ered common in real food webs (6–9). The cascade model assumes a hierarchical relationship between species that constrains the trophic role as consumer or resource: hierarchically higher species are always feeding on species with a lower ranking. This model cannot generate any loops, but explains some important properties of natural food webs such as the species richness among trophic levels (18).


24. Iterated simulation runs with randomly chosen parameter sets (\( f_r = 0.0 \) to 0.1, \( f_r = 0.0 \) to 1.0, \( s_r = 1.0 \)), initial abundances (\( X_i = 0.0 \) to 0.1) and linkage patterns, and measured the frequency of all species co-occurring (\( X_i > 1 \times 10^{-1} \) for all \( i \)) after sufficiently long periods (\( t = 10^4 \) by which time community persistence reaches an asymptote) in \( 10^4 \) runs. As in other models (1, 3–5), every species persists (with positive equilibrium biomass, \( r/s \) ) in the absence of interspecific interaction (food web being always persistent when \( C = 0 \)), they are either autotrophs or use external resources (35). This assumption avoids a confounding effect (36) of an increase in interspecific links decreasing the number of heterotrophic species with no potential diet present in the web.


**Phylogenetic Shadowing of Primate Sequences to Find Functional Regions of the Human Genome**

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Nonhuman primates represent the most relevant model organisms to understand the biology of *Homo sapiens*. The recent divergence and associated overall sequence conservation between individual members of this taxon have nonetheless largely precluded the use of primates in comparative sequence studies. We used sequence comparisons of an extensive set of Old World and New World monkeys and hominoids to identify functional regions in the human genome. Analysis of these data enabled the discovery of primate-specific gene regulatory elements and the demarcation of the exons of multiple genes. Much of the information content of the comprehensive primate sequence comparisons could be captured with a small subset of phylogenetically close primates. These results demonstrate the utility of intraprima sequence comparisons to discover common mammalian as well as primate-specific functional elements in the human genome, which are unattainable through the evaluation of more evolutionarily distant species.

Genomic sequence comparisons between distant species have been extensively used to identify genes and determine their intron-exon boundaries, as well as to identify regulatory elements present in the large noncoding fraction of the genome (1–3). This strategy has been successful in human-mouse comparisons, because the ~75 million years (My) of separation from their last common ancestor have provided sufficient time for a large fraction of nucleotides to have been exposed to considerable mutation and selection pressure. Although such comparisons readily identify regions of the human genome performing general biological functions shared with evolutionarily distant mammals, they will invariably miss recent changes in DNA sequence that account for uniquely primate biological traits.

As a consequence of their short evolutionary separation (apes 6 to 14 My, Old World monkeys 25 My, New World monkeys 40 My) (4), there is a paucity of sequence variation between humans and each of their nearest primate relatives. This lack makes it difficult to distinguish functional from passive conservation on the basis of pairwise comparisons, thus limiting the usefulness of such comparisons. However, the additive collective divergence of higher primates as a group (fig. S1) is comparable to that of humans and mice. This suggests that deep sequence comparisons of numerous primate species should be sufficient to identify important regions of conservation that encode functional elements. Phylogenetic footprinting (5, 6) has been used to identify highly conserved putative regulatory elements, exploiting alignments across numerous evolutionarily distant species. We developed a variant of phylogenetic footprinting, which we termed phylogenetic shadowing. In contrast to footprinting, phylogenetic shadowing examines sequences of closely related species and takes into account the phylogenetic relationship of the set of species analyzed. This approach enabled the localization of regions of collective variation and complementary regions of conservation, facilitating the identification of coding as well as noncoding functional regions.

We first examined the ability of this strategy to identify functional regions with precise locations within the human genome, such as intron-exon boundaries. The lack of clone-based libraries for multiple primate species limited us to sequencing orthologous regions from a large set of primates, using genomic DNA as template (7). The sole criterion used in the selection of the four different regions that we studied was that each should contain at least one annotated exon. The sequences were generated (8) for a set of 13 to 17 primate species that included those evolutionarily closest to humans, such as Old World and New World monkeys and hominoids, but not distant primates such as prosimians. The resulting sequences were analyzed to determine the likelihood ratio under a fast- versus slow-mutation regime for each aligned nucleotide site across all four regions analyzed (supporting online text) (8). This ratio represents the relative likelihood that any given nucleotide site was subjected to a faster or slower rate of accumulation of variation and is related to functional constraints imposed on each site. The corresponding likelihood ratio curves were used to describe the variation profile of the four genomic intervals analyzed.

