A Simple Method for Estimating the Strength of Natural Selection on Overlapping Genes

Xinzhu Wei and Jianzhi Zhang*

Department of Ecology and Evolutionary Biology, University of Michigan

*Corresponding author: E-mail: jianzhi@umich.edu.

Abstract

Overlapping genes, where one DNA sequence codes for two proteins with different reading frames, are not uncommon in viruses and cellular organisms. Estimating the direction and strength of natural selection acting on overlapping genes is important for understanding their functionality, origin, evolution, maintenance, and potential interaction. However, the standard methods for estimating synonymous (d_S) and nonsynonymous (d_N) nucleotide substitution rates are inapplicable here because a nucleotide change can be simultaneously synonymous and nonsynonymous when both reading frames involved are considered. We have developed a simple method that can estimate d_N/d_S and test for the action of natural selection in each relevant reading frame of the overlapping genes. Our method is an extension of the modified Nei-Gojobori method previously developed for nonoverlapping genes. We confirmed the reliability of our method using extensive computer simulation. Applying this method, we studied the longest human sense–antisense overlapping gene pair, *LRRC8E* and *ENSG00000214248*. Although *LRRC8E* (leucine-rich repeat containing eight family, member E) is known to regulate cell size, the function of *ENSG00000214248* is unknown. Our analysis revealed purifying selection on *ENSG00000214248* and suggested that it originated in the common ancestor of bony vertebrates.

Key words: synonymous substitution, nonsynonymous substitution, evolution.

Introduction

Overlapping genes generally refer to pairs of genes that overlap in their transcribed sequences. In this study, however, overlapping genes refer to pairs of genes that overlap in their protein-coding regions but use different reading frames. The first overlapping genes were discovered nearly 40 years ago in bacteriophage φ X174 (Barrell et al. 1976). Overlapping genes have since been found in numerous viruses and cellular organisms including multicellulars such as humans, and their functional importance has been demonstrated in some case studies (Giorgi et al. 1983; Normark et al. 1983; Chen et al. 1993; Veeramachaneni et al. 2004; Pavesi 2006; Chung et al. 2008; Dornenburg et al. 2010). In theory, two genes may overlap in one of the five possible phases (fig. 1), two being sense–sense (ss) and three being sense–antisense (sas). The sas11 phase, in which the second codon position in one gene faces the third codon position in the other gene (fig. 1), was reported to be the most common type (in prokaryotes), likely because this phase minimizes the mutual constraints of the protein sequences of the overlapping genes (Rogozin et al. 2002).

To study the functionality, origin, maintenance, and evolution of overlapping genes, it is often necessary to infer the direction and strength of natural selection acting on them. The standard approach for studying natural selection acting on protein-coding genes is by estimating the ratio between the rate of nonsynonymous nucleotide substitution (d_N) and that of synonymous nucleotide substitution (d_S). However, because a mutation may be simultaneously synonymous and nonsynonymous in overlapping genes, the commonly used methods for estimating d_S, d_N, and d_N/d_S are inapplicable. Several attempts have been made to estimate selection strengths in overlapping genes. Some authors treated a pair of overlapping genes as two nonoverlapping genes and calculated d_N/d_S for each gene independently using the standard methods (Yu et al. 2005; Pavesi 2006; Simon-Loriere et al. 2013). As pointed out long ago (Miyata and Yasunaga 1978), this approach is problematic, because a synonymous mutation to one of the overlapping genes may be nonsynonymous to the other gene and thus may be non-neutral. Realizing that the neutral expectation of d_N/d_S for each overlapping gene may not be 1, Nekrutenko et al. (2005) simply calculated d_N

SENSE-SENSE

ss12:
ORF1:	1-2-3-1-2-3-1
ORF2:	2-3-1-2-3-1-2

ss13:
ORF1:	1-2-3-1-2-3-1
ORF2:	3-1-2-3-1-2-3

SENSE-ANTISENSE

sas11:
ORF1:	1-2-3-1-2-3-1
ORF2:	1-3-2-1-3-2-1

sas12:
ORF1:	1-2-3-1-2-3-1
ORF2:	2-1-3-2-1-3-2

sas13:
ORF1:	1-2-3-1-2-3-1
ORF2:	3-2-1-3-2-1-3

Fig. 1.—Five phases of overlapping genes. Sense–sense overlap is abbreviated as "ss," whereas sense–antisense overlap is abbreviated as "sas." The two ss overlaps are equivalent if one switches the names of the two ORFs.

and d_S rather than their ratio, but they still applied a standard method directly to each overlapping gene. As such, the biological meanings of the estimated d_S and d_N are unclear. Rogozin et al. also noted the impact of one mutation on two genes and hence considered only sites that are 4-fold degenerate for one of the overlapping genes. Specifically, they were able to estimate d_N for each gene in gene pairs with the sas11 phase (Rogozin et al. 2002). But this method does not apply to all overlapping genes, and estimating d_S remains difficult (e.g., Rogozin et al. estimated d_S from nonoverlapping regions). Extending Goldman and Yang's (1994) method for nonoverlapping coding sequences, Sabath et al. (2008) developed a maximum likelihood (ML) method for simultaneous estimation of the selection intensity in each of the two overlapping genes. However, as currently implemented, the method cannot test whether d_N/d_S significantly differs from 1 for either gene (Sabath et al. 2008, 2012), rendering the utility of the method limited.

Here, we describe a simple method that estimates the selection strength of each of the two overlapping genes by separating the effects of each mutation on the two genes. Our method also estimates the associated variance, allowing a test of neutrality for each gene. We evaluate the performance of our method using computer simulation, and illustrate its utility by analyzing the human sas gene pair with the longest overlapping region.

