Scientists Create Atlas of the Human Genome Using Stem Cells
Embryonic stem cells are a unique resource as they can turn into any adult cell in our bodies. Their versatile nature puts them at the centre of attention in the fields of regenerative medicine, disease modelling and drug discovery.

In parallel to the discovery of human embryonic stem cells, another milestone in biology was completed with the sequencing of the human genome, and the identification of the entire set of genes responsible for our genetic identity. This finding has led to a new challenge of understanding the function of the genes in the human genome.

Now, scientists have generated an atlas of the human genome that illuminates the roles our genes play in health and disease.

Researchers from the Hebrew University of Jerusalem analysed virtually all human genes in the human genome by generating more than 180,000 distinct mutations. To produce such a vast array of mutations, they combined CRISPR-Cas9 screening with a new type of embryonic stem cells that was recently isolated by the same research group.

This new type of stem cells harbours only a single copy of the human genome, instead of two copies from the mother and father, making gene editing easier thanks to the need of mutating only one copy for each gene.
Scientists Have Discovered a New DNA Structure Inside Human Cells

Forget about the double helix – now, researchers have identified the existence of a new DNA structure that could play a crucial role in how DNA is expressed.

Their research, published in *Nature Cell Biology*, shows that a mere 9% of all the genes in the human genome are essential for the growth and survival of human embryonic stem cells, whereas 5% of them actually limit the growth of these cells.

The researchers were also able to analyse the role of genes responsible for all hereditary disorders in early human development and growth, showing how cancer-causing genes could affect the growth of the human embryo.

“This gene atlas enables a new functional view on how we study the human genome and provides a tool that will change the fashion by which we analyse and treat
cancer and genetic disorders,” said professor Nissim Benvenisty, MD, PhD, Director of the Azrieli Center for Stem Cells and Genetic Research and senior author of the study.

Another key finding of the study was the identification of a small group of genes that are uniquely essential for the survival of human embryonic stem cells but not to other cell types. These genes are thought to maintain the identity of embryonic stem cells and prevent them from becoming cancerous or turning into adult cell types.

“This study creates a new framework for the understanding of what it means to be an embryonic stem cell at the genetic level,” said Dr. Atilgan Yilmaz, PhD, postdoctoral fellow and a lead author on the paper.

“The more complete a picture we have of the nature of these cells, the better chances we have for successful therapies in the clinic.”
MORE FROM FRONT LINE GENOMICS

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Japanese Pharmaceuticals to Leverage Thermo Fisher’s Oncomine Dx Target Test
Thermo Fisher Scientific has signed agreements with Takeda Pharmaceuticals and Daiichi Sanyo that will enable the Japanese firms to leverage Thermo Fisher’s FDA approved Oncomine Dx Target Test for their clinical trials and drug development programs.

NEWS

Scripps Translational Science Institute Awarded $34 Million in NIH Funding
The Scripps Translational Science Institute has received over $34 million in renewed funding from the National Institutes of Health’s National Center for Advancing Translational Sciences to advance medical research and clinical care through genomic and digital technologies.

NEWS

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Researchers have identified a protein critical for the aggressiveness of T-cell leukaemia, a subtype of leukaemia that afflicts children and adults.

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A new method for diagnosing breast cancer involves a pill, which could do a better job distinguishing between benign and aggressive tumours.