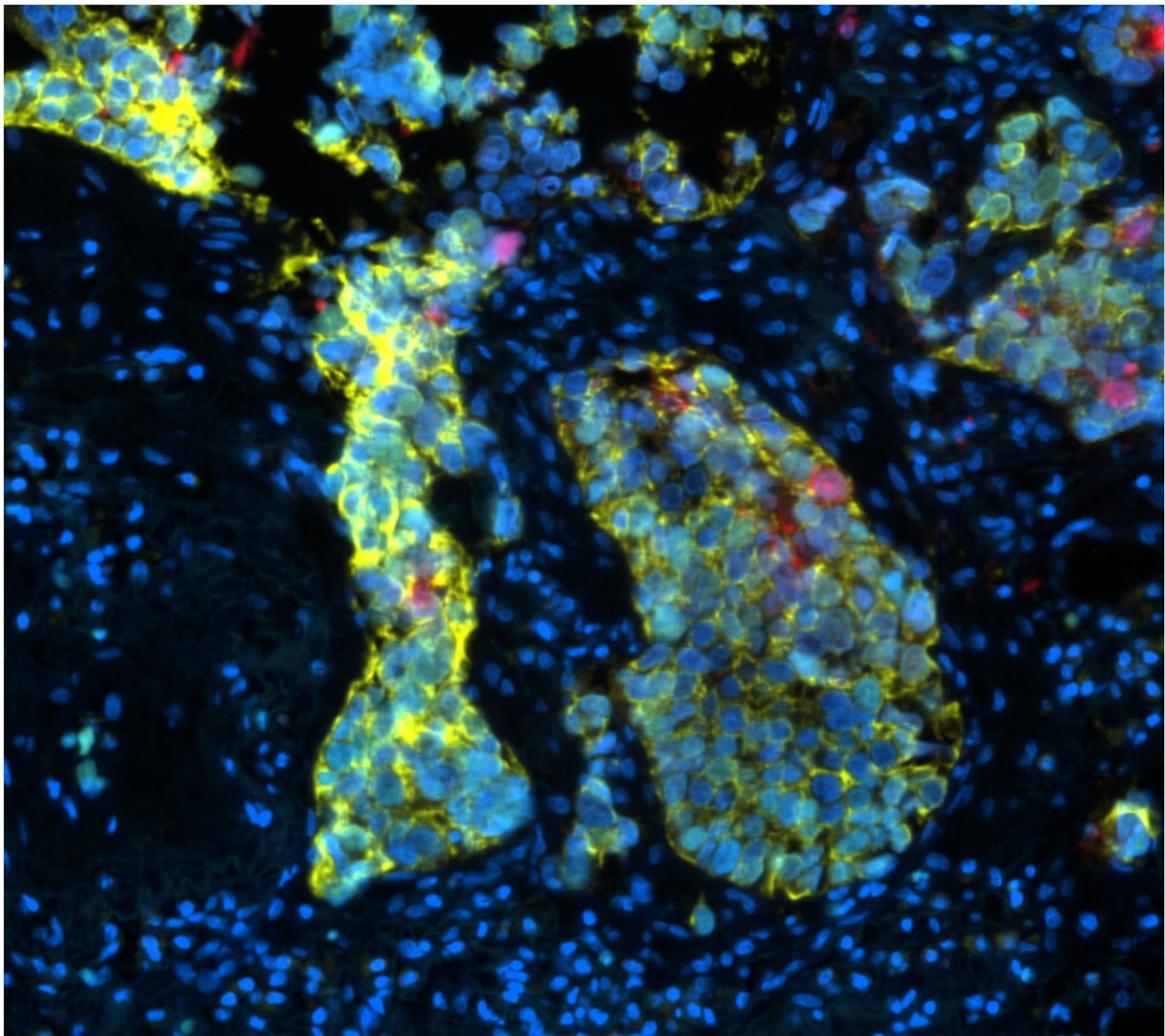


Tumour analysis following surgery could predict response to drug treatment

November 8 2017



Lung tumour explants from patients are treated with current standard therapy, Cisplatin, and novel drug X. Yellow represents tumour area, green represents

proliferating cells and red dying cells (top left). We can computationally detect all cells that are actively dividing and dying (top right). These data can then be used to calculate a patient specific response to standard of care drugs (Cisplatin), and novel targeted therapies (drug X) (bottom), allowing us to predict if novel drugs are better than current therapeutic options. Ki67 represents dividing cells, PARP represents proportion of dying cells. Credit: University of Leicester

A novel approach developed by researchers from the University of Leicester and the MRC Toxicology Unit could help to predict how well patients respond to drugs designed to fight various forms of cancer.