Materials and Methods

Computer Simulation

Our new method for estimating the selection strengths in overlapping genes is described in the Results section. Here, we describe the simulation used to evaluate the performance of our method. To generate a pair of overlapping genes, we set the following parameters: the overlapping phase, the length of the overlapping region l, the ratio (R) between the number of transitions and number of transversions, the distance (d) between two sequences defined by the expected number of substitutions per neutral site, selection strength on open reading frame 1 (ORF1) (ω_1), and selection strength on ORF2 (ω_2). We generated an ancestral sequence that contained overlapping ORFs by first randomly choosing sense codons for the first ORF and then removing all stop codons until no stop codon is found in each ORF. We then introduced mutations following Kimura's (1980) two-parameter model with a preset R. The fixation probability of a mutation is determined jointly by ω_1 and ω_2. Specifically, if the mutation is synonymous in both ORFs, its fixation probability is set to be a ($0 < a < 1$); if the mutation is synonymous to ORF1 but nonsynonymous to ORF2, its fixation probability is $a\omega_2$; if the mutation is synonymous to ORF2 but nonsynonymous to ORF1, its fixation probability is $a\omega_1$; if the mutation is nonsynonymous to both ORFs, its fixation probability is $a\omega_1\omega_2$. The parameter a must be small enough so that $a\omega_1$, $a\omega_2$, and $a\omega_1\omega_2$ are all smaller than 1. Under this scheme, both positive and negative selection can be simulated. When negative selection is simulated for both ORFs, a can take any value between 0 and 1, but we assigned 0.9 to a to decrease the computational time. When positive selection is simulated for ORF1 but negative selection is simulated for ORF2, $0.9/\omega_1$ was assigned to a. If both ORFs are under positive selection, $0.9/(\omega_1\omega_2)$ was assigned to a. Each ancestral sequence was evolved independently to produce two derived sequences, by either accepting or rejecting the randomly generated mutations. Simulation ended when the number of mutations introduced equals the preset number (dl/a). ω_1 and ω_2 were then estimated by comparing the two simulated derived sequences. The scripts used for simulating overlapping genes and for estimating ω were written with Perl and are available at http://www.umich.edu/~zhanglab/download.htm (last accessed January 8, 2015).

Case Study

Annotation for human protein-coding genes and sequences used in the selection analysis were downloaded from Ensembl GRCh37 (http://useast.ensembl.org/, last accessed January 8, 2015). Overlapping genes were identified by comparing exon start and end positions of each gene on the same chromosome. For example, if exon 2 of gene A starts at position 13,780 and ends at 13,942 on Chromosome 1, and exon 5 of gene B starts at 13,950 and ends at 13,820 on the same

chromosome, we can infer that these two genes form a pair of sas overlapping genes and that the overlapping region between the two exons has $(13942-13820+1)=123$ bp. The overlapping genes analyzed were identified from Ensembl annotations using a Python script. Sequences were aligned using an online version of ClustalW2 (http://www.ebi.ac.uk/Tools/msa/clustalw2/, last accessed January 8, 2015). Transition/transversion ratio was calculated using MEGA5 (Tamura et al. 2011). The protein expression levels were from ProteomicsDB at https://www.proteomicsdb.org/ (last accessed January 8, 2015) (Wilhelm et al. 2014). The GenBank accession numbers of LRRC8 genes analyzed are provided in supplementary tables S1 and S2, Supplementary Material online. We used MEGA5 to reconstruct the neighboring-joining tree of LRRC8 genes using protein p distances.

Results

A New Method for Estimating the Selection Strength in Overlapping Genes

Because most species use double-stranded DNA, one segment of DNA can harbor at most six different ORFs. However, very rarely do all six ORFs coexist. Even in cases where all six ORFs coexist, it is unclear whether all ORFs code for actual proteins (Menon et al. 1990). The simplest and most common overlapping coding regions harbor two different ORFs, which can be either on the same strand (ss overlap) or on opposite strands (sas overlap) (fig. 1). The two types of ss overlap are in fact equivalent, because they both have the third codon positions of one ORF facing the first codon positions of the other ORF (fig. 1). Here, we use the ss overlap as an example to describe our method, but the same applies to all overlaps between two ORFs.

Our method is an extension of the modified Nei-Gojobori (mNG) method for estimating d_S and d_N in nonoverlapping genes (Nei and Gojobori 1986; Zhang et al. 1998), but considers the complication that one mutation simultaneously affects two ORFs, often with different effects. Let us consider a pair of homologous DNA sequences (e.g., respectively from human and mouse) that harbor overlapping ORF1 and ORF2. Our method for quantifying the selection strength in ORF1 and that in ORF2 involves the following four steps.

In the first step, we classify human nucleotide sites in the overlapping region into four categories depending on the impacts of potential mutations on the two ORFs. The four categories are referred to as NN, NS, SN, and SS sites, respectively, where N stands for nonsynonymous and S stands for synonymous. That is, if all potential mutations at a site cause nonsynonymous changes in both ORFs, it is an NN site, and so on. A site may belong to multiple categories and be called, for example, 1/3 NN site and 2/3 NS site, if one-third of potential mutations at the site cause nonsynonymous changes in both ORFs and two-thirds of potential mutations at the site cause

nonsynonymous changes in ORF1 but synonymous changes in ORF2. When considering potential mutations, it is important to separate transitions from transversions because they typically have different mutation rates and have different probabilities of causing nonsynonymous changes (Zhang 2000). Let R be the ratio between the number of transitional mutations and that of transversional mutations and be estimated from external information (e.g., from nonoverlapping regions or other genes). Hence, we consider a fraction of $R/(1+R)$ mutations to be transitions and the rest transversions (Zhang et al. 1998) in determining to which of the above four categories a site belongs. For instance, if the transitional mutation at a site causes a synonymous change in both ORFs and the two transversional mutations both cause a synonymous mutation in ORF1 and a nonsynonymous mutation in ORF2, this site is counted as $R/(R+1)$ SS site and $1/(R+1)$ SN site. We then calculate the total number of sites in the human overlapping region belonging to each of the four categories. The corresponding values are also calculated for the mouse sequence, and the averaged value from the two sequences for each category (L_{NN}, L_{NS}, L_{SN}, and L_{SS}) will be used subsequently.

