



ELECTROGENICS LABORATORIES LTD

ACN 625 525 745

Next-Generation Radiation Dosimetry

Investment Overview

Series C Round In conjunction with

Raising AUD \$1.2M

(\$500k already secured - Overs will be Considered)

May 2025



Opportunity Snapshot

Key Definitions (Medical Physics)

Dosimetry: The science of measuring radiation, calculated through magnitude and specific calculations.

Radiation Oncology: A medical specialty using high levels of radiation to treat cancer, delivered through a Linear Accelerator (LINAC).

Interventional Radiology: Minimally invasive procedures guided by imaging, where precise radiation measurement is critical.

Key Market Insights

Global Need: Over 20 million new cancer cases annually, with 6 million patients treated using radiation therapy.

Current Challenges: Radiation dosimetry is costly, time-intensive, and lacks real-time accuracy, making it rare in mainstream use.

Significant Market Opening: MOSkin™ is as the clear option for replacement in market due to the recall of a major competitor

MOSkin™ Solution

Innovation: ELL's MOSkin™ dosimeter is fast, low-cost, and provides real-time, accurate radiation measurement—making dosimetry viable for mainstream use.

Market Position: MOSkin™ is fully patented (80% global market coverage) with successful trials in 20+ international institutions.

Unique Advantage: Competitors face market exit due to FDA recalls, positioning MOSkin™ as the clear market leader.

Recent Achievements

Grant Funding: Awarded a \$1.1m Industry Growth Program grant after detailed review by AusIndustry. This is 2nd federal Industry grant awarded to the Company.

Regulatory Pathway: *FDA 510k submission completed in March 2025, with market entry expected around July ~ Aug 2025.*

Engineering & Production Ready: Supply Chains enabled & QA certified

USA Distributors: *Identified and negotiations commenced.*

Current Medical Radiation Measurement Tools Are Outdated and Inefficient

Globally, there are 8,000 + Radiotherapy centres and over 15,000 Linear Accelerators, with 5,800 in the USA & China alone.



Competitor technologies have not changed in over 30 years



Expensive with large capex, servicing costs, and device disinfection



Risks patient safety with inaccurate dosage and delayed results



Completely out of touch with modern clinical pathways

Existing Dosimeters Have Major Flaws



Time consuming and **difficult** to calibrate and use,
1 ~ 3 hrs.



Require post processing which delays results for
hours or days and **cannot provide immediate**
feedback of radiation dose.



Don't measure the dose at skin depth, per an
internation standard.



Are **not single use**, therefore require disinfection
and cleaning.
Are **not Radiolucent**, blocking clinician vision.



Impedes LINAC patient throughput and lower
productivity.



Cumbersome for routine use, needing **large capital outlays** and **expensive radiation physicist skills**.

MOSkin™

Introducing MOSkin™

Simple, Immediate, Affordable, & Accurate Radiation Dosimetry

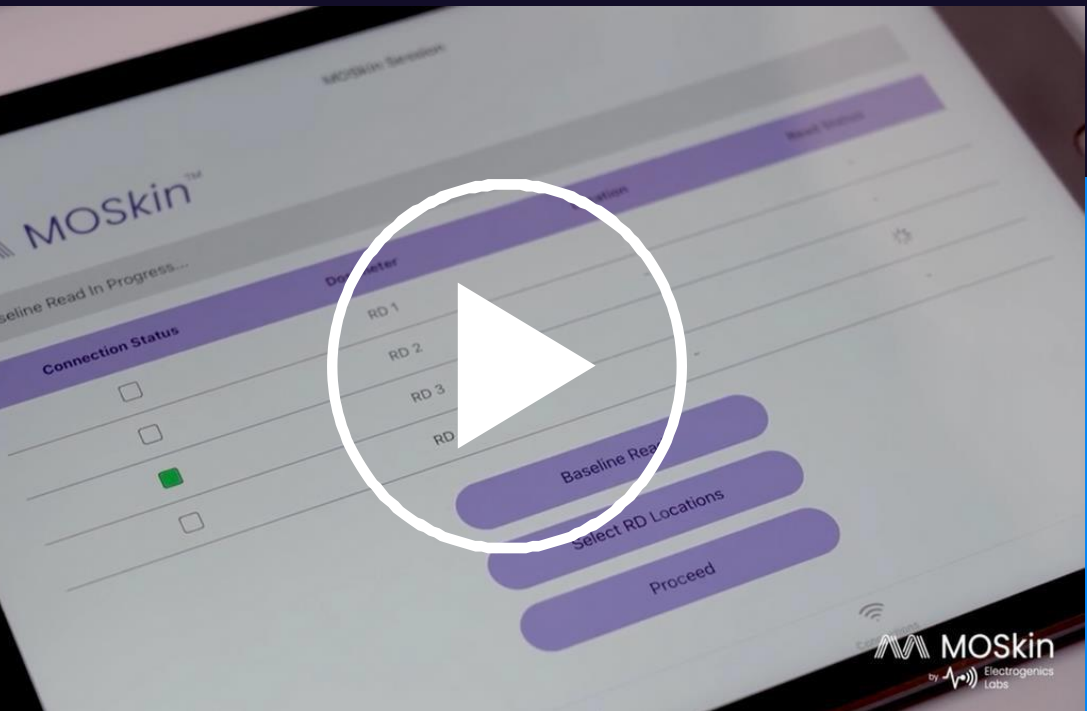
The MOSkin system materially improves clinical pathways, making use of the medical physics laboratory, redundant.



