



## TECHNOLOGY

# Treating Cancer with Specific DNA Secondary Structure Binding Compounds

## OVERVIEW

### Background

While the structure of DNA is most commonly associated with the double helix, research has shown that DNA can adopt many secondary structures for which the biological functions are still being determined. One secondary structure, the G-quadruplex, can form when the double helix unwinds during transcription. One common area for the formation of a G-quadruplex is at the end of DNA strands in a region called the telomere. Telomeres form at the end of DNA strands to protect them from degradation as a decrease in DNA strand length would eventually lead to cell death. Telomere length is maintained in the cell by an enzyme called telomerase, and over activity of telomerase is observed in ~85% of cancers. Recent research has shown that G-quadruplex formation in the telomere decreases telomerase activity, making stabilizing these formations a potential target for cancer treatment. G-quadruplex binding compounds are known, but many lack specificity for G-quadruplex formations and also bind duplex DNA structures, which can lead to increased toxicity during treatment. Therefore, a compound that specifically binds G-quadruplex formations could hold great potential as a new anti-cancer agent.

### Innovative technology

Researchers at the University of Maryland have discovered that a drug currently used to treat some forms of protozoan infection in animals binds to G-quadruplex formations. Through further testing, they were able to identify the structural components of the drug that were necessary to bind G-quadruplex formations. With this information, they synthesized derivatives that have increased specificity for G-quadruplex formations. These high specificity derivatives may prove to be useful anti-cancer agents.

## APPLICATIONS

- Cancer treatment

## ADVANTAGES

- Repurposed compound has known toxicity profile
- Potential new class of drugs for cancer treatment
- High specificity for G-quadruplex formations may decrease cellular toxicity

## CONTACT INFO

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## **Additional Information**

### **INSTITUTION**

University of Maryland, College Park

### **PATENT STATUS**

Pending

### **LICENSE STATUS**

Available for exclusive or non-exclusive license

### **CATEGORIES**

- Small molecules
- Repurpose Drug

### **EXTERNAL RESOURCES**

- [US Patent 10,130,625](#)

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