

Evolutionary regulation of the blind subterranean mole rat, *Spalax*, revealed by genome-wide gene expression

L. I. Brodsky*[†], J. Jacob-Hirsch*^{†§}, A. Avivi*, L. Trakhtenbrot*[§], S. Zeligson*[§], N. Amariglio*[§], A. Paz*, A. B. Korol*, M. Band[¶], G. Rechavi*[§], and E. Nevo*^{||}

*Institute of Evolution, University of Haifa, Mount Carmel, Haifa 31905, Israel; [†]Division of Pediatric Hematology/Oncology, Edmond and Lily Safra Children's Hospital and Sheba Cancer Research Center, The Chaim Sheba Medical Center, Tel Hashomer 52662, Israel; [§]Sackler School of Medicine, Tel Aviv University, Tel Aviv 69978, Israel; and [¶]The W. M. Keck Center for Comparative and Functional Genomics, University of Illinois at Urbana-Champaign, Urbana, IL 61801

Contributed by E. Nevo, June 16, 2005

We applied genome-wide gene expression analysis to the evolutionary processes of adaptive speciation of the Israeli blind subterranean mole rats of the *Spalax ehrenbergi* superspecies. The four Israeli allopecies climatically and adaptively radiated into the cooler, mesic northern domain (N) and warmer, xeric southern domain (S). The kidney and brain mRNAs of two N and two S animals were examined through cross-species hybridizations with two types of Affymetrix arrays (mouse and rat) and muscle mRNA of six N and six S animals with spotted cDNA mouse arrays. The initial microarray analysis was hypothesis-free, i.e., conducted without reference to the origin of animals. Principal component analysis revealed that 20–30% of the expression signal variability could be explained by the differentiation of N–S species. Similar N–S effects were obtained for all tissues and types of arrays: two Affymetrix microarrays using probe oligomer signals and the spotted array. Likewise, ANOVA and *t* test statistics demonstrated significant N–S ecogeographic divergence and region–tissue specificity in gene expression. Analysis of differential gene expression between species corroborates previous results deduced by allozymes and DNA molecular polymorphisms. Functional categories show significant N–S ecologic putative adaptive divergent up-regulation of genes highlighting a higher metabolism in N, and potential adaptive brain activity and kidney urine cycle pathways in S. The present results confirm ecologic–genomic separation of blind mole rats into N and S. Gene expression regulation appears to be central to the evolution of blind mole rats.

adaptive speciation | genome-wide expression | microarray | ecological stress | cross-species hybridization

Evolutionary change is primarily driven by ecological stress (1–3). Climatic stresses are instrumental in advancing the twin evolutionary processes of adaptation and speciation. Darwin recognized that “climate plays an important part in determining the average number of a species, and periodical seasons of extreme cold or drought seem to be the most effective of all checks” (4).

Evolutionary Ecological Genomics

The genomic revolution has linked molecular genomics with organismic evolution (3). The ongoing problems associated with speciation and adaptation (3) could fruitfully be explored by genome-wide gene expression analysis (5). How many genes, which ones, and, most importantly, what regulatory variations across the genome are possibly adaptive, associated with ecological stress, and affect adaptation and speciation? What genes and/or regulatory elements are associated with ecological speciation? Ecological and functional genomics (3) could elucidate some of the major yet unresolved evolutionary problems.

The Evolutionary Global Experiment of Subterranean Mammals

Subterranean mammals provide a long-term natural evolutionary experiment of mammals adapted to life underground. Subterranean mammals display mosaic evolution of regression, progression, and global convergence across all organizational levels from molecular to organismal (6). Their underground evolution originated in Middle Eocene to early Oligocene (45–35 million years ago) and continued stepwise in the Cenozoic period after climatic cooling and drying. The subterranean ecotope is characterized by simple, climatically stable, and specialized profiles but is subjected to severe stresses of darkness, energetics, infection, and hypoxia–hypercapnia (6).

