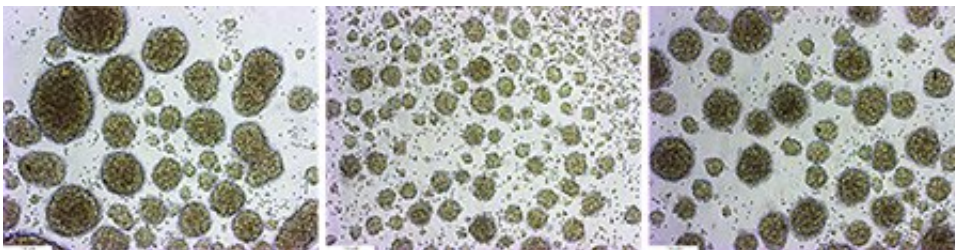


# Zika virus' key into brain cells ID'd, leveraged to block infection and kill cancer cells

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3D human brain organoids. Left: normal, uninfected. Center: infected with Zika virus. Right: infected with Zika virus and treated with cilengitide, which protects the cells from destruction by the virus. Credit: UC San Diego Health Sciences

Zika virus infection can stunt neonatal brain development, a condition known as microcephaly, in which babies are born with abnormally small heads. To determine how best to prevent and treat the viral infection, scientists first need to understand how the pathogen gets inside brain cells.

Employing different approaches to answer different questions, two research teams at University of California San Diego School of Medicine independently identified the same molecule— $\alpha v \beta 5$  integrin—as Zika [virus](#)' key to entering [brain stem cells](#).

In a pair of papers published January 16, 2020 by *Cell Press*, the

researchers also found ways to take advantage of the integrin to both block Zika virus from infecting [cells](#) and turn it into something good: a way to shrink [brain](#) cancer stem cells.

Integrins are molecules embedded in cell surfaces. They play important roles in cell adherence and communication, and are known to be involved in cancer progression and metastasis. Several other integrins are known entry points for other viruses, including adenovirus, foot-and-mouth disease virus and rotavirus, but  $\alpha\text{v}\beta 5$  was not previously known for its role in [viral infections](#).

## Finding the key

One team, led by Tariq Rana, Ph.D., professor and chief of the Division of Genetics in the Department of Pediatrics at UC San Diego School of Medicine and Moores Cancer Center, used CRISPR gene editing to systematically delete every gene in a 3-D culture of human glioblastoma (brain cancer) stem cells growing in a laboratory dish. Then they exposed each variation to Zika virus to determine which genes, and the proteins they encode, are required for the virus to enter the cells. The virus was—for the first time—labeled with green fluorescent protein (GFP) to allow the researchers to visualize viral entry into the cells.

Their study, published in *Cell Reports*, uncovered 92 specific human brain cancer stem cell genes that Zika virus requires to infect and replicate in the cells. But one gene stood out, the one that encodes  $\alpha\text{v}\beta 5$  integrin.

"Integrins are well known as molecules that many different viruses use as doorknobs to gain entry into human cells," Rana said. "I was expecting to find Zika using multiple integrins, or other cell surface molecules also used by other viruses. But instead we found Zika uses  $\alpha\text{v}\beta 5$ , which is unique. When we further examined  $\alpha\text{v}\beta 5$  expression in brain, it made

perfect sense because  $\alpha\text{v}\beta 5$  is the only integrin member enriched in neural stem cells, which Zika preferentially infects. Therefore, we believe that  $\alpha\text{v}\beta 5$  is the key contributor to Zika's ability to infect brain cells."

## Blocking Zika virus infection

The second study, published in *Cell Stem Cell*, was led by Jeremy Rich, MD, professor in the Department of Medicine at UC San Diego School of Medicine and director of neuro-oncology and of the Brain Tumor Institute at UC San Diego Health. Knowing that many viruses use integrins for entry into human cells, Rich's team inhibited each integrin with a different antibody to see which would have the greatest effect.

"When we blocked other integrins, there was no difference. You might as well be putting water on a cell," said Rich, who is also a faculty member in the Sanford Consortium for Regenerative Medicine and Sanford Stem Cell Clinical Center at UC San Diego Health. "But with  $\alpha\text{v}\beta 5$ , blocking it with an antibody almost completely blocked the ability of the virus to infect brain cancer stem cells and normal brain stem cells."

