Expression Quantitative Trait Locus Analysis Identifies Novel Genes for Statin Myopathy

Deepak Voora, MD

Study Hypothesis
Inhibitors of 3-hydroxy-3-methylglutaryl-CoA reductase (or statins) are the most common and effective medications used for the treatment and prevention of cardiovascular morbidity and mortality. Statins exert their beneficial effects primarily by inhibiting hepatic cholesterol synthesis. In general, statins are well-tolerated medications. However, a small but clinically significant number of patients taking statins develop statin myopathy, a range of clinical syndromes characterized by common muscle aches and pains to a rare but sometimes fatal condition of frank muscle necrosis called rhabdomyolysis. Although often described as benign, statin-induced side effects can significantly affect quality of life and are a frequent cause of premature drug discontinuation. As a consequence, millions do not receive the cardiovascular benefits of these life-saving drugs.

The reasons that some patients do (or do not) develop statin-induced side effects have been difficult to identify. Certain patient characteristics such as age, sex, and concomitant medications have been implicated in some. Genome-wide association studies have implicated specific genetic variants in the \( SLCO1B1 \) gene that underlie statin-induced side effects. However, many people without these genetic variants still go on to develop side effects, and many people who develop side effects have normal genotypes. Therefore, there are large gaps in our understanding surrounding not only the biology of statin-induced side effects but also useful biomarkers to identify individuals at highest risk for side effects. To overcome these gaps, Mangravite et al. combine these 2 concepts together to identify differential eQTLs or genetic loci that exert an influence on the magnitude or direction of gene expression changes in response to drug exposure. The underlying assumption is that genetically determined variation in drug responses can be revealed by differential changes in gene expression between those with different genotypes. Based on the known biology surrounding their top hit, the authors then tested for associations of their variants of interest with clinical responses to simvastatin therapy from 2 independent patient cohorts.

Principal Findings
In their gene expression and genetic data set of lymphoblastoid cell lines, 4590 genes with cis-eQTLs (defined as eQTLs within 1 MB of the transcription start/end site) were identified, and the expression of 5509 genes changed in response to simvastatin exposure. However, when the authors combined these analyses to identify transcripts that changed differently depending on genotype, they narrowed these large lists to only 6 loci that met criteria for a genome-wide significance level.

The top loci included a large group of variants near the \( GATM \) gene, and carriers of the minor allele had a decrease in \( GATM \) expression in response to simvastatin. Because of the known role of \( GATM \) in muscle physiology, the authors then hypothesized that these same variants may underlie the risk for statin-induced myopathy. Using data from 2 previous cohorts of cases of statin-induced myopathy, the authors confirmed that the same loci near \( GATM \) were protective for statin-induced myopathy in these patient cohorts.
Implications
The mechanisms by which statin induced reductions in GATM may alter the risk of statin-induced myopathy are not clear from this study. However, this work opens the possibility of using the differential eQTL approach in lymphoblastic cell lines as a general platform to understand the variability in drug responses. Beyond statins, one can envision testing entire panels of medications using banked cell lines to identify novel mechanisms that underlie variable drug responses or to compare drugs with different mechanisms for their effects on gene expression in different genetic backgrounds. By demonstrating the feasibility of this approach, this work should dramatically accelerate our understanding of the response to medications used to treat cardiovascular disease.

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Disclosures
None.

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