

# The human genome at ten

Nearly a decade on from the completion of the draft sequence of the human genome, researchers should work with the same intensity and focus to apply the results to health.

**T**he race to complete the first human genome sequence had everything a story needs to keep its audience enthralled — right down to a neck-and-neck sprint for the finish by two fierce rivals. In the end, the result was basically a tie. The rivals — the international, publicly funded Human Genome Project and the private, for-profit company Celera Genomics then based in Rockville, Maryland — jointly announced the completion of their draft sequences in June 2000 at a gala televised press conference attended by US President Bill Clinton and UK Prime Minister Tony Blair.

The White House press statement articulated the hope, felt by many, that this landmark achievement would “lead to a new era of molecular medicine, an era that will bring new ways to prevent, diagnose, treat and cure disease”.

This issue of *Nature* takes a look at the next chapter in the story — the first post-genome decade — then asks how the tale might unfold in the years to come.

For many scientists, the chronicle of that first decade is an intensely personal one. Not only were they inspired by this example of what researchers can do as group, but they found that the availability of the sequence shaped their lives and their research in ways they could not have predicted. On page 668, we share the experiences of some of those in thick of the race. But we also want to hear about yours: we would like you to take part in *Nature's* brief survey about the genome's impact at [www.nature.com/humangenomesurvey](http://www.nature.com/humangenomesurvey). The results will be published later this year.

The first post-genome decade saw spectacular advances in science. The success of the original genome project inspired many other ‘big biology’ efforts — notably the International HapMap Project, which charted the points at which human genomes commonly differ, and the Encyclopedia of DNA Elements (ENCODE), which aims to identify every functional element in the human genome. Dramatic leaps in sequencing technology and a precipitous drop in costs have helped generate torrents of genetic data, including more than two dozen published human genomes and close to 200 unpublished

ones (see page 670). Along the way, geneticists have discovered that such basic concepts as ‘gene’ and ‘gene regulation’ are far more complex than they ever imagined (see page 664).

But for all the intellectual ferment of the past decade, has human health truly benefited from the sequencing of the human genome? A startlingly honest response can be found on pages 674 and 676, where the leaders of the public and private efforts, Francis Collins and Craig Venter, both say ‘not much’. Granted, there has been some progress, in the form of drugs targeted against specific genetic defects identified in a few types of cancer, for example, and in some rare

inherited disorders. But the complexity of post-genome biology has dashed early hopes that this trickle of therapies would rapidly become a flood. Witness the multitude of association studies that aimed to find connections between common genetic variants and common diseases, with only limited success, or the discovery that most cancers have their own unique genetic characteristics, making widely applicable therapies hard to find.

Biotechnology companies, including Celera, deCODE Genetics in Reykjavik, Iceland, and Human Genome Sciences of Rockville, have had to rethink their optimistic assumption that selling human genetic information could turn a profit. Excitement over start-up companies offering personal genetic testing has withered just as fast, as it has become clear that their predictions have little actionable value (see page 680).

This gap between basic research and clinical application has not escaped the notice of funding agencies, many of which are now investing serious money in a bid to close it. The US National Institutes of Health, for example, has established a string of major clinical and translational science centres over the past few years, and in February it established a joint council with the Food and Drug Administration aimed at promoting translation. Similar efforts are under way elsewhere, such as those being rolled out by the UK Medical Research Council.

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## WHAT DO YOU THINK?

**What did the human genome sequence mean to you? Take part in *Nature's* online survey**  
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It is not yet clear whether such efforts will be enough for genomics. Given the biological complexities involved, applying knowledge of the human genome to health may well require a community-wide effort as determined and as systematic as was the project to sequence it in the first place.

That effort would need to solve a long-standing mismatch: the rapidly increasing ease of gathering genomic data versus the continuing difficulty of establishing what the genetic elements actually do. One intriguing experiment in high-throughput functional analysis appears on page 721, where the authors detail how they systematically disabled each of the 21,000 protein-coding genes in human cells and then captured and processed microscope images of the resulting cellular behaviour.

