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Long-lived flies

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Mutations that increase the [life span](#) of *Caenorhabditis elegans* encode components of the insulin/IGF signalling pathway. In the April 6 [Science](#), two papers describe mutations that link insulin signaling with longevity in *Drosophila melanogaster*. Clancy *et al.* report that homozygous null mutations in [chico](#), encoding an insulin receptor substrate protein, increased the female fly life span by up to 48% (*Science* 2001, **292**:104-106). They were able to demonstrate that the effects of *chico* on longevity could be separated from effects on fertility, stress resistance or body size. Tatar *et al.* show that the *Drosophila InR* (insulin-like receptor), which is a homolog of the *C. elegans daf-2* gene, also regulates adult longevity (*Science* 2001, **292**:107-110). Female flies with heteroallelic combinations of hypomorphic *InR* alleles lived up to 85% longer than controls. Tatar *et al.* provide evidence that the effects of *InR* are mediated by regulation of neuroendocrine function and juvenile hormone production. The conservation of insulin signalling pathways suggests that similar genes may affect longevity and [aging in mammals](#).

References

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