Explanatory Videos



Introducing the MOSkin™ system



Watch our video above to see the difference between MOSkin™ and competitive systems

Materially Improves Clinical Workflow

The MOSkin™ system improves facility outcomes by delivering accurate, real-time dosimetry at point-of-care, rather than after slow laboratory-based post processing.

Affordable



Easy to Use



Fast

MOSkin™



Clinical Benefits

- Can be **set up in minutes** by nurses or lab technicians (no longer needing specialist staff)
- **Real-time** radiation measurement to support clinicians.
- **Cost-effective**, disposable, and easy to use.
- Provides **instant results** directly at the treatment site.

Important Patient Benefits



Reduces risk of overdosing – skin burns + increased risk of secondary cancers



Reduced risk of underdosing – Potential for more visits and recurrence



Much quicker set up at the LINAC, reducing patient time in treatment

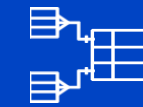
How MOSkin™ Works



Easy to use sensors



Immediate and accurate results



Integrates easily into existing systems

1. Initial Setup



2. Dose measured During Treatment



3. Immediate Data Transmission



4. Real-Time Results at the Bedside



Simplified Clinical Pathway



No disinfection factored in



Use of lower skilled medical staff

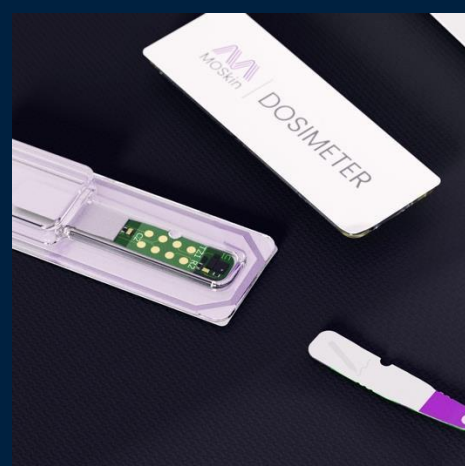


Faster recording and analysis of results

1. Register
Dosimeter At
Hub



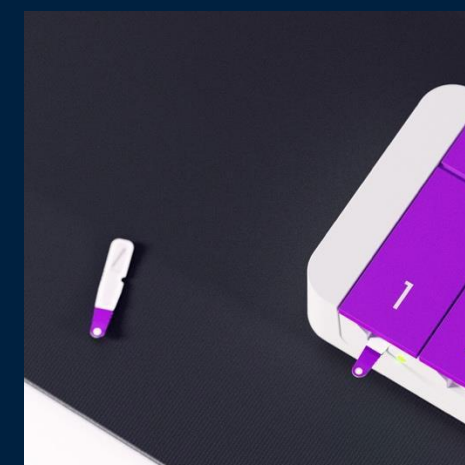
2. Stick on patient



3. Dose measured at
LINAC



4. Dosimeter placed
in hub and result
wirelessly
transmitted to
tablet



5. Immediate results
on ipad and
emailed for
medical records.



Why MOSkin™ Stands Out

- ◆ Superior to competitors in all key facets
- ◆ Largest competitor has exited the market due to an FDA recall, clearing the path for MOSkin™
- ◆ Fully patented and tested in over 20 global institutions

Feature	Old Technology	MOSkin™
Setup Time	1-2 hours per use	3 minutes
Results Delivery	Delayed (1.5+ hours after treatment)	Real-time, instant data during treatment
Accuracy	Moderate (70-85%)	High (>95% accuracy)
Usability	Requires specialist staff for calibration	Operable by nurses/technicians, minimal setup
Patient Safety	Limited monitoring during treatment	Real-time monitoring reduces patient & facility RISK
Cost per Use	High (due to reusable devices needing cleaning), etc	Low (single-use, disposable sensors)
Data Transfer	Wired, often delays in analysis	Wireless, instant data transfer

A Large & Underserviced Global Market



20 million new cancer cases annually



\$300M-\$400M annual market potential in Radiation Oncology



\$80M-\$100M annual market potential in Interventional Radiology

Proprietary Technology

- ◆ Patents are fully assigned to and owned by Electrogenics Laboratories Ltd.
- ◆ ELL has first rights to all new IP developed by UoW in this field.
- ◆ Further ELL IP in development
- ◆ Considerable key Trade Secrets around design & manufacturing not disclosed in the patent

Image shows patents granted in 3 regions, with international patent pending.



