

ASX Announcement

Racura Oncology to Present at TechKnow Invest Conference

29 July 2026

Racura Oncology Limited (“Racura”) is pleased to advise that CEO and Managing Director, Dr Daniel Tillett will be presenting at the TechKnow Invest Conference in Sydney on 29 July 2026 at 12.30pm. The TechKnow Invest Conference is being held at Hotel Swissotel, 68 Market Street, Sydney, Australia.

A copy of the presentation is attached to this announcement.

-ENDS-

About Racura Oncology

Racura Oncology (ASX: RAC) is a Phase 3 clinical-stage biopharmaceutical company with a mission to silence cancer.

Racura's lead asset, (E,E)-bisantrene, is a clinically derisked small-molecule anticancer agent that primarily acts by stabilising G4-DNA and RNA structures, driving potent silencing of c-MYC, a key cancer gene dysregulated in up to 70% of all cancers. (E,E)-bisantrene has shown therapeutic activity in cancer patients and has a well-characterised safety profile. Racura's discoveries have supported composition-of-matter IP filings that, if granted, could provide up to 20 years of patent protection for (E,E)-bisantrene.

Racura is advancing RC220, its proprietary formulation of (E,E)-bisantrene, to address significant unmet need across multiple MYC-driven oncology indications. The Company's clinical programs include the Phase 3 EMILI trial in acute myeloid leukaemia, the Phase 1a/b HARNESS trial in combination with osimertinib for EGFR-mutated non-small cell lung cancer, and the Phase 1a/b CPACS trial in combination with doxorubicin, where Racura aims to deliver both anthracycline cardioprotection and enhanced anticancer activity.

Racura has collaborated with Astex, Emory University, Purdue University, MD Anderson, Sheba City of Health, UNC School of Medicine, the University of Wollongong and the University of Newcastle. Racura is actively exploring partnerships, licence agreements and potential commercial merger and acquisition opportunities to accelerate global patient access to RC220. Learn more at www.racuraoncology.com.

For questions about this announcement or previous Racura Oncology announcements, please visit our [Interactive Announcements](#) page.

Racura encourages investors to go paperless by registering their email details with the Company's share registry, Automic Registry Services, at www.automicgroup.com.au.

Release authorised by

Daniel Tillett, CEO/MD

info@racuraoncology.com

Media Contact

Cherie Hartley +61 418 737 020

cherie.hartley@irdepartment.com.au



July 2026

Racura Oncology

TechKnow Invest Roadshow

Dr Daniel Tillett, CEO & MD

Important notice and disclaimer

The material in this presentation has been prepared by Racura Oncology Limited (ACN 149 318 749) (Company).

THIS IS NOT A PROSPECTUS

This presentation is not a prospectus, product disclosure statement or disclosure document for the purposes of the Corporations Act 2001 (Cth) (Corporations Act). It has not been lodged with the Australian Securities and Investments Commission, or otherwise.

Statements in this presentation are made only as of the date of this presentation unless otherwise stated and the information in this presentation remains subject to change without notice. No representation or warranty, express or implied, is made as to the fairness, accuracy or completeness of the information, opinions and conclusions contained in this presentation or any other information the Company or any other person otherwise provides to you.

This presentation does not constitute an offer to sell, or a solicitation of an offer to buy, securities in the United States. Any securities described in this presentation have not been, and will not be, registered under the US 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 transactions exempt from, or not subject to, registration under the Securities Act and applicable US state securities laws. This presentation may not be released to US wire services or distributed in the United States. The distribution of this presentation in other jurisdictions outside Australia may also be restricted by law and any such restrictions should be observed.