Spalacids and *Spalax ehrenbergi* Superspecies in Israel

The Spalacidae are Eurasian, primarily East Mediterranean, subterranean rodents representing extreme adaptations to life underground (figure 7.4b of ref. 6). Spalacids originated some 25 million years ago in southwest Asia Minor based on fossils and 40 million years ago based on molecules (6). The *Spalax ehrenbergi* superspecies adaptively radiated in the Near East and split into at least 12 allopecies. Four species evolved in Israel during the Pleistocene, adapting to four climatic regimes (7). The two northern Israeli species, *Spalax galili* ($2n = 52$) and *Spalax golani* ($2n = 54$), range in cool humid climate, whereas the central species, *Spalax carmeli* ($2n = 58$), ranges in warm humid climate, and the southern species, *Spalax judaei* ($2n = 60$), ranges in warm dry climate (figures 5, 11, and 12 in ref. 7). Generally, the complex extends along a southern gradient of increasing aridity with underground temperature increasing and humidity decreasing southwards, although mildly and without extremes as above ground. This ecological speciation into a trend of four species subdivided into older northern (*S. galili* and *S. golani*) and younger southern (*S. carmeli* and *S. judaei*) species has been documented by diverse studies of noncoding and coding genomic regions. These regions include chromosomes (7), allozymes (8, 9), mtDNA (10), random amplified polymorphic DNAs, minisatellites and chiasma frequency (11, 12), microsatellites (13), DNA hybridization (14), mitochondrial cytochrome *b* (15), the mitochondrial control region (16), erythropoietin (17), neuroglobin and cytoglobin (our unpublished observations), and the Clock and MOP3 genes (18). Here, we show that previous genetic and phenetic patterns are corroborated by genome-wide gene-expression analysis based on an average of 15,000 genes, highlight-

Conflict of interest statement: No conflicts declared.

Abbreviations: N, northern domain; S, southern domain; PC, principal component; PCA, PC analysis; PM, perfect match; MM, mismatch.

[†]L.I.B. and J.J.-H. contributed equally to this work.

^{||}To whom correspondence should be addressed. E-mail: nevo@research.haifa.ac.il.

© 2005 by The National Academy of Sciences of the USA

ing the remarkable importance of at least partly adaptive gene-expression regulation in *Spalax* evolution.

Materials and Methods

Animals Analyzed. We analyzed 16 animals, eight animals representing the cool mesic northern domain (N) [*S. galili* ($2n = 52$) from Dalton, Upper Galilee (seven animals), and *S. golani* ($2n = 54$) from Quneitra, Golan Heights (one animal)], and eight animals representing the warm xeric southern domain (S) [*S. carmeli* ($2n = 58$) from Mukhraka, Mt. Carmel (one animal), and *S. judaei* ($2n = 60$) from Lahav, northern Negev (seven animals)]. For the Affymetrix experiments, two N (one *S. galili* and one *S. golani*) and two S (one *S. carmeli* and one *S. judaei*) were used, with each animal on an individual chip. For the cDNA experiment, six N (six *S. galili* only) and six S (six *S. judaei* only) were used.

All were adult males of ≈ 150 g; they were captured in the field in July 2002 and housed in individual cages, each species in a separate room of the Institute of Evolution, University of Haifa. All animals were housed according to regulations conforming to animal handling protocols approved by the University of Haifa under controlled conditions of 22–24°C with seasonal light/dark hours on a diet of carrots. Animals were killed three to four months after capture (October 2002 and December 2004). Brain, kidney, and muscle tissues were removed from each animal, immersed in liquid nitrogen, and stored in -80°C until RNA processing. Brain and kidney mRNA of two animals from N and two animals from S were hybridized with Affymetrix rat and mouse oligonucleotide arrays. In addition, muscle mRNA of six animals from N and six animals from S were hybridized with a spotted mouse microarray.

Affymetrix Microarray Experiments. RNA from two Northern and two Southern individuals were analyzed with Affymetrix mouse MOE430A and rat RAE230A oligonucleotide arrays according to the manufacturer's instructions. Total RNA from each sample was used to prepare biotinylated target RNA by following the manufacturer's recommendations. Quality and quantity of starting RNA was confirmed by using agarose gel and OD measurement, respectively. After scanning, array images were assessed by eye to confirm scanner alignment and the absence of significant bubbles or scratches on the chip surface. 3'/5' ratios for GAPDH and β -actin were confirmed to be within acceptable limits, and BioB spike controls were present on all chips, with BioC, BioD, and CreX also present in increasing intensity. When scaled to a target intensity of 150 (Affymetrix MAS 5.0 array analysis software), scaling factors for all arrays were within acceptable limits, as were background, Q values, and mean intensities. Details of quality control measures are available from the National Center for Biotechnology Information or the Chaim Sheba Medical Center.