The team has developed a patient-relevant pre-clinical platform called 'explants', whereby tumour samples are taken from cancer [patients](#) immediately following surgery and drug responses are then assessed in situ.

The 'explant' approach is cheap, quick, and reproducible and has already given results that are more predictive of patient outcome than other methods.

The study, which was led by Professor Catrin Pritchard and Dr Howard Pringle from the University of Leicester Cancer Research Centre and Professor Marion MacFarlane, who is based within the MRC Toxicology Unit at Leicester, applied the 'explant' approach to lung cancer.

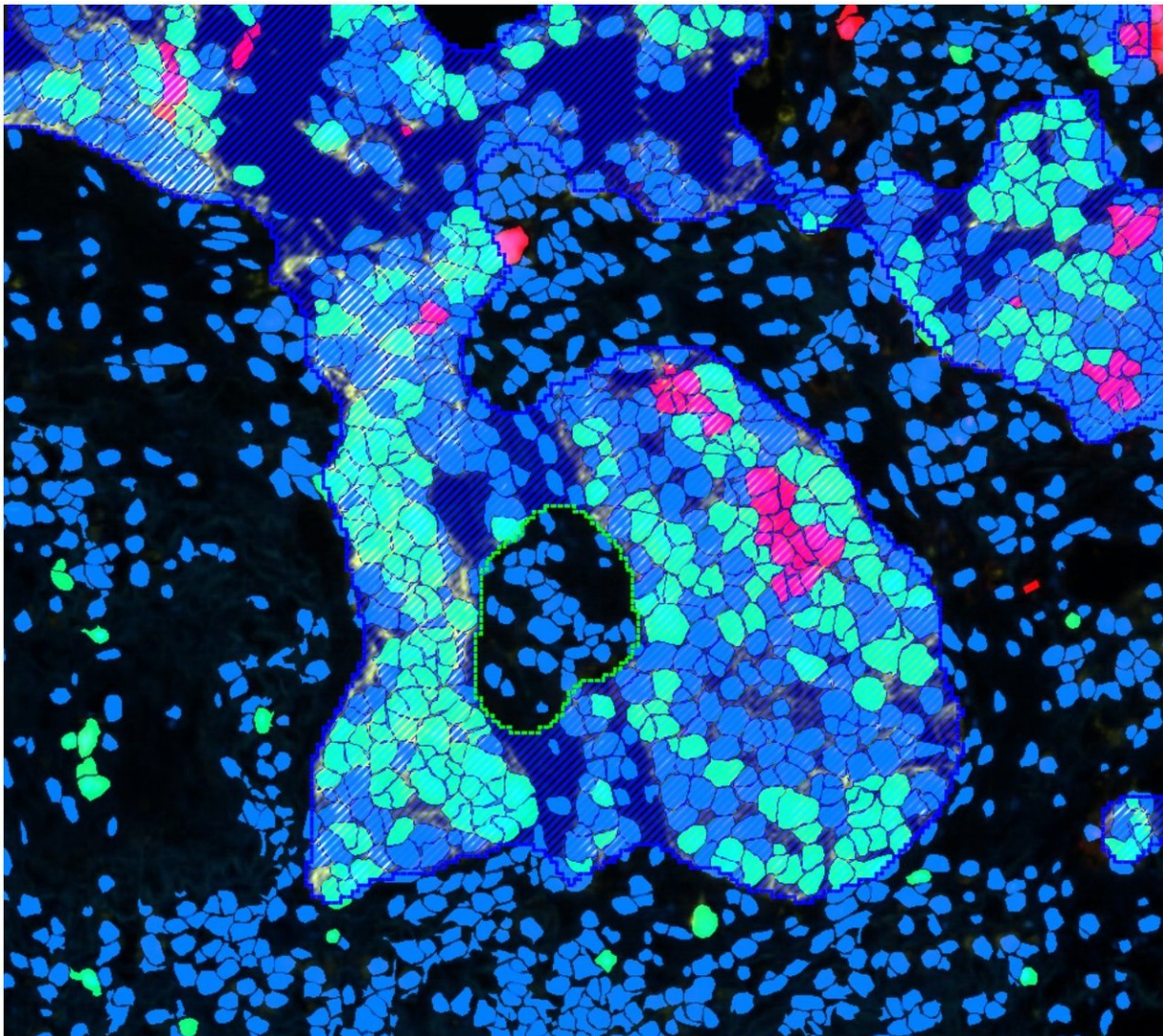
They found that the different cell types within a tumour are retained intact and that the way in which the tumour 'explants' respond to drugs predicts how well individual patients respond in the clinic.