The Details

- Expected FDA approval circa July ~ Aug 2025, following a 510(k) Filing in March
- Access to ~8,000 radiotherapy centres globally, with LINAC installations growing at a CAGR of 6%.
- Lower cost per use vs. legacy products due to single-use design and ease of setup.

A Lucrative & Scalable Business Model ...*Underpinned by Platform Technology, for Application in New Markets*



Consumables: Single-use MOSkin™ dosimeters (recurring revenue)



Capital Equipment: MOSkin™ Hubs (one-time purchases) + Utilising iPad



Software: Annual licenses (recurring revenue stream)



Fast Path to Market Entry

R&D, Product Development & supply chain enablement complete



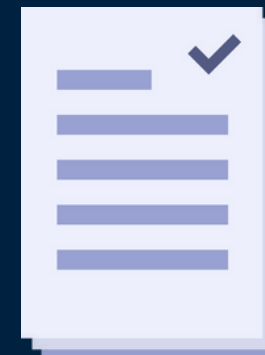
March 2025

510(k) submission complete with FDA as a Class II device.



April-July 2025

Engage and Establish USA distributors and fill supply chains with Product ready for Sales



July ~ Aug 2025

FDA 510(k) clearance anticipated by July ~ Aug, enabling immediate US market entry.



Q3 2025

Initial sales and distribution launch following FDA approval.



Q4 2025

Pursue EU Regulatory Approvals and distribution development activities .

Key Points:

- **FDA clearance expected circa July 2025**
- **Distributor-based model for rapid growth**
- **Positive Distributor negotiations started**
- **1st sales in Qtr. 3, CY 2025.**

1.

United States

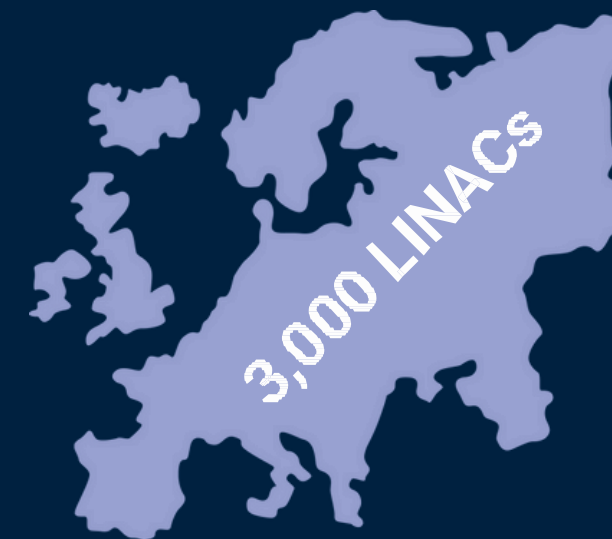
Primary initial target market due to high demand and straightforward regulatory pathway to FDA approval.



2.

Europe

Strategic European countries with high LINAC installations, including UK, Germany, France, and Italy.



Distribution Network

3.

Asia

Initial targets in Japan and China due to growing healthcare infrastructure and need for advanced dosimetry.



4.