NOT FINANCIAL PRODUCT ADVICE

No attempt has been made to independently verify the information contained in this presentation. The information in this presentation is of a general nature and does not constitute financial product advice, investment advice or any recommendation. Nothing in this presentation constitutes legal, financial, tax or other advice. The information in this presentation does not take into account your particular investment objectives, financial situation or needs, or those of any other person. You should make your own assessment of an investment in the Company and should not rely on this presentation. In all cases, you should conduct your own investigations and analysis of the financial condition, assets and liabilities, financial position and performance, profits and losses, prospects and business affairs of the Company and its business, and the contents of this presentation. You should seek legal, financial, tax and other advice appropriate to your jurisdiction.

THIS PRESENTATION DOES NOT CONSTITUTE AN OFFER OR ADVERTISEMENT

This presentation does not constitute an invitation, offer or recommendation to apply for or purchase Shares and does not contain any application form for Shares. This presentation does not constitute an advertisement for an offer or proposed offer of Shares.

NO LIABILITY

The Company has prepared this presentation based on information available to it at the time of preparation, from sources believed to be reliable and subject to the qualifications in this presentation. No representation or warranty, express or implied, is made as to the fairness, accuracy, completeness or correctness of the information, opinions and conclusions contained in this presentation. To the maximum extent permitted by law, none of the Company or its subsidiaries or affiliates or the directors, employees, agents, representatives or advisers of any such party, nor any other person accepts any liability for any loss arising from the use of this presentation or its contents or otherwise arising in connection with it, including without limitation, any liability arising from fault or negligence on the part of the Company or its subsidiaries or affiliates or the directors, employees, agents, representatives or advisers of any such party.

FORWARD-LOOKING STATEMENTS

This presentation may contain forward-looking statements that are subject to risk factors associated with an oncology company. Forward looking statements can be identified by the use of forward-looking terminology, including, without limitation, the terms “believes”, “estimates”, “anticipates”, “expects”, “predicts”, “intends”, “plans”, “goals”, “targets”, “aims”, “outlook”, “guidance”, “forecasts”, “may”, “will”, “would”, “could” or “should” or, in each case, their negative or other variations or comparable terminology. These forward-looking statements include all matters that are not historical facts. By their nature, forward-looking statements involve known and unknown risks, uncertainties and other factors because they relate to events and depend on circumstances that may or may not occur in the future and may be beyond the Company’s ability to control or predict which may cause the actual results or performance of the Company to be materially different from the results or performance expressed or implied by such forward-looking statements. Forward looking statements are based on assumptions and are not guarantees or predictions of future performance. No representation is made that any of these statements or projections will come to pass or that any forecast result will be achieved, nor as to their accuracy, completeness or correctness. Similarly, no representation is given that the assumptions upon which forward looking statements may be based are reasonable.

Corporate snapshot

Racura Oncology is an ASX-listed, late-stage clinical biopharmaceutical company

Key metrics (\$A)

ASX code	RAC
Share price	\$2.09 ¹
Market capitalisation	\$409.8m ¹
Cash at bank	\$34.3m ²
Debt	Nil
Enterprise value	\$375.5m ¹
Shares on issue	196,068,940 ¹
Options on issue	10,201,193 ¹

1. As at 27 July 2026
2. As at 30 June 2026

Racura 12-month trading history¹



Current funding

- On 29 May 2026, Racura concluded a fully underwritten option issue and intuitional placement which has provided an additional \$17.8m million since 31 March 2026
- Racura is fully funded to undertake all trials and other programs into 2028 from existing funds

Why invest in Racura Oncology?

Investment opportunity for Racura has fundamentally changed in recent times



Clinically validated drug

Bisantrene used in >50 clinical trials. Established safety and anticancer activity.



Improved drug - RC220

Reformulated for better administration. 20 years of patent protection.



20yr composition of matter patent filed

Covers the active isomer, (E,E)-bisantrene discovered by Racura research team.



G4-DNA & RNA binder

MOA identified to be G4-DNA & RNA binding. Targets the key cancer growth regulator MYC.



Many value drivers

Significant news flow throughout 2026 & 2027. Three open label clinical trials reporting.



Investor & patient focused

Ethos, communication and funding strategies.

