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# Mouse duplicate genes are as essential as singletons

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**Duplicate genes in mouse are widely thought to have functional redundancy, and to be less essential than singleton genes. We analyzed nearly 3900 individually knocked out mouse genes and discovered that the proportion of essential genes is ~55% in both singletons and duplicates. This suggests that mammalian duplicates rarely compensate for each other, and that the absence of phenotypes in mice deficient for a duplicate gene should not be automatically attributed to paralogous compensation.**

## Duplicates, singletons and redundancy

Duplicate genes occur in all organisms [1], especially in multicellular eukaryotes [2]. Because gene duplication is the primary source of new genes [3], there is enduring interest in understanding the function of each duplicate gene [4,5]. However, early mouse studies that ‘knocked out’ duplicate genes revealed only mild or even no phenotypes [6,7], prompting the hypothesis that many mouse duplicates are functionally redundant and, therefore, that it would be difficult to discern the function of each copy by knocking out individual genes [8–10]. This view was reinforced when genome-wide gene deletion experiments showed that 12.4% of duplicates, compared with 29.0% of singletons, are essential to the viability or fertility of the yeast *Saccharomyces cerevisiae* [11] (Figure 1a). Similarly, in the nematode *Caenorhabditis elegans*, 2.3% of duplicates, but 7.6% of singletons, show lethal phenotypes in genome-wide knock-down experiments by RNA interference (RNAi) [12] (Figure 1a). However, the presumption that removing a mouse duplicate gene generates milder phenotypes than removing a singleton gene was based on anecdotal evidence and has not been systematically verified. Because of the expense and effort required to generate knockout mice and the potential value of such studies in understanding and treating human diseases, this verification is important

because it could substantially affect the design and interpretation of mouse knockout experiments.

## Proportion of mouse essential genes

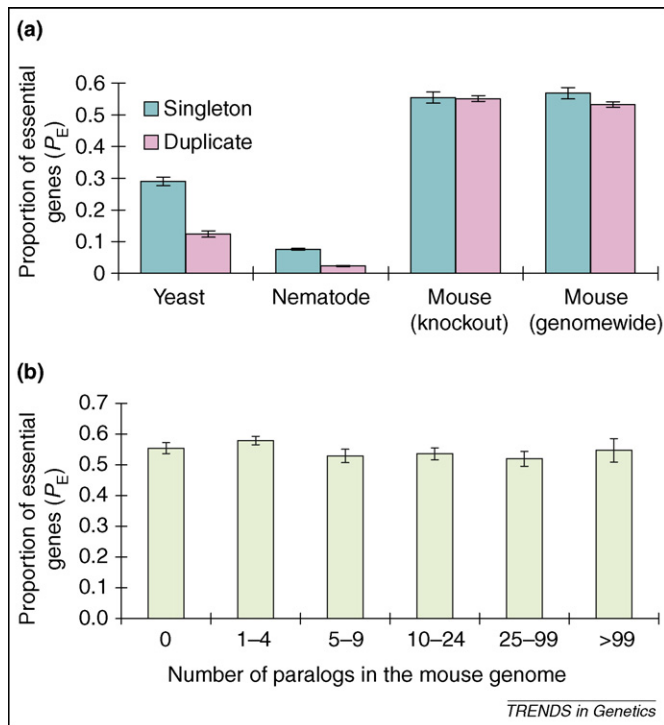
We examined the presumption that mouse duplicates are functionally redundant using a list of 3872 genes that have been individually knocked out from the mouse genome. Because there are numerous different mutant phenotypes and it is not easy to compare their severities, we separated all phenotypes into two categories based on the phenotype annotation by Mouse Genome Informatics (MGI 3.51; <http://www.informatics.jax.org>). If the deletion of a gene leads to either lethality before reproduction or sterility (i.e. fitness reduces to 0), the gene is referred to as essential (see [Methods in supplementary material online](#)). All other genes are considered as nonessential, because they are not essential to viability or fertility. With this classification, our dataset includes 2136 essential and 1736 non-essential genes. We also classified the 3872 genes into 3087 duplicate genes (Table S1 in [supplementary material online](#)), which have at least one duplicate in the genome, and 785 singleton genes (Table S2 in [supplementary material online](#)). Unexpectedly, we found that the proportion of essential genes ( $P_E$ ) is not significantly different between duplicate genes (55.1%) and singleton genes (55.4%) ( $P = 0.89$ ;  $\chi^2$  test; Figure 1a). Use of different criteria to define duplicate and singleton genes does not change this result qualitatively (Table S3 in [supplementary material online](#)). There is also no difference in  $P_E$  among genes belonging to small gene families and those belonging to large families (Figure 1b). These results differ from yeast and nematode genomic studies (Figure 1a) and contradict the widely held view that mouse duplicate genes are functionally redundant.

