Homo sapiens under Neutral Evolution

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Received March 3, 2014; Revised April 26, 2014; Accepted April 28, 2014
J-STAGE Advance published date: May 3, 2014

People often would like to think themselves as special existence. In studies on organismal evolution, unexpectedly many researchers believe in contribution of positive selection for evolution of characters which made us human. However, most of evolution at genomic level is neutral process, and this is also true for evolution at protein level. There is no exception in evolution toward human lineage. I would like to show various examples on this point, including studies of my group, and would like to confirm natural phenomenon that most of evolution which produced Homo sapiens was neutral. It should also be noted that majority of this paper is from “Introduction to Evolutionary Genomics” written by Saitou (Saitou N. Berlin: Springer; 2014).

Key words: evolutionary genomics, natural selection, human evolution

Neutral Evolution as Default Process of the Genome Changes

It is now established that the majority of mutations fixed during evolution are selectively neutral, as amply demonstrated by Kimura (1) and by Nei (2). Reports of many genome sequencing projects routinely mention neutral evolution in the 21st Century. I thus discuss neutral evolution as one of the basic processes of genome evolution in this paper.

Neutral evolution is characterized by the egalitarian nature of the propagation of selectively neutral mutants. Mutation is the ultimate source of diversity of organisms. If a mutation occurring in some gene modifies gene function, there is a possibility of heterogeneity in terms of number of offsprings. This is the start of natural selection. However, some mutations may not change gene function, and although they are somewhat different from parental type DNA sequences, mutants and parental or wild types are equal in terms of offspring propagation. We meet the egalitarian characteristic of the selectively neutral mutants. If all members of evolutionary units, such as DNA molecules, cells, individuals, or populations, are all equivalent, the frequency change of these types are dominated by random events. It is therefore logical that randomness is the most important factor in organismal evolution.

Our World is Finite

Randomness also comes in when abiotic phenomena are involved in organismal evolution. Earthquakes, volcanic eruptions, continental drifts, meteorite hits, and many other geological and astronomical events are not the outcome of biotic evolution, and they can be considered to be stochastic from organismal point of view.

Before proposal of the neutral theory of evolution by Kimura (3), randomness was not considered as the basic process of evolution. Systematic pressure, particularly natural selection, was believed to play the major role in evolution. This view is applicable if the population size, or the number of individuals in one population, is effectively infinite. However, the earth is finite, and the number of individuals is always finite. Even this whole universe is finite. This finiteness is the basis of the random nature of organismal evolution.

Basic Concept of Natural Selection

The fundamental source of evolution is mutation, or any change of genome sequences. Therefore, natural selection is tightly connected with the effect of mutations. If mutations are highly deleterious, they will soon disappear though natural selection. Before Darwin (4) proposed the possibility of natural selection as a creative power of evolution, natural selection in modern term has been considered as the mechanism to keep status quo, as initially created by the divine power. Elimination of deleterious mutations is now called ‘negative’ selection after Kimura (1). When the conservative nature of this process is stressed, it is called ‘purifying’ selection (5). To keep the current genetic entity thorough elimination of deleterious mutations is the core of the “struggle for existence”.

Most of mutations become extinct simply by chance, and only a small fraction exists for a long evolutionary

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time. If we focus on these mutations that contribute to evolution for a certain period, say 1000 generations, the majority is selectively neutral, and only a small fraction is kept through natural selection that favors advantageous mutations. Charles Darwin, together with Alfred Russel Wallace, proposed this type of natural selection as the mechanism of evolution in 1858. This process is therefore sometimes called ‘Darwinian’ selection, but nowadays ‘positive’ selection, after Kimura (1), is commonly used. Negative and positive selection sandwich the selective neutrality, the dominant situation in evolution.

Natural selection is tightly connected with two concepts: adaptation and fitness. Although ‘adaptation’ is widely used in many books on evolution, it is often not quantitatively defined, and we should be careful for using this word in the context of natural selection unless clearly defined. This word is also used to describe various characters of many organisms without testing whether those characters were products of positive selection. For example, ‘adaptive radiation’ has often been used on the evolution of many mammalian lineages as if adaptation caused by positive selection is the main factor for radiation. Recent studies, however, suggest that the geographical isolation caused by the continental drift seems to be the main cause of mammalian radiation. Any evolutionary biologist should have a discreet attitude on the usage of ‘adaptation’.