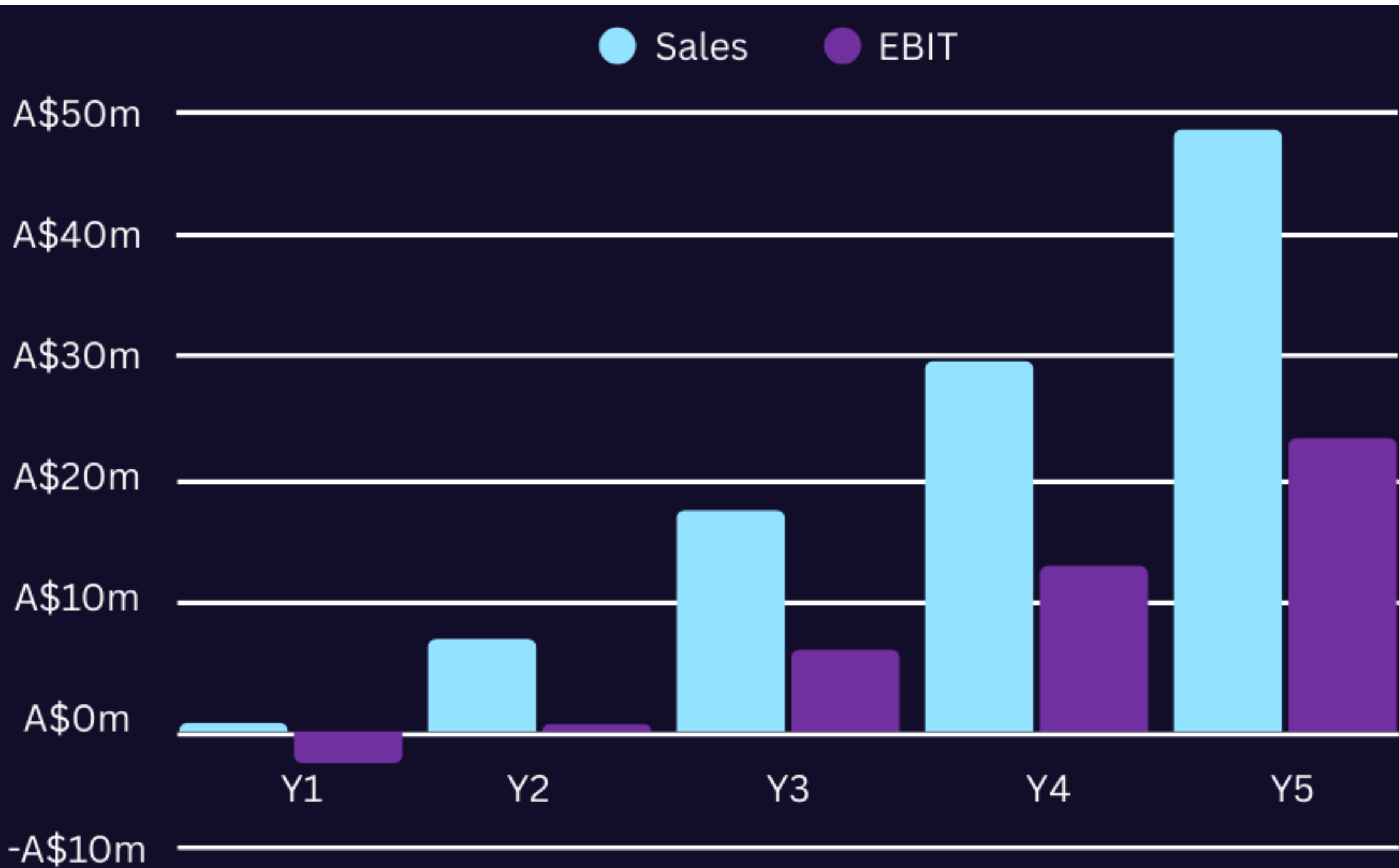
Australia & NZ

Existing clinical partnerships and trial locations make this a target market for expansion.



Out GTM strategy is indirect via in-country channel partners

Linear Growth & Profitability



Break even expected in Year 2

Revenue growth driven by market penetration, replacing old technologies, and competitor withdrawal

IPO or Exit Planned Q4 2026

45% + EBIT by year 3

The Projected Numbers

P&L Projection - Existing Market Only*						
Year (Base Year 2025)	1 - (6mo)	2	3	4	5	Total
Sales						
Sensors Sold (MOSkin™ Units) (rounded)	9,000	92,000	240,000	450,000	750,000	
Software Licenses Active (Units) rounded	70	810	2,250	4,140	7,000	
Hub Sales (New Customers)	70	740	1,450	1,890	2,850	
LINAC Population (6% CAGR) Rounded	15,000	15,900	17,000	18,100	19,200	
Market Share of LINAC machines	0.47%	5.09%	13.25%	22.91%	36.42%	
Total Sales	699,000	7,458,000	17,889,000	29,931,000	48,694,000	AUD\$104,671,000
Royalty/License fees #	48,000	543,000	1,244,000	2,066,000	3,298,000	
COGS	353,000	2,212,000	4,557,000	6,998,000	11,341,000	
Gross Profit	298,000	4,703,000	12,089,000	20,867,000	34,055,000	AUD\$72,012,000
Gross Margin %	43%	63%	68%	70%	70%	
Other income R&D rebates etc.	319,000	224,000	215,000	269,000	438,000	
Expenses						
Sales & Marketing Expense	216,000	766,000	1,189,000	1,713,000	2,539,000	
R&D Engineering & Technical Expense	950,000	746,000	716,000	898,000	1,461,000	
Tech Support & SW Maintenance	69,000	654,000	1,456,000	2,411,000	3,615,000	
Corp Overheads	1,960,000	2,153,000	2,327,000	2,730,000	3,154,000	
Total Expenses	3,195,000	4,319,000	5,688,000	7,752,000	10,769,000	AUD\$31,723,000
Expenses as a % of Sales	457%	58%	32%	26%	22%	
EBIT	-2,577,000	608,000	6,616,000	13,383,000	23,725,000	AUD\$41,755,000
EBIT % of Sales	-369%	8%	37%	45%	49%	NPV 14.5 ***
EBIT NPV ^{14.5}	-2,363,970	480,065	4,496,363	7,828,681	11,945,671	AUD\$22,386,811
			Potential Valuation ##		3x EBIT NPV ^{14.5}	AUD\$ 67,160,000
*Projections are for the existing market only and do not take into account that the enhanced MOSkin™ functionality, low cost and convenience that will most likely result in a considerably expanded market as oncologists and surgeons specify the use of dosimeters in more and more cases.						
# Royalty & License Fees are subject to expiry of the US patents in 2032						

