

***eShadow*: a tool for comparing closely related sequences**

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ABSTRACT

Primate sequence comparisons are difficult to interpret due to the high degree of sequence similarity shared between such closely related species. Recently, a novel method, *phylogenetic shadowing*, has been pioneered for predicting functional elements in the human genome through the analysis of multiple primate sequence alignments. We have expanded this theoretical approach to create a computational tool, *eShadow*, for the identification of elements under selective pressure in multiple sequence alignments of closely related genomes, such as in comparisons of human to primate or mouse to rat DNA. This tool integrates two different statistical methods and allows for the dynamic visualization of the resulting conservation profile. *eShadow* also includes a versatile optimization module capable of training the underlying Hidden Markov Model to differentially predict functional sequences. This module grants the tool high flexibility in the analysis of multiple sequence alignments and in comparing sequences with different divergence rates. Here, we describe the *eShadow* comparative tool and its potential uses for analyzing both multiple nucleotide and protein alignments to predict putative functional elements. The *eShadow* tool is publicly available at <http://eshadow.dcode.org/>.

Introduction

Cross-species sequence comparisons between distantly related genomes, such as those of humans and rodents, have been instrumental in defining evolutionarily conserved elements with critical biological roles, whether they function as coding exons (Gilligan *et al.*, 2002; Pennacchio *et al.*, 2001), regulatory elements (Ghanem *et al.*, 2003; Loots *et al.*, 2000; Nobrega *et al.*, 2003), or microRNAs (Lim *et al.*, 2003). Four widely used tools intended for generating genomic alignments are web-accessible and augmented by easy-to-use graphical interfaces: *Blast2Seq* (Tatusova & Madden, 1999), *PipMaker* (Schwartz *et al.*, 2000), *Vista* (Mayor *et al.*, 2000) and *zPicture* (<http://zpicture.dcode.org>). Also, several genome browsers provide an effortless access to precomputed genome-scale sequence alignments for any region in the human genome. These include *Ensembl* (Hubbard *et al.*, 2002), the *ECR Browser* (<http://ecrbrowser.dcode.org/>), *Pip Dispenser* (Schwartz *et al.*, 2003), *Vista Browser* (Couronne *et al.*, 2003) and the *Genome Browser at UCSC* (Karolchik *et al.*, 2003).

However, a deeper understanding of the biology and the evolution of *Homo sapiens* will require comparisons not only to distantly related genomes, such as rodents and fishes, but also to our closest relatives, the great apes. In such comparisons, it is very challenging to extract statistically significant differences since the genomes of humans and their primate relatives are very similar at the nucleotide level (>90%) (Anzai *et al.*, 2003; Britten, 2002; Hellmann *et al.*, 2003; Silva & Kondrashov, 2002). Available comparative sequence analysis tools and methodologies have, in general, been developed to analyze more distant evolutionary relationships and are not fine-tuned to analyze recent evolutionary events. Such tools are not sensitive enough to allow for meaningful comparisons involving recent segmental duplications in the human genome (Bailey *et al.*, 2002), dynamically evolving clusters of paralogous genes such as the zinc finger transcription factor families (Shannon *et al.*, 2003), or slowly diverging

genomic intervals such as the HOX-clusters (Balavoine *et al.*, 2002). But most importantly, in light of the emerging sequence of the chimpanzee genome, we require new resources that will permit meaningful comparisons between humans and other primates. Such tools will advance our ability to extract functional information from primate comparisons and expand our current understanding of human health and biology.

Recently, a novel approach, *phylogenetic shadowing*, was developed to compute and statistically evaluate conservation profiles of multiple sequence alignments from closely related species. This statistical method permitted the accurate prediction of exons and transcriptional regulatory elements in human-primate comparisons, and validated the use of this approach for deciphering primate-specific functional DNA sequences (Boffelli *et al.*, 2003). Based on the successful use of the *phylogenetic shadowing* method we have created a publicly accessible automated tool, *eShadow* that applies this strategy to the analysis of any closely related sequences. Also, *eShadow* extends the phylogenetic shadowing approach to include the analysis of multiple protein alignments, and to reduce the number of species required for the identification of functional elements. The *eShadow* tool incorporates two distinct approaches for finding functional elements - *Hidden Markov Model Islands* (HMMI) and *Divergence Threshold* (DT) scans of multiple (or pairwise) sequence alignments. Here, we report the computational design and algorithms underlining the *eShadow* tool, and we suggest several applications including the analysis of (1) coding exons, (2) noncoding elements, and (3) protein domains. We show that by overlapping HMMI predictions with a distribution of open reading frames (ORF) and fully conserved splice sites, *eShadow* can also be used to highlight regions with coding potential. Finally, we demonstrate how this program is trainable, and highly flexible to be generalized to human/baboon pairwise comparisons.

