

# Tracking the 'social networks' of genes disrupted in complex diseases

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Credit: Valdo Peixoto and Daniel Marbach

Your personal risk of developing complex diseases such as diabetes, depression or cancer is influenced in part by genetic variants, that is, letters in your DNA sequence that differ between people. These variants disrupt networks of interacting genes in different tissues of your body, two studies published in *Nature Methods* and *PLoS Computational Biology* found. Innovative software tools allowed the scientists to construct accurate "maps" of gene networks for about 400 different human cell and tissue types, ranging from immune cells to brain tissues, whereas previous studies were limited to just one or few tissues. Each of these networks describes hundreds of thousands of regulatory interactions among thousands of genes, giving the first global view of the "control system" of diverse cells and tissues.

The team found that genetic variants disrupt components of these networks in disease-specific tissues, giving new insights on disease mechanisms which may lead to targeted treatments that are more effective and have fewer side-effects for the patient. The project was spearheaded by researchers from the newly

formed Department of Computational Biology (DCB) at the University of Lausanne (UNIL) and the SIB Swiss Institute of Bioinformatics, in collaboration with researchers from the University Hospital of Lausanne (CHUV) and the Broad Institute of MIT and Harvard.

Advances in genome-sequencing and related technologies have given rise to large studies that compare genetic variants between healthy people and people with a given condition. These studies have successfully identified thousands of genetic variants that are linked to different diseases. However, the mechanisms by which these variants influence disease processes remain mostly unknown, which is currently hindering progress to developing better diagnostic tests and personalized treatments for patients.

Daniel Marbach, David Lamparter and Prof. Sven Bergmann (SIB, UNIL), in collaboration with Prof. Zoltán Kutalik (SIB, CHUV, UNIL) and Prof. Manolis Kellis (MIT), have now mapped networks of interacting genes that are perturbed by disease variants with unprecedented resolution across hundreds of human cell types and tissues.

## Poorly understood regulatory regions

"The challenge is that over 90% of disease variants lie outside of genes, in regions of the genome that are still poorly understood" Marbach says. "These regions can have regulatory functions, which are sometimes disrupted by genetic variants. Things get even more complicated as the regulatory relationships may vary between different tissue types. For example, a certain gene may activate another one in the liver, but not in the heart."

The team thus tackled the ambitious task of creating accurate "maps" of the regulatory networks that control the activity of genes in a given [tissue](#). Data from an international research consortium (FANTOM) coupled with novel analysis techniques

allowed them to create the largest collection of such perturbations across complex diseases, *Nature* networks to date, describing the regulatory interactions among over 19,000 genes in close to 400 human cell types and tissues. The study thus gives the first comprehensive view of the regulatory systems across a broad range of cells and tissues. *Methods* (2016). [DOI: 10.1038/nmeth.3799](https://doi.org/10.1038/nmeth.3799)

Provided by Swiss Institute of Bioinformatics

### **Taking inspiration from social networks**

The underlying hypothesis was that genetic variants may impact genes that are connected within regulatory networks of tissues that are specific to certain diseases. To test their hypothesis, the researchers employed techniques similar to those applied to social networks to gain information about users on the basis of their interconnections. But in this case, they used biological networks to gain information about genes that can lead to diseases.

In a large study including genetic data for diverse neurodegenerative, psychiatric, immune-related, cardiovascular and metabolic disorders, the researchers found that disease variants often affect groups of genes that were densely interconnected within regulatory networks, confirming their hypothesis. Moreover, these affected [network](#) components pinpointed with remarkable precision cell types or tissues that are implicated in disease processes. "For example, people with schizophrenia were found to have genetic variants that perturb interacting genes in brain tissues that are responsible for cognitive and emotional behavior, while genetic variants associated with obesity impact [genes](#) that interact in tissues of the intestinal system" Marbach says.

### **Towards better targeted treatments**

"Our work shows that accurate maps of gene networks for different tissues will be of tremendous value to advance our understanding of how diseases start and progress, which is essential to design targeted treatments and to identify patient groups that respond to these treatments in a personalized medicine setting," concludes Prof. Bergmann.

**More information:** Daniel Marbach et al. Tissue-specific regulatory circuits reveal variable modular

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