

# Researchers discover why pancreatic cancer cells are so aggressive

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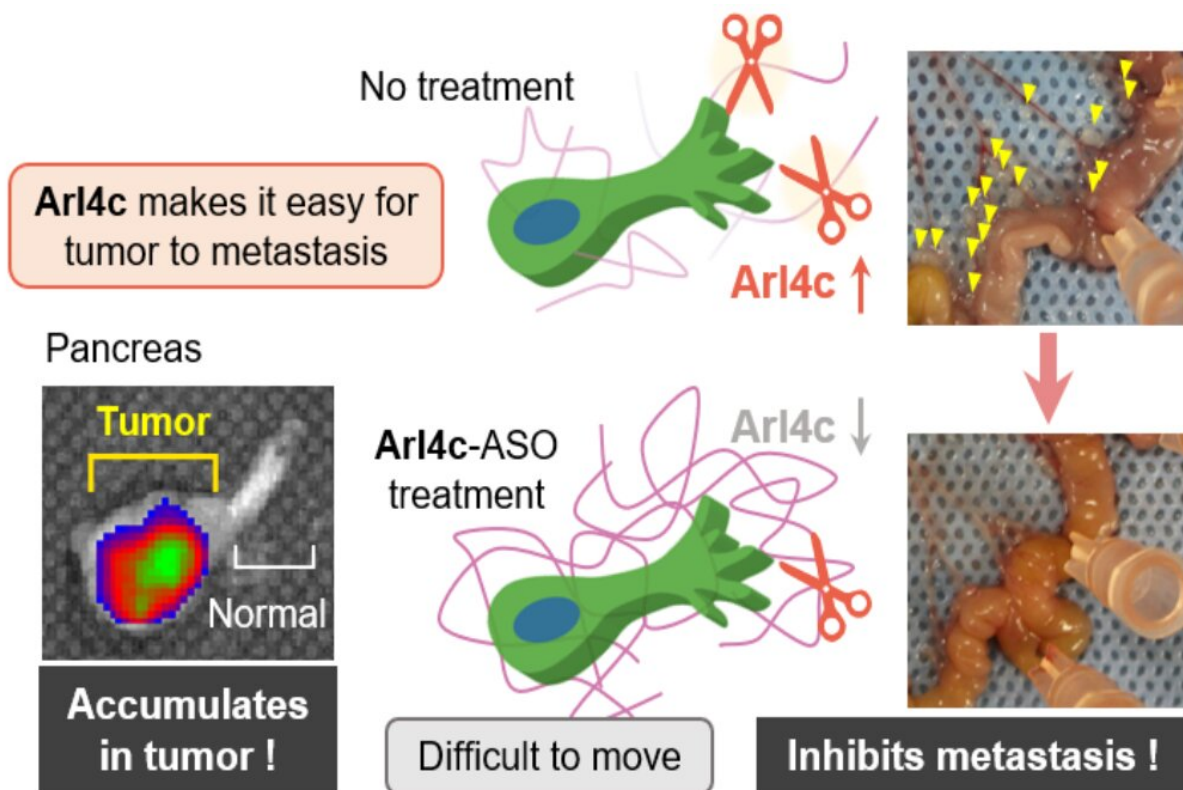


Fig.1 Antisense oligonucleotide (ASO) targeting Arl4c inhibits pancreatic cancer metastasis. Credit: Akikazu Harada

Pancreatic cancer is very aggressive and, unfortunately, has a poor prognosis with a 5-year survival of only 5%. Most deaths related to pancreatic cancer are due to metastasis, which occurs when the tumor

invades other organs. Treatments to stop tumor metastasis, which will save thousands of lives, are still missing. In this context, understanding the molecular mechanisms that cause pancreatic cancer metastasis is the first step to developing effective targeted treatments.

In a study published in *eLife*, researchers from Osaka University revealed a previously unknown mechanism for metastasis of [pancreatic cancer](#) that could be used as a therapeutic target.

First, the study analyzed human pancreatic [cancer](#) tissues and demonstrated that a small signaling protein called ARL4C is overexpressed in pancreatic cancer patients. Initial results regarding the function of this protein suggested that it could be involved in the migratory and invasive abilities of pancreatic cancer cells. To investigate this, and to confirm the location of ARL4C in invading pancreatic cancer cells, the researchers designed an elaborate experiment that simulated the invasion of cancer cells in the human body. They created a 3D culture device that could monitor cancer cells invading into the surrounding collagen gel which they used to observe the invasion of living cells containing fluorescently labeled ARL4C through a microscope.

"We discovered that ARL4C localized to so-called invasive pseudopods—functionally analogous to, but structurally distinct from invadopodia—at the cell surfaces," explains Akikazu Harada, lead author of the study. Invadopodia are well-characterized foot-like extrusions arising from a cell's ventral surface that cancer cells use to invade other tissues. Conversely, invasive pseudopods are longer and have a larger diameter than invadopodia, and are extended from the front end of the cell. "In these pseudopods ARL4C recruited another protein called IQGAP1—that is also highly expressed in numerous cancers including pancreatic cancer—which transported an enzyme called MMP14 to the pseudopods allowing the cancer cell to break into and invade the

collagen gel or extracellular matrix."