RESULTS

***eShadow* Computational Design and Visualization Scheme**

eShadow is an interactive computational tool for aligning, visualizing, and evaluating evolutionarily conservation profiles in multiple and pairwise nucleotide or protein alignments of closely-related sequences. The tool works best in alignments between sequences characterized by divergence rates less than the average human vs mouse neutral substitution rate of 0.46 substitutions per site (Waterston *et al.*, 2002). *eShadow* analysis proceeds in three major steps: (1) generating multiple sequence alignments (MSAs), (2) visualizing MSAs as percent variation plots, and (3) statistically evaluating MSAs to detect regions of high conservation (Figure 1).

eShadow generates MSAs using the multiple aligner program *ClustalW*. This tool creates alignments by first constructing a phylogenetic tree based on sequence similarity, and then follows with successive pairwise alignments in the order provided by the tree. *ClustalW* is able to align multiple closely related sequences (such as human/baboon/chimp or mouse/rat genomic sequences, which are $\geq 80\%$ identical) up to 200kb in length. While performing protein alignments, *ClustalW* weighs amino acid substitutions according to the divergence rates of the sequences being aligned, and assigns residue-specific gap penalties which are adjusted locally, resulting in increasing or decreasing the score depending on the potential secondary structure of the protein (Chenna *et al.*, 2003; Thompson *et al.*, 1994).

eShadow visualizes MSAs as percent variation (or mismatch) plots. The percent of mismatched nucleotides or amino acids is calculated in a sliding window of user-defined length; where the percent identity y , in a window size w , centered at a given position x , is plotted at the (x,y) coordinate. The peaks and valleys of the conservation plot correspond to regions of low and high variation, respectively, and 0% variation signifies 100% sequence identity in the MSA. To increase plasticity for the visual representation of the data, *eShadow* uniquely allows users to

interactively choose the base organism, modify parameters and annotation files, features absent from all other available multiple or pairwise sequence alignment tools.

The *eShadow* analytical module implements two different statistical methods of scanning MSAs to detect slow-mutating regions: (1) *Hidden Markov Model Islands* (HMMI), and (2) *Divergence Threshold* (DT). While Hidden Markov Model methods are extensively used to detect functional elements in raw genomic and protein sequences as well as in sequence alignments, DT methods are typically used for the analysis of conservation plots. HMMI implementations include gene prediction (Burge & Karlin, 1997; Krogh, 1997), CpG island localization (Takai & Jones, 2002), noncoding RNAs identification (Rivas & Eddy, 2001), protein domain (Truong & Ikura, 2002) and protein fold predictions (Bienkowska *et al.*, 2000). We used a two-state HMMI to analyze conservation profiles and to predict conserved (slowly diverging) regions. This method does not utilize a sliding-window, rather it analyzes the underlying distribution of matches and mismatches in the alignments. The DT method distinguishes conserved elements based on their length and the level of complete nucleotide/amino acid identity. In contrast to the HMMI approach, DT analysis is performed by scanning the alignment summary for regions corresponding to the number of matches x in a sliding window of predefined length y . The DT method is employed by most pairwise alignment programs and is probably the most commonly used approach for biologists to define evolutionary conservation (Schwartz *et al.*, 2000).

The analytical component of the *eShadow* tool also contains several optional features: (1) an open reading frame (ORF) detection block and (2) an optimization module to assist during the characterization of conservation patterns across alignments. By superimposing ORF predictions, *eShadow's* HMMI detection module identifies nucleotide regions with high potential to code for proteins, possibly differentiating coding from noncoding conserved elements. The optimization

module allows the user to train the program to identify parameters that would distinguish features similar in nature to a set of known elements (exons or regulatory elements) from the background noise. We have implemented three complementary optimization methods into the *eShadow* tool: (1) *Baum-Welch* (Durbin R., 1998), (2) *Maximum Likelihood* (Durbin R., 1998), and (3) *Golden Section Search* (Press W.H., 1988).