In the second step, we classify all nucleotide differences between the two sequences into four categories: NN, NS, SN, and SS. That is, if a difference is nonsynonymous in both ORF1 and ORF2, it belongs to the NN group, and so on. When a nucleotide difference is in isolation, meaning that in neither ORF is there another difference in the same codon as the focal difference, the classification is straightforward. But when a codon (in either ORF) harbors two or more differences, the situation becomes complicated, because to determine the categories of the multiple differences, one has to consider all possible evolutionary pathways that can give rise to the observed nucleotide differences. In the case of nonoverlapping ORFs, there are two equally shortest evolutionary pathways between a pair of codon sequences with two differences (e.g., to evolve from AAA to AGG, one can go through AAG or AGA) and six equally shortest pathways when it harbors three differences (Nei and Gojobori 1986). For overlapping ORFs, however, one may need to consider a lot more pathways, because a codon in ORF1 overlaps with a codon in ORF2, which overlaps with another codon in ORF1, and so on. Thus, we need to find a segment of DNA in which each codon (defined by both ORFs) has multiple nucleotide differences with the exception of the codon at each end of the segment (fig. 2). When this segment has a total of m nucleotide differences between the pair of homologous sequences, a total of $m!$ pathways should be considered, each of which contains a unique order of m nucleotide changes. For each pathway, we count the number of nucleotide changes belonging to each of the four categories (NN, NS, SN, and SS). We average these numbers across all open pathways, which are pathways with no intermediate sequences that contain stop codons. An example is provided in supplementary figure S1, Supplementary Material online. After classifying all

Human CTTCATGGCCAGCTG

Mouse CTTCTAGCGCTCCTG

Fig. 2.—Determining the shortest overlapping region for mutational pathway consideration. Shown is an example of the ss overlap. Codons in ORF1 are marked with lines above the sequences, whereas codons in ORF2 are marked with lines below the sequences. Differences between the two species are in black, whereas identical nucleotides are in grey. The boxed region is the shortest region for mutational pathway consideration.

nucleotide differences between the pair of homologous sequences into the four categories, we count their numbers (M_{NN}, M_{NS}, M_{SN}, and M_{SS}, respectively).

In the third step, we calculate the proportion of sites with nucleotide differences by $p_{NN} = M_{NN}/L_{NN}$, $p_{NS} = M_{NS}/L_{NS}$, $p_{SN} = M_{SN}/L_{SN}$, and $p_{SS} = M_{SS}/L_{SS}$ for NN, NS, SN, and SS sites, respectively. The Jukes–Cantor formula (Jukes and Cantor 1969) may be used to correct for multiple hits. For instance, the number of nucleotide substitutions per site at NN sites can be estimated by $d_{NN} = -\frac{3}{4}\ln(1 - \frac{4p_{NN}}{3})$; d_{NS}, d_{SN}, and d_{SS} can be similarly estimated. Here, we used the Jukes–Cantor correction instead of more complex corrections such as Kimura's (1980) two-parameter model or Tamura and Nei (1993) model, because overlapping regions are usually so short that the variance of a distance estimate would be large under complex corrections (Nei and Kumar 2000).

In the fourth step, we propose that the strength of natural selection acting on ORF1 be estimated by $\omega_1 = d_{NN}/d_{SN}$ and that acting on ORF2 be estimated by $\omega_2 = d_{NN}/d_{NS}$. This formulation is based on two assumptions. First, synonymous mutations are neutral. Although not all synonymous mutations are neutral due to their potential impacts on DNA–protein interaction, pre-mRNA splicing, mRNA folding, translational efficiency, translational accuracy, and other aspects of cell biology (Chamary and Hurst 2005; Pagani et al. 2005; Warnecke and Hurst 2007; Qian et al. 2012; Park et al. 2013; Yang et al. 2014), most synonymous mutations may be considered largely neutral when compared with nonsynonymous mutations, especially in species with small effective population sizes (Li 1987; Ohta 1992). Second, the two overlapping genes do not have genetic interaction, such that the probability that a mutation gets fixed is the product of the probability with which it gets fixed in the absence of ORF1 and the probability with which it gets fixed in the absence of ORF2. This assumption implies that 1) NN-type mutations and SN-type mutations have comparable average effects on ORF2 and 2) NN-type mutations and NS-type mutations have comparable average effects on ORF1. Hence, ω_1 can be estimated by d_{NN}/d_{SN} and ω_2 can be estimated by d_{NN}/d_{NS}. In theory, we could also estimate ω_1 by d_{NS}/d_{SS} and estimate ω_2 by d_{SN}/d_{SS}. But, such estimates are usually subject to large sampling errors, because with the exception of the sas12 overlap that has a sizeable fraction of SS sites (fig. 1), overlapping regions

typically have few SS sites. Thus, unless otherwise noted, we do not use d_{SS} in this study. It is sometimes of interest to compare the selective pressures acting on the two overlapping genes. For this purpose, we can compute ω_1/ω_2, which equals d_{NS}/d_{SN}.

To calculate the variances of d_{NN}, d_{NS}, d_{SN}, and d_{SS}, the commonly used bootstrap method (Nei and Kumar 2000) is inapplicable because of the difficulty in bootstrapping codons from one ORF while maintaining the other ORF. We therefore extend an approximate analytical method previously developed for estimating the variances of d_S and d_N in the Nei–Gojobori method (Nei 1987), which is known to be quite accurate (Ota and Nei 1994). Following this method, we calculate the variance of d_{NN} by $\text{Var}(d_{NN}) = \text{Var}(p_{NN})/(1-4p_{NN}/3)^2$, where the variance of p_{NN} is given by $\text{Var}(p_{NN}) = p_{NN}(1-p_{NN})/L_{NN}$. Variances of d_{NS}, d_{SN}, and d_{SS} can be similarly estimated. Standard deviations (SDs) of d_{NN}, d_{NS}, d_{NS}, and d_{SS} are then estimated by taking the square root of their variances, respectively. The hypothesis of neutral evolution of ORF1 can be tested by a Z-test of the equality between d_{NN} and d_{SN}. That is, we can conduct a Z-test using $Z = (d_{NN} - d_{SN})/(\text{Var}(d_{NN}) + \text{Var}(d_{SN}))^{1/2}$. Similarly, the neutral evolution hypothesis for ORF2 can be tested by a Z-test of the equality between d_{NN} and d_{NS}. We can also test if the strengths of natural selection acting on the two ORFs are equal by a Z-test of the equality between d_{SN} and d_{NS}.