Rich's team followed up by inhibiting  $\alpha\text{v}\beta 5$  in a glioblastoma mouse model with either an antibody or by deactivating the gene that encodes it. Both approaches blocked Zika virus infection and allowed the treated mice to live longer than untreated mice. They also found that blocking the  $\alpha\text{v}\beta 5$  integrin in glioblastoma tumor samples removed from patients during surgery blocked Zika virus infection.

Rana's team also blocked  $\alpha\text{v}\beta 5$  in mice, treating them daily with cilengitide or SB273005, two experimental cancer drugs that target the integrin. Six days after Zika virus infection, the brains of their drug-treated mice contained half as much virus as mock-treated mice.

"The neat thing is that these findings not only help advance the Zika virus research field, but also opens the possibility that we could similarly block the entry of multiple viruses that use other integrins with antibodies or small molecule inhibitors," Rana said.

Rana and team are now engineering a mouse model that lacks  $\alpha v\beta 5$  integrin in the brain—a tool that would allow them to definitively prove the molecule is necessary for Zika viral entry and replication.

## Leveraging Zika to treat brain cancer

Rich is a neuro-oncologist who specializes in diagnosing and treating patients with glioblastoma, a particularly aggressive and deadly type of brain tumor. When he first saw how the Zika virus shrinks brain tissue, it reminded him of what he hopes to achieve when he's treating a patient with glioblastoma. In 2017, he and collaborators [published a study](#) in which they determined that Zika virus selectively targets and kills glioblastoma stem cells, which tend to be resistant to standard treatments and are a big reason why glioblastomas recur after surgery and result in shorter patient survival rates.

Rich's latest study helps account for the virus' preference for glioblastoma stem cells over healthy [brain cells](#). The  $\alpha v\beta 5$  integrin is made up of two separate subunits— $\alpha v$  and  $\beta 5$ . The team found that glioblastoma stem cells produce a lot of both the  $\alpha v$  subunit (associated with stem cells) and  $\beta 5$  subunit (associated with cancer cells). Together, these units form the  $\alpha v\beta 5$  integrin, which, the team discovered, plays an important role in glioblastoma stem cell survival. Those high levels of  $\alpha v\beta 5$  integrin also help explain why, in the study, glioblastoma stem cells were killed by Zika virus at much higher rates than normal stem cells or other brain cell types.

"It turns out that the very thing that helps cancer cells become aggressive

cancer stem cells is the same thing Zika virus uses to infect our cells," Rich said.

To see how this might play out in a more realistic model of human disease, Rich's team partnered with an expert in human brain disease modeling—Alysson Muotri, Ph.D., professor at UC San Diego School of Medicine, director of the UC San Diego Stem Cell Program and a member of the Sanford Consortium for Regenerative Medicine, and team. Pinar Mesci, Ph.D., a postdoctoral researcher in Muotri's lab, generated a new brain tumor model, where human glioblastoma tumors were transplanted into human brain organoids, laboratory "mini-brains" that can be used for drug discovery. The researchers discovered that Zika virus selectively eliminates glioblastoma [stem cells](#) from the brain organoids. Inhibiting  $\alpha\beta5$  [integrin](#) reversed that anti-cancer activity, further underscoring the molecule's crucial role in Zika virus' ability to destroy cells.

Now Rich's team is partnering with other research groups to perform targeted drug studies. In addition to searching for drugs to block Zika virus, as Rana's group is doing, Rich is interested in genetic modifications to the virus that could help better target its destruction to brain cancer cells, while leaving healthy cells alone.

"While we would likely need to modify the normal Zika virus to make it safer to treat brain tumors, we may also be able to take advantage of the mechanisms the virus uses to destroy cells to improve the way we treat glioblastoma," Rich said. "We should pay attention to viruses. They have evolved over many years to be very good at targeting and entering specific cells in the body."

Zika virus was perhaps best known in 2015-16, when a large outbreak affected primarily Latin America, but also several other regions of the world. While that particular epidemic has passed, Zika virus has not

gone away. Smaller, local outbreaks continue and this past summer, the first few cases of native Zika virus infection were recorded in Europe. Scientists warn Zika could continue to spread as climate change affects the habitat range of the mosquito that carries it. The virus can also be transmitted from pregnant mother to fetus, and via sexual contact. More than half of all people on Earth are at risk for Zika virus infection, and there is no safe and effective treatment or vaccine.

**More information:** *Cell Reports* (2020).

[www.cell.com/cell-reports/full ... 2211-1247\(19\)31491-3](http://www.cell.com/cell-reports/full...2211-1247(19)31491-3)

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