Applications for the *eShadow* tool

eShadow has been designed for comparing sequences with relatively small interspecies divergence rates, preferably classified to the same class or order. The detection of such elements is very difficult by currently available computational means, and is critical for molecularly distinguishing biological functions unique to small clades of organisms (Cooper *et al.*, 2003). Unlike most available comparative sequence analysis tools, *eShadow* can be dynamically adjusted and trained to accommodate evolutionary relationships as distant as human and mouse or as close as two primates. However, other tools such as *Pipmaker* and *VISTA* are more effective in pairwise analysis of distantly related sequences and are specifically designed for such applications (Schwartz *et al.*, 2000; Mayor *et al.*, 2000). The particular analytical scope of the *eShadow* tool is to assist in the discovery of elements distinctly shared by classes of organisms tightly clustered on the same branch of the evolutionary tree. Here, we illustrate three major applications for the *eShadow* tool: (1) identification of coding exons, (2) prediction of conserved noncoding elements and (3) evaluation of protein domains in alignments.

Detecting Coding Exons

The two statistical methods implemented into the *eShadow* tool were tested for the ability to detect conserved elements across 4 genomic intervals and to accurately predict the known coding regions corresponding to 5 exons from 4 different genes (*ApoB*; *Plasminogen*; *LXR-alpha*;

CETP). These 4 genomic intervals have been extensively sampled across multiple primate species and currently represent the only dataset for which there are 13 to 16 unique primate sequences available (Boffelli *et al.*, 2003). Even though the set of sequenced primates slightly varied for each tested region, the available sequences for each genomic interval spanned the primate phylogeny evenly, resulting in similar substitution rates per human base pair in all tested cases (Supplement Table S1). We optimized the parameters for each analytical method to detect regions with slow-mutation rates and found a set of common parameters that identified all the exons present in these genomic intervals. The HMMI method detected all exons [parameters: $eS=0.85$; $eF=0.77$; $T=0.1$ (Details in Supplementary Materials; Figure S1)] while the DT method missed the shortest exon [parameters: % variation=15; length=100bp] (Figure 2).

Despite the tremendous advances achieved in DNA sequencing technologies, obtaining the sequence of dozens of closely related vertebrate-sized genomes is still not a practical goal. We therefore asked whether single human/primate pairwise alignments might contain enough information to distinguish conserved (slow-mutating) from neutral (fast-mutating) regions. This task is highly intricate since the substitution rate decreases significantly when switching from an MSA to a pairwise alignment (Supplement Figure S2). Figure 3 illustrates *eShadow*'s ability to predict the ApoB exon from a single human/primate [*Allouatta seniculus*] pairwise alignment (Figure 3C) as accurately as it can be predicted from a human/mouse (Figure 3A) or a primate MSA (Figure 3B). Similar results were obtained in 54 other exons analyzed in human/baboon alignments (Table 1). These results suggest that if properly analyzed, single human/primate pairwise alignments have the potential to be as informative for exon identification as human/rodent alignments are.

Detecting Conserved Noncoding Elements

Since humans and rodents share the majority of their protein coding genes (Waterston *et al.*, 2002) it has been hypothesized that most of the phenotypic differences between clades of mammals are attributed to differences in noncoding sequences. In some cases, these differences may involve substantial changes in regulatory sequences that have occurred during the ~80 Myrs separating the rodent and human lineages (Dermitzakis & Clark, 2002; Scemama *et al.*, 2002), limiting H/M comparisons to functions that are more globally shared by mammals and vertebrates. The *phylogenetic shadowing* approach has been shown to be suited for the identification of lineage-specific noncoding conserved elements through comparisons of several closely related primate genomic regions (Boffelli *et al.*, 2003). Since this approach is limited by the amount of sequence data available for species tightly-clustered on the same branch of the phylogenetic tree, we tested *eShadow*'s ability to recapitulate conservation patterns when the number of input sequences is reduced from >10 different species to only 2 or 3. We also limited our analysis to organisms evolutionarily close to humans and in the primate lineage— such as baboon and chimp (genome sequences expected to be generated in the foreseeable future).