Performance of the New Method in Estimating the Selection Strengths in Overlapping Genes

To examine the performance of the new method, we conducted extensive computer simulation of overlapping genes of each phase. The overlapping region had 3,000 nucleotides, and the simulation was repeated 100 times under each parameter set. We used exceptionally long overlapping regions to minimize the sampling error such that potential biases of our estimators became more readily detectable. We start by describing the results obtained under the ss overlap. We first examined the situation that both overlapping genes are under purifying selection. We fixed $\omega_1 = 0.2$ and $\omega_2 = 0.5$ and studied how the distance between a pair of homologous sequences affects the accuracy of estimation (fig. 3A), where the distance is defined by the expected number of substitutions per neutral site between the two homologous sequences (i.e., the expected value of d_{SS}). We found that the mean ω_1 estimate and the mean ω_2 estimate are both slightly greater than their true values, and this excess in the estimated ω value appears unrelated to the distance. This bias may be due to the fact that we simulated sequence evolution using Kimura's two-parameter model, but estimated d_{NN}, d_{NS}, and d_{SN} using the Jukes–Cantor correction, which is known to undercorrect multiple hits in this scenario. When ω_1 and ω_2 are lower than 1, d_{SN} and d_{NS} are greater than d_{NN}, making the undercorrection more severe for the former than the latter
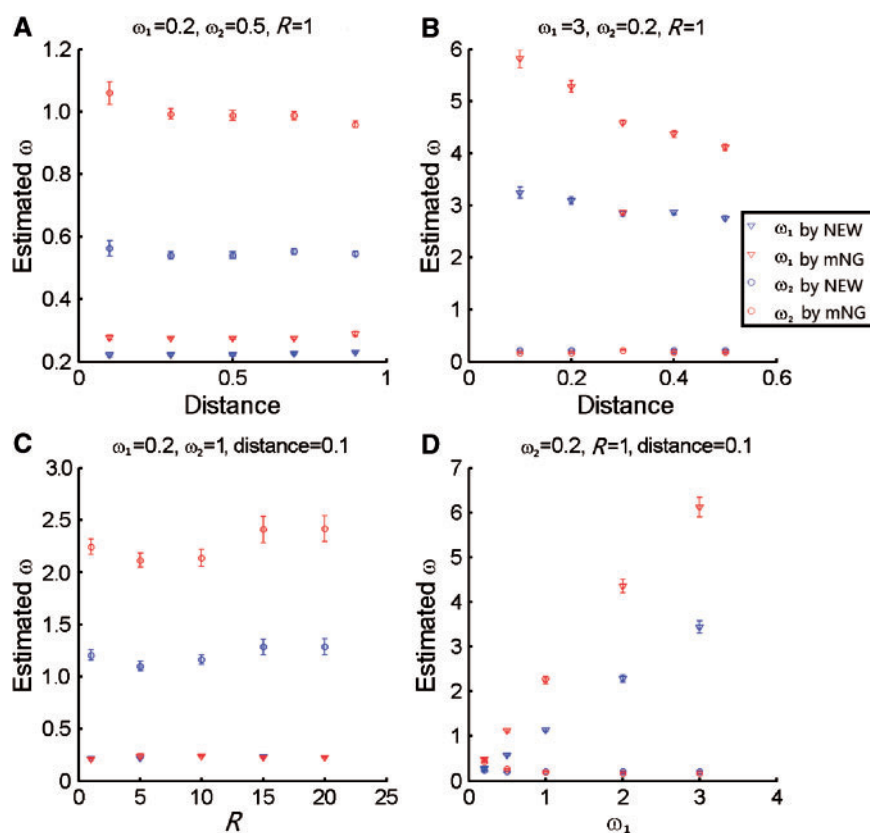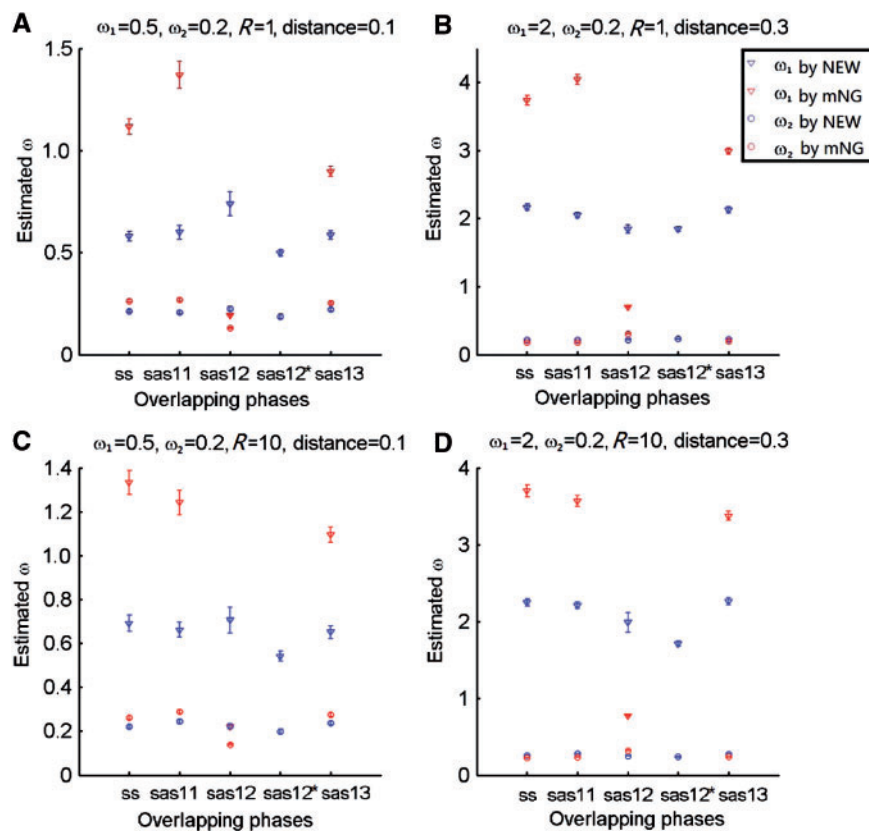
FIG. 3.—Performances of the new (NEW) method and modified Nei–Gojobori (mNG) method in estimating the selection intensities (ω_1 and ω_2) on overlapping genes. Shown are results from computer simulations of overlapping genes with the ss overlap. Each symbol represents the mean from 100 replications under a given parameter set, and error bars show the standard error. In each panel, the common parameters are listed above the panel, whereas the varying parameter is shown on the x axis. Distance is defined as the expected number of nucleotide substitutions per neutral site between the two sequences under comparison.

and the resultant ω_1 and ω_2 upward biased. Nevertheless, the biases appear to be generally lower than 10%. In contrast, if we estimate ω_1 and ω_2 by the mNG method without considering the mutual influences between overlapping genes, the estimates are much higher than their respective true values (fig. 3A). This is because some synonymous mutations to one ORF are nonsynonymous to the other ORF and hence have been removed by purifying selection, causing overestimation of ω_1 and ω_2. Because the true $\omega_1 < \omega_2 < 1$, ω_2 is overestimated to a larger extent than ω_1 (fig. 3A).