Capital Strategy & Offer Overview

Offer Details	Information
Raise Amount (Series C Round)	A\$1.2M
Price per Share	A\$0.065
Issue Instrument	Ordinary Shares
Pre-Money Valuation	A\$12M*
Target Liquidity Event Horizon	Q4 2026

This Round



Series C Round Initiation

USE OF FUNDS: Initial setup for US market entry pending FDA 510(k) clearance.
A \$1.2M @ A\$0.065 per share



FDA Submission

510k Class IIa.
FDA submission lodged in March of 2025



FDA Clearance & Market ENTRY

Expect FDA 510(k) clearance in July~Aug of 2025.
Prepare for market entry and establish early sales channels.



Initial Sales & Distribution

Launch initial sales Q3 CY 2025, pre-IPO if required.
Supports commercial readiness.



Possible Liquidity Event

Planned for Q4 2026 Est.
A\$0.25-A\$0.30 +per share
based on market conditions
IPO or Trade sale

EV ~A\$12M*
A\$ 6.5c

1

March 25

2

March 25

3

July 2025

4

Q3 2025

5

Q4 2026

**Next Raise A\$2M ~A\$3m @
A\$0.18-A\$0.20**

*Estimated range of EV at Round ^a Contingent on Capital raise and market conditions

Pro Forma Capital Structure

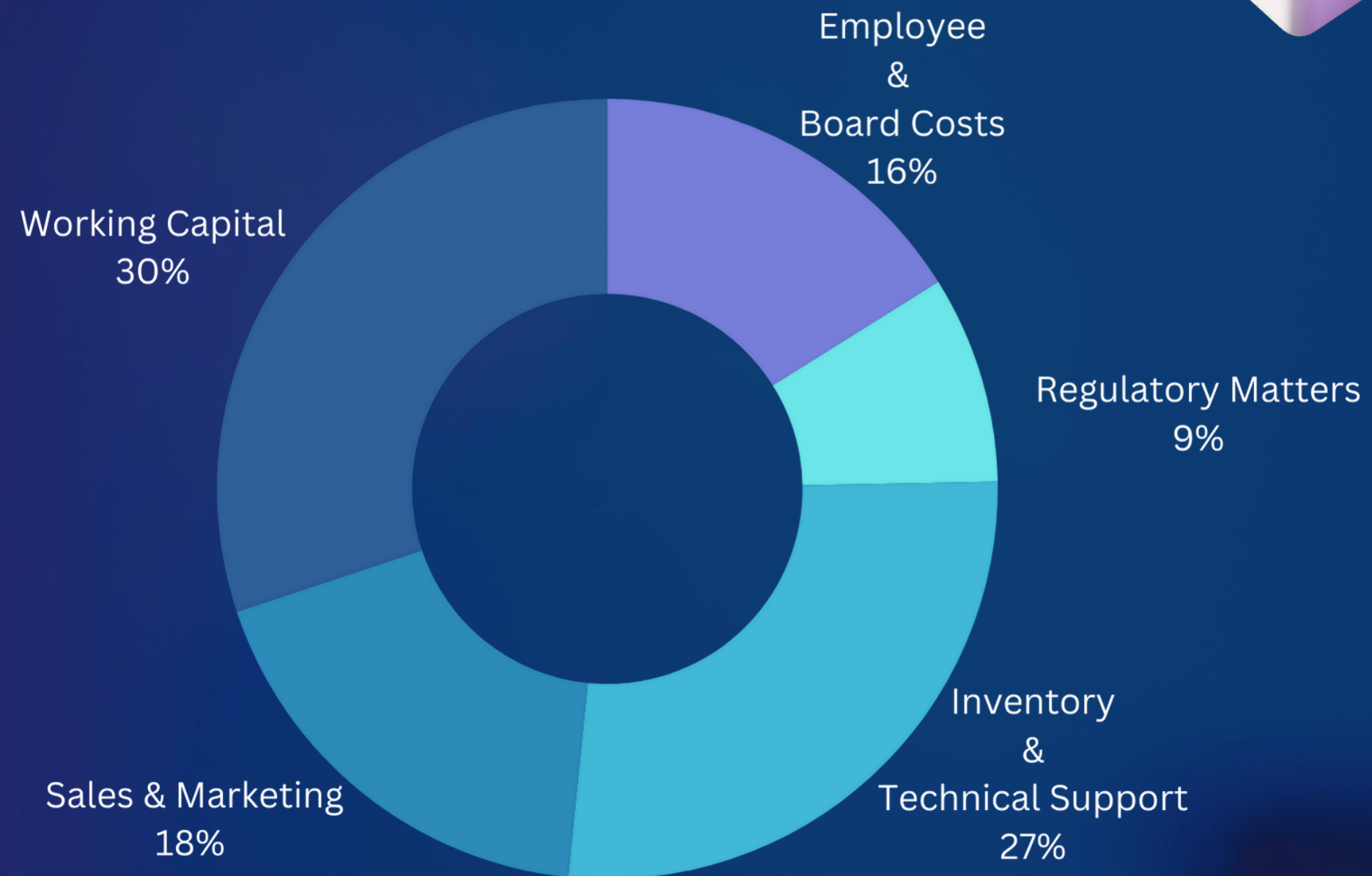
Pro Forma Capital Table – Post Series C Raise		
Shareholders	Shares	Holding
University of Wollongong	11,475,000	5.66%
Founders (3 people)	43,861,254	21.63%
ELL Trust shares & Board & Management	30,631,811	15.1%
Service Providers and Advisors	2,049,996	1.01%
Other shareholders from A & B rounds (93 shareholders)	96,306,133	47.49%
Current Round Series C @ 6.5 cents per share	18,461,538	9.10%
TOTAL Shares	202,785,732	100.00%