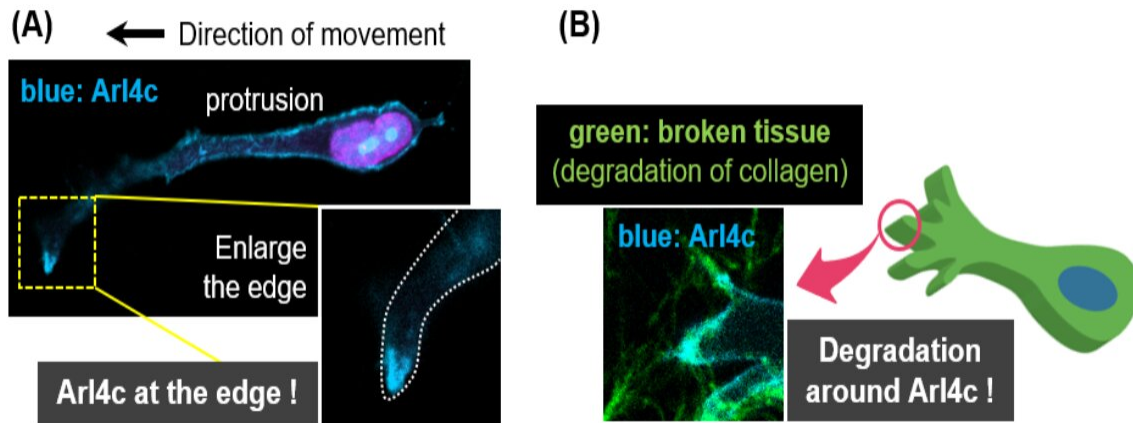


Fig.2 Unique localization of Arl4c. Credit: Akikazu Harada

Once this novel mechanism was revealed, the researchers wanted to harness this new knowledge for therapeutic purposes. Specifically, they adopted an oligonucleotide therapy approach using antisense oligonucleotide or ASO. These are short molecules of single-stranded DNA that act inside the cell to influence—in this case, block—protein production. "An ASO targeted against ARL4C was able to suppress the lymph node metastases of pancreatic cancer [cells](#) implanted into the pancreas of an immunodeficient mouse," says Akira Kikuchi, senior author of the study. If ARL4C is blocked, the [cancer cells](#) are less aggressive and less likely to spread. Although only preliminary, these findings open up promising novel therapeutic avenues for this extremely aggressive cancer and shed light on its mechanism of metastasis.

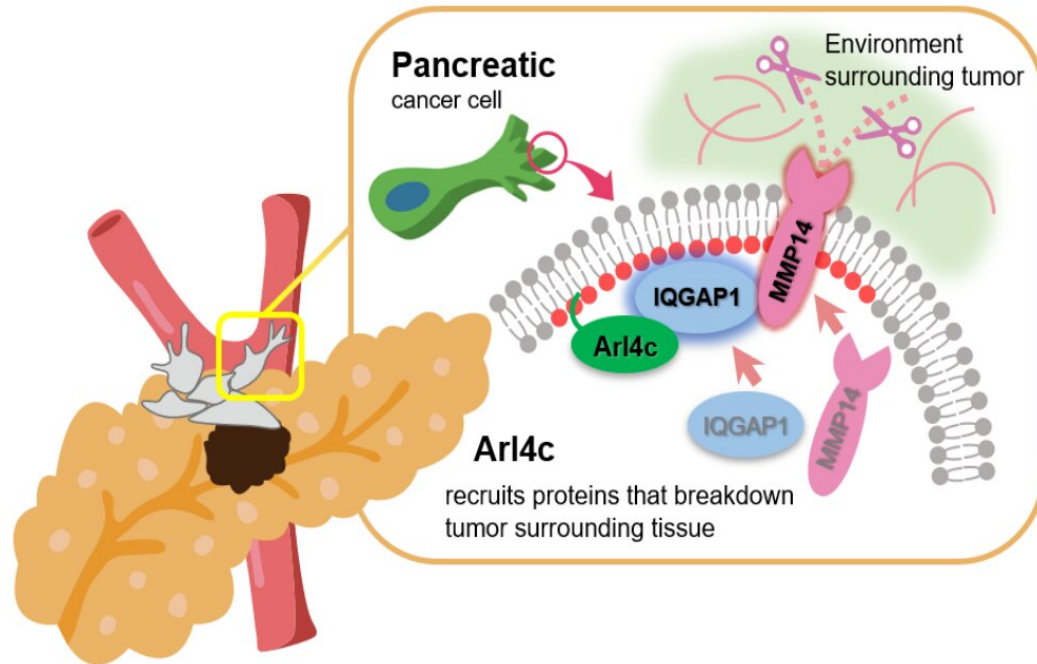


Fig.3 Novel mechanism of pancreatic cancer metastasis. Credit: Akikazu Harada

**More information:** Akikazu Harada et al, Localization of KRAS downstream target ARL4C to invasive pseudopods accelerates pancreatic cancer cell invasion, *eLife* (2021). DOI: [doi.org/10.7554/eLife.66721](https://doi.org/10.7554/eLife.66721)

Provided by Osaka University

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