² currently certified and further products expected to be submitted in next 12 months



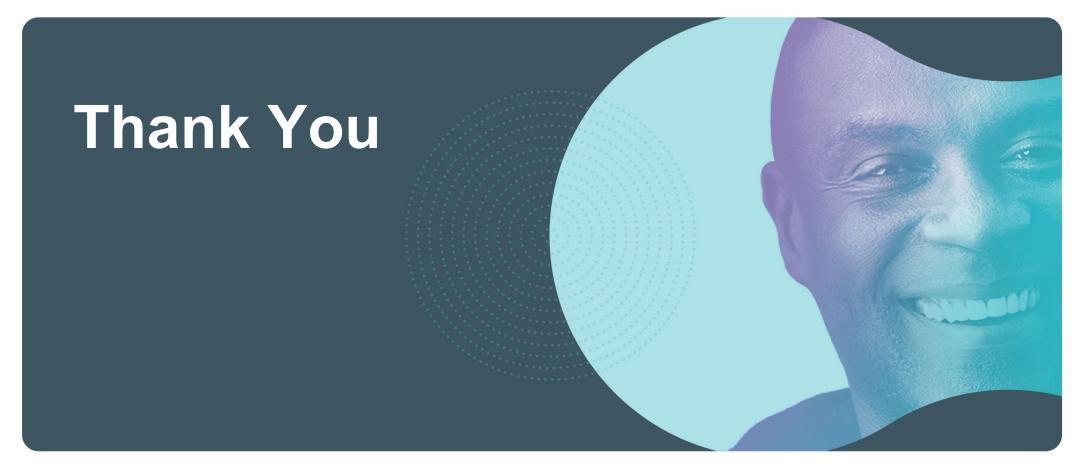
Europe

• Looking to commence CE certification with the view of entering the European market with our novel genetic risk tests in CY2021

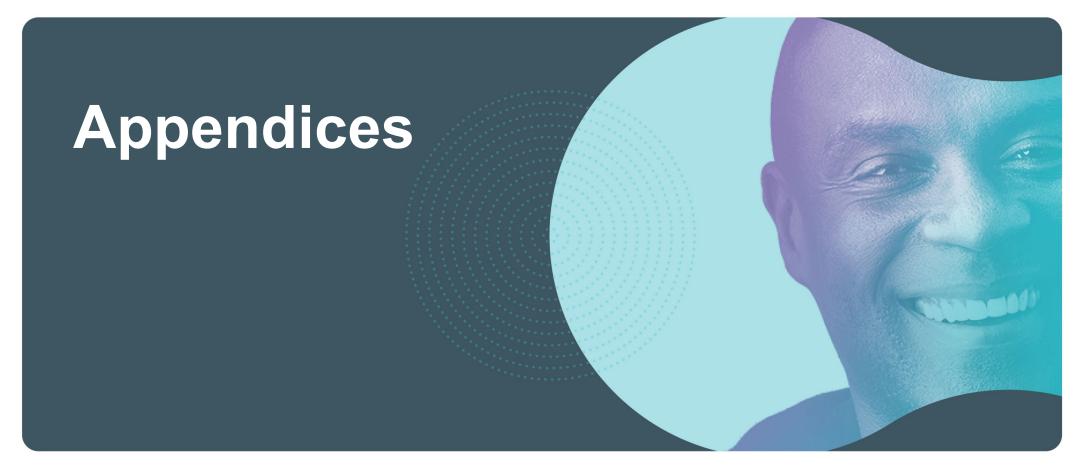
- GeneType for Breast Cancer certified for sale via online sales platform
 GeneType for Breast Cancer and Colorectal Cancer certified for sale via online sales platform

Genetic Technologies Empowering a healthier life!





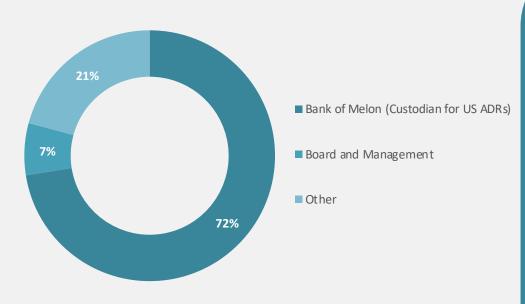




Corporate Overview







Dual Listed on the ASX and Nasdaq

Financial Information

Share price (AUD) as at 5 March 2021	0.9c
ADR price (USD) as at 4 March 2021	\$4.16
Ord Share on Issue (M) ¹	9,012
ASX 52-week trading (low/high)	0.3/1.4c
Nasdaq 52-week trading (low/high)	\$1.41/10.30
Market Cap (A\$M/\$USM)	85.63/62.50
Cash (31 December 2020)	\$16.4m
Debt (31 December 2020)	nil