cDNA Microarray Experiments. Muscle tissue from 12 individual animals, six S [*S. judaei* ($2n = 60$) from Lahav] and six N [*S. galili* ($2n = 52$) from Dalton], were directly compared by using mouse cDNA microarrays representing 15,247 clones from the National Institute on Aging 15K libraries (19, 20) as well as positive and negative controls. We added to the cDNA arrays 32 *Spalax* sequences for the following genes: *Epo*, *Epo-R*, *VEGF*, *VEGF-R(FLK)*, *VEGF-R(FLT)*, *HIF1A*, *HuR*, *Ngb*, *CYTG*, *Mdm2*-promoter, *Apaf1*-promoter, *p53*, *CARP*, *HPSE*, *LDHa*, *GT1*, *CLOCK*, *MOP3*, *Per1*, *Per2*, *Per3*, *Cry1*, *Cry2*, *Tim*, *OPN4*, *ADCYAPI*, *MAOA*, *LIS*, *DCX*, *CRYAB*, and *FGF-R3*. Description of the National Institute on Aging 15K microarray printing parameters and format are available from the authors upon request. For all comparisons, 10 μg of *Spalax* muscle total RNA was reverse-transcribed according to indirect aminoallyl incorporation protocol and labeled with Alexa Fluor 555 or Alexa Fluor 647 dyes (Invitrogen). Slides were hybridized overnight at 42°C, washed, and scanned with an Axon GenePix 4000B microarray scanner (Molecular

Devices, Sunnyvale, CA). Detailed protocols describing labeling and hybridization are available from the authors upon request. All slide images were analyzed by using GENEPIX PRO 6.0 software. Analyzed slide images were manually edited, and aberrant spots were flagged for exclusion in downstream analysis.

Quality Control and Normalization of Microarray Data. The initial Affymetrix hybridization probe expression signals (CEL files) were preprocessed separately for each set of hybridizations per array type (mouse or rat). Because all hybridizations were cross species and, presumably, some mouse or rat oligomer probes could be incompatible for hybridization with *Spalax* mRNA, we applied probe-based detection (21, 22) of N- and S-discriminating genes rather than a gene-based analysis. This approach is also expected to be generally more accurate because only a subset of each probe set may be suitable for analysis because of putative gene splice variants in the mRNA sample and possible slide artifacts (23).

The quality control analysis of the Affymetrix slides was performed by using techniques described in ref. 23. The major criterion for detection of slide artifacts was the deviation from uniformity of the distribution for log-transformed perfect match (PM) probe signals of high and low intensity over the microarray geometry. As a result of the quality control analysis, some condensed areas of high and low probe signals ("patterns") were detected on slides, and sometimes their artifactual nature was very obvious (*Supporting Materials and Methods* and Fig. 3, which are published as supporting information on the PNAS web site).

Statistically significant (23) local artifactual patterns were cleaned. Corresponding probe pairs were removed from all mice and rat microarray hybridization sets. In total, 15% of the probe pairs were excluded from the subsequent analysis. Exclusion of local regions across the chip did not significantly affect the results because probe pairs of each probe set are uniformly distributed over the microarray geometry.

Considering that in cross-species hybridizations PM and mismatch (MM) probes may potentially differ in sequence from hybridized mRNA (21, 22), both probe signals were included in the analysis. However, we relied primarily on the PM signals, assuming they had higher sequence similarity. Log-transformed probe signals of each sample were quantile-normalized (24). Normalization was performed separately for PM and MM signals. The goal of the normalization was to equalize distributions of probe log-signals for all samples within the set.

The six spotted array hybridizations were normalized by nonlinear LOWESS regression (25). The slide artifacts for spotted array hybridizations were checked with the procedure described in ref. 23.

ANOVA Model for Log-Signal Gene Expression Values. We used a general additive ANOVA model for the influence of effects of two factors (ecogeography and tissue) and their interaction on expression log-signal of a probe (see *Supporting Materials and Methods*). The effects of the model were estimated through general linear model regression (26). The significance of the estimated effect for individual probe profile (the vector of probe signals across all samples) was detected through t -distributed statistic (26). The ANOVA estimation was used to detect genes with significant geography–tissue interaction effect.