Next, we examined the situation that one overlapping gene is under positive selection ($\omega_1 = 3$), while the other is under purifying selection ($\omega_2 = 0.2$). We again found the mean estimates of ω_1 and ω_2 by our method to be close to their respective true values, for all levels of distance considered (fig. 3B). When the mNG method is used, ω_2 is slightly underestimated (fig. 3B), likely because some synonymous mutations to ORF2 are beneficial to ORF1 and are fixed by positive selection. In contrast, ω_1 is grossly overestimated by mNG (fig. 3B), for the reason mentioned in the previous paragraph.

We next examined the impact of the transition/transversion ratio R on estimates of ω_1 and ω_2 when their true values are 0.2 and 1, respectively (fig. 3C). We found both ω_1 and ω_2 slightly overestimated. This becomes moderately severe for ω_2 when $R \geq 10$, probably due to the aforementioned undercorrection of multiple hits by the Jukes–Cantor formula that is more serious when R gets higher. The mNG method performs similarly well as the new method in estimating ω_1 (fig. 3C), likely because of the lack of any selection on ORF2. But ω_2 is grossly overestimated by mNG (fig. 3C). Because ORF2 itself is not under any selection, the above phenomenon must be due to the fact that synonymous mutations to ORF2 are more likely than nonsynonymous mutations to ORF2 to be deleterious to ORF1.

We next varied ω_1 from 0.2 to 3.0 while keeping ω_2 at 0.2. We found estimates of ω_1 and ω_2 by our method to be generally reliable (fig. 3D). By contrast, ω_1 is consistently and grossly overestimated by mNG, whereas ω_2 is overestimated when $\omega_1 < 1$ and underestimated when $\omega_1 > 1$, as expected (fig. 3D).

Fig. 4.—Performances of the new (NEW) method and modified Nei–Gojobori (mNG) method in estimating the selection intensities (ω_1 and ω_2) on simulated overlapping genes of various phases indicated on the x axis. Each symbol represents the mean from 100 replications under a given parameter set, and error bars show the standard error. In each panel, the parameters are listed above the panel, whereas different overlapping phases are shown on the x axis. The results for sas12* are estimates using SS sites (i.e., $\omega_1 = d_{NS}/d_{SS}$ and $\omega_2 = d_{SN}/d_{SS}$) under the sas12 phase.

In addition to the ss overlap, we also examined the three sas overlapping phases with different parameter sets. We found that our method generated reliable results under all phases (fig. 4). In contrast, the mNG method can make grossly wrong estimates, and the direction and extent of the error depend on ω_1, ω_2, and the specific overlapping phase (fig. 4). For phase sas12, third codon positions in ORF1 overlap with third codon positions in ORF2. Consequently, the fraction of SS sites is higher than that in other phases, allowing the possibility of estimating natural selection using SS sites. We thus also estimated ω_1 by d_{NS}/d_{SS} and estimated ω_2 by d_{SN}/d_{SS} for phase sas12 (see sas12* in fig. 4). The results showed that these estimates are either similar to or slightly better than those using NN sites (see sas12 in fig. 4).

Because the analytical formulas for SDs are approximate, we used computer simulation to investigate their accuracies. For the ss phase, we examined the reliabilities of the analytically computed SD(d_{NN}), SD(d_{NS}), and SD(d_{SN}), but could not examine SD(d_{SS}) because of the paucity of SS sites. We conducted 100 simulation replications under each set of parameters. We then compared the SD among the 100 d_{NN} values

obtained and the mean of SD(d_{NN}) analytically calculated using the data from each simulation. The same was done for d_{NS} and d_{SN}. We found the analytically calculated SD values to be overall similar to the simulation observations, with statistically insignificant differences (fig. 5).

Evolutionary Analysis of the Human Gene Pair with the Longest Sense–Antisense Overlapping Region

