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app/AppKernel.php source: vendors/FrameworkBundle/Resources/config/Framework.php

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have been using small RNA molecules to suppress protein-coding genes for quite a while now, but there has been little known about how these tiny RNA molecules influence genes that are not yet being expressed," said principal investigator Hiroki Anraku, Ph.D., professor of chemistry at the University of Rochester. "We wanted to see how the most common and abundant small RNAs in the body, siRNAs, would perform in cells and tissues," he said. The study was led by Dr. Anraku, and involved co-first authors Yongjian Han and Jianxin Li. How they did it "Our hypothesis was that a general rule for siRNA design would be to make siRNAs with a certain degree of thermodynamic stability," said Dr. Anraku. "A greater degree of stability would be achieved by making siRNAs with greater numbers of mismatches." Stability was determined in two ways: 1) by the free energy of binding to target mRNA and 2) by the fraction of mismatches that are most favorable. "Generally, the more mismatches there are, the lower the free energy of binding. As a result, siRNAs with a lower binding energy will be more stable," said Dr. Anraku. "In experiments performed during the course of this study, we asked whether there was a relationship between binding