We analyzed 53kb from the Wingless-type MMTV Integration Site Family, Member 2 gene locus (*WNT2*), which has been deeply sequenced in many species, including chimps and baboons in addition to humans and mice (Thomas *et al.*, 2003). H/M comparisons identified all translated and untranslated (UTR) exons, included in a collection of 62 Evolutionary Conserved Regions (ECRs; ≥ 100 bps and $\geq 70\%$) (Figure 4). We addressed whether *eShadow* primate-specific comparisons can possibly recapitulate the H/M conservation patterns, as well as identify additional primate-specific conserved elements through the use of human/baboon/chimp (H/B/C) comparisons. The *eShadow* HMMI module was first trained on the *WNT2* exons, and the optimized parameters were used to analyze the conservation pattern across the entire 53 kb genomic interval.

While the H/B/C alignment demonstrated ~12% nucleotide variation, the conservation pattern clearly distinguished regions with different evolutionary rates. HMMI predictions (0.98/0.90/0.005) identified 26 ECRs, including all the *WNT2* coding exons. While this number (26) is approximately three times less than the corresponding number for H/M ECRs (62), in general, strings of smaller H/M ECRs were incorporated within larger H/B/C ECRs, such that 68% (42) H/M ECRs were recapitulated in H/B/C ECRs. On a per base pair basis, the sum of all human nucleotides classified as highly conserved either in rodent (H/M) or primate (H/B/C) comparisons were highly similar spanning 15kb in H/B/C and 17kb H/M alignments. Four primate-specific ECRs lacked a highly homologous counterpart in the mouse ortholog. These elements could either represent regions that did not accumulate enough mutations throughout primate evolution due to chance alone, or could be primate specific elements. Computationally distinguishing between these two possibilities is not yet feasible; rather the true biological relevance of these lineage-specific elements must be determined experimentally.

To evaluate the specificity and sensitivity by which the *eShadow* tool is able to recapitulate H/M conservation profiles from a single human/primate alignment, we analyzed a test set of four completely finished baboon BACs spanning ~677kb that were syntenic to contiguous regions in both the human and mouse genomes (Table 1). This analysis was performed using fixed HMMI parameters trained on the *WNT2* region (0.98/0.90/0.005). These regions exhibited strong correlations between the H/B HMMI predictions and the ECRs present in the H/M alignments. We estimated the sensitivity of recapitulating H/M conservation patterns by HMMI modeling of H/B alignments to be ~59.3% and the specificity ~77.6% (Table 1). To provide a measure of H/B exon-detection sensitivity we calculated the number of exons identified by the HMMI approach across these four baboon BACs. Exons were scored as “detected” if it contained a partial or a full overlap with the HMMI prediction. 62% (54/87) of

all exons were detected by this approach. That also corresponds to 83% (25/30) of the long exons (>150bps) suggesting that the *eShadow* approach works more efficiently for detecting longer elements (Table 1). These results imply that similar analysis may be used to define primate specific elements in the human genome in an unbiased manner by comparing a minimum number of different primate sequences and therefore suggest the use of the *eShadow* tool for computationally identifying primate-specific elements when the genome sequence of additional primates become available.

Identifying Conserved Protein Domains

Discovering deleterious mutations within candidate genes is fundamental to elucidating the genetic basis of human disorders. For most genes, the significance of any particular amino acid change is mostly unknown and requires comprehensive structural and functional studies. Multi-species protein alignments (MPA) can provide valuable information about the phylogenetic relationships between species and identify evolutionarily constraints in regions that are central to structural and biochemical interactions (Cline *et al.*, 2002). Evaluating evolutionary rates at specific sites through the use of likelihood-ratio tests (LRTs) has been extensively used to characterize amino acid rate changes likely to underlie functional constraints on proteins (Knudsen *et al.*, 2001). Evolutionary rate analysis complements existing approaches for the identification of conserved residues. Despite the intuitive correlation between conserved residues and functionally significant protein domains, distinguishing conservation associated with genuine biological interactions solely resulting from the shared phylogeny is a very difficult task (Pollock & Taylor, 1997).

One application for the *eShadow* tool includes the analysis of MPAs to detect protein domains under selective pressure using HMMI predictions. This strategy is particularly promising since HMM profiling is one of the most successful strategies for detecting statistically

significant regions of protein homology (Madera & Gough, 2002) and is already implemented by homology-based protein motif search tools such as the *Pfam* database. The *eShadow* tool can also be used to visualize the distribution of nonsynonymous amino acid changes within MPA. To illustrate this we have mapped all cystic fibrosis (CF) missense mutations documented in the *Cystic Fibrosis Mutation Database* to the CFTR MPA built from seven different species and correlated their distribution with the *eShadow* HHMI predictions. 80% of the documented CF missense mutations (32/40) were enclosed by regions of high protein homology as identified by HHMI (0.77/0.65/0.1) (Figure 5). At the same time, 95% the most common CF disease causing alleles (16 CF missense mutations; 3 single amino acid deletions) were found to be present in HHMI predicted domains.