The Statistical Significance of Gene Regulation. For the Affymetrix array experiments, the statistical significance of ANOVA effects for a given gene were evaluated by Wald statistics (27). First, for each probe profile the t test statistic (see Eq. 2 in *Supporting Materials and Methods*) of the effect estimation was calculated as a measure of normalized deviation (including direction) of the factor combination-specific log-signal from the probe log-signal average. The null hypothesis assumes that a significant individual

effect (t test) would appear randomly for probe profiles across the entire set of probes of the set of hybridizations. For a specific gene, the null hypothesis assumes that all probes within the gene probe set demonstrate effects of factors by chance (see *The Independence of Probe Signals Within Probe Set* in *Supporting Materials and Methods*; see also Figs. 4–6, which are published as supporting information on the PNAS web site). Afterward, two Wald statistics (the sum of squares of t statistics) were calculated for every gene effect (N–S effect, tissue effect, and their interaction): one for all probes with positive effects and another one for all negative effect probes. Under the null hypothesis, these are χ^2 -distributed (27). The P value of the ratio of these two χ^2 statistics, scaled by corresponding degrees of freedom (F test), reflects the significance of the effect for a gene in the corresponding direction.

The same Wald approach was applied for detection of N–S differentiating genes per tissue. Considering that fold changes detected by the Affymetrix experiments were rather small (≈ 1.5) even for the best discriminating probes and because the t test significance of the effect for an individual probe was based on only four animals, we used the above mentioned probe set “voting” approach to determine whether an individual gene was differentially expressed in N and in S and whether there is a specificity of expression under some geography–tissue combination. The N–S discriminating power of a gene was estimated through voting of its probe set t -test values either “for” or “against” regulation in a given direction. Two Wald statistics were calculated for each gene: with N up-regulated probes of the gene probe set and with S up-regulated probes of the same gene. If one Wald statistic was significant and the other was not, we considered that gene to be significantly up-regulated in the direction of the significant Wald statistic. To flag a gene as significantly differentially expressed, a single Wald test P value (calculated according to all 11 probes of the probe set) needed to pass the 10% false discovery rate (28) significance level.

For spotted cDNA arrays, genes with significant N–S differences of expression signals (N–S effect) were detected with a t test taking into consideration three direct and three dye-swapped hybridizations. Because all 12 RNA samples represent individual animals (six N and six S), all direct vs. all swapped hybridizations were paired to estimate the N–S effect that is not confounded by dye. The nine estimations of N–S effect are slightly interdependent. Indeed, the estimation of every one of nine N–S effects is based on a pair of direct and dye-swapped hybridizations. Each pair has six related pairs: three pairs with the same first hybridization and three pairs with the same second hybridization. This weak interdependence of nine N–S difference estimations decreases the effective number of independent members for the χ^2 statistic of the corresponding t test. The Satterthwaite’s formula was used to calculate the effective number of degrees of freedom for the variance estimation according to interpenetrating subsamples (29).

Results

After quality control filtering of the Affymetrix signals, we analyzed a data set that contained 430,509 probe pairs representing 17,941 genes and 308,042 probes representing 14,708 genes on the mouse and rat GeneChips, respectively. For the mouse cDNA, 16,416 spots were analyzed, including control spots.

The process of data analysis was carried out by using a multistep approach. First, a hypothesis-free principal component (PC) analysis (PCA) was done to detect what factors explain the major trends in the overall variability. Next, we investigated the N vs. S differences as these stem from the PCA analysis of two types of Affymetrix microarray data (PM and MM probe oligomer signals separately for the mouse and rat) and from spotted array data (Fig. 1a). Finally, we characterized the effect of N vs. S species for all three tissues at the individual probe level (Affymetrix hybridiza-