## Potential data biases

### Protein sequence divergence

Because most of the mouse gene knockouts were generated by individual laboratories for different purposes, rather than by a genome-wide systematic effort, it is important to consider potential biases in the data that might have led to

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**Figure 1.** Proportion of essential genes ( $P_E$ ) among singletons and duplicates. (a) Contrary to observations from yeast (*Saccharomyces cerevisiae*) and nematode (*Caenorhabditis elegans*), mouse (*Mus musculus*) duplicates and singletons are equally essential. Error bar indicates one standard error. The yeast results, based on gene deletion experiments, are from ref. [11], whereas the nematode results, based on RNAi knockdown experiments, are from ref. [12]. The mouse results labeled 'knockout' are based on the gene knockout data, whereas those labeled 'genomewide' are predicted from the knockout data after correction for potential data biases (see main text). (b) Essentiality of a mouse gene does not depend on how many paralogs it has in the genome (0, singletons;  $\geq 1$ , duplicates). Error bar indicates one standard error.

overestimation of the  $P_E$  for duplicates. In both yeast [11] and nematode [12],  $P_E$  increases as the protein sequence divergence between the knocked-out gene and its closest paralog increases, presumably because functional redundancy between duplicate genes decreases as the protein sequence divergence increases [11,12]. If this relationship also exists in mouse, and if mouse geneticists have been avoiding targeting genes with close paralogs, our sample of duplicates might have been biased for a greater  $P_E$  compared with the  $P_E$  of randomly chosen duplicate genes in the genome. To examine this possibility, we measured the protein sequence divergence between a gene and its closest paralog by the proportion of amino acid positions that differ between the two proteins ( $p$ -distance) (see [Methods in supplementary material online](#)). We plotted the frequency distribution of the  $p$ -distance for the duplicate genes in the knockout gene dataset and for all duplicates in the mouse genome. Indeed, duplicates in the knockout dataset are clearly underrepresented in low  $p$ -distance ranges, compared with all duplicates in the genome ( $P < 10^{-168}$ ,  $\chi^2$  test; [Figure 2a](#)). However, contrary to the observations in yeast [11] and nematode [12], mouse duplicates show a weak negative correlation between  $p$ -distance and  $P_E$  (Spearman rank correlation coefficient  $\rho = -0.063$ ,  $P < 10^{-3}$ ; see [Figure 2a](#) for binned results). Thus, the underrepresentation of knockout duplicates with low  $p$ -distances does not lead to an overestimation of  $P_E$  for duplicates.

