

fix could delay the launch by 4 months, says Denny Holt, manager for the Hubble mission at NASA's Johnson Space Center in Houston.

Holt notes, however, that there's only an "outside chance" that the old software would have to be scrapped, causing such a major delay. So far, tests have not uncovered Y2K-related problems. The analysis, however, won't be completed until next month.

Hubble has had only three working gyroscopes since January and another could fail at any time. In announcing plans for the repair mission last March (SN: 3/27/99, p. 203), the agency had pushed for the earliest launch possible, an October date. Wiring problems discovered in the shuttle, however, forced the

mission to be rescheduled to December.

In choosing the earliest launch date, the agency hoped to avoid what NASA chief scientist Edward J. Weiler called a "science emergency." Each month that the telescope lies dormant, about \$20 million in operational costs would go to waste.

To save time, NASA decided last spring to use the same software that successfully navigated the shuttle during a Hubble repair mission 2 years ago. They had estimated that testing the old software for Y2K problems would have added another 6 months to the planning time, Holt says. If modifying the program now proves necessary, he says, the experience acquired in certifying Y2K compatibility for a newer version of shuttle software should make that task easier. —R. Cowen

Chimps outdo people in genetic diversity

If variety is the spice of life, then chimpanzees come loaded with genetic seasoning. Common chimps, with their three subspecies, exhibit far more diversity along a particular stretch of DNA than people do, a new study finds. Moreover, it reveals a surprisingly close genetic connection between common and pygmy chimps.

The results reflect a tight evolutionary relationship, nurtured by frequent interbreeding, among the different chimp groups, report geneticists Svante Pääbo, Henrik Kaessmann, and Victor Wiebe of the Max Planck Institute for Evolutionary Anthropology in Leipzig, Germany.

Broad genetic consistency among subspecies of these close human relatives supports the notion that chimp groups' unique behaviors in different regions develop through teaching and imitation (SN: 6/19/99, p. 388) rather than through genetic determination, the scientists say.

Until now, genetic studies of chimps have focused on mitochondrial DNA, which is inherited only from the mother. The German team instead studied the X chromosome, which is inherited from both parents. Chemical changes and rearrangements occur rarely in the section examined, thus enhancing efforts to reconstruct ancient evolutionary relationships.

The team determined the chemical arrangement of this DNA segment in 30 common chimps of three subspecies—12 central African chimps, 17 western African chimps, and 1 eastern African chimp. The scientists also tested five members of the pygmy chimp species.

Overall, the common chimps exhibited about four times as much diversity in this genetic region as a group of 70 people did, Pääbo's team reports in the Nov. 5 SCIENCE. Differences between individuals in a single chimp subspecies often exceeded those between common chimps and pygmy chimps.

The two chimp species may thus have taken different evolutionary directions



Chimps' genetic diversity: X chromosome tests mark the spot.

relatively recently, the scientists contend. They calculate that this split occurred about 930,000 years ago. Prior estimates, based on more changeable sections of DNA, had placed the species' division at around 2.5 million years ago.

Further research will examine whether gorillas and orangutans display the abundant diversity of chimps or the narrower genetic range of people, the team says.

The scientists hold that people's relatively low genetic variation has implications for how they evolved. "The simplest explanation is that at some rather recent point in the past, humans were few in numbers," asserts Pääbo. "That point could have been the genetic origin of modern humans."

Pääbo and other researchers have similarly argued, using analyses of mitochondrial DNA, that modern humans arose only about 100,000 years ago. This interpretation of the evidence has proven controversial, however (SN: 2/6/99, p. 88).

Since evolutionary processes have yielded chimp subspecies whereas modern humans fall within a unified species, it's not surprising that chimps harbor more genetic diversity than humans do, remarks geneticist Alan R. Templeton of Washington University in St. Louis.

He adds that lesser human diversity doesn't show, as the German team argues, that a decimated human population in the Stone Age depleted human genetic variation. —B. Bower

Soy slows growth of prostate cancers

Men who eat soy-rich diets face a lower risk of deadly prostate cancer, epidemiological studies have indicated. Animal experiments now suggest how soy defends the prostate. The legume induces suicide among cancer cells and limits their spread, scientists report.

In a pair of 6-month-long studies, Göran Hallmans of the University of Umeå in Sweden and his colleagues implanted cancer cells under the skin of rats and hairless mice. Then, they fed the rodents diets deriving one-third of their calories from protein. Some animals got protein from soy, which is rich in plant estrogens known as isoflavones. Whole-grain rye, which contains large amounts of lignans—another family of plant estrogens—provided the protein for others. The final group consumed milk casein, an estrogen-free protein.

Both the soy- and rye-based diets reduced the growth of tumors, compared with the growth of cancers in casein-fed animals. Moreover, only the soy and rye diets induced apoptosis—or natural, programmed death—in the implanted tumor cells, notes coauthor Herman Adlercreutz of the University of Helsinki.

Ordinarily, tumor cells' failure to undergo normal aging and apoptosis contributes to their uncontrolled growth. Adlercreutz says that his team's data represent "the first time it has been shown diet can induce apoptosis."

Jin-Rong Zhou of Harvard Medical School in Boston described related data from mice. His team had placed human cancer cells into the animals' prostates. As tumors began to grow, some mice received diets rich in soy or supplemented with genistein, soy's primary isoflavone. The rest ate casein-based chow.

After 10 weeks, Zhou reports, tumors in casein-fed mice were about twice as big and twice as likely to spread as those in animals fed the soy or genistein.

Zhou notes that most primary cancers aren't lethal; it's their dissemination to other organs that kills. His data indicate that soy retards the cancers' growth and spread at least in part by inhibiting the body's production of blood vessels to supply nutrients to the prostate tumors.

Both Adlercreutz and Zhou reported their findings Monday in Washington, D.C., at the Third International Symposium on the Role of Soy in Preventing and Treating Chronic Disease.

Soy has garnered plenty of media attention for hints that it might cut the risk of breast cancer, says conference chairman Mark Messina, a Port Townsend, Wash., nutritionist. "I think we can now make the case that the prostate data are more impressive," he notes. —J. Raloff