DISCUSSION

We have developed a computational web-based tool, *eShadow*, that is highly proficient in performing *phylogenetic shadowing* analysis for closely related nucleotide and protein sequences. *eShadow* amplifies the information content from pairwise or multiple alignments by combining independent mutations present in each different lineage, and detects regions with the lowest cumulative density of mutations through the use of two different statistical methods, DT and HHMI. This tool also includes a parameters-optimization module for the HHMI model that can be amended to any particular evolutionary history underlying the input sequences and trains the program to predict conserved elements in a wide variety of alignments. Unlike other available tools that analyze conservation across alignments using static parameters, *eShadow*, permits for dynamic modifications of all parameters and picture settings creating conservation plots in real-time.

eShadow can be used to detect coding exons, protein domains and conserved noncoding elements. While *eShadow* identifies exons, exon-intron boundaries are not exactly delineated;

therefore this tool provides a good starting point for transcript analysis that could benefit from external gene prediction information. When protein alignments are analyzed *eShadow* can be used to highlight protein domains that are conserved across multiple species and may be involved in vital biochemical processes such as protein-protein contacts or DNA binding. We have also indicated how amino acid mutational analysis can be superimposed on HHMI predictions in MPAs and this analysis can be used to evaluate missense mutations.

We have shown that *eShadow* can recreate most of the information obtained from human/mouse alignments when human/baboon/chimp or human/baboon alignments are analyzed. A 53kb three-way primate alignment analysis for the WNT2 locus recovered 68% of the human/mouse conserved elements, as well as identified several primate-specific conserved elements. While the functional significance of these lineage-specific sequence elements is presently unknown, we speculate that they may potentially represent sequences that underlie noncoding functions shared by primates but not by other mammals. Regulatory modifications of conserved genes have been proposed to define the major molecular differences that set different organisms phenotypically apart (Boffelli *et al.*, 2003), suggesting one potentially very interesting application for *eShadow* that cannot currently be performed by any other publicly available computational tool. Although we have focused on primate sequence alignments, *eShadow* can be tuned to align closely related sequences from any species. In addition, *eShadow* may be uniquely applicable to other problems including alignments between recent segmental duplications. Such duplications can often generate new functional gene copies that do not have true orthologs in other species and are therefore not amenable to standard cross-species comparative analysis (Bailey *et al.*, 2002; Shannon *et al.*, 2003). *eShadow* therefore adds an important set of capabilities to the current comparative genomics toolkit, providing unique

access to species- and lineage-specific elements throughout sequenced genomes from any evolutionary clade.

METHODS

Hidden Markov Model Islands

We used a two-states HMM method to predict slow diverging regions in MSA. We modeled the distribution of matches and mismatches assuming that they correspond to two mutation states [slow and fast], with different probabilities of emitting a match (eS and eF in slow- and fast-mutation states, respectively), and under the assumption that the probability T of transferring from state to state is equal in both directions (Supplement Figure S1). The emission probability of a state relates to the average sequence conservation of that particular state, while the transfer probability is inversely proportional to the average length of regions in the alignment. *Viterbi algorithm* was used to predict the underlying distribution of slow-mutation states.

Parameters Optimization

Baum-Welch (Durbin R., 1998), *Maximum Likelihood* (Durbin R., 1998), and *Golden Section Search* (Press W.H., 1988) were used for optimizing parameters. Only *Maximum Likelihood* and *Golden Section Search* utilize user-provided annotation files (base sequence) as a training dataset. *Maximum Likelihood* scans MSAs and sets emission probabilities equal to the observed number of matches per length of sequence in a particular state; the transition probability is estimated as a number of transitions from state to state divided by the total length of the base sequence. The guided *Baum-Welch* optimization tests different paths through the sets of slow- and fast-mutation states and identifies the set that maximizes the likelihood probability for the HMMI. This method does not require annotation (it is used as an initial guess, if available) to calculate HMMI parameters, instead it scans through all the possible regions of slow-mutation and identifying regions and HMMI parameters that will maximize the log-likelihood of the model given the distribution of matches and mismatches. *Golden Section Search* (Brent's method) is a one-dimensional approach for determining the minimum (or optimum) of a