### Expression divergence

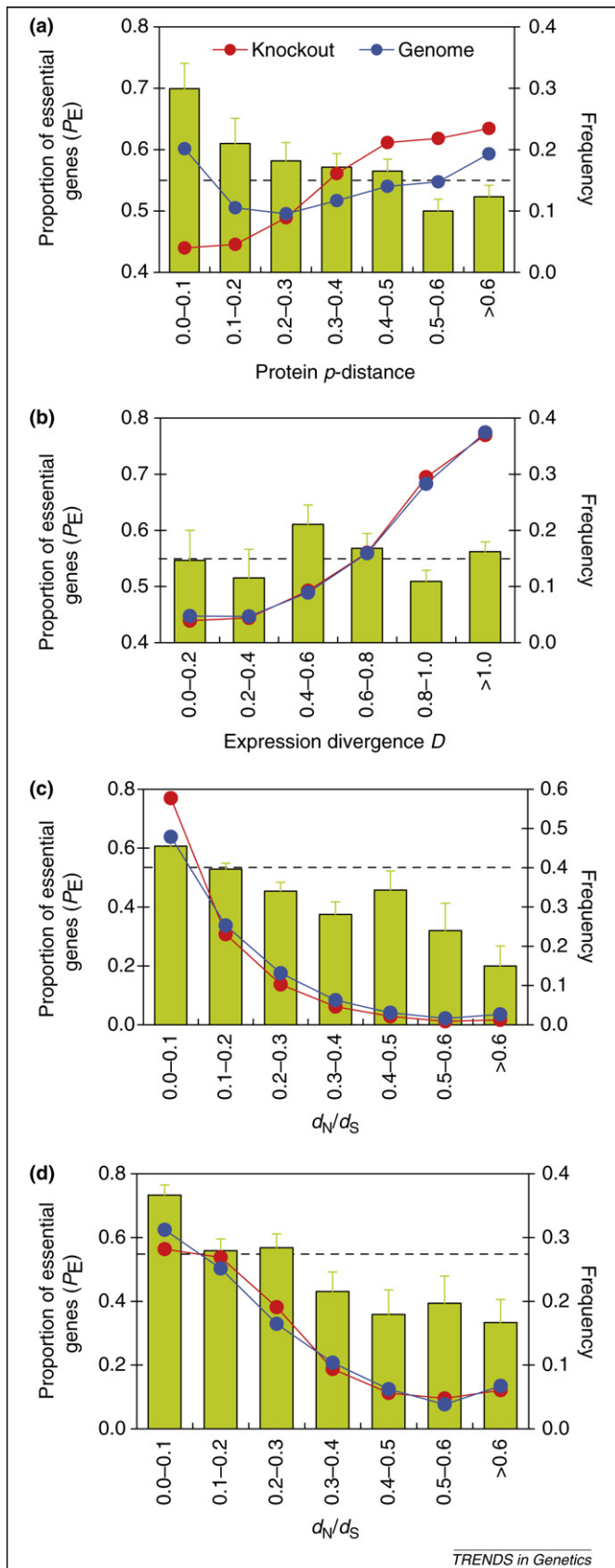
It is possible that duplicate genes with similar expression patterns have also been disfavored as knockout targets, because they might be more likely than those with dissimilar expression patterns to compensate for each other. We used  $D = 1 - r$  to measure the temporal and spatial expression-profile divergence between a duplicate gene and its closest paralog, where  $r$  is the Pearson correlation coefficient between microarray expression signals of the two genes across 61 mouse tissues at different developmental stages (see [Methods in supplementary material online](#)). However, we did not observe a significant correlation between  $D$  and  $P_E$  ( $\rho = -0.0031$ ,  $P > 0.5$ ). Furthermore, there was no difference in the distribution of  $D$  between the knockout data and the entire genome ( $P > 0.5$ ,  $\chi^2$  test; see [Figure 2b](#) for binned results). Thus, there is no need to correct the  $P_E$  estimate for duplicates here.

### Evolutionary conservation

It is also possible that genes chosen for 'knockout' experiments tend to be evolutionarily more conserved than other genes in the genome, because conserved genes might be involved in fundamental cellular processes that are of particular interest to biologists. A recent mouse study showed that the rate of sequence evolution is on average 25% lower in essential genes than in nonessential genes [13]. Thus, if genes targeted for knockout differ from the genomic average in evolutionary rate, a bias in  $P_E$  might result. We measured evolutionary conservation of each mouse gene by computing the ratio of the number of nonsynonymous substitutions per site ( $d_N$ ) to the number of synonymous substitutions per site ( $d_S$ ) between the gene and its rat ortholog (see [Methods in supplementary material online](#)). A low  $d_N/d_S$  indicates that a gene is evolutionarily conserved. Mouse genes without one-to-one rat orthologs were not considered. As expected,  $P_E$  decreases as  $d_N/d_S$  increases, for both duplicates ( $\rho = -0.17$ ,  $P < 10^{-18}$ ; [Figure 2c](#) for binned results) and singletons ( $\rho = -0.26$ ,  $P < 10^{-11}$ ; [Figure 2d](#) for binned results). Interestingly, duplicate genes targeted for knockout have lower  $d_N/d_S$  values than the genomic average ([Figure 2c](#)), whereas the opposite is true for singletons ([Figure 2d](#)), suggesting that  $P_E$  has been overestimated for duplicates but underestimated for singletons. Using the  $P_E$  estimate in each  $d_N/d_S$  bin for duplicates ([Figure 2c](#)), we estimated that a randomly picked duplicate gene from the mouse genome has an expected  $P_E$  of 53.2% ([Figure 1a](#)). Similarly, we estimated that a randomly picked singleton gene from the mouse genome has an expected  $P_E$  of 56.8% ([Figure 1a](#)). These two numbers are not significantly different from each other ( $P = 0.10$ ,  $\chi^2$  test).