Use of Funds

How we will spend \$1.2M

Primary Allocation:

- Regulatory Matters
- Inventory build-up
- Employee & Board Costs
- Sales & marketing
- Working capital



Experienced Leadership with Proven Expertise



**David Rundell,
CEO**

Extensive experience within the Medical Device and Radiopharmaceutical industries with an exceptional track record.

Previously 9 years as the CEO of Laudauer's Nanodot Dosimetry business in Australia and NZ, substantially growing Laudauer's Dosimetry business. He is an expert in the commercial sphere of medical physics.



**Geoff Neilson,
Non-Executive
Director**

Over 30 years in Medical Devices including and senior VP level roles at ResMed across multiple business functions including Product Development, Commercial, and Supply Chain Management roles



**Mario Pennisi AM,
Non-Executive
Director**

Over 30 years in life sciences, experienced in commercialisation and strategic leadership. Director of several successful healthcare businesses in Australia and the USA.

Foundation career in Health tech including Mayne Health.



**Dr. Arthur
Brandwood,
Non-Executive
Director**

40 years in MedTech, expert in regulatory and product commercialisation.

Senior roles in the TGA and adviser to international regulators over many years.



**Geoff Marshall,
Chair**

Deep experience in medical and non-medical sectors. Former big 4 Accounting Partner, investment banker, and founding MD of Nanosonics Ltd.

Previous roles include COO of Mayne Health Private Hospital Group and founder of several start ups.

Specialist Advisors, Key Team & Partners



**Prof. Michael Jackson,
Medical Advisor, MD**

25 years in MedTech and business development, ex-VP at ResMed.



**Prof. Anatoly Rozenfeld,
Scientific Advisor.**

World renowned leader in radiation therapy instrumentation, founder of the Centre for Medical Radiation Physics.



Matthew Harrington, Product Development & Supply

20+ years in MedTech and CleanTech, former ResMed product lead.



Brad Tvedt, Sales & Marketing

Senior sales leader, previously at ResMed and Philips Healthcare, scaled sales from start-up to multi-million revenue.



**Grant Palmer,
QA/RA/Clinical**

30 years in regulatory affairs, clinical research, and quality assurance; experience with global market approvals.

GTM Partners



Recap



Strong Market Need

- Immediate demand for accurate radiation dosimetry following regulatory induced competitor exit.
- Over 20 million new cancer cases annually; 6 million treated with radiation therapy.



Regulatory & Market Readiness

- Minimal regulatory risk with FDA approval expected within 4 ~ 5 months from submission in March 25 (Q1 CY 2025).
- Successful trials in over 20 international institutions testing product reliability.



Compelling Business Model

- Recurring revenue from single-use dosimeters and software licenses.
- Proven 'razor/razorblade' model, maximising customer lifetime value (est. \$60k-\$100k per LINAC over 10 years)



Experienced Leadership

- Highly skilled team with decades of MedTech expertise and strong R&D backing from leading institutions (UoW & CMRP)



Attractive Investment Potential:

- Conservative valuation with potential for 2-3x uplift post-FDA approval.
- Near-term liquidity event planned via a trade sale or IPO, anticipated Q4 2026



Solid IP Portfolio

- Fully owned and patented in all significant global markets.