MYC: Oncology's largest untapped opportunity

Treating MYC dysregulation is a huge opportunity across numerous cancer types

70%

of all cancers have elevated MYC activity

MYC is a potent accelerator of cancer growth

- ✓ Discovered in 1982
- ✓ Controls genes involved in cell growth

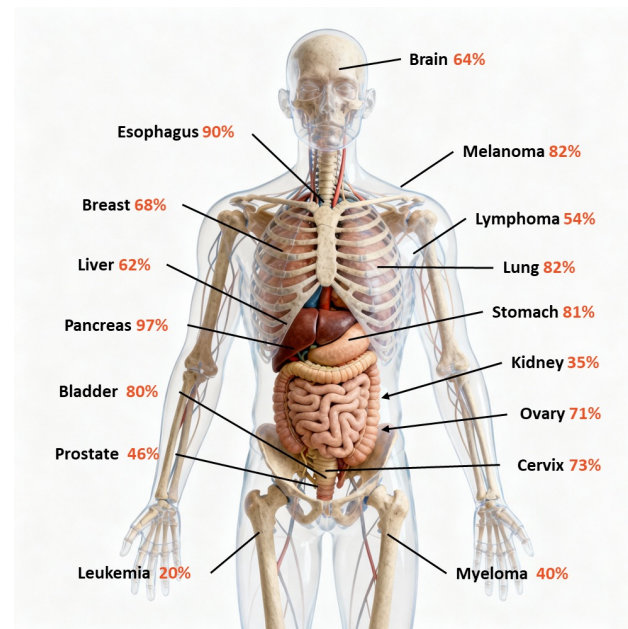
Difficult to target - considered undruggable

- ✓ No conventional binding pocket as protein is unstructured
- ✓ No clinically effective MYC inhibitors even after decades of research

Broad oncology applicability

- ✓ Elevated MYC across multiple tumour types
- ✓ More than 1 million new cancers each year in the USA

Major cancers with elevated MYC activity¹



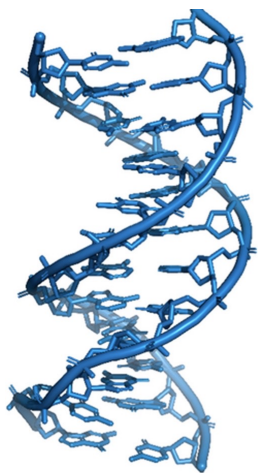
MYC inhibition has broad clinical and commercial value across most cancer

(E,E)-bisantrene

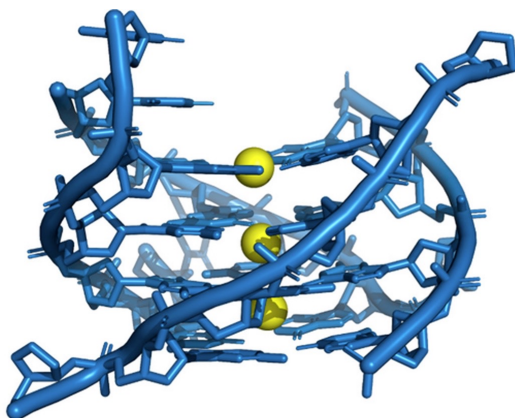
Binds and stabilises G-quadruplex DNA, silencing the c-MYC oncogene¹