### Gene essentiality, functional compensation and genetic robustness

Estimating the proportion of essential genes ( $P_E$ ) in a genome is not an easy task. For example, the nematode  $P_E$  values shown in [Figure 1a](#) are certainly underestimates because of frequent failure of RNAi to suppress gene expression completely [14], although the relative magnitudes of the  $P_E$  values for singletons and duplicates are probably correct



**Figure 2.** Factors that might influence the proportion of essential genes ( $P_E$ ) in the mouse knockout data. **(a)** Relationship between  $P_E$  in duplicate genes and the protein  $p$ -distance between the target gene and its closest paralog in the genome. **(b)** Relationship between  $P_E$  in duplicate genes and the expression divergence between the target gene and its closest paralog. **(c)** Relationship between  $P_E$  in duplicate genes and the evolutionary conservation of the gene, measured by the ratio of the nonsynonymous ( $d_N$ ) to synonymous ( $d_S$ ) nucleotide distances

[12]. In the case of mouse, a recent random *N*-ethyl-*N*-nitrosourea (ENU) mutagenesis targeting a 50 million-nt genomic region showed that 13.7% of mouse genes cause embryonic lethality when mutated [15]. The corresponding number from our knockout data is 14.0%. This comparison reconfirms the reliability of our estimate of mouse  $P_E$ . It is interesting that  $P_E$  seems to increase with organismal complexity, because it is  $\sim 7\%$  in *Escherichia coli* [16], 17% in yeast [17,18] and 55% in mouse, when the organisms were examined under relatively benign laboratory conditions. Apparently, complex organisms devote a greater fraction of their genes to ensure viability and fertility than simple organisms do.

Our analyses of the phenotypic data from  $\sim 15\%$  of genes in the mouse genome show that singletons and duplicates are equally likely to be essential to the organism. A similar result has been independently obtained by Liang and Li [19]. These findings strongly suggest that functional compensation between duplicates is rare in mouse, refuting the anecdote-based perception. In addition, there is no positive correlation of  $P_E$  and the protein sequence divergence between duplicates, providing further evidence for our conclusion. These findings differ from observations in yeast [11] and nematode [12] (Figure 1a). One possible reason is that the expression patterns of duplicates might diverge more rapidly in mouse than in yeast and nematode [20], because of the existence of many more cell types and developmental stages in mouse than in the other two species. Such expressional differences make functional compensation between duplicates unlikely even if the protein sequences of the duplicates are similar. However, we did not observe lower  $P_E$  for duplicates with lower expression divergence (Figure 2b), suggesting that the rapid expression divergence is probably not the main reason.

The second possible reason is that factors determining gene duplicability might be different in different species. It has been shown in yeasts that nonessential genes tend to duplicate, rendering  $P_E$  lower for duplicates than for singletons [21]. The absence of this mechanism in mouse [19] might explain our observation here. Regardless, our finding of low functional redundancy between mouse duplicate genes shows that there is no reason to avoid targeting duplicate genes in knockout experiments unless there are technical difficulties. Furthermore, the absence of expected phenotypes in mice deficient of a duplicate gene should not be automatically attributed to functional compensation from its paralogs.

Genetic robustness against null mutations seems to be a universal phenomenon of all living systems [22]. Such robustness can arise from the existence of duplicate genes with similar function and expression or from other reasons collectively known as ‘distributed robustness’ [22,23]. For

between the target gene and its rat ortholog. **(d)** Relationship between  $P_E$  in singleton genes and the evolutionary conservation of the gene, measured by  $d_N/d_S$  between the target gene and its rat ortholog. In each panel, the green bars indicate  $P_E$ , corresponding to the left y-axis. Error bars show one standard error. The red and blue lines show the frequency distributions of genes from the knockout dataset and the entire genome, respectively, and the values are indicated on the right y-axis. The horizontal black dotted line shows the observed  $P_E$  among singleton knockout genes. None of these potential biases alter the relationship between  $P_E$  of duplicates and  $P_E$  of singletons.

example, even in a metabolic network where no two enzymes catalyze the same reactions, some enzymes can be removed without rendering the network nonfunctional [22]. Our results indicate that duplicate genes have a negligible role in the genetic robustness of mouse. Furthermore,  $P_E$  of singleton genes is much greater in mouse than in yeast, indicating that the distributed robustness is also lower in mouse than in yeast.

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#### Supplementary data

Supplementary data associated with this article can be found, in the online version, at [doi:10.1016/j.tig.2007.05.006](https://doi.org/10.1016/j.tig.2007.05.006).

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