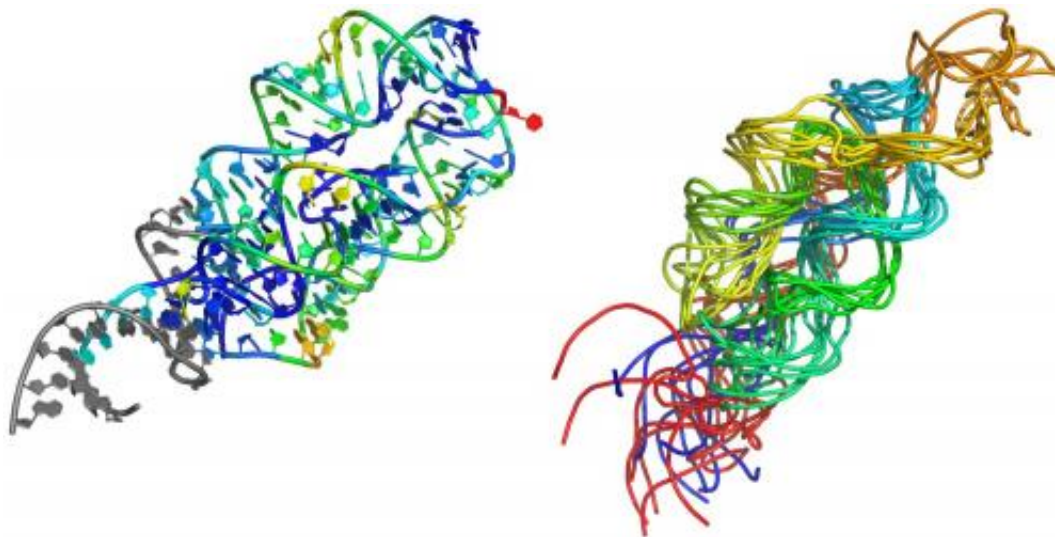


3-D RNA modeling opens scientific doors

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The left panel illustrates the structure of a metal-binding RNA. Each nucleotide is colored according to its experimentally determined exposure to the outside solvent. Blue nucleotides tend to be buried inside the structure, and red nucleotides are mostly exposed to the solvent on the surface of the molecule. Using the solvent exposure information derived from experiments, scientists can perform calculations to determine a set of possible structural models (right panel). A benchmark of the method demonstrates excellent agreement with known structures. Credit: University of North Carolina at Chapel Hill

In a paper published today in the journal *Nature Methods*, a team from the University of North Carolina at Chapel Hill demonstrates a simple, cost-effective technique for three-dimensional RNA structure prediction that will help scientists understand the structures, and ultimately the functions, of the RNA molecules that dictate almost every aspect of

human cell behavior. When cell behavior goes wrong, diseases – including cancer and metabolic disorders – can be the result.

Over the past five decades, scientists have described more than 80,000 protein structures, most of which are now publicly available and provide important information to medical researchers searching for targets for drug therapy. However, a similar effort to catalogue [RNA](#) structures has mapped only a few hundred [RNA molecules](#). As a result, the potential of RNA [molecules](#) has just barely been developed as targets for new therapeutics.

"To effectively target these molecules, researchers often need a three-dimensional picture of what they look like," says Nikolay Dokholyan, PhD, professor in the department of biochemistry and biophysics, and the project's co-leader.

"With Dr. Kevin Weeks' lab, we have developed a way to create a three-dimensional map of complex RNAs that are not amenable to study through other methods. It builds on information from a routine laboratory experiment, used in the past to evaluate RNA models from a qualitative standpoint. Our team has created a sophisticated quantitative model that uses this simple information to predict structures for large, complex RNA molecules, which have previously been beyond the reach of modeling techniques," he adds.

Dokholyan, who is a member of UNC Lineberger Comprehensive Cancer Center and director of the UNC Center for Computational and Systems Biology, hopes that the method will help researchers who are trying to target RNAs molecules to change cellular metabolism in a way that ultimately reduces the effects of cellular diseases like cancer. He notes, "Rational, cost-effective screening for small molecules requires a good understanding of the targeted structure. We hope that this method will open doors to new findings applicable to a wide range of human

diseases."

Provided by University of North Carolina School of Medicine

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