G-quadruplexes or G4

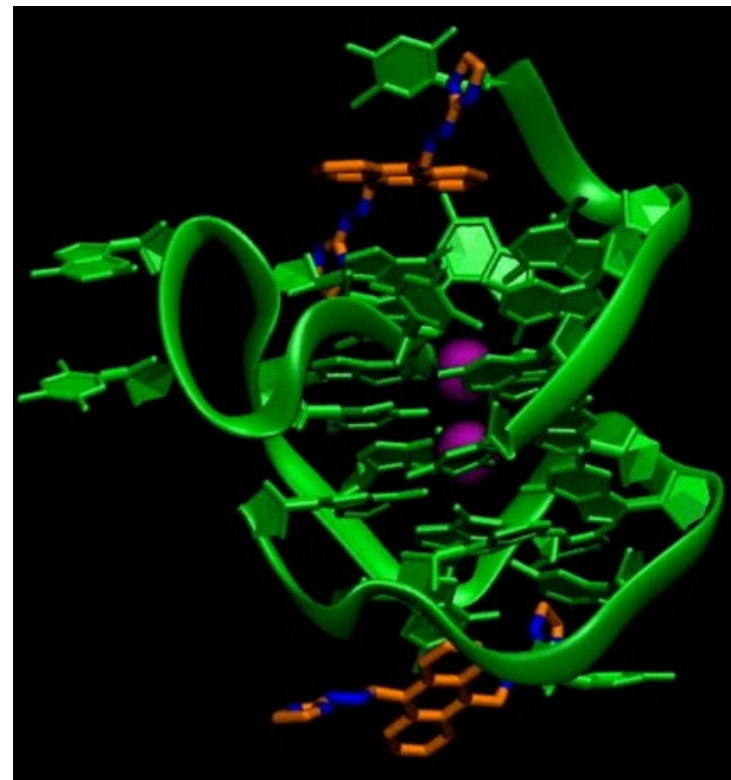
Key regulatory structures in DNA (& RNA) recognised by transcription and translation factors to regulate cell growth and division



Double stranded DNA
(double helix)

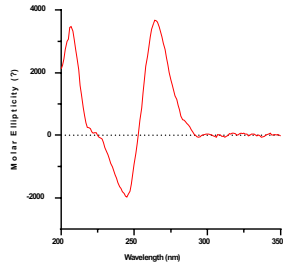


G-quadruplex (G4) DNA



Scientific support for c-MYC gene silencing activity

(E,E)-bisantrene stabilises MYC G-quadruplex (G4) and turns off expression of c-MYC gene