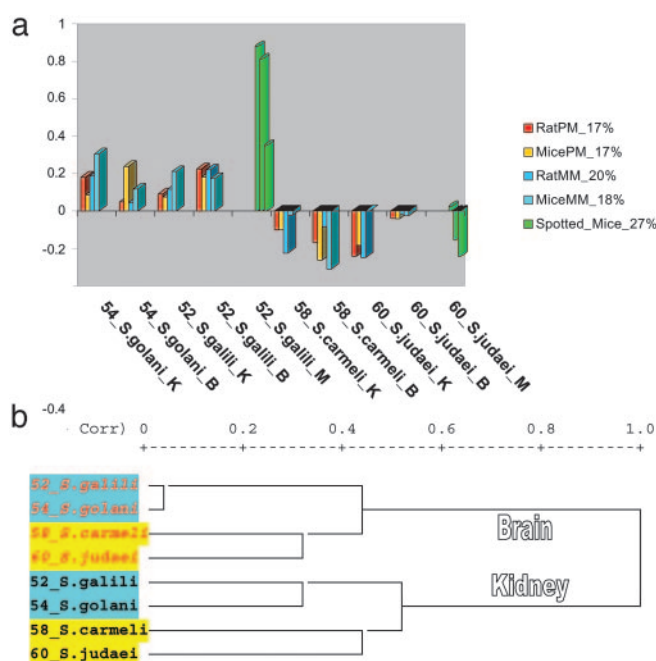


Fig. 1. PCA and cluster analysis of probe oligomer (Affymetrix arrays) and gene signals (spotted array). Profiles of the second PC of PM and MM log signals for mouse and rat Affymetrix microarray experiments and mouse spotted-array experiment. (a) PC2 reflects the general signal difference between N and S animals for two types of Affymetrix microarrays [PM (red bars) and MM (blue bars) signals] and for the spotted array experiment (green bars). This PC2 component explains 20–30% of total signal variability. (b) The dendrogram of correlation distances between Affymetrix hybridizations (average linkage) after the cleaning of artifacts. All nonpatterned probe oligomers with average signals of >300 according to all eight hybridizations were taken. Clear N–S (*S. galili* plus *S. golani* vs. *S. carmeli* plus *S. judaei*) separation is seen for both tissues as in all other evolutionary discrimination of *Spalax* species based on diverse molecular markers and morphological criteria (figures 16, 17, 19, 20–22, 23 a and b, 31, and 33 in ref. 7).

tions) or at the gene level (spotted arrays). The last step of our analysis was a “supervised approach,” for which we distinguished genes with significant N–S expression divergence within each individual tissue by estimating differences on spotted arrays and the correlated differential behavior of probe sets for Affymetrix arrays. Also, genes with geography–tissue specificity in expression were detected by ANOVA.

PCA. The general picture of PM and MM probe log signals according to the PCA (30) demonstrated similar patterns for the variability of Affymetrix array signals across the two series of hybridizations. This similarity supports the hypothesis of small differences between PM and MM probes in cross-species hybridizations.

The first PC of all Affymetrix hybridizations (mouse and rat arrays) reflected the difference between tissues. PC1 of spotted hybridizations is related to the variability of individual slides. In all cases, PC1 covers 40–50% of total signal variability.

Notably, PC2 reflected an important pattern of signal variability for all sets of hybridizations (Affymetrix and spotted arrays) and accounts for gene expression divergence between N and S species (Fig. 1a). This PC explains 20–30% of the total signal variability.

Genes with Ecogeography–Tissue Specificity in Their Expression. The ANOVA analysis detected ≈ 100 genes with ecogeography–tissue specificity in their expression (Data Set 1, which is published as supporting information on the PNAS web site). We

dian rhythmicity, and aggressive behavior (figures 61–63 in ref. 7) all display clinal patterns associated with increasing climatic stress toward the desert. Remarkably, aggression increases northward and “pacifism” increases southward (pp. 127–132 in ref. 7). Notably, the 5-hydroxytryptamine (serotonin) receptor expression associated with aggression levels is significantly up-regulated (our unpublished results) in *S. judaei*, possibly relating to its more pacifistic behavior in xeric environments.

Two antagonistic biological trends occur in *S. ehrenbergi* superspecies in Israel; metabolism increases northward, but relative brain size and encephalization, underlying complex behavioral patterns, increase southward and climax in *S. judaei* ($2n = 60$) (tables 3 and 4 a–c in ref. 7), (i.e., one of the S species) the smallest species in size (figure 25 in ref. 7). This extraordinary pattern is presumably adaptive, reflecting the Bergmann rule and driven by natural selection. Larger relative brain size in the southern *S. judaei* appears to be associated with increasing aridity stress and climatic unpredictability. The more significantly N-up-regulated kidney genes (Table 2 and Fig. 7) display the higher metabolism in N. Behavioral traits like memory and orientation tend to increase in S. Microarray results support the previously described physiological and behavioral opposite N–S trends (7).

