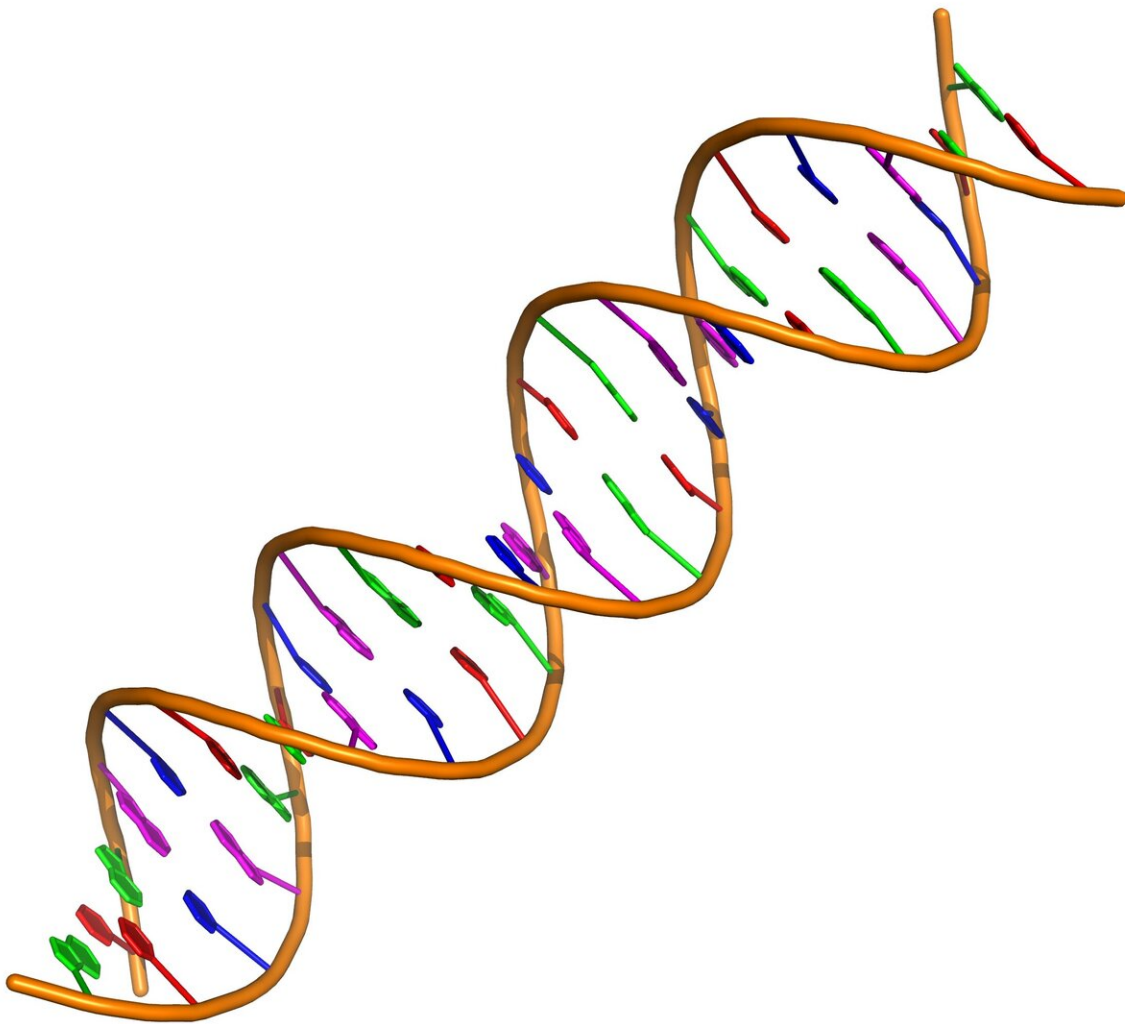


DNA tags enable blood-based tests to assess cancer treatment outcomes

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A double stranded DNA fragment. Credit: Vcpmartin/Wikimedia/ CC BY-SA

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Cell-free DNA (cfDNA) shed into the blood was discovered in the late 1940s but with rapid advances in genomics and computational analytics in just the past few years, researchers at Georgetown Lombardi Comprehensive Cancer Center now believe that studying tags, or modifications to this type of DNA, may lead to a better understanding of how to assess, and possibly modulate, treatment approaches for cancer and other diseases. Their perspective, drawn from a review of studies to date, appears July 27 in *Frontiers in Genetics*.

During [cell death](#), which is a normal part of [tissue](#) regeneration, cfDNA is shed from tissue. The shed cfDNA can be isolated from a [blood sample](#), thus providing a reading of cell death across the body in both normal and [cancer cells](#) without the need for taking invasive biopsy samples.

"Taking tumor tissue biopsies is a hit or miss process and is usually not a good representation of the whole tumor or its spread," says Anton Wellstein, MD, Ph.D., professor of oncology and pharmacology at Georgetown Lombardi and corresponding author for this article. "Using blood, or liquid biopsies, on the other hand, provides a homogeneous representation of cfDNA that is being shed from all types of [cells](#)."

The scientists note that short fragments of DNA and chemical modifications to those fragments, known as [methyl groups](#), help tell researchers what cell type the respective snippet of DNA came from because these methylation patterns are unique to specific cell types. By using cfDNA to compare damage to cells from various forms of treatment with undamaged normal cells from the same tissue, the researchers can analyze a treasure-trove of data about how the cells in a

tissue are affected by treatments and other external forces. This knowledge could be key in assessing if a therapy is effective and what adverse effects it may cause.

"Fine-tuning these applications of cfDNA analysis is challenging and requires in-depth approaches, both at the genome sequencing level and computationally," explains Megan Barefoot, a MD/Ph.D. student in the Wellstein lab at the Cancer Center and lead author of the article.

"Methylated cfDNA has opened a new and minimally invasive way to detect damage to cells in the body as there are often hundreds of methyl markers per cell that can mark, very specifically, where the cells came from, much like a barcode scanner at a grocery checkout tells the store the identity of a particular product. Combined biological and computational analyses make deciphering these methylation patterns/molecular barcodes possible so that researchers can trace the origins of cfDNA."

The end-result of these analyses aids investigators in determining the tissue of origin of a [cancer](#), for example, and also allows researchers, when comparing damaged cells to healthy cells, to see where the damage originated, especially if it was due to a certain type of treatment.

"This approach can be applied to any therapy that will impact tissue equilibrium by causing cells in tissues to become damaged and die, including chemotherapy, radiation, and immunotherapy. This review really helps set the stage for our future research efforts," concludes Wellstein. "My lab is very actively pursuing methods and technologies that further refine analyses of methylated cfDNA. We believe these efforts are affordable and will soon become standard in labs and they should make a difference in advancing the understanding and treatment of many cancers."

More information: Megan E. Barefoot et al, Detection of Cell Types

Contributing to Cancer From Circulating, Cell-Free Methylated DNA, *Frontiers in Genetics* (2021). [DOI: 10.3389/fgene.2021.671057](https://doi.org/10.3389/fgene.2021.671057)

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