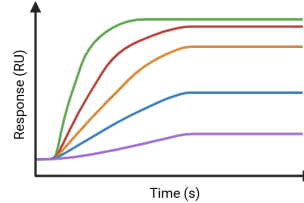
G4 Melting

Bisantrene stabilises c-MYC G4



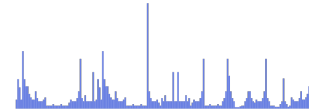
Molecular Dynamics

Predicts binding of bisantrene to c-Myc G4



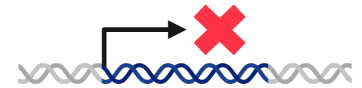
Surface Plasmon Resonance

Measures affinity of bisantrene for c-MYC G4



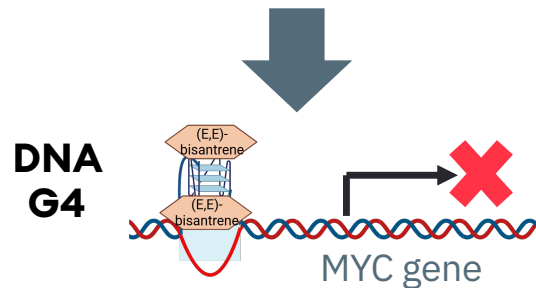
¹H-NMR

Shows how bisantrene binds to c-MYC G4



Gene Expression Analysis

Consistent downregulation of MYC regulated genes



MYC inhibitors in the clinic

RC220 competitors are all at early stage and have limited clinical activity

Drug	Developer	Mechanism of action	Clinical activity
MYC Direct (protein/gene)			
RC220	Racura Oncology	G-quadruplex stabiliser	Phase 3 AML, Phase 1 NSCLC, Phase 1 solid tumours; 40% CR/PR in AML
OMO-103	Peptomyc S.L.	MYC-MAX inhibitor (Omomyc miniprotein)	Phase 2 (osteosarcoma); 8/22 SD, no CR or PR
IDP-121	IDP-Pharma	Stapled peptide direct inhibitor of MYC	Phase 1/2 in MM; No clinical efficacy data released
PC-002	Cotherra Bio	De-ubiquitinase inhibitor	Phase 2 lymphoma; 1/13 CR, no PR
LMP744	Gibson Oncology	G-quadruplex stabiliser	Phase 1/2 GBM; 2/40 PR, no CR
CX-5461	Senhwa Biosciences	G-quadruplex stabiliser	Phase 1b BRCA1/2 or HR deficient solid tumours; 4/40 PR, 11/40 SD
WBC100	Hangzhou Weben	MYC-specific PROTAC degrader	Phase 1 in MYC+ solid tumours; No clinical efficacy release
Other pathways			
WP1066	Moleculin Biotech	p-STAT3 inhibitor (impacts MYC transcription)	Phase 2 GBM with radiotherapy
GP2250	Persevere Tx	Inhibits MYC and NFκB	Phase 1 pancreatic
PMR-116	Pimera Tx/ ANU	Inhibits pathway downstream of MYC	Phase 1 in MYC+ solid tumours



Clinical Programs

Clinical pipeline

Three clinical programs in AML, lung cancer and solid tumours

Cancer indication	Total addressable patients ¹	Phase 1	Phase 2	Phase 3	Comment
AML Acute myeloid leukaemia ²	US/EU: 50,000	 Phase 3			40% response rate in two Phase 2 trials ³ Bisantrene approved in 1988 in France
EGFRm lung cancer EGFR Non-small lung cancer (NSCLC) in combination with osimertinib (TKI) ²	China: 445,000 US/EU: 95,000	 Phase 1			Osimertinib 2025 sales >US\$7.2 billion ⁴
CPACS Cardioprotection & anticancer synergy in combination with doxorubicin in solid tumours ⁵	Global: 4 million	 Phase 1			Patients have been safely dosed with RC220 and RC220+Dox ⁵

1. Estimated number of patient per year for each indication. US – USA & Canada; EU - Europe

2. ASX Announcement (17 Nov 2025)

3. ASX Announcement (30 Jul 2024)

4. AstraZeneca Annual Report 2025; Janssen Annual Report 2025; Hansoh Annual Report 2025; Shanghai Allist Annual report 2025

5. ASX Announcement (20 Oct 2025)

Shared mechanism, multiple pathways to market

RC220's shared MYC mechanism supports multiple oncology programs



RC220

Program 1: Acute myeloid leukaemia (EMILI)

Phase 3

Monotherapy

US\$3.5B+

Global AML market size (2024)

MYC represents the lead validation pathway for RC220's MYC-targeting mechanism

*Clinical foundation for all programs.
Provides early MYC data insights.*

Program 2: EGFRm lung cancer (HARNESS)

Phase 1

In combination with osimertinib (Tagrisso)

US\$7.2B+

Global osimertinib annual sales (2025)

MYC-mediated resistance emerges following osimertinib (AstraZeneca's top selling cancer drug) treatment – supporting combination potential

MYC involved in resistance to many other drug classes.

Program 3: Cardioprotection & anticancer (CPACS)

Phase 1

In combination with doxorubicin

US\$1.5B+

Global doxorubicin (generic) market size

Anthracyclines remain widely used oncology backbones, but cardiotoxicity limits treatment intensity – RC220 aims to improve anthracycline anticancer activity and provide cardioprotection

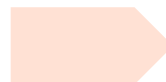
RC220 reduces doxorubicin toxicity, enabling more doses to be given.

