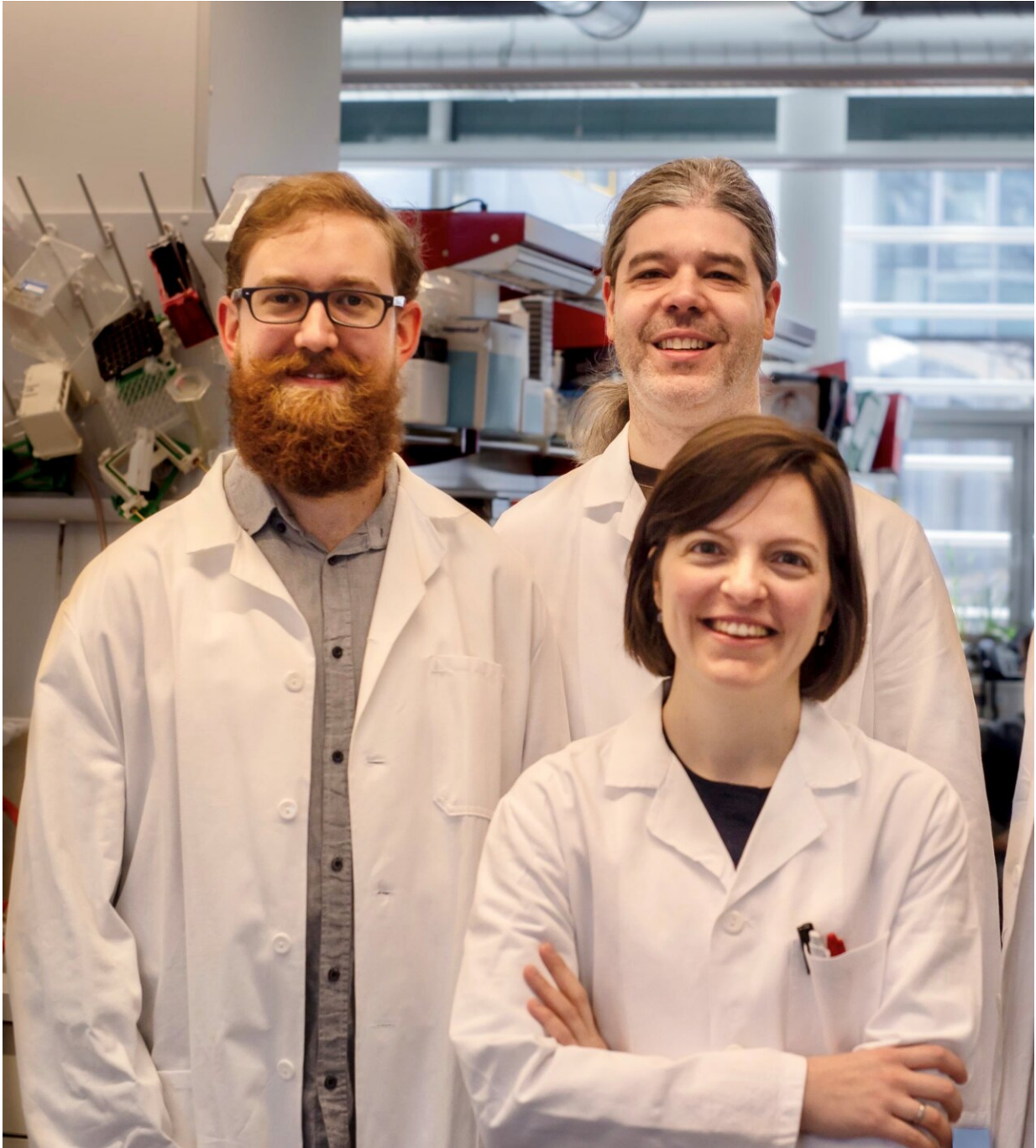


Blood test detects childhood tumors based on their epigenetic profiles

May 28 2021



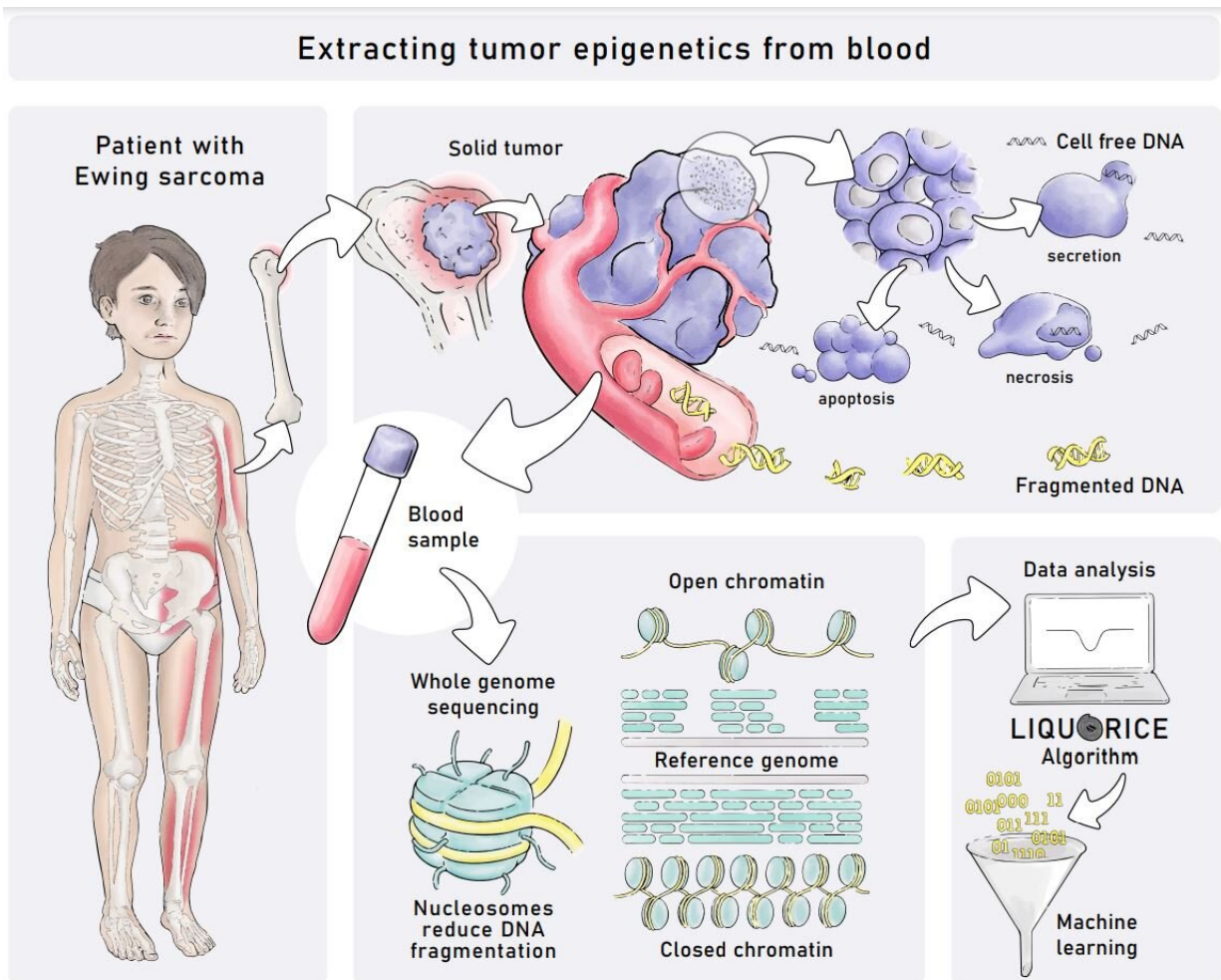
Peter Peneder, MSc, Adrian Stuetz, PhD, and Eleni Tomazou, PhD, develop a liquid biopsy analysis based on tumor epigenetics. Credit: St. Anna Children's Cancer Research Institute

A new study exploits the characteristic epigenetic signatures of childhood tumors to detect, classify and monitor the disease. The scientists analyzed short fragments of tumor DNA that are circulating in the blood. These "liquid biopsy" analyses exploit the unique epigenetic landscape of bone tumors and do not depend on any genetic alterations, which are rare in childhood cancers. This approach promises to improve personalized diagnostics and, possibly, future therapies of childhood tumors such as Ewing sarcoma. The study has been published in *Nature Communications*.

