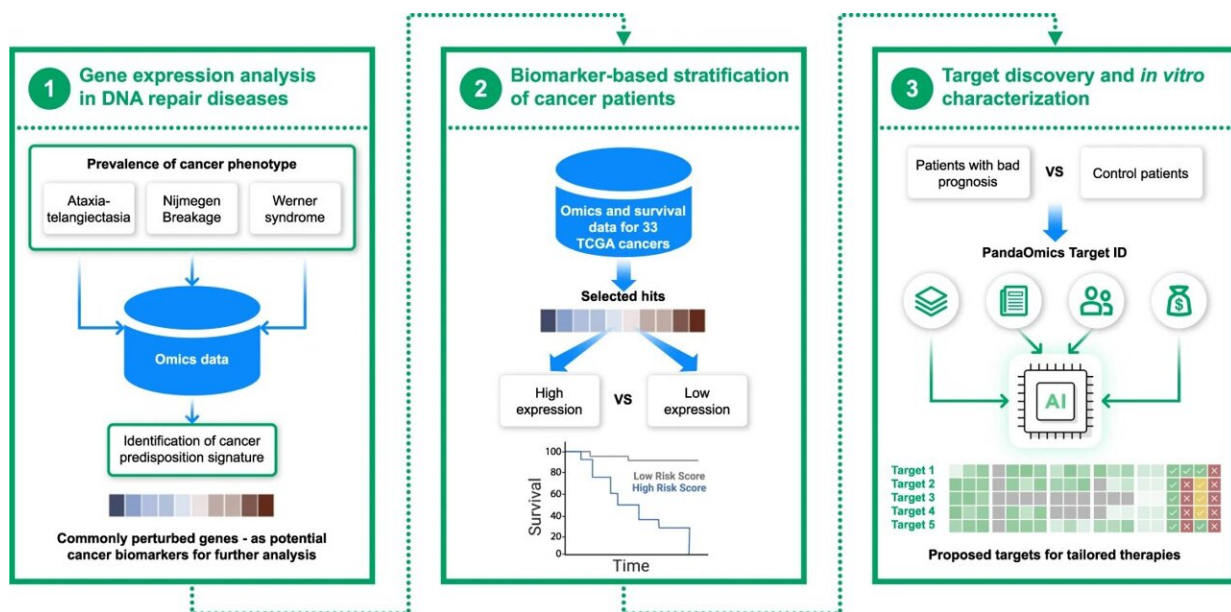


Gray hair and wrinkles at an early age lead researchers to potential new treatment for rare cancer

March 8 2023



Schematic representation of the PandaOmics application for a rapid biomarker discovery and target characterization in cancer. Gene expression signatures have been examined in DNA repair diseases with high cancer predisposition (1), followed by the analysis of the most significantly perturbed genes as potential biomarkers stratifying cancer patients based on their survival rates (2). The group of patients with low survival outcomes have been further used for identification of potential therapeutic candidates for cancer treatment via PandaOmics Target ID approach (3). Credit: *Cell Death & Disease* (2022). DOI: 10.1038/s41419-022-05437-w

Sarcomas are cancer tumors found in the bones, muscles or fatty tissue. They are a rare type of cancer seen in only one percent of cancer patients, and are complex and difficult to treat.

However, a new study may have found a new treatment that can help the sickest [sarcoma](#) patients.

"We have learned that sarcoma patients whose [cancer cells](#) have a high expression of the cep135 protein are worse off. But inhibiting a gene called plk1 also inhibits growth of the sarcoma cells, and this suggests that we can target the treatment of the sickest sarcoma patients," says Associate Professor Morten Scheibye-Knudsen from the Center for Healthy Aging at the Department of Cellular and Molecular Medicine, who is responsible for the new study.

Methods for identifying sarcoma patients' prognoses are already available, as are different forms of treatment. But the new study has identified a new method.

What are sarcomas?

There are two main types: bone sarcomas and soft tissue sarcomas (found in muscles, [fatty tissue](#), connective tissue, blood vessels and neurilemma).

Sarcomas affect one percent of [cancer](#) patients. In Denmark, around 45 people are diagnosed with bone sarcomas each year and 220 with soft tissue sarcomas. Adults diagnosed with bone sarcomas have a 60-percent five-year survival rate, while adults diagnosed with non-bone sarcoma have a 50-70-percent five-year survival rate.

"This is a new way of stratifying and possibly a new and better way of treating sarcomas. And the introduction of yet another method is always

good news to patients. Because no two cancers are alike. Ideally, treatment should always be tailored to the individual patient," Morten Scheibye-Knudsen stresses.

He hopes other researchers with access to the necessary test facilities will study his results in more detail and eventually design a new treatment. If the method turns out to work, he believes a new treatment may be available to patients in five to 10 years.

Gray hair, wrinkles and loss of fatty tissue at an early age

Morten Scheibye-Knudsen and his colleagues started out by studying patients suffering from the rare neurological disorders Werner's syndrome, Nijmegen breakage syndrome and Ataxia-telangiectasia syndrome.

These patients experience symptoms of early aging, such as gray hair, wrinkles and loss of fatty [tissue](#)—and they have a high risk of developing cancer at an early age.

"Age-associated diseases such as cancer is one of my main areas of interest as a researcher at the Center for Healthy Aging. As we grow older, a lot of things happen to the body, and determining causality can be difficult. But in people suffering from e.g. Werner's syndrome it is easier to see which genes are responsible for which processes. This gives us a molecular handle, so to speak," says Morten Scheibye-Knudsen.

In order to establish why these patients develop cancer at an early age, the researchers compared gene expressions across the three disorders. Here they worked together with the company Insilico Medicine, whose large Pandaomics platform made it possible to identify gene mutations in thousands of different disorders. It turned out that *cep135* is a common

denominator for the cancer genes of the three disorders.

"This made us study the gene expressions of various cancers, and we learned that cep135 is associated with high mortality in i.a. sarcoma, but also in bladder cancer. Sarcoma was particularly interesting, as many Werner's syndrome patients develop sarcoma," explains Morten Scheibye-Knudsen.

Finally, the researchers sought to find ways to inhibit the sarcoma. Cep135 is not a useful target, as it is a so-called structural protein, which are difficult to target. Instead, the researchers learned that by inhibiting the plk1 gene they were able to target the sarcoma.

"The study indicates that we can use [genetic diseases](#) that exhibit accelerated aging to identify new treatment targets. In this study, we investigated cancer, but the method can in principle be used for all age-related diseases such as dementia, cardiovascular diseases and others," says Morten Scheibye-Knudsen.

The work is published in the journal *Cell Death & Disease*.

More information: Garik V. Mkrtychyan et al, High-confidence cancer patient stratification through multiomics investigation of DNA repair disorders, *Cell Death & Disease* (2022). [DOI: 10.1038/s41419-022-05437-w](https://doi.org/10.1038/s41419-022-05437-w)

Provided by University of Copenhagen

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