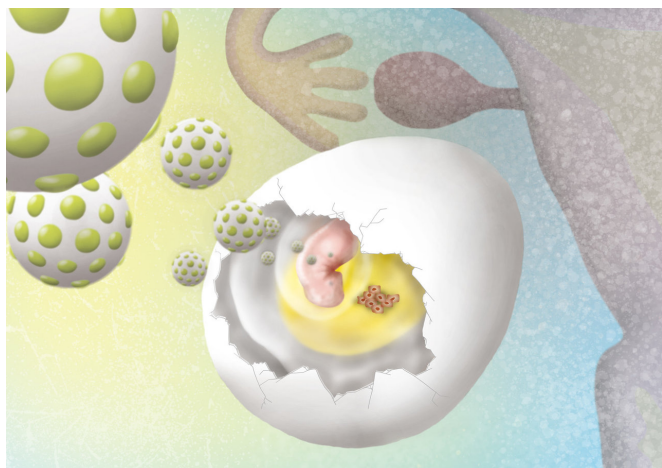


A major step towards individualized cancer therapy

4 June 2018



The team opened a small window in eggshells and implanted ovarian tumour cells on top of the vascular membrane surrounding 10-day-old chicken embryos. Credit: Izumi Mindy Takamiya

Fuyuhiko Tamanoi of Kyoto University's Institute for Integrated Cell-Material Sciences (iCeMS) and colleagues in the U.S. succeeded in establishing a powerful and convenient model to analyse human cancer.

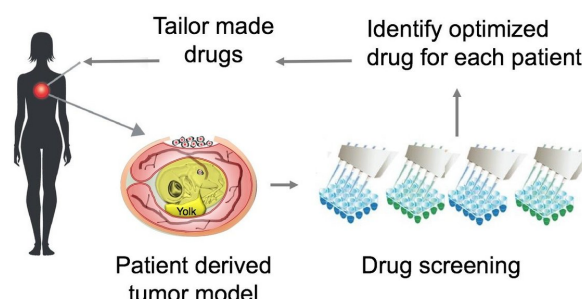
The researchers developed a [chicken egg tumour model](#) in which cultured [ovarian cancer cells](#) are transplanted on top of the membrane that surrounds a 10-day-old [chicken](#) embryo. An ovarian tumour forms on top of the membrane within three days of transplantation.

The team had similar results when they used ovarian tumour samples taken directly from patients, showing that their chicken egg model provides a convenient system for replicating human cancer. This conclusion is supported by their detailed characterization of the tumour, demonstrating that it possesses all major cancer features. "We were surprised when the tumour was

formed in three days," says Tamanoi."

This is very rapid, considering that it takes weeks to do the same with mice. We can start using this model to test for anti-cancer drugs tailored to each cancer patient's needs. The process can be completed within one week," he says. This is a major step toward individualized medicine for cancer patients.

Tamanoi's team, in collaboration with colleagues in France and Saudi Arabia, also developed a new type of biodegradable silica nanoparticle called biodegradable PMO, which is only 200 nanometres in size. The nanoparticles were loaded with the anti-cancer drug doxorubicin and were tested on human ovarian tumour established in the chicken egg.



Researchers developed a faster and cheaper way to test cancer drugs to develop patient-specific treatments. Credit: Kyoto University iCeMS

The biodegradable PMO carrying doxorubicin quickly eliminated the human ovarian tumours without affecting other organs in the chicken embryo. When a smaller amount of the drug, not enveloped in the nanoparticles, was injected in the

egg, severe organ damage ensued. This indicates that the team's nanoparticles prevent anti-cancer drug side effects due to their ability to directly target the [tumour](#).

The chicken egg model has several advantages over existing models, such as mouse models, for testing anti-[cancer](#) therapies. The tumours form much more rapidly on the chicken embryonic membranes than in mice due to the rich nutrient environment and the incomplete immune system at this stage of embryonic development. Fertilized chicken [eggs](#) are also less expensive to use than immune-compromised mice making the model suitable for high throughput experiments.

More information: Binh Thanh Vu et al. Chick chorioallantoic membrane assay as an in vivo model to study the effect of nanoparticle-based anticancer drugs in ovarian cancer, *Scientific Reports* (2018). [DOI: 10.1038/s41598-018-25573-8](https://doi.org/10.1038/s41598-018-25573-8)

Provided by Kyoto University

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