

# ScienceScope

**Reaching Out** The recent India-Pakistan summit may have ended in a diplomatic stalemate, but it did produce new joint research and training opportunities that the Indian government hopes will provide a back channel for improving relations between the two nuclear neighbors. Prime Minister Atal Bihari Vajpayee earlier this month unveiled a plan to award 20 scholarships to Pakistani students to attend Indian technical institutions and invited Pakistani academics to visit "as guests of the government." The focus would be on education, health, environment, and gender equity.

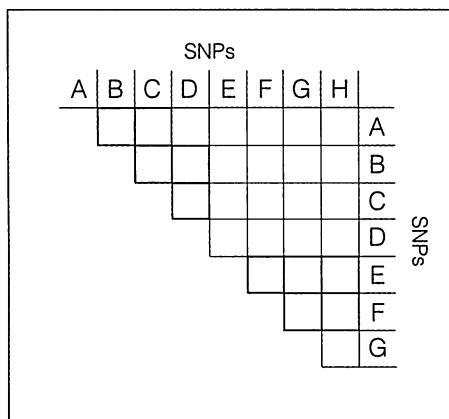
"There is an enormous amount of misinformation about India in Pakistan," says Pavagada Venkata Indresan, former president of the Indian National Academy of Engineering. "But once young minds can be trained at India's top technical institutions, a more positive message would certainly go out." Pakistan reacted cautiously, with a government spokesperson calling the idea "peripheral" to diplomatic efforts.

**Bioprospecting Under the Microscope** The National Park Service is starting a court-ordered environmental study of revenue-sharing agreements with firms that make profitable discoveries in national parks. Officials plan to kick off the bioprospecting assessment next month after getting public comment on the issues the study should address.

After Yellowstone National Park signed a 1997 profit-sharing deal with a San Diego biotech, bioprospecting opponents sued, charging that commercial activities in the parks violated federal law (*Science*, 13 March 1998, p. 1624). A federal judge suspended the deal earlier this year, ruling that Yellowstone could make such agreements, but that it first had to study their environmental impacts. In a 25 June *Federal Register* notice, the service said it would study bioprospecting impacts at all 384 of its parks in order to formulate consistent policy.

Critics want the study to take a careful look. By allowing exploitation, "these types of agreements fundamentally change the spirit of what the parks are about," says Joseph Mendelson III of the International Center for Technology Assessment, a Washington-based group that participated in the earlier legal challenge. Comments on the study's scope are due 10 August, with a draft expected in the fall.

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**Identifying blocks.** Hypothetical SNPs A through D are correlated (red) and probably constitute a haplotype. SNPs E through G are also fellow travelers.

Only in the past year or so have genome researchers realized that a haplotype map might be feasible. Until now, they have focused mainly on identifying DNA variations called single-nucleotide polymorphisms (SNPs)—sites along the genome at which individuals differ by just one base—to use in tracking disease genes. And although they've had some success recently in identifying genes associated with diabetes and the gastrointestinal ailment Crohn's disease, it's been an arduous, expensive process (see p. 593).

Because there's roughly one SNP for every 1000 bases of DNA, there might be a huge number of SNP patterns over a given stretch of sequence. If, for example, a 50,000-base sequence contains 50 SNPs, those SNPs could, in theory, come in  $2^{50}$  different variations. Computer simulations suggested that haplotype blocks—DNA stretches containing the same SNP pattern—would stretch only 3000 to 8000 bases—too short to make a haplotype map worth the trouble for tracking disease genes. But the reality turned out to be much more promising.

As several teams reported at the meeting, haplotype blocks are at least 10 times longer than predicted, and there are a relatively small number at each chromosomal position. For some sequences of 50,000 bases, for example, just four or five patterns of SNPs—that is, four or five different haplotypes—might account for 80% or 90% of the population. "It didn't have to be this way," said Eric Lander of the Whitehead Institute for Biomedical Research/MIT Center for Genome Research in Cambridge, Massachusetts. But because it is, instead of trying to correlate each of the 50 SNPs with disease, researchers can restrict their studies to SNPs that differentiate the few common patterns. This should ease their work by cutting down on the amount of DNA they have to scan to identify disease genes.

