## SCIENCE'S COMPASS



LETTERS

In considering the preservation of species DNA it is noted that "microorganisms, not animals, represent the bulk of the world's phylogenetic diversity and are also in need of preservation in collections." The complexities and triumphs of distinguishing real but very rare or very subtle results from statistical fluctuations with Bayesian analysis are discussed. The benefits and shortcomings of Old World monkeys versus chimpanzees as subjects for analysis in any future primate genome project are examined. And it is observed that the study of artificial genetics has the potential to unite researchers who are on one side or the other of the traditional divide between "natural history" and the "physical sciences."

### Microorganisms Should Be High on DNA Preservation List

The Policy Forum "DNA banks for endangered animal species," by Oliver A. Ryder and colleagues (*Science*'s Compass, 14 Apr., p. 275) generated comment from Phillip A. Morin with response from the authors concerning the need to preserve cells and DNA from threatened and endangered species (Letters, 4 Aug., p. 725). However, it is poorly recognized that microorganisms, not animals, represent the bulk of the world's phylogenetic

diversity and are also in need of Green non-sulfur lections, especially those from habitats under threat of environmental degradation (1). Microorganisms may also be considered endangered in cases where they exist in obligate symbiotic associations with endangered plant or animal species (2). The world's network of microbial culture collections

work of microbial culture collections do a magnificent job with limited resources, but these preserved cultures represent only a tiny fraction of the microbial species present in the environment as most microorganisms are not readily culturable with established techniques (3). For example, there are estimated to be at least 36 major lineages (divisions) of the domain *Bacteria* (4), of which only four are even

moderately well represented in culture collections (see the figure). This limited representation is due to most of new divisions having been recognized by direct molecular methods not involving cultivation. Indeed, many of these divisions do not have even a single representative species preserved in culture collections [e.g., see (5)]. Therefore, the concept of ex situ conservation of microbial biodiversity should be extended beyond pure cultures to habitat samples.

Culture collections could be a valuable tool for the task of preserving samples of endangered habitat reference material or DNA derived from it. Samples could include those from potentially ephemeral extreme environments such as hot springs, Low G+C gran Fibrobacte Marine group A OP Green sulfur Dictvoglomus Cytophanal ermus/Deinococcus Spirochetes 00,, 0.10 Archaea

**Microbial tree of life.** Phylogenetic tree of the domain *Bacteria*, based on comparative analysis of 16*S* ribosomal RNA gene sequences, showing currently recognized major divisions. Divisions moderately well represented in culture collections are shown in red. [Adapted from (4).] Scale bar indicates changes per nucleotide.

acid mine drainage sites, and submarine hydrothermal vents, as well as from unique habitats such as rainforest soil, coral reef invertebrate tissue, oligotrophic lake water, and microbial mats in living stromatolites.

Preservation of habitat reference material could begin with selection of habitat sam-

ples that would ensure that all known microbial lineages of life are represented. It would then be possible to preserve reference samples containing at least one representative of all known divisions of Bacteria and Archaea. This project would be a worthy initiative for support in the International Biodiversity Observation Year 2001-2002 and could be coordinated with existing similar research funded by the National Science Foundation and the National Air and Space Administration, such as Life in Extreme Environments (6) and microbial observatories (7). The resulting material would provide an invaluable scientific resource for comparison and testing of hypotheses concerning microbial diversity, physiology, and evolution, in addition to storing essential reference material in case of future habitat loss.

### John A. Fuerst Philip Hugenholtz

Department of Microbiology and Parasitology, University of Queensland, Brisbane, Queensland 4072, Australia. E-mail: fuerst@biosci.uq.edu.au and philiph@biosci.uq.edu.au

#### References

- N. R. Pace, Science 276, 734 (1997); C. R. Woese, Proc. Natl. Acad. Sci. U.S.A. 95, 11043 (1998); A. T. Bull, A. C. Ward, M. Goodfellow, Microbiol. Mol. Biol. Rev. 64, 573 (2000).
- 2. J.T. Staley, Curr. Opin. Biotechnol. 8, 340 (1997).
- R. I. Amann, W. Ludwig, K.-H. Schleifer, *Microbiol. Rev.* 59, 143 (1995).
- P. Hugenholtz, B. M. Goebel, N. R. Pace, J. Bacteriol. 180, 4765 (1998).
- M. A. Dojka, J. K. Harris, N. R. Pace, Appl. Env. Microbiol. 66, 1617 (2000).
- 6. http://www.nsf.gov/pubs/2000/nsf0037/nsf0037. htm
- 7. http://www.nsf.gov/pubs/2000/nsf0021/nsf0021. htm

# **Figuring the Odds**

Bayesians are not surprised that results at several standard deviations are often spurious, as Charles Seife points out in his News Focus report "CERN's gamble shows perils, rewards of playing the odds" (29 Sept., p. 2260). The significance levels he discusses overstate the evidence, often by a great deal (1). Is a tossed coin that gave 60 heads and 40 tails fair? In the sidebar, "A Greek letter, demystified" (p. 2261), it is incorrectly stated that this happens only 2% of the time with a fair coin. In fact, the probability of obtaining exactly 60 heads and 40 tails is 0.011 (from the binomial distribution). What was apparently meant was that the probability of obtaining 60 or more heads out of 100 is 2%, the one-sided P-value (but this is actually 2.8%).

The *P*-value is a poor measure of the evidence against "no bias." If a biased coin averages 60% heads, the probability of obtaining 60 heads and 40 tails is 0.081, so on this evidence the odds against "no bias" are at most 0.081/0.011 or 7.5:1, not 35:1 or 50:1. In addition, this computation is done under the extreme assumption that