To illustrate the utility of our method, we searched for an appropriate pair of overlapping genes from Ensembl for detailed analysis. We found that Ensembl annotates most ss overlapping genes with different reading frames as alternative splicing (Curwen et al. 2004), greatly underestimating the prevalence of ss overlapping genes. We thus focused on sas overlapping and identified the longest sas overlapping coding region in the human genome, containing 732 bases. The involved genes are *LRRC8E* (leucine-rich repeat containing eight family, member E) and an uncharacterized gene with an Ensembl Gene ID of *ENSG00000214248*. The structure of this gene pair (fig. 6A) shows that the entire 243 amino acid
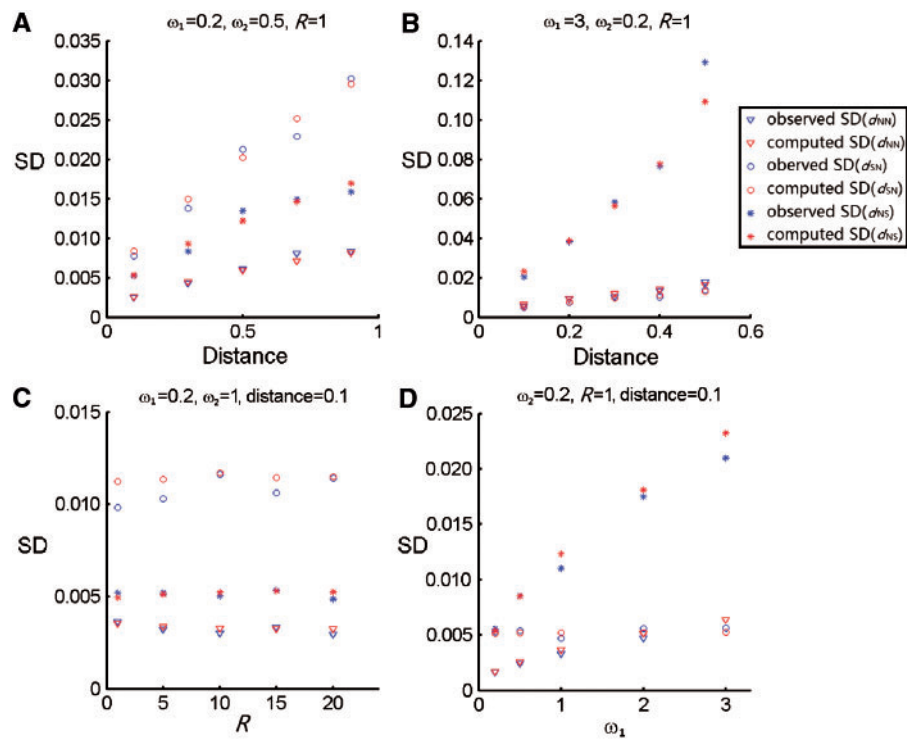
Fig. 5.—Performance of the new method in estimating the SD of d_{NN}, d_{NS}, and d_{SN}. Shown are the results from computer simulations of overlapping genes with the ss overlap. The analytically computed SD, averaged across 100 replications, is shown by red symbols, whereas the actual SD, observed from the 100 simulation replications, is shown in blue. In each panel, the common parameters are listed above the panel, whereas the varying parameter is shown on the x axis. Using 400 bootstrap samples of the 100 replicates under each parameter set, we derived a frequency distribution of the observed SD. We found that the mean computed SD is within the central 95% of the frequency distribution of the observed SD under all parameter sets examined.

coding region of *ENSG00000214248* lies within the second exon of *LRRC8E*, with the sas12 overlapping phase. It was recently discovered that LRRC8E functions as an essential component of the cell volume-regulated anion channel VRAC (Voss et al. 2014), but whether *ENSG00000214248* encodes a functional protein and what its function is are unknown.

We found from the recently published human proteomic data (Wilhelm et al. 2014) that *ENSG00000214248* is not only transcribed but also translated in coronary sinus and blood platelet (Fig. 6B). The protein expression sites of *ENSG00000214248* and those of *LRRC8E* overlap in blood platelet but are otherwise distinct (fig. 6B). The expression levels of the two proteins are generally comparable (fig. 6B). We acquired the sequences of the orthologous genes of human *ENSG00000214248* and *LRRC8E* from the macaque genome sequence. Using our method, we estimated the ω values for the two genes in the overlapping region as well as the ω in the nonoverlapping region of *LRRC8E*. R was estimated to be 3.61 from the nonoverlapping region of *LRRC8E* using Kimura's (1980) two-parameter model. We found that the overlapping region and the nonoverlapping region of *LRRC8E* have been under similar levels of purifying selection, with ω = 0.08 and 0.09, respectively. The ω for

ENSG00000214248 is 0.20, significantly lower than the neutral expectation of 1 (P < 0.002, two-tail Z-test), suggesting that this uncharacterized gene has been under purifying selection at least since the divergence between human and macaque. For the overlapping region, we used SS sites in the above estimation of ω values for *ENSG00000214248*, because there was no substitution at NS sites.

Because *ENSG00000214248* is entirely within *LRRC8E*, we traced the origin of *ENSG00000214248* by examining its presence in *LRRC8E* of various species. We were able to identify *LRRC8E* in all bony vertebrate genome sequences available at Ensembl and NCBI, but not in shark, lamprey, or any invertebrate genome. Interestingly, we also identified the ORF of *ENSG00000214248* within *LRRC8E* in most bony vertebrates, including zebrafish (fig. 6C). Apparently, *ENSG00000214248* already existed in the common ancestor of bony vertebrates, but was pseudogenized several times in subsequent evolution (fig. 6C). Because *LRRC8E* is a member of the *LRRC8* family that contains five genes in human, we reconstructed the phylogeny of this gene family (supplementary fig. S2, Supplementary Material online) to investigate if *ENSG00000214248* originated before *LRRC8E*. We discovered that the closest relative to *LRRC8E* is *LRRC8C*, which can be found in bony vertebrates and shark. However, the