In all regions examined, the exon-containing sequences displayed the least amount of cross-species variation, in agreement with the constraint imposed by their functional role (Fig. 1, A to D). A limited number of short regions of minimal variation similar to the exon-containing sequences appeared in the likelihood plots.
These regions, however, were all less than 50 base pairs (bp) long, making them unlikely to be candidate exons for alternative splicing (the vast majority of internal exons predominantly range between 50 and 200 bp in length) (9). These regions may represent previously unidentified regulatory elements. In agreement with observations from other parts of the genome (10), the four sequenced regions have evolved at different rates, as indicated by their differing absolute likelihoods. Comparison of human-mouse versus the multiple-primate visualization plots (Fig. 1, a to d) yields similar results, with exons showing the highest level of conservation in all regions studied. The primate sequence comparisons illustrate the effectiveness of phylogenetic shadowing in yielding a precise identification of the exon boundaries. In the case of the cholesterol ester transfer protein gene (Fig. 1, B and b), the human-mouse comparison is unavailable, because this gene has been inactivated in the mouse. This inactivation represents an extreme example of the frequently encountered situation where the mouse genome, owing to the lack of meaningful alignments between human and mouse sequences, fails to localize functional elements in the human genome (11).

We next analyzed the sequence data for the four regions studied to assess the contribution of additional species to the discriminative power of phylogenetic shadowing and to identify the most informative minimum subset of species (8). The sequences from the most informative subset of only four to seven species, depending on the genomic locus, were calculated to be sufficient to capture ~75% of the total available discriminative power of this approach (Table 1). We were able to unequivocally identify the position of exon 3 of liver X receptor–α from the five most informative species (Homo, Saguinus, Colobus, Callicebus, Allenopithecus), as indicated by the likelihood curve (Fig. S2). Comparison with Fig. 1C shows that additional species only marginally improve this plot. As would be predicted, the species included in the set maximizing the discriminative power of phylogenetic shadowing include representatives of the different clades that compose the primate phylogenetic tree and therefore constitute the least related species in the set studied (Fig. S1).

To investigate the ability of phylogenetic shadowing to discover primate-specific gene-regulatory elements, we studied apolipoprotein (apo) [apo(a)], a recently evolved primate gene whose orthologous distribution is limited to Old World monkeys and hominoids (12). Defining the regulatory sequences determining apo(a) expression levels is of considerable biomedical relevance, because high plasma levels of this protein serve as an important cardiovascular disease risk predictor (13, 14). The sparse distribution of this gene among mammals precludes classical comparative genomic approaches. Sequence comparison of the apo(a) locus in humans, chimps, and baboons revealed a region of extreme conservation of ~1.6 kb adjacent to the transcription start site, surrounded by ~8 kb of reduced conservation on either side, owing to the different patterns of repetitive-element insertion in the three species. We sequenced and analyzed this 1.6-kb region in 18 Old World monkeys and hominoids (8). Phylogenetic shadowing revealed regions of varying conservation similar to that observed for the intron-exon regions described previously (Fig. 2A). In addition to the region containing the first exon of apo(a) (region E) (15), the remaining regions with the lowest variation include the apo(a) promoter’s TATA box (region C9) and a critical hepatocyte nuclear factor (HNF)–1α transcription factor binding site (region C10), both of which have been functionally described (16).

Because these elements were located immediately upstream of the apo(a) promoter, we
of the protein-bound oligonucleotides relative to the mobility of the pure oligonucleotides (Fig. 2B, lower bands). Conversely, oligonucleotides from nonconserved regions showed only weak or no protein binding. Overall, binding of oligonucleotides from conserved regions with hepatic nuclear proteins was more than six times as strong as binding of oligonucleotides from regions with lower conservation (Fig. 2C, regions C1 to C10.2 versus N1 to N7).

Consistent with the prediction that conserved regions interact preferentially with DNA binding proteins is the prediction that these regions are involved in transcriptional regulation activity. To explore this possibility, we developed an assay for in vitro enhancer activity based on transient transfection of a human liver cell line. The 10 most conserved regions, as well as the 7 least conserved regions, were individually deleted from the whole 1.6-kb region (8). These constructs were compared with the intact 1.6-kb region construct for their ability to drive the expression of a luciferase reporter gene. Three independent experimental determinations reproducibly indicated that conserved regions had a much larger functional impact than did nonconserved regions in these reporter-gene expression assays (8). The deletions of the conserved sequence elements reduced the expression of the full reporter by 25 to 55% (17), with the exception of the construct carrying the deletion of region 6 (Fig. 3). In contrast, deletions of all but one of the seven nonconserved regions had a minimal effect on the expression of the full reporter construct, reducing its levels by a median value of only 4%.

The analysis of closely related primates facilitates the discovery of functional elements specific to primates as well as elements shared with evolutionarily distant mammals. The high absolute degree of similarity minimizes ambiguity in the computation of the multiple alignment, which in turn greatly facilitates the subsequent construction of the phylogenetic tree (18). Furthermore, the generalization of gene-finding algorithms to multiple organism annotation is simplified, which results in more accurate predictions (supporting online text). The facility of sequence alignments and the possibility of comparative assembly for nonhuman primates, with human as the reference sequence, have important
practical relevance, because they considerably diminish the depth of sequence coverage required for an organism to be informative in annotating the human genome.