In contrast, ‘fitness’ is quantitatively defined in population genetics theory. There are two kinds of fitness; absolute and relative. Because the basis of evolution is the change of genetic constituents of organisms, natural selection should be discussed in terms of differential rates of reproduction among genotypes. The absolute fitness is the mean number of offsprings for one particular genotype. If this is larger than 1, the genotype is expected to increase its offspring if the effect of random genetic drift is negligible. Consideration of the absolute fitness is important, for we can directly discuss the temporal change of population size. It should be noted that the absolute fitness may change without mutations, if the environmental condition changes.

We are often interested in the relative success of one genotype to other ones, and consider the relative fitness. The relative fitness of one particular genotype is usually set to be 1, and that of other genotypes are expressed using selection coefficients. Genotypes are identical with alleles in haploid organisms, and the relative fitness for allele \( i \) may be written as \( 1 + s_i \), where \( s_i \) is selection coefficient for allele \( i \). Let us assume that the relative fitness of allele 0 is 1 \((s_0 = 0)\). If \( s_i (i > 0) \) is positive, allele \( i \) is advantageous compared to allele 0 and is positively selected, while allele \( i \) is deleterious compared to allele 0 and is negatively selected when \( s_i \) is negative. Allele \( i \) is selectively neutral with allele 0 when \( s_i = 0 \).

### Positive Selection for Ape and Human Genes

We now move to discuss positive selection. Because the author is familiar with studies on primates, let us first consider positive selection in primates. Hughes and Nei (6,7) showed that both MHC class I and class II genes experienced positive selection, probably of the overdominant type, through the comparison of synonymous and non-synonymous nucleotide substitutions (Dn/Ds test). This test is quite powerful to detect positive selection based on the neutral theory, and many studies were conducted to detect such selection. However, one drawback of this test is that it can detect only a limited type of genes under positive selection, and to obtain a statistically significant result, we need to have many substitutions. Therefore, a protein gene must have many variable amino acids, a pre-condition that may be satisfied in protein-coding genes that are involved in the immune system, such as MHC. If a single amino acid change was responsible in enhancing fitness, however, it would be difficult to detect it though the Dn/Ds test.

The majority of protein-coding genes have lower non-synonymous substitutions than synonymous ones, and only a small proportion is under positive selection. Although most of the genes were found to have experienced positive selection thorough the Dn/Ds test, the FOXP2 gene studied by Enard et al. (8) and Zhang et al. (9) did not show significantly higher Dn than Ds.

The FOXP2 gene was initially found in mouse (10), and its highly homologous human ortholog was shown to be responsible for hereditary orofacial dyspraxia associated with dysphasia (11). Interestingly, two nonsynonymous nucleotide substitutions at codon 304 (ACC:Thr⇒AAC:Asn) and at codon 326 (AAT:Asn⇒AGT:Ser) occurred after the human and chimpanzee divergence, and the DNA polymorphism study of modern human populations suggested that the FOXP2 gene region experienced a selective sweep after mutations occurred within modern humans (8,9). However, one of the two amino acid substitutions occurred in the human lineage after the divergence of the human-chimpanzee common ancestor also occurred in the carnivores (9). It is therefore not clear if these amino acid changes were truly responsible for the emergence of language. There was a report that FOXP2 of Neandethals also had these two amino acid substitutions (12). If this finding is true, two nonsynonymous substitutions occurred in the common ancestor of modern humans and Neandethals. Coop et al. (13) suggested two alternative scenarios for this; low rates of gene flow between modern humans and Neanderthals or contamination of modern human DNA in the putative Neanderthal genome. Mouse FOXP2 gene knockin experiment by Fujita et al. (14) and Enard et al. (15) both showed some phenotypic differences in vocalization. Konopka et al.
Pollard et al. (17,18) found a series of short DNA sequences which are highly conserved in vertebrates but show accelerated evolution only in human, and named them HARs (human accelerated regions). Prabhakar et al. (19) and Bird et al. (20) also conducted similar genome-wide studies. Prabhakar et al. (21) found one such sequence from the human genome, HACNS1 which includes 119-bp HAR2, and showed it to act as a limb bud enhancer with enhanced limb enhancer activity specifically in human. This change was caused by 13 human-specific substitutions within that region, and they interpreted that accumulation of these positively selected substitutions created multiple novel transcription factor binding sites (gain-of-function) and that the deposition of those facilitated the human-specific enhanced activity (21).