GBE
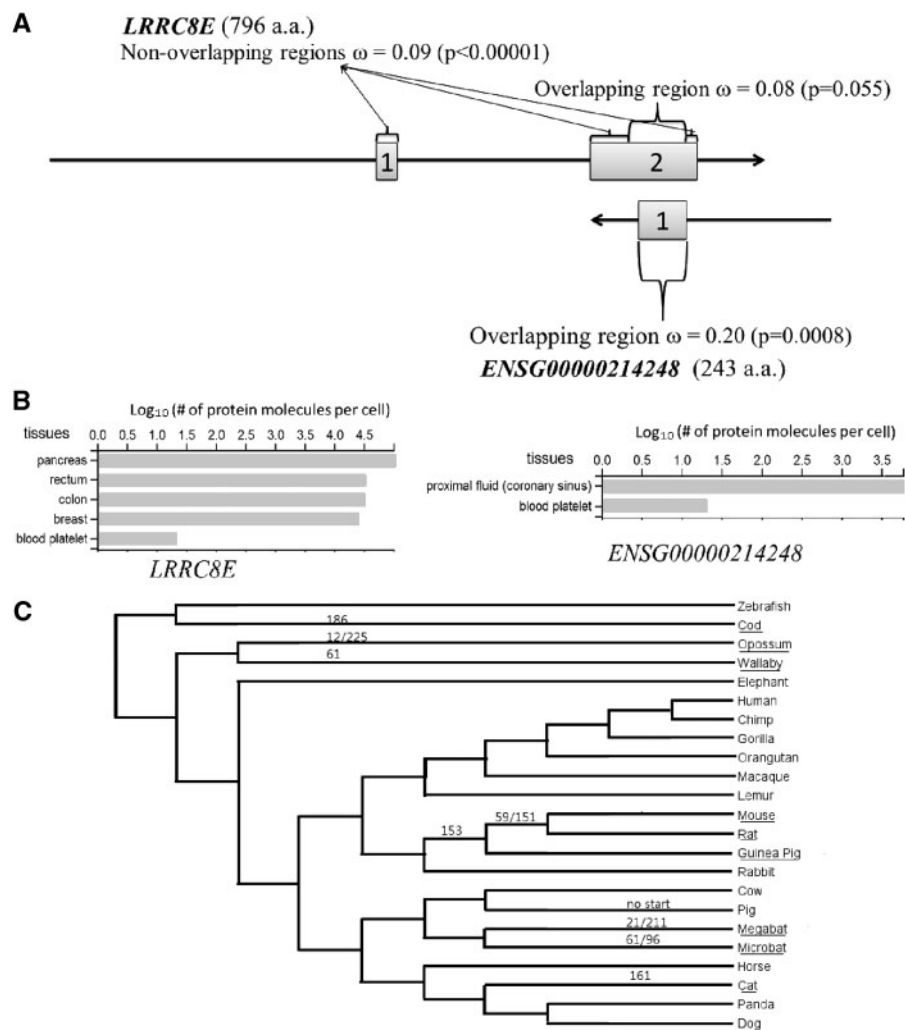
FIG. 6.—Evolution of the overlapping genes *LRRC8E* and *ENSG00000214248*. (*A*) The structures of the sas overlapping (sas12) genes of *LRRC8E* and *ENSG00000214248*. The ω values are estimated by comparing the human and macaque orthologs, with *P* values indicating the probabilities with which the null hypothesis of ω = 1 is true. (*B*) Protein expression levels of *LRRC8E* and *ENSG00000214248*. Median protein intensities from multiple samples, based on ProteomicsDB (Schwanhausser et al. 2011; Wilhelm et al. 2014), are shown for each tissue. (*C*) Evolution of *ENSG00000214248*. Species in which the ORF for *ENSG00000214248* is broken are underlined. Numbers on branches show the amino acid positions of premature stop codons. Branches are not drawn to scale.

presumable *ENSG00000214248* reading frame in *LRRC8C* contains several premature stop codons in each species examined (human, macaque, mouse, rat, zebrafish, and shark), suggesting that the common ancestor of *LRRC8C* and *LRRC8E* did not contain *ENSG00000214248*. Thus, the antisense reading frame probably originated in *LRRC8E* shortly after the birth of *LRRC8E* from the duplication of *LRRC8C*.

Discussion

Overlapping genes have been identified in many species and are particularly common in bacteria and viruses (Normark et al. 1983; Veeramachaneni et al. 2004), but their evolutionary studies have been hampered by the inapplicability of the standard methods for inferring natural selection acting on overlapping genes. We developed a simple method to estimate the selection strength on each of the overlapping ORFs and demonstrated the reliability of our method by computer simulation. Our method allows testing whether an overlapping gene is under natural selection and hence can be used to identify functional genes from hypothetical overlapping reading frames, as was demonstrated in the example of *ENSG00000214248*.

To more readily detect potential biases of our method, we simulated long overlapping regions (3,000 sites). In reality, however, overlapping regions are much shorter. We also

performed simulations using overlapping regions of 750 sites and 300 sites, respectively (supplementary fig. S3, Supplementary Material online), based on the parameters used in figure 3A and B. When the overlapping region is short and the distance is low, many sequences had no substitution in NS sites or SN sites, making our method inapplicable. For cases where our method did work, the mean ω estimates were reasonably good, although the standard errors were large, as expected (supplementary fig. S3, Supplementary Material online). Thus, accurately estimating ω values of short overlapping regions remains challenging unless the divergence between the two taxa compared is high. On the basis of current annotations of eukaryotic genomes, there are not many overlapping genes that have long evolutionary histories. However, as in the example studied, although the orthologs of human *ENSG00000214248* are present in many vertebrates, they have not been annotated outside primates. It is likely that much more overlapping genes and long-lasting overlapping genes than currently annotated exist. Overlapping genes are prevalent in viral genomes. Many viruses have high mutation rates, allowing the use of our methods even for relatively short overlapping regions.

Sabath et al. (2008) noted that the ML method they developed does not perform well under low distances (mean sequence divergence across sites < 8%). To examine if our method suffers from the same problem, we compared the two methods using the parameters in figure 3A and B. The results showed that the two methods are similar in their sensitivity to distance (supplementary fig. S4, Supplementary Material online). However, under both negative (supplementary fig. S4a, Supplementary Material online) and positive (supplementary fig. S4b, Supplementary Material online) selection, our method outperforms the ML method in terms of the accuracy of the ω estimates.

Although we introduced our method in the context of estimating the selective strength using interspecific comparisons, our method may also be applied to intraspecific data or comparisons between intraspecific and interspecific data. For instance, let us use D_{NN}, D_{NS}, D_{SN}, and D_{SS} to denote the numbers of the four types of substitutions in a pair of overlapping genes, respectively, and use P_{NN}, P_{NS}, P_{SN}, and P_{SS} to denote the corresponding numbers of the four types of polymorphisms, respectively. We can conduct a selection test similar to the McDonald–Kreitman test (McDonald and Kreitman 1991) for ORF1 by comparing D_{NN}, D_{SN}, P_{NN}, and P_{SN}, because D_{NN}/P_{NN} equals D_{SN}/P_{SN} under the null hypothesis of neutrality. Similarly, we can test selection in ORF2 by comparing D_{NN}, D_{NS}, P_{NN}, and P_{NS}. In addition to studying overlapping genes, our method can also be applied to the study of the functionality of certain alternative splicing. Alternative splicing is generally demonstrated by the existence of various transcripts from a gene, but the existence of a transcript is not a proof that the transcript is functional. For splice variants using alternative reading frames, our method may be used to test if the

alternative reading frame has been under natural selection, which would support the functionality of the splice variant.

In summary, we believe that our development of a simple method for estimating the selective strengths on overlapping genes will facilitate researches toward understanding the origin, evolution, and functionality of overlapping genes.

Supplementary Material

Acknowledgments

Literature Cited

Barrell BG, Air GM, Hutchison CA 3rd. 1976. Overlapping genes in bacteriophage phiX174. Nature 264:34–41.

