true in his national epidemiological study. The reasons for dysrhapic developmental accidents may lie in the genotype, in the pre- or postconceptional environment, or in both. If so, is there a case for studying preventive vitamin supplementation in mothers after the occurrence of a minor developmental mishap like the one reported?

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References

Vogel and Motulsky refer to this explicitly on page 449 of their text Human Genetics (1st ed), and implicitly in putting Dürer’s woodcut of Adam and Eve on the front cover. But that ‘giant saltation’ was, surely, a one-off event. It is difficult to conceive of mini-saltations (hops and skips?) occurring at a rate of several per generation, and which so reorder the genome that the cerebral phenotype is improved. Speculation is fine (and fun), but one must resist the temptation to go overboard.

Otherwise, I found this an interesting and useful paper, particularly in respect of the data concerning the phenotypes of children with de novo apparently balanced rearrangements.

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Tracheo-oesophageal dysraphism

Sir,

I am alarmed by Dr David’s wish (J Med Genet 1984;21:74) to get his hands on ‘whoever first used the awful term tracheo-oesophageal dysraphism’ and wish to disclaim responsibility, though I cannot remember where I came across the phrase when I first used it (Lancet 1980;ii:80). I thought ‘dysraphism’ (defective fusion) was an acceptable way of getting around the clumsiness of ‘tracheo-oesophageal fistula and/or oesophageal atresia’. What would Dr David suggest?

As for the ‘probably spurious’ increase in NTD in sibs of children with TED, both negative studies cited by Dr David came from low frequency areas, and Baird’s study did not measure the number of sibs but only failed to find a sib with NTD reported to the BC registry. Drs Ilyina and Lurie (J Med Genet 1984;21:73–4) place our study in the ‘low incidence’ group, but in fact Montreal had a frequency of over three per 1000 during the time of the study. The figure for ‘Canada’ given is for British Columbia, a low incidence area. Before giving up the idea of an increase in NTD in sibs of children with TED, I would like to see negative data from a high incidence area.

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