However, a GC-biased gene conversion may be an alternative explanation for fixation of such mutations causing loss-of-function in a repressor element within HACNS1 (22–24). GC-biased gene conversion is a consequence of DNA double strand break repair between homologous chromosomal regions, and the alleles from one chromosome are converted to the other with a bias of A or T to G or C (25). Neutral or even deleterious alleles could be fixed by GC-biased gene conversions (26). It is possible that the 13 human-specific substitutions were caused by a GC-biased gene conversion and resulted in a disruption of repressor function of the 81 bp region (loss-of-function), which may eventually enhance the activity of human HACNS1.

To evaluate the function of HACNS1, Sumiyama and Saitou (27) performed transgenic mouse assay by using the HACNS1 construct lacking the 13 human-specific substitutions. The deleted construct showed similar enhancer activity to the intact human HACNS1. This result suggests that the function of HACNS1 is not an activating enhancer, but rather a disrupted repressor. If so, loss-of-function in the HACNS1, possibly via a GC-biased gene conversion, not via positive selection, played an important role in human-specific evolution.

Kryukov et al. (28) reported that the selective pressure affecting the evolution of regulatory elements in the hominid lineage was significantly relaxed compared with that of the rodent lineage. Keightley et al. (29) suggested that regulatory elements in hominids may be diverging at a neutral evolutionary rate. All these studies discussed in this section revealed the difficulty in detecting evidence of positive selection in one lineage. We therefore should be careful for any study which jumped to a conclusion of accelerated evolution without carefully examining an alternative neutral evolution scenario.

Detection of Positive Selection through Genome-wide Searches

Let us now discuss on positive selection on modern humans. Adaptation to a high altitude environment has long been an interesting subject in human genetics. Recently, three independent genome-wide studies (30–32) found that the EPAS1 gene showed the highest differentiation between Han Chinese and Tibetans who are believed to adapted to high altitude. This gene codes a transcription factor HIF–2α involved in the induction of genes regulated by oxygen. The EGLN1 gene, which encodes HIF-prolyl hydroxylase 2, was the second-best differentiated gene. EGLN1 catalyzes the post-translational formation of 4-hydroxyproline in HIF–α proteins.

Moreno-Estrada et al. (33) compared ~11,000 human genes with their orthologs in chimpanzee, mouse, rat, and dog, and found 11 genes as showing the signatures of positive selection on the human lineage through a branch-site likelihood method (34). These genes were then analyzed for signatures of recent positive selection using SNP data in modern humans. One SNP every 5–10 kb inside each candidate gene and up to around 30 kb in both upstream and downstream flanking regions, plus additional SNPs around 200 kb in both flanking regions were selected, and a total of 223 SNPs were typed for 39 worldwide populations from the HGDP-CEPH diversity panel (35). They also analyzed 4,814 SNP data distributed along 2 Mb centered on each gene from the HGDP-CEPH panel. Through examination of allele frequency spectrum, population differentiation, and the maintenance of long unbroken haplotypes, they found signals of recent adaptive phenomena in only one gene region. The signal of recent positive selection came from a neighboring gene CD5, which codes a transmembrane receptor expressed in the T-cell surface (33). This careful study suggests that most of positively selected genes in modern humans are involved in the immune system. It is not surprising, for our ancestors have suffered many sorts of infectious diseases, and the human immune system-related genes confronted the battle against bacteria, virus, or parasitic eukaryotes. When we discuss positive selection on modern humans, we should consider natural selection on the interaction with other organisms, before attempting to apply other types of natural selection such as sexual selection.

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