## Ice particles and the greenhouse

 $S_{IR}$  – Within the framework of a suitable model the properties of the Earth's greenhouse can be calculated from tables of absorptivities of the greenhouse gases1. We find that, taking the Earth's albedo to be 0.30, the Earth's effective infrared temperature with no greenhouse would be about 255 K. Adding the present-day greenhouse gases raises the Earth's mean temperature to about 292 K, in agreement with experience. Doubling the present atmospheric CO<sub>2</sub> content would raise the mean temperature by a further 1.5 °C without including a knock-on-effect from water vapour. The latter is what one cares to make it. A typical increase of a factor of 2 gives a rise of about 3 °C.

Even slight variations of circumstances that have been kept constant could alter the result of this investigation; the albedo is one such factor. Increasing the Earth's albedo from 0.30 to 0.315 would conceal the doubling of  $CO_2$ . The albedo depends on clouds in the troposphere, and on the nature



The backscatter mass exclusion coefficient of spherical ice particles averaged over the solar spectrum and the sunlit hemisphere, as a function of particle radius<sup>2</sup>.

of the Earth's surface - oceans, deserts, grasslands, forests, crops, ice and snow.

Small ice crystals in the mesosphere also significantly affect the albedo. The figure shows the average mass coefficient for the exclusion of solar radiation due to backscattering by spherical ice grains uniformly distributed over a sunlit hemisphere2. The computations are based on Mie theory combined with integrations over a hemisphere and an averaging with respect to the solar spectrum. We note that for particles with radii of a few tenths of a micrometre, a typical value of the mass exclusion coefficient is about 4,000 cm<sup>2</sup>  $g^{-1}$ , the result being insensitive to the precise particle size. Taking the mass loading of the particles over the whole Earth to be  $10^{12} \xi g$ (where  $\xi$  is a parameter that ranges from about 0.1 to 10), the albedo is increased by  $10^{12} \xi \times (4,000) / 5 \times 10^{18} = 8 \times 10^{-4} \xi$  (the surface area of the Earth being  $\sim 5 \times 10^{18}$ cm<sup>2</sup>). A value of  $\xi$  of order 10 would therefore yield a cooling of the Earth sufficient to cancel out a doubling of the atmospheric CO<sub>2</sub> content. Although such a value of  $\xi$  is not currently operative, the total water vapour content of the high atmosphere would, if condensed, be far more than sufficient. The measured mid-stratospheric water vapour content is indeed in excess of  $10^{15}$  g. Conditions for the condensation of water vapour into particles appear to be delicate<sup>3</sup>, suggesting that no great change would be needed to increase  $\xi$  substantially. Episodes of extensive ice-crystal condensation could be triggered by either terrestrial or extra-terrestrial

## **African fishes**

Sir — Meyer *et al.* conclude <sup>1</sup> that morphological diversification in flocks of cichlid fishes in Lake Victoria occurs without much molecular evolution. This agrees with our own data on hybrids between different species of cichlid fish from Lake Victoria, which can be bred easily over several generations<sup>2–4</sup>. In addition to the molecular data on mitochondrial DNA, our results show the genetic compatibility for the entire genome of some Lake Victoria 'haplochromines'.

But we disagree with Meyer et al.'s second conclusion that Lake Malawi species are more closely related to those of Lake Victoria than to those of Lake Tanganyika. We tested by hybridization the genetic compatibility of Astatotilapia burtoni, a riverine species from the Lake Tanganvika basin, with Astatotilapia nubilus, a riverine species from the Lake Victoria basin. The fertility of the hybrids between these species suggests that they are genetically almost as compatible as haplochromines from the Lake Victoria basin<sup>2,3</sup>. Moreover, we also tried to breed hybrids between the monotypic genus Astatoreochromis alluaudi and several haplochromines of Lake Victoria. According to Meyer et al., this species should be more closely related to Lake Victoria haplochromines than to those of Lake Tanganyika. We succeeded, however, in breeding only one hybrid population between Astatoreochromis alluaudi and Astatotilapia 'black lividus' (a so far undescribed species of Lake Victoria). These hybrids reproduced, but the eggs deteriorated before hatching. This is the only evidence for sterile hybrids that we have obtained so far. Astatoreochromis alluaudi is also the only species in Lake Victoria that stands out in its protein characteristics5.

We believe that Meyer *et al.* should include molecular data from the riverine Lake Tanganyika haplochromine *A. burtoni* in their analysis. Their second conclusion will then be that some haplochromines of different lakes arose from more closely related riverine species; in other words, that segregation had started before the lakes formed. The major problem of the species flocks of the great African lakes is clearly how they have evolved and are maintained in sympatry,

factors connected with the supply of condensation nuclei. Perhaps this is what happened during the Little Ice Age.

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while obviously relying exclusively on premating barriers — a significant ethological problem in its own right<sup>4</sup>.

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MEYER *ET Al.* REPLY — Inferring relationships from patterns of hybridization is risky. The hybridization studies involving *Astatotilapia burtoni* males and *Astatotilapia nubilis* females produced only lethal hybrids<sup>2</sup>. Females of *A. burtoni* mated rarely with males of *A. nubilis*, but produced viable hybrids<sup>2</sup>. Numerous studies have shown that hybrid incompatibility, although usually correlated



Evolutionary tree based on comparisons of part of the control region of mtDNA. The region sequenced and the methods used are described by Meyer et al.<sup>1</sup>. Bootstrap values appear above the two internal branches that are statistically significant (a-b and c-d). The distance from any node to the tips of the descendant branches are roughly proportional to the average number of base substitutional differences between species united by that node. The root of the tree (node d) was determined by using Julidochromis as an outgroup (see ref. 1). The two sequences of Astatotilapia have been deposited in Gen Bank with accession numbers X58151 (A.c.) and X58152 (A.b.).

with phylogenetic distance, is not an accurate metric for building trees and time scales.

In anticipation of questions about Astatotilapia's phylogenetic status, one of us (A. M.) sequenced the most variable part of the control region of mitochondrial DNA from two key species, A. burtoni and A. calliptera (a non-endemic cichlid from Lake Malawi). Three Astatotilapia species from the Lake Victoria basin (A. nubilis, A. elegans and A. piceatus) had already been tested<sup>1</sup>. The figure shows a tree relating the two new sequences to those of the species flocks in

Howard, J. N., Garing, J. S. & Walker, R. G. in *Handbook of Geophysics and Space Environments* (ed. Valley, S. L.) (McGraw Hill, 1965).