Key takeaway

MYC dysregulation is implicated across all three target pathways, supporting broader applicability of RC220

Extreme value from clinical success with difficult targets

RAS provides a compelling precedent for MYC – value of success with major “undruggable” targets

Company	Target	Stage at peak (all pre-revenue)				Peak market value	Outcome/ Status
		1/1b	2	3	Approved		
 MIRATI THERAPEUTICS®	KRAS G12C					US\$12B	2023 - Acquired for \$US4.8bn +\$1bn earn out  Bristol Myers Squibb
	Pan-RAS / KRAS					US\$40B	2026 - Takeover speculation
	Pan-RAS / KRAS					US\$6B	2026 - Raised US\$260m

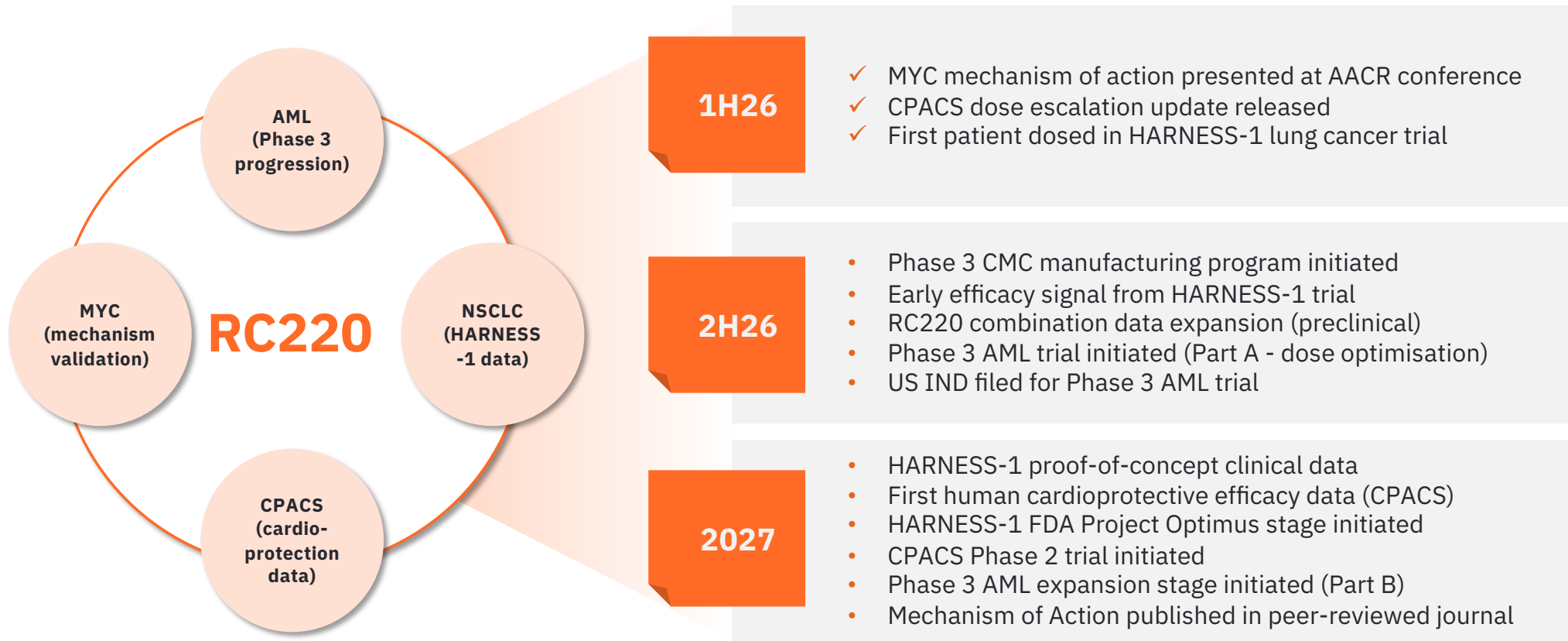
Key takeaway

RAS was undruggable for decades – changing this has generated significant value

RC220 is the only clinically efficacious MYC-targeting drug and is at Phase 3

Multiple clinical and strategic catalysts through 2027

Upcoming milestones across AML, NSCLC, cardioprotection and mechanism validation





Contact

DANIEL TILLET

CEO & MD

 +61 2 8051 3043

 daniel.tillett@racuraoncology.com

JANE LOWE

Investor & Media Relations

 +61 411 117 774

 jane.lowe@irdepartment.com.au