^{1.} American Depository Receipts (ADRs) are interchangeable via custodian Bank of Mellon with 70% of stock held in the USA 2.600 shares on the ASX equate to 1 ADR (American Depository Receipts) in the USA which are interchangeable via custodian Bank of Mellon

Financial Overview



- Strong proforma cash position of A\$24 million to provide runway for commercialisation and further product development
- Cash burn of \$1.48 million a decrease on prior quarter (Q1 FY21: \$1.88 million) as a result of a reduction in administrative costs despite increased products in development and focus on product distribution and marketing
- Strong interest from US based institutional investors as evidenced by completion of post quarter end US\$6.56 million capital raise on the 25th January used to:
 - Support the introduction and distribution of its new products in the United States and Europe
 - Reimbursement studies for the polygenic risk tests;
 - Implementation of its consumer-initiated testing platforms;
 - Preparation for its COVID-19 PRS Test;
 - Introduction of germline testing division;
 - General product research and development; and
 - For general working capital and potential acquisitions.

\$A '000s	30 Sept 2020	31 Dec 2020	Change
Net operating cashflow	(1,875)	(1,481)	(21%)
Payments for Research and Development	438	358	(18%)
Cash	18,095	16,435	(9%)

Our board and management





Mr. Peter Rubinstein
BSc, BEc, LLB
Chairman - Non – Executive Director



Dr. Lindsay WakefieldMBBS
Non – Executive Director



Mr Nick Burrows

B.Com, FAICD, FCA, FGIA, FTIA, F Fin
Non – Executive Director



Dr. Jerzy "George" MuchnickiMBBS
Executive Director & Chief Medical Officer



Simon MorrissGAICD
Chief Executive Officer



Richard AllmanBSc, PhD, Microbiology
Chief Scientific Officer

Defined Terms



Common Complex Diseases (CCP) – A complex disease is caused by the interaction of multiple genes and environmental factors. Complex diseases are also called multifactorial. Examples of common complex diseases include cancer and heart disease.

Polygenic Risk Score (PRS) - A polygenic risk score tells you how a person's risk compares to others with a different genetic constitution. However, polygenic scores do not provide a baseline or timeframe for the progression of a disease. For example, consider two people with high polygenic risk scores for having coronary heart disease.

Serious Disease Risk (SDR) - Risk associated with acquiring COVID-19 and requiring hospitalisation withs its associated morbidities and mortalities.

Germline Testing – Germline testing is done on cells that do not have cancer. It is done to see if a person has a gene mutation that is known to increase the risk of developing cancers and other health problems. This test uses cells (such as blood or skin cells) that do not have any cancer cells. Germline mutations can sometimes be passed down from parents.

Clinical Laboratory Improvement Amendments (CLIA) - Regulates laboratory testing and require clinical laboratories to be certified by the Center for Medicare and Medicaid Services (CMS) before they can accept human samples for diagnostic testing

National Association of Testing Authorities (NATA) - the authority responsible for the accreditation of laboratories, inspection bodies, calibration services, producers of certified reference materials and proficiency testing scheme providers throughout Australia. It is also Australia's compliance monitoring authority for the OECD Principles of GLP. NATA provides independent assurance of technical competence through a proven network of best practice industry experts for customers who require confidence in the delivery of their products and services.

Next Generation Sequencing (NGS) – Next-generation sequencing (NGS), also known as high-throughput sequencing, is the catch-all term used to describe a number of different modern sequencing technologies. These technologies allow for sequencing of DNA and RNA much more quickly and cheaply than the previously used Sanger sequencing, and as such revolutionised the study of genomics and molecular biology.

Laboratory Developed Tests (LDT) – A type of in vitro diagnostic test that is designed, manufactured and used within a single laboratory.

Consumer Initiated Tests (CIT) - laboratory testing that is initiated by the consumer without a physician order but reviewed and communicated back to the consumer via a physician.

Direct to Consumer (DTC) — laboratory testing that is initiated by the consumer without a physician order. The results are reported back directly to the consumer.

Genome Wide Association Studies (GWAS) - an approach used in genetics research to associate specific genetic variations with particular diseases. The method involves scanning the genomes from many different people and looking for genetic markers that can be used to predict the presence of a disease. Once such genetic markers are identified, they can be used to understand how genes contribute to the disease and develop better prevention and treatment strategies.