No one is sure yet why the genome is so blocky, but geneticists mention two candi-

date explanations. During the meiotic divisions that give rise to sperm and eggs, the two copies of each chromosome sometimes swap stretches of DNA, or recombine. If, for some unknown molecular reason, some parts of the chromosome are less likely to recombine than others, some stretches of DNA will be conserved as blocks while others change rapidly across generations.

Population bottlenecks apparently contribute as well. As Kenneth Kidd of Yale University and others have found, there is a greater diversity of haplotypes in people in Africa, where humans first arose, than in other populations. In addition, descendants of people who settled in Asia carry a somewhat different set of haplotypes from those who settled Europe.

And that's what raises the ethical issues discussed at the meeting. Researchers are still figuring out how to construct the haplotype map—what pilot studies to run, for instance, and how to standardize the definition of a haplotype. ("This notion of a block is a little hazy," said Leonid Kruglyak of the Fred Hutchinson Cancer Research Center in Seattle to much laughter.) But their most pressing problem may be whether to include ethnic or geographic identifiers on DNA samples. Such identifiers were stripped from the samples used to create the human genome sequence and SNP maps. But as Collins points out, if certain haplotypes are more common in some ethnic groups than others, haplotype mappers run the risk of missing distinctive patterns of DNA that might predict disease susceptibility in some populations but not others.

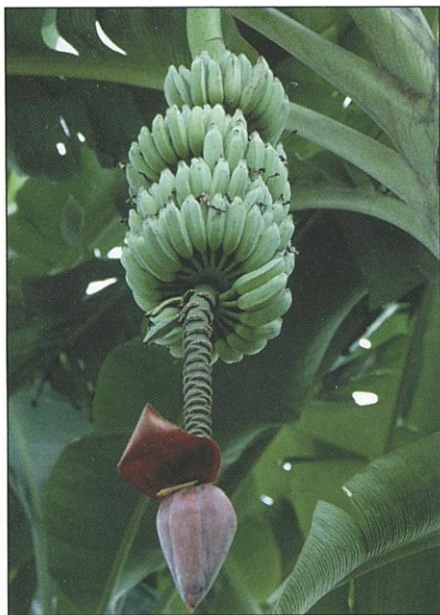
Summing up the meeting, Collins said that there was a "consensus" that the project "would have considerable medical value" and is worth pursuing. He solicited volunteers for two working groups that in the upcoming weeks will do the heavy lifting—a scientific steering committee to nail down working definitions of haplotypes and set priorities for pilot studies, and a second group to keep an eye on social and ethical issues.

—LAURA HELMUTH

## GENOME RESEARCH

### DNA Sequencers To Go Bananas?

Among scientists, the banana gets little respect. It's one of the most popular fruits on Earth and the developing world's fourth most important food crop, yet only a handful of labs are working on it. Now, a group of researchers is hoping to put the banana (*Musa*) on the scientific map. On 19 July, an international consortium announced that it hopes to sequence the entire banana genome, perhaps as early as 2006. The announcement of the multimillion-dollar effort



**To be revised.** International group seeks disease-resistant banana genes.

has already raised the banana's public profile, but some plant scientists say the project won't bear much scientific fruit.

The Global *Musa* (Banana) Genomics Consortium says it expects the project to take anywhere between 5 and 10 years, depending on the degree of accuracy desired, and cost up to \$7 million a year. Members of the consortium have promised \$2 million a year for the duration of the project, says geneticist Emile Frison, chair of the nonprofit International Network for the Improvement of Banana and Plantain and coordinator of the sequencing effort. The chief sponsor to date is the French agricultural institute CIRAD, which has pledged \$1.5 million over the next 5 years.

The consortium has already begun work but is still looking for funding. To help with fund raising, it engaged a public relations firm and announced the project's launch last week, noting that better bananas would help reduce hunger and poverty in the developing world. Frison explains that it would have been "a handicap in raising resources" not to have sought publicity.