nonlinear function, and is used to fine-tune the HMMI parameters by cyclically optimizing the probability parameters until full convergence is achieved. Every parameter optimization is performed while the other two variables are fixed. This is achieved by sampling the surface and identifying the local minimum of the penalty function. We define the scoring function S as a discrepancy in the location of the HMMI predictions and the locations of functional regions (G):

$$S = L_G + L_{HMM} - 2L_{G \cap HMM},$$

where L_G , L_{HMM} , and $L_{G \cap HMM}$ are the lengths of guiding regions and HMMI predictions, and the length of their overlap, respectively. An exact fit for the HMMI predictions and guiding regions will zero the scoring function S , while any discrepancies will increase it. This method iterates and converges to the point of the local minimum, due to the discrete nature of the scoring function. Usually there are several local extremum points in the three-dimensional surface of HMMI probability parameters in *Baum-Welch* and *Golden Section Surface* optimizations. The implemented procedure converges to one of the extremum points depending on the input parameters. Therefore, when HMMI generates no predictions or there is a large discrepancy between the predicted and the expected elements, parameters need to be modified accordingly.

Open Reading Frame (ORF) Detection

ORFs are identified in all six reading frames, excluding the ones <60 bp in length, and are illustrated as gray bars. Stop codons from all sequences are collapsed onto the reference sequence. All ORFs internally spanned by HMMI predictions >75% in length are preserved, identifying the most probable frame of translation for each HMMI. At the final step every frame is truncated in order to confirm the standard AG-GT exon-intron splice-site (Burset *et al.*, 2000). AG and GT dinucleotides are required to be evolutionary conserved in all the species throughout the alignment and the pair closest to HMMI edge is used to define the exon boundaries.

Independently generated ORF and HMM predictions that overlap with each other are demarcated by different colors: red (positive strand); blue (negative strand) (Figure 3C).

Output option

The text output option of the *eShadow* tool provides the user with a detailed summary for the alignment and exhaustive information about the detected slow-mutating regions. This module calculates the number of mismatches in all the pairwise sequence comparisons for the base sequence versus each different sequence and compares it to the whole MSA. This serves as an estimate of evolutionary divergence between different species as well as it characterizes the substitution rate in the MSA. The second part of the summary data consists of a module calculating the discriminative power introduced by each additional species used in the multiple sequence alignment. The program presents the tabulated data identifying the n most distant species, where n varies from 2 to the maximum number of species in the initial alignment, by analyzing all the possible combinations of n species and extracting one with the highest substitution rate per base sequence (Supplementary Figure S2). This is done by a sequential grouping of sequences that introduce the largest number of mutations into the MSA. Every group of n sequences that is presented is the optimal group that has the maximum possible amount of mutations in all the possible sets of groups of n sequences.

This section also contains a report of all the slow-mutating regions identified by the selected methods, providing the coordinates of the predicted elements in the base sequence and indicating their parameters. HMMI predictions also contain scores that reflect confidence (statistically evaluating predicted regions versus background noise for a given set of parameters). Every predicted slow-mutating interval I of the HMMI collects a score $S(I)$, which reflects a log likelihood probability of this interval not to be a fast-mutating region:

$$S(I) = \sum_{i \in I} \log \left(\frac{P(m_i = \text{slow} | A)}{P(m_i = \text{fast} | A)} \right),$$

where the summation is done across all the base pairs in the interval and $P(m_i = k | A)$ is the posterior probability of the observed state k at the position i in the alignment A , which has a known structure of complete matches/mismatches. We prioritized longer conserved intervals over the shorter ones by not normalizing interval scores based on length.

Table 1. Evaluating *eShadow*'s performance on recapitulating human/mouse conservation patterns in human/baboon alignments. Sequences for the WNT2 region were obtained by excising ~53 kb from the ~2 Mb region [NISC Comparative Sequencing Program has sequenced for the CFTR locus sequencing project (Thomas *et al.*, 2003)]. 4 baboon BACs spanning CECR, PCQAP, SNAP29, and TCF4 gene loci were downloaded from NCBI (Acc# AC091672, AC128639, AC129881, AC113267). Human and mouse sequences were obtained from UCSC database, coordinates and alignments are available in Supplementary Materials.

