Reproductive factors are crucial in the aetiology of breast cancer – a reply

Sir,

The letter by Professor Khuder raises several points of potential importance. Our statistical analysis correlating *Mus* species and human breast cancer (HBC) incidence is described as simplistic, ignoring social, cultural and demographic variables. Thus, it may suffer from ecological bias, due to the effect of hormonal promoters on the development of HBC. The greatest influence would likely be associated with fecundity, which is best reflected in the world statistics on 'total fertility rate' (TFR) (US Bureau of the Census, Report WP/98, World Population Profile (1998) US Government Printing Office: Washington DC, 1999).

TFR was evaluated as a potential confounder of the association of *M. domesticus* geography with human breast cancer incidence. For our sample of 39 countries (less two regions, Hawaii and 'circumpolar Inuit' for want of data), we analysed the reported 1990 (or 1998, where lacking) TFR for correlation with the world age-standardized incidence rate (WASIR) for female breast cancer (as in Stewart et al, 2000). The expected negative correlation of WASIR with TFR [$R = -0.327$, $P = 0.048$] was found. However, across Europe there was no difference in TFR between lands of *M. domesticus* and lands of other mice (mean TFR $1.656 \pm \text{SD } 0.368$, vs 1.657 ± 0.346 , $P = 0.993$).

Internationally, excluding Europe, there was a higher reported TFR in *M. domesticus* lands (TFR 2.875 ± 0.822 vs 2.371 ± 0.971 , $P = 0.244$). Overall, the crude difference in mean WASIR due to *M. domesticus* lands is +15.6, accounting for 38.3% of the observed variation in this sample. The TFR-adjusted difference in mean WASIR is +17.4, accounting for 48.4% of variation, both highly statistically significant ($P < 0.001$). Thus, in addressing Professor Khuder's concern about reproductive factors by adjusting for TFR, the association of WASIR with lands of *M. domesticus* was strengthened.

The report by McCredie et al (1999) on the incidence of HBC in Maori and non Maori women emphasizes that all parameters suggesting a lower incidence of HBC were seen in Maori women in a highly significant fashion, lower educational level, lower socio-economic status, lower age at first full term pregnancy, high parity and longer duration of breast-feeding. Despite this, the incidence of HBC in Maori women before the age of 54 is twice that of non-Maori women in New Zealand. Could this reflect a greater exposure of the Maori to *Mus domesticus* which occurs in both urban areas and native forests in New Zealand (King, 1982)?

In the paper by Chie et al (1995), no data on the incidence of HBC in white women is given in the text. White women form a minuscule proportion of the female population of Taiwan. The use of oral contraceptives in Asian women migrating to the USA, adjusted for age, ethnicity, study area, years since migration,

family history of HBC and age at first full-term birth was not associated with increased risk of breast cancer (Ursin et al, 1999). The low incidence of HBC in Spanish women and Hispanic women living in the US is a fact. Genetic susceptibility to MMTV was not addressed in our paper, although it has been well studied in mice (Ross et al 1997; Golovkina, 2000).

In summary, adjustment of our analysis for a possible ecologic bias related to fecundity and hormonal influence on breast cancer increases the statistical significance of our reported association. We agree with Professor Khuder that one should seek a correlation in breast-cancer risk with more direct measures of contact and potential exposures to mice, such as local mouse population levels, or occupational exposures such as in farming (Khuder et al, 1998), or in laboratory work with experimental handling of mice (Dion et al, 1986). Some areas of the world do have wide fluctuations in *M. domesticus* population levels due to epizootic diseases, or climatic variations. One must keep in mind that the MMTV is the proposed cause, and that *M. domesticus* would be a surrogate of MMTV exposure. The actual risk will depend on the likely modes of MMTV transmission, exposure, and the burden of infectious MMTV in the resident mouse population.

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