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elections set for January, some analysts are predicting that he may agree to a power-sharing alliance with opposition leader Benazir Bhutto in which he remains head of state but gives up his role as head of the army, with Bhutto assuming the role of prime minister (for the third time).

Such a scenario could have uncertain consequences for science and education. Bhutto's previous stint as prime minister (1993–96) was characterized by low investment in research and higher education. She was also much criticized by the scientific community for failing to

appoint a minister for science.

"For 20 years, Pakistani universities have been quite indifferent to national politics."

Leaders of Pakistan's scientific community are meeting this week at the Academy of Sciences for the Developing World in Trieste, Italy, for a seminar originally designed

to showcase the country's progress in science, technology and higher education. Much of the meeting is now likely to be devoted to speculation on what will happen after the election.

Among the speakers will be Atta-ur-Rahman, chair of Pakistan's Higher Education Commission and a professor of chemistry at the University of Karachi. He told *Nature* that an incoming government had a responsibility not to throw away the past: "I hope the new government will continue on the present path. We have made huge advances in the past several years and it would be a great shame if this were lost."

Ehsan Masood

strategy is that they argued it at the individual level. It certainly drove down the estimates of how much this was going to cost them," says Garret FitzGerald, chair of pharmacology at the University of Pennsylvania in Philadelphia, whose mechanistic studies in humans and mice first pointed to the cardiovascular risks of Vioxx and other drugs in its class.

The Vioxx cases are in stark contrast to the lawsuits brought by people who took Fen-Phen, a popular diet-drug combination withdrawn in 1997. The signature, and relatively uncommon, heart-valve damage caused in those cases left drug-maker Wyeth liable for damages of more than \$21 billion.

Indeed, for scientists and clinicians, the main message of the settlement may be that reliable, personalized lawsuit victories remain as elusive as personalized medicine itself, and for many similar reasons.

Meredith Wadman

HIV vaccine may raise risk

An HIV vaccine made by Merck that failed in clinical trials may have increased the susceptibility of some trial participants to the AIDS virus, researchers reported last week. The findings have left scientists grappling with the problem of how to handle future trials of vaccines that use similar strategies to stimulate an immune response.

The trial was suspended in September, when preliminary analyses showed that the vaccine failed to protect participants against HIV (see *Nature* 449, 390; 2007). The vaccine consisted of three HIV genes carried by a weakened viral vector. Neither the HIV genes nor the vector, made from a common-cold virus called adenovirus 5, were capable of causing infection.

But it seems that trial participants who had a pre-existing immunity to adenovirus 5 were more likely to become HIV-positive if they received the vaccine. Among male volunteers who had high levels of antibodies against the adenovirus, 21 of 392 became infected in the vaccinated group, but only 9 of 386 were infected in the placebo group. The vaccine did not affect infection rates in men with no adenovirus-5 immunity.

The trial was not designed to examine the relationship between adenovirus-5 immunity and HIV infection, and researchers cannot say whether the result is statistically significant. But they can't ignore the findings, says Anthony Fauci, director of the National Institute of Allergy and Infectious Diseases in Bethesda, Maryland. The start of another HIV-vaccine trial has been pushed back several months to allow more time for scientific review, he says.

For now, researchers are trying to determine whether the vector is to blame or whether those with high adenovirus-5 immunity are at risk for unrelated reasons. Immunologists say it is also theoretically possible that vaccination in those with pre-existing immunity may have triggered an immune response that transiently boosted production of activated CD4 cells — the very immune cells that HIV infects.

That theory leaves several questions unanswered. For example, it is not known how long after vaccination the activated CD4 cells would have remained in the mucosal surfaces where HIV first gains entry, says Fauci.

Researchers are nevertheless

concerned. One possible solution is to limit future trials of adenovirus-5-based vaccines to participants with very low levels of immunity to the vector, says Gary Nabel, director of the Vaccine Research Center at the National Institutes of Health in Bethesda, Maryland. But doing so would close off the trials to a large proportion of the population they aim one day to help: more than 80% of sub-Saharan Africans have significant levels of adenovirus-5 immunity. In addition to the planned HIV-vaccine trial, an Ebola vaccine in development also relies on an adenovirus-5 vector.

But at this stage in vaccine development, researchers are just looking for a proof of concept rather than a vaccine that can be put directly to use, says Lawrence Corey, lead investigator of the HIV Vaccine Trials Network. Alternatives to adenovirus 5 are being developed, and may be substituted once the efficacy of a vaccination strategy has been shown. Nevertheless, vaccine developers will face challenging decisions in light of the new data, he says: "Translating uncertain data into policy is always a difficult issue."

Heidi Ledford

Hopes for a future vaccine against the devastating Ebola virus may have been set back by the results of a clinical trial of an HIV vaccine.

J.-M. BOUJ/AF