Gene Locus	Sensitivity	Specificity	Common region length**	Detected coding exons	
				all	> 150 bps
WNT2	67.9%	61.1%	52.8kb	5/5	4/4
CECR	55.7%	82.5%	63.4kb	7/20	5/7
PCQAP	51.1%	89.8%	123.4kb	13/20	8/8
SNAP29	46.0%	89.6%	125.1kb	19/27	7/10
TCF4	61.4%	56.7%	137.8kb	10/11	1/1
Average	59.3%	77.6%	100.5kb	54/87 (62%)	25/30 (83%)

**Common region lengths are defined as regions of homology (kb) flanked by at least one ECR and one HMMI prediction. HMMI parameters used: 0.98/0.90/0.005.

Figure 1. Schematic dataflow of the eShadow program.

Figure 2. Predicting exons in multiple primate alignments. *eShadow* exon prediction and conservation plots for Apo-B (1.4kb) (A), Plasminogen (1.2kb) (B), LXR-alpha (1.35kb) (C), and CETP (1kb) (D) loci using multiple primate alignments. HMMI predictions (0.85/0.77/0.1) are in beige and DT regions (15/100) are in green. The locations of known exons are depicted as red bars. Sequences are available at NCBI, acc# AY190030-AY190042, AY192729-AY192785.

Figure 3. Predicting exons in single primate pairwise alignments. *eShadow* analysis for Apo-B (1.4kb) region using human/mouse (a), human and 14 primate sequences (b) and human/*Allouatta seniculus* sequences (c); 50 bps sliding window was utilized to smoothen the conservation profile. Right panel shows human/mouse (d) and human/ *Allouatta seniculus* (e) sequence comparisons plotted with standard 50% to 100% thresholds and 100 bps sliding window. Exons (blue) and evolutionary conserved regions (ECRs) (red) are indicated in (d) and (e) zplots (<http://zpicture.dcode.org>). HMMI predictions (beige), ORF predictions (gray), exon annotations (red) and exon predictions (yellow) are visualized in *eShadow* plots (a-c). Percent variation- y axis, size in bp- x axis. (a-c). Percent identity- y axis, size in kb- x axis (d,e).

Figure 4. *eShadow* analysis for the WNT2 region. Human/mouse conservation plot (A) compared to human/baboon/chimp *eShadow* conservation plot (B). Human/mouse alignments were generated and visualized by the zPicture program (<http://zpicture.dcode.org>), using standard parameters ($\geq 100\text{bp}$; $\geq 70\%$) and conserved elements corresponding to exons (blue), UTRs (yellow), intronic (pink) and intergenic (red) elements are indicated (A). Human/baboon/chimp alignment plot (B) depicting regions of conservation (purple) and HMMI predictions (beige). y

axis corresponds to- percent identity (A) and percent variation (B). x axis corresponds to size in base pairs (A,B).

Figure 5. *eShadow* application in multiple protein alignment analysis. Human, baboon, cow, sheep, mouse, rat, and rabbit CFTR proteins were aligned using *eShadow* and highly homologous regions were predicted using the HMMI module (0.77/0.65/0.1). Mutations known to cause cystic fibrosis (<http://www.genet.sickkids.on.ca/cftr>) are mapped to the CFTR MPA and annotated as red tick marks.

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<i>Cystic Fibrosis Mutation Database</i>	http://www.genet.sickkids.on.ca/cftr/
<i>ECR Browser</i>	http://ecrbrowser.dcode.org/
<i>Ensembl</i>	http://www.ensembl.org/
<i>eShadow</i>	http://eshadow.dcode.org/
<i>Human Genome Browser at UCSC</i>	http://genome.ucsc.edu/
<i>Pip Dispenser</i>	http://bio.cse.psu.edu/genome/hummus/
<i>PipMaker</i>	http://bio.cse.psu.edu/pipmaker/
<i>Pfam database</i>	http://www.sanger.ac.uk/Software/Pfam/index.shtml/
<i>Vista</i>	http://www-gsd.lbl.gov/VISTA/VistaInput.html
<i>Vista Browser</i>	http://pipeline.lbl.gov/
<i>zPicture</i>	http://zpicture.dcode.org/

Input File

- 1. human
DUTRIN00000
- 2. human
P00000000
- 3. ortho
P00000000

ClustalW Alignment

```
human    DUTRIN00000  
human    P00000000  
ortho     P00000000  
          ..  
          ..
```

Annotation of Conserved Regions

Divergence
threshold

TMM
islands

ORF
prediction

Visualization & Output



1-100 conserved
300-400 conserved