Evaluation of our data suggests that sequence from as few as four to six primate species in addition to humans is sufficient for the identification of a large fraction of functional elements in the human genome, many of which are likely to be missed by human-mouse comparisons. With the goal of annotating the human genome, these studies indicate that the generation of a limited amount of sequence from a few additionally selected primate genomes will provide information previously unavailable from comparisons of humans with more evolutionarily distant species.

References and Notes
7. Genomic DNA from the following species was analyzed in this study: Pan troglodytes (*), Pan paniscus (*), Macaca mulatta (*), Macaca nemestrina (*), Macaca fascicularis (*), Ateles geoffrey (*), Callicebus moloch (*), Aotus trivirgatus (*), Callithrix jacchus (*), Hylobates syndactylus (*), Hylobates lar (*), Hylotes lamar (*), Pygathrix nemaeus (*), Trachypithecus frisvani (*), Presbytis entellus (*), Nasalis larvatus (*), Atelates geoffrey (*), Callithrix jacchus (*), Aotus trivirgatus (*), Callithrix (*), and Saguinus labiatus (*).
8. For a complete list of species, see Table S1.
15. This region was used to learn the mutation rates for more evolutionarily distant species.
17. Deletion of region 9, containing the promoter’s TATA box, reduces expression by only 30%, likely attributable to the presence of one of the several cryptic TATA boxes further upstream.
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21. Supporting Online Material
www.sciencemag.org/cgi/content/full/299/5611/1391/DC1

Materials and Methods
SOM Text
Figs. S1 and S2
References
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EDEM As an Acceptor of Terminally Misfolded Glycoproteins Released from Calnexin
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Terminally misfolded proteins in the endoplasmic reticulum (ER) are retrotranslocated to the cytoplasm and degraded by proteasomes through a mechanism known as ER-associated degradation (ERAD). EDEM, a postulated Man8BB-binding protein, accelerates the degradation of misfolded proteins in the ER. Here, EDEM was shown to interact with calnexin, but not with calreticulin, through its transmembrane region. Both binding of substrates to calnexin and their release from calnexin were required for ERAD to occur. Overexpression of EDEM accelerated ERAD by promoting the release of terminally misfolded proteins from calnexin. Thus, EDEM appeared to function in the ERAD pathway by accepting substrates from calnexin.

The endoplasmic reticulum (ER) possesses a quality-control mechanism that discriminates correctly folded proteins from misfolded proteins (1). The misfolded proteins are dislocated to the cytosol through the translocon and degraded by the ubiquitin-proteasome system known as ER-associated degradation (ERAD) (1–4). It has been established that formation of the mannose–N-acetylglucosamine Man8GlcNAc2 isomer B of N-linked oligosaccharides is the critical luminal event as a signal for ERAD (3–5). EDEM is an ER stress-inducible membrane protein with homology to α-mannosidase but lacking mannosidase activity. Overexpression of EDEM greatly accelerates ERAD, most likely through recognition of the substrate-bound Man8B structure, because the mannosidase inhibitor kifinensine abolished the effects (3). However, the precise role of EDEM in the quality-control machinery is unknown.

In an attempt to search for proteins that work in concert with EDEM, we focused on calnexin (CNX), an ER protein with a potential role in the ERAD pathway (10–12). Although CNX is an abundant transmembrane component with an affinity to monoglucosylated oligosaccharides (13, 14), its role in ERAD remains controversial (15–17). We first assessed whether CNX can physically interact with EDEM. Immunoprecipitation of cell lysates with antibodies to CNX and hemagglutinin (HA) revealed that pulse-labeled EDEM-HA was coprecipitated with the antibody to CNX (Fig. 1A). Similarly, immunoprecipitation followed by immunoblotting of cells expressing EDEM-myc also showed that EDEM-myc bound to CNX (Fig. 1B). The observed association was not a result of the chaperone function of CNX, because (i) EDEM remained in the CNX complex even after 1 hour of chase (Fig. 1A), and (ii) EDEM was still coprecipitated with CNX even after cells were treated with castanospermine, an inhibitor of glucosidase, which is known to prevent the binding of substrate proteins to CNX.

Next, we tested whether EDEM interacts with calreticulin (CRT), a soluble homolog of CNX that is present in the ER lumen (13, 14) (Fig. 2A). EDEM-HA was not coprecipitated with the antibody to CRT (Fig. 2B), suggesting that there was no interaction between EDEM and CRT. Although CNX and CRT appear to share similar functions, CNX contains a transmembrane region that CRT lacks (Fig. 2A), suggesting that the transmembrane region might be responsible for the binding of EDEM to CNX. CRT that anchored to the ER membrane by fusing it with the membrane region of CNX (CRT-CNXXm) was revealed to coprecipitate with EDEM-myc (Fig. 2C). On the contrary, soluble CNX without its cytoplasmic and transmembrane segments (FS-CNXX) was not coprecipitated with EDEM-myc (Fig. 2D).