Kidney and Brain Function Based on GO Classification. How does the genome-wide gene expression underline *Spalax* adaptations to underground life? The S kidney is characterized by extraordinary urinary concentration ability under protein and salt loads (pp. 71–73 in ref. 7). Only the kidney of *S. judaei* ($2n = 60$) produced hyperosmotic urine adapted to the desert succulent and halophyte plants, pushing speciation southward (pp. 71–77 and figure 35 in ref. 7). Genes of the GO categories protein signaling, acid metabolism, and amine transport characterize the presumed adaptive uniqueness of the *Spalax* desert kidney, associated with increasing ecological stresses of aridity and climatic unpredictability (pp. 35 and 173–175 in ref. 7). Intriguingly, the up-regulated genes in the relatively larger brain of *S. judaei* in S relate to the GO category phosphate metabolism and are

significantly overrepresented regarding all LocusLink rat genes. These genes include those in the GO categories phosphorous metabolism, phosphorylation, protein modification, and protein amino acid phosphorylation (see Data Set 5).

Conclusions and Prospects

This study highlights the fact that allelic diversity is not the only pattern that ecologically differentiates N from S *Spalax*. Most remarkably, genome-wide gene expression regulation in speciation and adaptation seems to be of paramount importance in *Spalax* evolution. This regulatory impact resides primarily in the noncoding genome (e.g., random amplified polymorphic DNAs, amplified fragment-length polymorphisms, variable number tandem repeats, simple sequence repeats), which has been repeatedly implicated in diverse environmental, climatic, and respiratory stresses. Climatic selection seems to be instrumental in driving the noncoding and coding *Spalax* genome to presumably adapt to life underground and speciate in a N–S trend along a southward gradient of increasing aridity stress. Environmentally stressed regulation of genome-wide gene expression in *Spalax* and probably more generally in other organisms across life appears to be the hallmark of the ecological evolutionary process of speciation and adaptation.

We plan to deeply explore the nature of the divergent genes in *Spalax* and unravel their functional significance. Future studies also could elucidate the interaction of potential adaptive evolution of *Spalax* genomics, proteomics, and phenomics to local and regional stresses underground by focusing on sequencing and SNP of stress genes and their evolutionary environmental regulation (e.g., ref. 30).

We thank Eshel Ben Jacob and Yossi Hillel for reviewing the paper and Harris Lewin and Lena Feinstein for comments that helped improve the paper. This work was supported by an Arison family donation (to the Center of DNA Chips in Pediatric Oncology, The Chaim Sheba Medical Center) and by the Ancell Teicher Research Foundation for Genetics and Molecular Evolution.