Chamary JV, Hurst LD. 2005. Evidence for selection on synonymous mutations affecting stability of mRNA secondary structure in mammals. Genome Biol. 6:R75.

Chen HS, et al. 1993. The woodchuck hepatitis virus X gene is important for establishment of virus infection in woodchucks. J Virol. 67:1218–1226.

Chung BY, Miller WA, Atkins JF, Firth AE. 2008. An overlapping essential gene in the Potyviridae. Proc Natl Acad Sci U S A. 105:5897–5902.

Curwen V, et al. 2004. The Ensembl automatic gene annotation system. Genome Res. 14:942–950.

Dornenburg JE, Devita AM, Palumbo MJ, Wade JT. 2010. Widespread antisense transcription in *Escherichia coli*. MBio 1:e00024–e00010.

Giorgi C, Blumberg BM, Kolakofsky D. 1983. Sendai virus contains overlapping genes expressed from a single mRNA. Cell 35:829–836.

Goldman N, Yang Z. 1994. A codon-based model of nucleotide substitution for protein-coding DNA sequences. Mol Biol Evol. 11:725–736.

Jukes TH, Cantor CR. 1969. Evolution of protein molecules. In: Munro HN, editor. Mammalian protein metabolism. New York: Academic Press. p. 21–132.

Kimura M. 1980. A simple method for estimating evolutionary rates of base substitutions through comparative studies of nucleotide sequences. J Mol Evol. 16:111–120.

Li WH. 1987. Models of nearly neutral mutations with particular implications for nonrandom usage of synonymous codons. J Mol Evol. 24:337–345.

McDonald JH, Kreitman M. 1991. Adaptive protein evolution at the Adh locus in *Drosophila*. Nature 351:652–654.

Menon NK, et al. 1990. Cloning and sequencing of a putative *Escherichia coli* [NiFe] hydrogenase-1 operon containing six open reading frames. J Bacteriol. 172:1969–1977.

Miyata T, Yasunaga T. 1978. Evolution of overlapping genes. Nature 272:532–535.

Nei M. 1987. Molecular evolutionary genetics. New York: Columbia University Press.

Nei M, Gojobori T. 1986. Simple methods for estimating the numbers of synonymous and nonsynonymous nucleotide substitutions. Mol Biol Evol. 3:418–426.

Nei M, Kumar S. 2000. Molecular evolution and phylogenetics. New York: Oxford University Press.

Nekrutenko A, Wadhawan S, Goetting-Minesky P, Makova KD. 2005. Oscillating evolution of a mammalian locus with overlapping reading frames: an XLalphas/ALEX relay. PLoS Genet. 1:e18.

Normark S, et al. 1983. Overlapping genes. Annu Rev Genet. 17:499–525.

Ohta T. 1992. The nearly neutral theory of molecular evolution. Ann Rev Ecol Syst. 23:263–286.

Ota T, Nei M. 1994. Variance and covariances of the numbers of synonymous and nonsynonymous substitutions per site. Mol Biol Evol. 11: 613–619.

Pagani F, Raponi M, Baralle FE. 2005. Synonymous mutations in CFTR exon 12 affect splicing and are not neutral in evolution. Proc Natl Acad Sci U S A. 102:6368–6372.

Park C, Chen X, Yang JR, Zhang J. 2013. Differential requirements for mRNA folding partially explain why highly expressed proteins evolve slowly. Proc Natl Acad Sci U S A. 110:E678–E686.

Pavesi A. 2006. Origin and evolution of overlapping genes in the family Microviridae. J Gen Virol. 87:1013–1017.

Qian W, Yang JR, Pearson NM, Maclean C, Zhang J. 2012. Balanced codon usage optimizes eukaryotic translational efficiency. PLoS Genet. 8: e1002603.

Rogozin IB, et al. 2002. Purifying and directional selection in overlapping prokaryotic genes. Trends Genet. 18:228–232.

Sabath N, Landan G, Graur D. 2008. A method for the simultaneous estimation of selection intensities in overlapping genes. PLoS One 3: e3996.

Sabath N, Wagner A, Karlin D. 2012. Evolution of viral proteins originated de novo by overprinting. Mol Biol Evol. 29:3767–3780.

Schwanhausser B, et al. 2011. Global quantification of mammalian gene expression control. Nature 473:337–342.

Simon-Loriere E, Holmes EC, Pagan I. 2013. The effect of gene overlapping on the rate of RNA virus evolution. Mol Biol Evol. 30:1916–1928.

Tamura K, Nei M. 1993. Estimation of the number of nucleotide substitutions in the control region of mitochondrial DNA in humans and chimpanzees. Mol Biol Evol. 10:512–526.

Tamura K, et al. 2011. MEGA5: molecular evolutionary genetics analysis using maximum likelihood, evolutionary distance, and maximum parsimony methods. Mol Biol Evol. 28:2731–2739.

Veeramachaneni V, Makalowski W, Galdzicki M, Sood R, Makalowska I. 2004. Mammalian overlapping genes: the comparative perspective. Genome Res. 14:280–286.

Voss FK, et al. 2014. Identification of LRRC8 heteromers as an essential component of the volume-regulated anion channel VRAC. Science 344:634–638.

Warnecke T, Hurst LD. 2007. Evidence for a trade-off between translational efficiency and splicing regulation in determining synonymous codon usage in Drosophila melanogaster. Mol Biol Evol. 24: 2755–2762.

Wilhelm M, et al. 2014. Mass-spectrometry-based draft of the human proteome. Nature 509:582–587.

Yang JR, Chen X, Zhang J. 2014. Codon-by-codon modulation of translational speed and accuracy via mRNA folding. PLoS Biol. 12: e1001910.

Yu P, Ma D, Xu M. 2005. Nested genes in the human genome. Genomics 86:414–422.

Zhang J. 2000. Rates of conservative and radical nonsynonymous nucleotide substitutions in mammalian nuclear genes. J Mol Evol. 50:56–68.

Zhang J, Rosenberg HF, Nei M. 1998. Positive Darwinian selection after gene duplication in primate ribonuclease genes. Proc Natl Acad Sci U S A. 95:3708–3713.

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