The effort would also require even more imaginative ways to visualize and draw meaning from the flood of genomic and molecular data (see page 678). It would require interdisciplinary teams that can provide know-how not just in research, but in the issues such as intellectual property, informed consent, finance and regulatory bureaucracy that are needed to keep discoveries moving through the agonies of clinical development.

Perhaps it would even require an explicit challenge to foster fervour and competition. For example, the X-Prize Foundation of Playa Vista, California, is offering a prize of US\$10 million to the first group to sequence 100 human genomes in ten days. Why shouldn't some organization offer a reward and kudos for, say, the next genetics-based cancer drug to go from basic genomic data to approved therapy within ten years? True, real-world therapies are unlikely to have metrics as clear as those in genome sequencing, in which progress can be measured in base pairs read per day and there is a well-defined end point. But the need is no less urgent, and the collective will to reach such goals should be no less intense.

More than anything, the race to sequence the human genome proved that researchers comfortable in their individual pursuits are capable of incredible cohesion, focus and breathtaking speed. They rewrote the research rule-book, broke with the conventions of individual academic goals and left the public with the sense that anything is scientifically possible. The ten years since have brought astounding technological and intellectual advances. But ten years from now, when the story of the genome's first two decades is being told, it should include equally astounding applications to human health. ■

## A new row to hoe

The time is right to revitalize US agricultural research.

In nations where food is plentiful, it is easy to take that abundance for granted. In the United States, for example — a country rich in corn fields and pasturelands, and where shops overflow with cheap produce — agricultural research has languished for years under comparatively low budgets and disorganized funding priorities. In the 2009 economic stimulus bill, for example, the National Science Foundation received a \$3-billion boost and the National Institutes of Health got \$10 billion — but the Department of Agriculture's internal research programme was allocated just \$176 million, all of which was restricted to improving facilities.

It is heartening, then, that the administration of President Barack Obama has begun a much-needed overhaul of the nation's agricultural-research programme. The groundwork was laid in June 2008, when US Congress mandated the creation of a National Institute of Food and Agriculture (NIFA) within the agriculture department. Under the leadership of Roger Beachy, whom Obama appointed in September last year, the new agency has taken over, expanded and revitalized the department's long-standing competitive grants programme, now called the Agriculture and Food Research Initiative.

Last week, NIFA released its eagerly anticipated first call for grant proposals, and the changes are indeed dramatic. In striking contrast to the smaller, two-year individual grants that were the mainstay of the programme, NIFA's offerings now include a series of five-year 'coordinated agricultural projects' of up to \$45 million for collaborative, interdisciplinary research. NIFA has defined its funding priorities much more clearly than its predecessor did, and it has aligned them with a series of 'societal challenges' that include climate change, bioenergy

and childhood obesity. The agency has also declared that the success of the programmes will be measured not just by scientific publications, but by real progress towards solving these challenges, such as reducing the amount of energy, nitrogen fertilizer and water used in agriculture by 10% by 2030. It has also introduced new fellowship and outreach programmes in an effort to stimulate the dwindling pipeline of young talent entering the field.

The overall intent, says Beachy, is to raise the status of the nation's agricultural research, hopefully drawing attention and support from the nation's lawmakers in the process. With just \$262 million available for the first round of grants, the Agriculture and Food Research Initiative is still financially constrained compared with other US science agencies. But the structure of its competitive grants programme is an important step towards maximizing the impact of the funds that it does have.

The transition has not pleased everyone. News of Beachy's appointment caused a stir among opponents of genetically modified crops, who noted that he has voiced support for such crops in the past. Beachy has maintained that NIFA will also support small farms and traditional crop-breeding programmes, and the first call for proposals does seem to be doing this. But the strength of that support will become clear only after the grants have been made.

Meanwhile, many long-time agricultural researchers are feeling unsettled by the abruptness of the changes. Some are dismayed to find that their favourite funding programmes have been cut. Others find the new structure too prescriptive, and worry that it leaves little room for creative approaches. These are legitimate concerns, and NIFA should follow through with its promise to solicit input from the community before it crafts next year's proposals.

Nevertheless, the community should seize this opportunity to tackle big problems. Growing pains are inevitable, but the shake-up has the potential to rejuvenate the field at a time when its talents are desperately needed. ■