- Hoffman, A. R. & Parsons, P. (1991) *Environmental Stress* (Oxford Univ. Press, Oxford).
- Nevo, E. (1998) *Heredity* **81**, 591–593.
- Nevo, E. (2001) *Proc. Natl. Acad. Sci. USA* **98**, 6233–6240.
- Darwin, C. (1859) *On the Origin of Species by Means of Natural Selection* (Murray, London).
- Brown, P. O. & Botstein, D. (1999) *Nat. Genet.* **21**, 33–37.
- Nevo, E. (1999) *Mosaic Evolution of Subterranean Mammals, Regression, Progression and Global Convergence* (Oxford Univ. Press, Oxford).
- Nevo, E., Ivanitskaya, E. & Beiles, A. (2001) *Adaptive Radiation of Blind Subterranean Mole Rats* (Backhuys, Leiden, Germany).
- Nevo, E. & Shaw, C. R. (1972) *Biochem. Genet.* **7**, 235–241.
- Nevo, E., Filippucci, M. G. & Beiles, A. (1994) *Heredity* **72**, 465–487.
- Nevo, E., Honeycutt, R. L., Yonekawa, H., Nelson, K. & Hanzawa, N. (1993) *Mol. Biol. Evol.* **10**, 590–604.
- Ben-Shlomo, R., Fahima, T. & Nevo, E. (1996) *Isr. J. Zool.* **42**, 317–326.
- Nevo, E., Ben-Shlomo, R., Beiles, A., Ronin, Y. I., Blum, S. & Hillel, J. (1995) in *Gene Families: Structure, Function, Genetics Evolution*, eds Holmes, R. S. & Lim, H. A. (World Scientific, Singapore), pp. 55–70.
- Karanth, P. K., Avivi, A., Beharav, A. & Nevo, E. (2004) *Biol. J. Linn. Soc.* **83**, 229–241.
- Catzefflis, F. M., Nevo, E., Ahlquist, J. E. & Sibley, C. G. (1989) *J. Mol. Evol.* **29**, 223–232.
- Nevo, E., Beiles, A. & Spradling, T. (1999) *J. Mol. Evol.* **49**, 215–226.
- Reyes, A., Nevo, E. & Saccone, C. (2003) *Mol. Biol. Evol.* **20**, 622–632.
- Shams, I., Avivi, A. & Nevo, E. (2004) *Proc. Natl. Acad. Sci. USA* **101**, 9698–9703.
- Avivi, A., Albrecht, U., Oster, H., Joel, A., Beiles, A. & Nevo, E. (2001) *Proc. Natl. Acad. Sci. USA* **98**, 13751–13756.
- Tanaka, T. S., Jaradat, S. A., Lim, M. K., Kargul, G. J., Wang, X., Grahovac, M. J., Pantano, S., Sano, Y., Piao, Y., Nagaraja, R., et al. (2000) *Proc. Natl. Acad. Sci. USA* **97**, 9127–9132.
- Kargul, G. J., Dudekula, D. B., Meng, Y. K., Saied, K. L., Jaradat, A., Tanaka, T. S., Carter, M. G. & Ko, M. S. H. (2001) *Nat. Genet.* **28**, 17–18.
- Ji, W., Zhou, W., Gregg, K., Yu, N. & Davis, S. (2004) *Nucleic Acids Res.* **32**, e93.
- Gautier, L., Cope, L. M., Bolstad, B. M. & Irizarry, R. A. (2004) *Bioinformatics* **20**, 307–315.
- Brodsky, L., Leontovich, A., Shtutman, M. & Feinstein, E. (2004) *Nucleic Acids Res.* **32**, E46.
- Bolstad, B. M., Irizarry, R. A., Astrand, M. & Speed, T. P. (2003) *Bioinformatics* **19**, 185–193.
- Yang, Y. H., Dudoit, S., Luu, P., Lin, D. M., Peng, V., Ngai, J. & Speed, T. P. (2002) *Nucleic Acids Res.* **30**, e15.
- Draper, N. R. & Smith, H. (1981) *Applied Regression Analysis* (Wiley, New York).
- Agresti, A. (1990) *Categorical Data Analysis* (Wiley, New York).
- Benjamini, Y. & Hochberg, Y. (1995) *J. R. Stat. Soc. Ser. B* **57**, 289–300.
- Qian, J. (1998) *Proc. Survey Res. Methods Section, Am. Stat. Assoc.*, 704–708.
- Alter, O., Brown, P. & Botstein, D. (2000) *Proc. Natl. Acad. Sci. USA* **97**, 10101–10106.
- Billington, C. J., Briggs, J. E., Harker, S., Grace, M. & Levine, A. S. (1994) *Am. J. Physiol.* **266**, R1765–R1770.
- McLaughlin, T., Hindges, R., Yates, P. A. & O’Leary, D. D. (2003) *Development (Cambridge, U.K.)* **130**, 2407–2418.
- Kang, F., Purich, D. L. & Southwick, F. S. (1999) *J. Biol. Chem.* **274**, 36963–36972.
- Chuang, J. Z., Yeh, T. Y., Bollati, F., Conde, C., Canavosio, F., Caceres, A. & Sung, C. H. (2005) *Dev. Cell.* **9**, 75–86.
- Dennis, G., Sherman, B. T., Hosack, D. A., Yang, J., Gao, W., Lane, H. C. & Lempicki, R. A. (2003) *Genome Biol.* **4**, R60.
- Shams, I., Avivi, A. & Nevo, E. (2005). *Comp. Biochem. Physiol.* **142**, 376–382.
- Gurnett, A. M., O’Connell, J. P., Harris, D. E., Lehmann, H., Joysey, K. A. & Nevo, E. (1984) *Protein Chem.* **3**, 445–454.
- Ashur-Fabian, O., Avivi, A., Trakhtenbrot, L., Adamsky, K., Cohen, M., Kajakaro, G., Joel, A., Amariglio, N., Nevo, E. & Rechavi, G. (2004) *Proc. Natl. Acad. Sci. USA* **101**, 12236–12241.