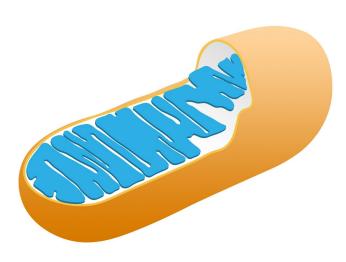


How dark matter of the genome interacts with mitochondria—and affects the fate of cancer

10 February 2021, by Delthia Ricks



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If an opinion poll were taken for the most popular component of mammalian cells, the result probably would be a 50-50 split: Half of respondents would likely vote hands down for the nucleus and the other half for the mitochondria—a decision that in no way diminishes or casts aspersions on the other hardworking constituents of cells.

Both the nucleus and mitochondria are critical membrane-bound organelles, cellular components with highly specialized roles. There are many other organelles, each with a specific role in the cell. The nucleus, only one per cell, is home to deoxyribonucleic acid—DNA— the spiraling master molecule of life. By contrast, there can be upward of 2,500 mitochondria per cell, which have their

own, more primitive DNA loop, encoded with instructions only for <u>energy production</u>. These bean-shaped fuel factories are the only organelles—other than the nucleus—to be endowed with their own unique genetic material.

In emerging genomics research, a growing cadre of scientists has begun tapping into the <u>organelle</u>'s connection to something far more mysterious: the so-called dark matter of the genome—long noncoding RNA—and the influence that these sequences have on <u>mitochondrial functions</u>. These RNA transcripts have an influence in health and disease, new studies are demonstrating.

Long non-coding RNAs get their name from their extraordinary lengths, most of which exceed 200 nucleotides. And as the moniker implies, this form of RNA doesn't code for proteins. Scientists have identified more than 50,000 non-coding RNAs of all lengths that are transcribed in the human genome. Most are 200 nucleotides long or longer and they exist between—or after— coding sequences.

Stunningly, these transcripts were once considered genomic junk, but increasingly they are being studied as the dark matter that holds sway over certain functions of multiple organelles, including mitochondria. These sequences of RNA possess enhancers and promoters that affect the expression of coding genes. Some activate or repress oncogenes, which means they play a role in certain cancers. Because these molecules have become the subject of intense research, new findings are helping to peel away much of the mystery about them. Long non-coding RNA—IncRNA—also possesses transposable elements—so-called jumping genes—which underscores how translocations and insertions into coding regions can and do occur.



In China, scientists who are conducting genomic research in the College of Life Sciences at Zhejiang as the Krebs cycle, but additionally is called the University in the city of Hangzhou, are among the global investigators exploring the association between mitochondrial functions and IncRNA (pronounced "linc" RNA).

A mitochondrion (singular) is a veritable energyproducing factory, providing copious amounts of adenosine triphosphate, ATP. Heart cells are chock In terms of cancer, GAS5, negatively correlates full of mitochondria because of the extraordinary energy demands required to pump blood. Unlike the nucleus, whose DNA is a spiraling helix, a mitochondrion has a single DNA loop, similar to bacterial DNA—and for good reason. Scientists theorize that 1.5 billion years ago mitochondria were primordial purple (non-sulfur) bacteria that insinuated themselves into primitive cells that were hungry for energy.

To shed light on the intimate relationship between mitochondrial function and IncRNA, the team also took stock of the indispensable roles that all organelles play in cells and the influences that this dark matter of the genome has on multiple organelles.

"Organelles are microscopic semi-organs involved in many <u>cellular processes</u>, including several important metabolic reactions, energy production, cellular signaling and cell growth," Dr. Linjie Sang and colleagues wrote in the journal Nature Metabolism.

It was from their research involving the mitochondrial association with IncRNA that Sang and colleagues made a startling discovery—they uncloaked the activity of a tumor suppressor molecule known as growth-arrest-specific 5, or more simply, GAS5. The team discovered its association with "mitochondrially localized IncRNA." GAS5 apparently helps maintain cellular energy homeostasis, the researchers found.

"Mechanistically, energy-stress-induced GAS5 modulates mitochondrial tricarboxylic acid flux," Sang wrote, noting this modulation occurs by disrupting metabolic enzyme sequences in the tricarboxylic acid cycle.

The tricarboxylic acid cycle is also famously known TCA and the citric acid cycle. Regardless of which name is used, the cycle is a series of chemical reactions that occur in all aerobic organisms to generate energy through the oxidation of acetate. The acetate comes from dietary fats, proteins and carbohydrates.

with levels of its associated mitochondrial metabolic enzymes in tumors and benefits the overall survival of individuals, particularly those with breast cancer, Sang and colleagues found.

"Our detailed annotation of subcellular IncRNA distribution identifies a functional role for IncRNAs in regulating cellular metabolic homeostasis, highlighting organelle-associated IncRNAs as potential clinical targets to manipulate cellular metabolism and diseases," the scientists concluded about their research.

While the findings illuminate a complex association between mitochondria and IncRNA, the research is likely just the beginning of discovering broader relationships between IncRNAs and mitochondria as well as IncRNAs and other organelles.

In addition to the nucleus and mitochondria, other major cellular organelles include the Golgi apparatus, which modifies and transports proteins. It is often referred to as the "post office" of cells. The endoplasmic reticulum is another organelle. It's involved in the synthesis, folding and transport of proteins. The lysosomes have the job of breaking down large molecules—macromolecules—into their component parts. Proteins, for example, are reduced to their individual amino acid components.

"Organelles use specialized molecules to regulate essential cellular processes," Sang explained, referring to IncRNA. However, systematically elucidating the sub-cellular distribution and function of IncRNAs in cellular homeostasis and human diseases, has not yet been fully realized, Sang emphasized.

The Zhejiang University scientists and their collaborators in the United States and elsewhere in



the world are also examining these long stretches of non-coding RNA in connection with other cellular organelles. They have identified a "diverse and abundant subcellular distribution" of organelles associated with IncRNAs that range from mitochondria, to lysosomes to the endoplasmic reticulum.

More information: Linjie Sang, et al. Mitochondrial long non-coding RNA GAS5 tunes TCA metabolism in response to nutrient stress, *Nature Metabolism* (2021) www.nature.com/articles/s42255-020-00325-z

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