Two dozen research centers expect to participate in the project, including The Institute for Genomic Research, a large-scale sequencing facility in Rockville, Maryland. Sequencing pros and banana experts met in Alexandria, Virginia, from 17 to 19 July to draw up a protocol and select a banana type for sequencing. Only specialists are likely to be familiar with their choice—*Musa acuminata calcutta 4*, a wild, nonedible variety native to southern India. According to Frison, *calcutta 4* was chosen in part because of its resistance to black sigatoka, a devastating fungus that reduces worldwide banana yields by as much as 50%. Network re-

searchers, who plan to make sequence data freely available as they are produced, say they hope the variety's resistance traits can be used to improve commercial bananas.

The banana would be the third major plant genome sequenced, after the wild mustard *Arabidopsis*, completed by a public consortium in 2000, and rice, undertaken by a private company and a public consortium. With 500 million to 600 million base pairs, the banana would be the biggest plant genome cracked yet.

But some members of the plant genomics community are skeptical of the project. They say bananas are intractable in the laboratory, which will make it hard to take advantage of the genomic data. "You can count the number of people working on banana on one hand," says Chris Somerville, a pioneer in *Arabidopsis* research who works at Stanford University. "The sequence is of limited utility if no one is working on the plant."

Backers of the project acknowledge that the ranks of banana geneticists are thin but point to recent accomplishments such as the identification of black sigatoka resistance genes. They argue that the banana is a very important crop even though it is not ideal for genetic research. "Just because something you sequence isn't a perfect model organism doesn't mean that its genome isn't useful," says University of Chicago plant geneticist Daphne Preuss.

—JOSH GEWOLB

## NEW FACILITIES

### Funding Backlog at NSF Sets Off Free-for-All

Elbowing your way to the head of a line that has been forming for years isn't polite. This year, however, it could be a winning strategy for several groups seeking major new research facilities from the National Science Foundation (NSF). The latest evidence came earlier this month in separate votes by both House and Senate spending panels on increasing NSF's \$4.4 billion budget.

The panels approved overall increases for NSF in 2002 of 9.4% and 5.5%, respectively—more than the 1.5% the Bush Administration requested but less than the 15% sought by research advocates. Those middle-of-the-road numbers hide some controversial decisions on individual projects, however. The Senate panel included funds for an underground laboratory in South Dakota, whereas the House

panel approved money for an expanded search for neutrinos at the South Pole and a high-altitude research plane. None of these projects were included in NSF's budget request to Congress. Although Congress must still reconcile those bills this fall, the trend is clear: Pushiness pays off.

NSF's budget has traditionally been nearly free of congressional earmarks—projects not requested by the agency and often advanced by politicians on the urgings of their constituents. But a decision in February by the Bush Administration to eliminate any so-called "new starts" from the foundation's capital budget, combined with congressional rejection of two projects contained in last year's budget request, has created a backlog of pent-up demand. That delay has prompted some researchers whose projects are stuck in NSF's pipeline to plead their cases to Congress.

The result is undermining the orderly process for setting priorities on big projects. The process begins with scientific reviews, runs through approval by the National Science Board (NSB), NSF's governing body, and culminates with a decision to include an item in the agency's budget request to Congress. Any major deviation, say most scientists, invites chaos and a misallocation of limited resources. "NSF [decision-makers] can see the big picture, and they set their funding priorities based on what's best for the entire community," says Anne Meltzer, a seismologist at Lehigh University in Bethlehem, Pennsylvania. She's involved with one of the delayed initiatives, a 10-year, \$350 million geosciences project called EarthScope that includes a network of seismic monitoring stations and a 4-kilometer hole drilled into the San Andreas fault. Although "it's all good research," says Meltzer about the projects Congress has favored, "it's very disappointing when people don't respect the process."

The science board hasn't even been briefed on one of the major projects that Congress wants the agency to take on: building a national underground laboratory in an old gold-mine shaft in South Dakota (*Science*, 15 June, p. 1979). Last week, the Senate panel that funds NSF included \$10 million to examine the feasibility of the \$280 million project, which was submitted to NSF for scientific review only last month. Propo-



**Cold cash.** Congressional funding may help scientists working on a neutrino detector at the South Pole greatly expand their experiment.

CREDITS: (TOP TO BOTTOM) CGIAR, AMANDA