A study led by scientists from St. Anna Children's Cancer Research Institute (St. Anna CCRI) in collaboration with CeMM Research Center for Molecular Medicine of the Austrian Academy of Sciences provides an innovative method for "liquid biopsy" analysis of childhood tumors. This method exploits the fragmentation patterns of the small DNA fragments that tumors leak into the [blood stream](#), which reflect the unique epigenetic signature of many [childhood cancers](#). Focusing on Ewing sarcoma, a bone [tumor](#) of children and young adults with unmet clinical need, the team led by Eleni Tomazou, Ph.D., St. Anna CCRI, demonstrates the method's utility for tumor classification and monitoring, which permits close surveillance of [cancer](#) therapy without highly invasive tumor biopsies.

In tumors, cancer [cells](#) constantly divide, with some of the cancer cells dying in the process. These cells often release their DNA into the blood stream, where it circulates and can be analyzed using genomic methods such as high-throughput DNA sequencing. Such "so-termed liquid biopsy" analyses provide a minimally invasive alternative to conventional tumor biopsies that often require surgery, holding great promise for personalized therapies. For example, it becomes possible to check frequently for molecular changes in the tumor. However, the use of liquid biopsy for childhood cancers has so far been hampered by the fact that many childhood tumors have few genetic alterations that are

detectable in DNA isolated from blood plasma.



Extracting tumor epigenetics from blood. Credit: Tatjana Hirschmugl

Exploiting tumor-specific epigenetic patterns

Cell-free DNA from dying tumor cells circulates in the blood in the form of small fragments. Their size is neither random nor determined solely by the DNA sequence. Rather, it reflects how the DNA is

packaged inside the [cancer cells](#), and it is influenced by the chromatin (i.e., complex of DNA, protein and RNA) structure and epigenetic profiles of these cells. This is very relevant because epigenetic patterns—sometimes referred to as the "second code" of the genome—are characteristically different for different cell types in the human body. Epigenetic mechanisms lead to changes in gene function that are not based on changes in the DNA sequence but are passed on to daughter cells. The analysis of cell-free DNA fragmentation patterns provides a unique opportunity to learn about the epigenetic regulation inside the tumor without having to surgically remove tumor cells or even know whether and where in the body a tumor exists.

"We previously identified unique epigenetic signatures of Ewing sarcoma. We reasoned that these characteristic epigenetic signatures should be preserved in the fragmentation patterns of tumor-derived DNA circulating in the blood. This would provide us with a much-needed marker for [early diagnosis](#) and tumor classification using the liquid biopsy concept," explains Dr. Tomazou, Principal Investigator of the Epigenome-based precision medicine group at St. Anna CCRI.

Machine learning increases sensitivity

The new study benchmarks various metrics for analyzing cell-free DNA fragmentation, and it introduces the LIQUORICE algorithm for detecting circulating tumor DNA based on cancer-specific chromatin signatures. The scientists used machine-learning classifiers to distinguish between patients with cancer and healthy individuals, and between different types of pediatric sarcomas. "By feeding these machine learning algorithms with our extensive whole genome sequencing data of tumor-derived DNA in the blood stream, the analysis becomes highly sensitive and in many instances outperforms conventional genetic analyses," says Dr. Tomazou.

When asked about potential applications, she explains, "Our assay works well, we are very excited. However, further validation will be needed before it can become part of routine clinical diagnostics." According to the scientists, their approach could be used for minimally invasive diagnosis and, but also as a prognostic marker, monitoring which patient responds to therapy. Additionally, it might serve as a predictive marker during neoadjuvant therapy (i.e., chemotherapy before surgery), potentially enabling dose adjustments according to treatment response. "Right now, most patients receive very high doses of chemotherapy, while some patients may be cured already with a less severe therapy, which would reduce their risk of getting other cancers later in life. There is a real medical need for adaptive clinical trials and personalized treatment of bone tumors in children."

More information: Multimodal analysis of cell-free DNA whole genome sequencing for pediatric cancers with low mutational burden, *Nature Communications* (2021). [DOI: 10.1038/s41467-021-23445-w](https://doi.org/10.1038/s41467-021-23445-w)

Provided by St. Anna Children's Cancer Research Institute

Citation: Blood test detects childhood tumors based on their epigenetic profiles (2021, May 28) retrieved 4 May 2024 from

<https://medicalxpress.com/news/2021-05-blood-childhood-tumors-based-epigenetic.html>

<p>This document is subject to copyright. Apart from any fair dealing for the purpose of private study or research, no part may be reproduced without the written permission. The content is provided for information purposes only.</p>
--