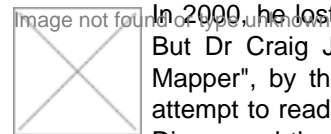


## Reader of the 'Book of Life'

10 April 2003 | News



In 2000, he lost narrowly to the then US president-elect, George W Bush, the title "The Time Man of the Year." But Dr Craig J Venter's influence will go beyond the immediate happenings of our life. For, called "Gene Mapper", by the Time magazine, his scientific work in completing the mapping of the human genome is an attempt to read the secrets of the humanity itself. This 56-year-old biochemist from University of California, San Diego, and then the chief of Celera Genomics, may have shared the fame with a government scientist Francis

Collins when they shared the podium at the White House in June 2000 to announce their simultaneous mapping of the human genome. But the world knew it was Venter's "shotgun sequencing method" that made it all possible. One of the most cited biologists of modern times, Craig J Venter was in New Delhi in March to speak at the Millennium Summit -III organized by ASSOCHAM. In an interview with Executive Editor N Suresh, Venter unraveled some of the secrets of human genome

### " You are not what your genes are"

#### **What is the progress in the identification of the functions of human genes?**

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For each species, for each set of genes, there is a precise set of environmental conditions that impact them. You are certainly not what your genes are. When we are studying the genetic code, we are only studying at best half the question. Humans

have 100 trillion cells and they work together in mass with genes. So it will be some time before we even understand the basic functions, let alone how all of them interact effectively with the environment.

### **Where does this leave the theory of genetic determinism which has been popular for decades?**

Each human has roughly two to three million of genetic code that are different from another person. But most of these codes probably have no biological significance whatsoever. They may have just some forensic and tracking applications, and maybe useful as mapping tools. Less than a percent of these codes occur in genes or regulatory regions. This means that at the biological level, at the gene level, we are all virtually identical twins. This clearly means that your life not determined by your genes. The fraction that is really different in a biological and meaningful way is such a tiny percentage it is stunning. The smaller number of genes and the similarity of human genes to that of a mouse hopefully put the nail in the coffin of the genetic determinants.

### **What is the next big thing after the genome mapping?**

We have some 100 trillion cells in each of our bodies. This all interact with each other in many complex ways and produce a variety of proteins. With the information about the human gene sequence, it is possible now to measure these proteins. Because we can get at every protein in the genome. For the known useful genes of around 50,000 there are some 200,000 to one million proteins. We are building a large-scale protein facility to do roughly one million protein sequences per day. We are looking at using these proteins as markers for specific diseases.

The genome is important because with mass spectroscopy sequencing, the proteins get blown apart into small fragments and we can compare those sequences back with databases. Till now most of these did not match anything on the databases and so interpretation was difficult. Now every one of these will have a match and we can rapidly determine the sequence of the proteins in the cells and in the blood.

### **Does genomic information provide clues to our past?**

Many scientists believe that evolution is adding on genetic information and adding on complexity. But we find that most human pathogens probably started from a much more complex organism and threw out genetic material during evolution. We tried to find out whether we could come out with a molecular definition of life. To test this we knocked out 200 or so genes from *Mycoplasma pneumoniae* and observed whether we still have a living organism. And with *Mycoplasma genitalium* (a small microbe found in the human genital area) we asked the question whether all its genes were necessary for life. And we found that the 200 extra genes in *pneumoniae* were completely dispensable and about 200 genes in *genitalium* too appeared dispensable.

We found out three stunning things. First of the 300-odd genes, 103 were completely new to science. We think we know a lot about biology, yet in this most minimal cell, we had no idea about what one third of the genes do, except that if we knock them out, the cell dies.

Two, we could not come up with a molecular definition of life. We found that life is context-sensitive, that is, the environment that the cell is in is equally important to any component of the genetic code. Third, the Darwinian evolution wasn't just random errors in the genetic code. With *haemophilus* and every pathogen we have worked on, we found there was preprogramming in the genetic code to cause change in the structure of specific genes. Essentially a pathogen fools the immune system of its host. And each pathogen has a different way of doing it.

Essentially, the human genome contains a molecular clock as repeated sequences of letters between genes and mutates over time. Measuring the rate of these changes would some day allow us to predict the time at which it happened. This may help to pinpoint the epoch in which the traits that make us unique humans emerged.

### **You have been talking about the development of personalized medicine. What are these?**

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For the first time, we can look at this genetic variation by chromosome. For example, you can discover genetic variation in the genome of individuals who have an increased risk of myocardial infarction. Scientists are using this information to find linkage to disease. The pharma industry is using this to find ways to improve clinical trials and drug effects. This leads to the development of personalized medicine.

Recently, there was a type II diabetes drug that had to be taken off the shelves because one out of 10,000 people had a severe liver toxicity to this drug. If we can find simple tests that predict toxicity, it will have a huge impact. Most drugs work only on 30 to 50 percent of the population but are given to everybody because in theory they are relatively safe and doctors

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We have to look closely at our own evolution and that of vertebrates. From the simple drosophila (fruit fly) to humans, the evolution has taken place in about 600 million years. And just four to five gene groups only expanded in this period. For example, humans have an immune system, while the fruit fly doesn't have a significant immune system. All the genes associated with our immune system expanded during this period. When we compare with homeostasis, all things associated with our vascular system expanded during this period. But we have virtually identical gene set with mice and other vertebrates. The key to our uniqueness are things like the transcription pattern in gene regulation, that turn on different sets of response to environmental conditions. Among all human chromosome, the chromosome 19 has the largest number of genes.

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### **In which areas of biotechnology should India invest?**

The best investment is to improve the quality of education and research work. Keep Inder Verma (a well known geneticist at Salk Institute, USA) in India itself.

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