

# Georg Winter wins the 2019 Eppendorf Award for Young European Investigators



Georg Winter receives his Eppendorf Award from jury chair Reinhard Jahn.

PHOTOS: EMBL PHOTOLAB, HEIDELBERG, GERMANY.

**Georg Winter, a chemical biologist at the CeMM Research Center for Molecular Medicine of the Austrian Academy of Sciences in Vienna, Austria, talks to science writer Geoff Marsh about his work on protein degradation. He describes his research on the notorious drug thalidomide, aiming to turn it into a 'chemical knock-out' for cancer. This year's ceremony took place at the EMBL Advanced Training Centre in Heidelberg, Germany, on 27 June, 2019.**

**Geoff Marsh:** Congratulations for winning this year's prize. You must be chuffed.

**Georg Winter:** It's a huge honour, and a very humbling experience.

**GM:** The work you're being celebrated for started with a key paper in 2015 in *Science*, detailing a new technique for targeting specific proteins for degradation. How does that work?

**GW:** It's a relatively simple principle. We started out by making small molecules that have two 'arms', where one arm binds to a protein of interest, and the second binds to the protein degradation machinery, which is active in every cell. We call them 'bifunctional molecules', and we use them to reprogram the degradation machinery to eliminate a protein of interest.

**GM:** The drug you use to bind those two components is a phthalimide, which has a contentious history.

**GW:** Yes. The first phthalimide that you'll have heard of is called thalidomide. It was marketed for morning sickness and as a sedative [in the 1950s], and it was taken by many pregnant women. Unfortunately, it turned out that if women took it in the first trimester of pregnancy, their babies were born with limb abnormalities. That particular molecule was withdrawn from the market, unfortunately only after around ten thousand babies were born

with limb defects. For many decades, nobody wanted to touch thalidomide or its analogues. But it then became clear that they have very interesting anticancer activities.

**GM:** How does this phthalimide work?

**GW:** In 2010, a Japanese research group identified its cellular binding partner. This turned out to be a protein called cereblon which is a so-called E3 ligase. This is one of the proteins engaged in the cellular degradation machinery. Once we knew that thalidomide binds to this E3 ligase, the question was whether we could use this binding event to reprogram the machinery — by conjugating thalidomide to another small molecule that would bind to a protein of interest? Once that happens, we have these 'heterobifunctional' molecules that bind to both the degradation machinery and the protein of interest, bring the two together, and redirect the degradation activity to the target protein.

**GM:** In that 2015 paper, your target protein was one essential for the growth and survival of cancer, and you tested it in mice. How did that go?

**GW:** It was one of the key findings of the paper. We were already very excited to see that the molecule worked in a cancer-cell line in a dish. But the real test is in an animal model, to check whether it had some therapeutic potential. We were amazed by how well it worked. These mice already had a tumour growing. We took out the tumour and stained and quantified it for levels of a protein called BRD4. A couple of hours after a single treatment with this particular molecule, levels of the protein had decreased in the tumour; when we treated those tumour-bearing mice every day, the tumour growth was profoundly reduced.



Georg Winter discussing his research with Geoff Marsh, Nature Research.



Left to right: Axel Jahns<sup>1</sup>, Wilhelm Plüster<sup>1</sup>, Laura Machesky<sup>2</sup>, Georg Winter, Maria Leptin<sup>2</sup> and Reinhard Jahn<sup>2</sup>

<sup>1</sup>Eppendorf AG  
<sup>2</sup>Eppendorf Award Jury

We slowed down the tumour growth by degrading a protein relevant to that cancer.

**GM:** Theoretically, could this phthalimide conjugation technique work for any protein target for which there is a ligand?

**GW:** That is a fair statement, and for some proteins we might even have ligands without knowing it. So far, we've only really cared about ligands that change protein function. But now, binding of the ligand can be entirely inconsequential — we just use it as an anchor to induce closeness to the protein degradation machinery.

Often in cancer research we're driven by genetic studies. For instance, we know that if we knock out a particular gene, we slow down cancer growth. But this particular gene might code for a protein with three different functionalities. Some of these functionalities might look like great targets for a small molecule ligand because they have been inhibited before. But it can be that inhibiting one functionality does not equate to knocking out or deleting the gene. We can now revisit all these cases because our chemical perturbation of degrading the protein is much closer to deleting it genetically than the conventional pharmacological way to just inhibit a protein's function.

**GM:** So it's kind of like a 'chemical knockout'?

**GW:** Very much so.

**GM:** What do you think should be the next protein targets that would help the fight against cancer, using this technique?

**GW:** We now have the tools to tackle some transcription co-activators and regulators of gene control. Ultimately, we should aim to target transcription factors for degradation. These will be the proteins with the highest bar, because for many of them we don't have defined binding pockets to help us predict the ligand. But this will be the challenge and there are already strategies out there to tackle these limitations.

**GM:** How close is this technology to readiness for human trials?

**GW:** Not directly with the technology of phthalimide conjugation, but there have been some with other closely related heterobifunctional molecules that hijack another part of the protein degradation machinery and have similar functionality. There is one biotech company from Newhaven called Arvinas. They have started trials in humans to degrade a transcription factor, called the androgen receptor, in prostate cancer. This is, to the best of my knowledge, the first clinical trial. The entire field is eagerly waiting for the results.

**GM:** Finally, what's next for you?

**GW:** We are very much interested in pushing this technology to the next limit and eventually being able to degrade proteins that we can't bind. I'll tell you how we plan to do this the next time we meet.

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## ABOUT THE AWARD (EST. 1995)

Presented in partnership with *Nature*, the Eppendorf Award for Young European Investigators recognizes outstanding work in biomedical science. Besides a prize money of € 20,000, it provides the opportunity for European researchers to showcase their work and communicate their research to a scientific audience.

The winner is selected by an independent jury of scientists under the chairmanship of Reinhard Jahn, Director Emeritus at the Max Planck Institute for Biophysical Chemistry in Göttingen, Germany. Eppendorf and *Nature* do not influence the selection.

For more information see:  
[www.eppendorf.com/award](http://www.eppendorf.com/award)

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We invite biological and biomedical researchers with an advanced degree, not older than 35 years who work in Europe to apply for the 2020 Eppendorf Award. Applications will be accepted from 1 October 2019 and the deadline for entries is 15 January 2020. The prize ceremony will take place at the EMBL Advanced Training Centre (ATC) in Heidelberg, Germany, on 25 June 2020 in conjunction with the Young European Investigators Conference 2020.



To find out more about Georg Winter's work, visit: <https://cemm.at/research/groups/georg-winter-group>