● JOIN US IN TRANSFORMING RADIATION DOSIMETRY

[Click here or scan QR Code
for More Information Or
Apply for shares](#)



Contact us for more information about this opportunity

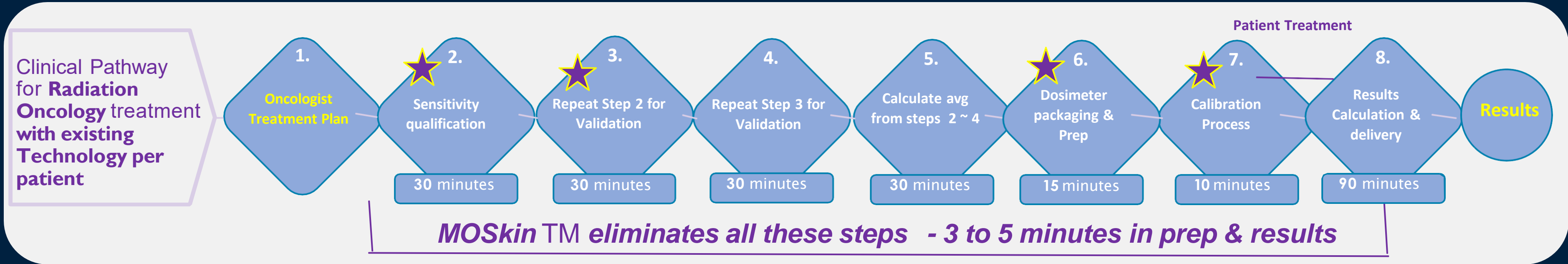
Geoff Marshall
Chair, Electrogenics
+61 414 272 833
geoff.m@electrogenicslabs.com


Geoff Neilson
Director, Electrogenics
+61 424 750 844
geoff.neilson@electrogenicslabs.com

Gavan Farley
Senior Corporate Advisor, Novus Capital
+61 420 520 300
gavan.farley@novuscapital.com.au

APPENDIX

Detailed Clinical Pathway & Competitor Analysis



	A\$ Capital Outlay	A\$ Service cost~10yrs	A\$ Cost for Sensors^	Prep time before Dose	Time to results	Max Patients /day/ #	Typical Accuracy	Application Coverage	Realtime	WED Std
Existing Technology TLD Sensors	Ave \$60K	Ave \$350K	\$30-\$60	~2.5 hrs	>1.5hrs	5~10	70~80%	~80%	NO	NO
Existing Technology DIODES 	Ave \$20K	Ave \$100K	\$1.5-\$2.5K	~1 hr	Real Time	12~15	75~85%	~70%	YES	NO
MOSkinTM	Ave \$12K	\$50K**	\$35	~3 mins*	Real time / Instant	20~25	>95%+	99%+	YES	YES

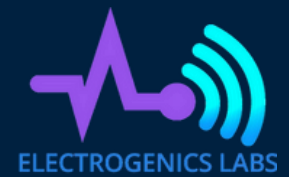
^ Per Sensor. Sensors can be used multiple fractions for same patient

* Assumes pre-calibrated

** Annual Software license over 10 yrs.

= Linear accelerator, uses electricity to generate high energy beams of electrons

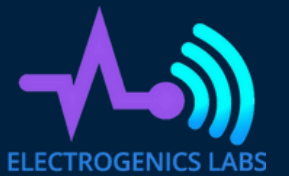
Fully Patented and Tested at More than 20 Specialist Clinics & Hospitals Around the World



- MOSkin™ has been **tested and validated** in Australia & internationally in >20 hospitals & clinics
- Tested **on over 2,000** patients and simulations
- Over **40 published** scientific papers
- A subject of **21 PhDs**
- **> 500 references** in Google scholar
- Over **\$10 million** in equity, cash and grants invested to date
- St George Public Radiation Oncology unit has been using the Company's prototypes under ethics committee approval for **over 12 months**, recently expressing interest to extend the use of prototypes in preference to existing in market technology

INTERNATIONAL	AUSTRALIA
Argon National Lab, USA	St George Cancer Care Centre, Prostate Cancer Institute, St George Hospital, Sydney
Mass General, Boston, USA	St Vincent's Hospital, Sydney
Wisconsin University Hospital, USA	Liverpool Hospital
Brookhaven National Lab, USA	Calvary Mater Hospital – Newcastle, Australia
University of Malaya Medical Centre, Kuala Lumpur, Malaysia	Royal Adelaide Hospital
Italian National Institute of Tumours, Milan, Italy	Illawarra Cancer Care Centre
CERN Switzerland	Perth Hospital
KEK Japan	Prince of Wales Hospital
Sun Yat-Sen University Cancer Centre, Guangzhou China	Mater Hospital, Newcastle
Radiation Dosimetry Lab, Federal University of Pernambuco, Brazil	Peter MacCallum Cancer Centre
City University Hong Kong Hospital	Geneva Hospital

Important Notice



This presentation has been prepared in connection with a proposed investment in Electrogenics Laboratories Ltd (ACN 635 525 745) (**Company**). It contains general background information only which is current at the date of this presentation unless otherwise specified. It contains selected summary information and does not purport to be all-inclusive, comprehensive or to contain all of the information that may be relevant, or which a prospective investor may require in evaluations for a possible investment in the Company. Investors should not rely solely on the information contained in this presentation and must satisfy themselves as to the accuracy of all such information. This presentation has been prepared based on information available at the time of preparation and is subject to change without notice. In receiving this presentation, each recipient agrees to the following terms and conditions, including any modifications to them. No person is under any obligation to update this presentation after its release.