Professor Pritchard explained: "Despite the huge level of investment into the development of new anti-cancer drugs by academia and the

pharmaceutical industry, most do not deliver patient benefit in the clinic.

"We carried out this study to test whether our 'explant' approach gives patient-relevant data with regard to standard-of-care chemotherapy and provide strong evidence that this is indeed the case for lung cancer.

"If incorporated into drug development procedures, then the 'explant' platform has an opportunity to make a huge contribution to the anti-[cancer drug development](#) pipeline."



Lung tumour explants from patients are treated with current standard therapy, Cisplatin, and novel drug X. Yellow represents tumour area, green represents proliferating cells and red dying cells (top left). We can computationally detect all cells that are actively dividing and dying (top right). These data can then be used to calculate a patient specific response to standard of care drugs (Cisplatin), and novel targeted therapies (drug X) (bottom), allowing us to predict if novel drugs are better than current therapeutic options. Ki67 represents dividing cells, PARP represents proportion of dying cells. Credit: University of Leicester

Professor MacFarlane added: "This work is important as it will reduce the attrition rates of cancer [drug](#) development and improve the efficiency and speed with which anti-cancer drugs are incorporated into the clinic to give patient benefit. This will also substantially reduce the costs of developing novel anti-cancer drugs.

"We are very excited by the potential of our platform and have already expanded our approach to other cancer types such as breast cancer, mesothelioma and colorectal cancer with excellent results in each of these disease types."

The study was published in *Cancer Research* earlier this year.

In recognition of the potential of the platform, Cancer Research UK Therapeutic Discovery Laboratories previously co-funded method development and following this successful pilot study the team has now completed a research agreement with both Cancer Research UK Therapeutic Discovery Laboratories and LifeArc to test a range of novel [anti-cancer drugs](#), attracting investment totalling >£500K.

Tim Hammonds, Deputy Director of Drug Discovery at Cancer Research UK Therapeutic Discovery Laboratories, said: "Access to

clinical tissues for our [drug discovery](#) projects should provide vital information on the mechanisms and possible application of new therapeutics. We are delighted to be working in this collaboration."

Dr Justin Bryans, Executive Director, Drug Discovery at LifeArc, said: "We are very excited about this collaboration. The ability to test experimental drugs against explants will provide valuable information and enable us to expedite the [discovery](#) of new cancer therapies. In addition, if we can use this platform to inform oncologists of the appropriate treatment for their patients then this could transform the way we treat [cancer](#)."

This has allowed the establishment of a 'Cancer Explant Facility' located in the University's Robert Kilpatrick Clinical Sciences Building at the Leicester Royal Infirmary Site. This facility is also being supported by the University's Leicester Drug Discovery and Diagnostics Centre (LD3), and the Leicester Precision Medicine Institute (LPMI).

Alasdair Gaw, the Commercial Director of both LD3 and LPMI, commented: "The development of this pre-clinical platform is a great example of how a collaborative approach based on excellent scientific research can positively impact clinical practice to provide patient benefit and typifies why Leicester is a leading centre for precision medicine."

More information: Ellie Karekla et al. Ex Vivo Explant Cultures of Non-Small Cell Lung Carcinoma Enable Evaluation of Primary Tumor Responses to Anticancer Therapy, *Cancer Research* (2017). [DOI: 10.1158/0008-5472.CAN-16-1121](https://doi.org/10.1158/0008-5472.CAN-16-1121)

Provided by University of Leicester

Citation: Tumour analysis following surgery could predict response to drug treatment (2017, November 8) retrieved 25 April 2024 from <https://medicalxpress.com/news/2017-11-tumour-analysis-surgery-response-drug.html>

This document is subject to copyright. Apart from any fair dealing for the purpose of private study or research, no part may be reproduced without the written permission. The content is provided for information purposes only.