This presentation is provided for general information purposes only. This presentation does not constitute an offer, invitation, solicitation or recommendation with respect to the purchase or sale of securities in the Company in any jurisdiction. It is not a prospectus, product disclosure statement, pathfinder document or any other type of public offer disclosure document for the purposes of the *Corporations Act 2001* (Cth) (**Corporations Act**) and has not been, and is not required to be, lodged with the Australian Securities and Investments Commission. It should not be relied upon solely by the recipient in considering the merits of the Company or the acquisition of shares in the Company.

This presentation does not constitute investment or financial advice. It is not intended to be used as the basis for solely making an investment decision, nor is it intended to constitute legal, tax, accounting or other advice. This presentation has been prepared without taking account of any person's individual investment objectives, financial situation or particular needs. Before making an investment decision, the recipient should consider its own financial situation, objectives and needs, and conduct its own independent investigation and assessment of the contents of this presentation, including obtaining investment, legal, tax, accounting and such other advice as it considers necessary or appropriate. Any references to or explanations of legislation, regulatory issues, benefits or any other legal commentary (if any) are indicative only, do not summarise all relevant issues and are not intended to be a full explanation of a particular matter. The information in this presentation has been obtained from and based on sources believed by the Company to be reliable. Past performance is not an indication of future performance.

This presentation may contain forward-looking statements, guidance, forecasts, estimates, prospects, projections or statements in relation to future matters that may involve risks or uncertainties and may involve significant items of subjective judgement and assumptions of future events that may or may not eventuate (**Forward Statements**). Forward Statements can generally be identified by the use of forward-looking words such as "anticipate", "estimates", "will", "should", "could", "may", "expects", "plans", "forecast", "target" or similar expressions. Forward Statements including indications, guidance or outlook on future revenues, distributions or financial position and performance or return or growth in underlying investments are provided as a general guide only and should not be relied upon as an indication or guarantee of future performance. To the extent that certain statements contained in this presentation may constitute 'forward-looking statement' or statements about forward-looking matters, then the information reflects the Company's (and no other party's) intent, belief or expectations as at the date of this presentation. No independent third party has reviewed the reasonableness of any such statements or assumptions. None of the Company, its related bodies corporate and their respective officers, directors, employees, advisers, partners, affiliates and agents (together, the Company Parties) represent or warrant that such Forward Statements will be achieved or will prove to be correct or gives any warranty, express or implied, as to the accuracy, completeness, likelihood of achievement or reasonableness of any Forward Statement contained in this presentation. Except as required by law or regulation, the Company assumes no obligation to release updates or revisions to Forward Statements to reflect any changes. Recipients should form their own views as to these matters and any assumptions on which any of the Forward Statements are based and not place reliance on such statements. All dollar values are in Australian dollars (\$) or A\$) unless otherwise stated.

An investment in the Company's securities is subject to known and unknown risks, many of which are beyond the control of the Company, including factors and risks specific to the industry in which the Company operates as well as general economic conditions, prevailing exchange rates and interest rates and conditions in financial markets. The Company does not guarantee any particular rate of return or the performance of the Company, nor does it guarantee any particular tax treatment. Prospective investors should make their own enquiries and investigations regarding all information in this presentation, including the assumptions, uncertainties and contingencies which may affect future operations of the Company and the impact that different future outcomes may have on the Company. By accepting this presentation, the recipient agrees to keep permanently confidential all information that it contains. It should not be made available to any other third party without the prior written approval of the Company.

The distribution of this presentation to persons or in jurisdictions outside Australia may be restricted by law and any person into whose possession this document comes should seek advice on and observe restrictions. Failure to comply with such restrictions may violate securities law. No party other than the Company has authorised, permitted or caused the issue, submission, dispatch or provision of this presentation, or takes any responsibility for, or makes or purports to make any statement, representation or undertaking in this presentation and there is no statement in this presentation that is based on any statement by any other party. No person, either as a director, partner or in the employment of the Company has any authority to make, imply, or give any representation or warranty whatsoever in relation to the information contained in this presentation. None of the Company Parties take any responsibility for any information in this presentation or any action taken by you on the basis of such information.

To the maximum extent permitted by law, the Company Parties: exclude and disclaim all liability, including (without limitation) any liability for fraud or negligence, for any expenses, losses, damages or costs incurred either as a result of the information in this presentation being inaccurate or incomplete in any way for any reason, or otherwise arising in connection with this presentation; and make no representation or warranty, express or implied, as to the currency, accuracy, reliability or completeness of information in this presentation.