Resource

Analysis of Multiple Genomic Sequence Alignments: A Web Resource, Online Tools, and Lessons Learned From Analysis of Mammalian SCL Loci

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Comparative analysis of genomic sequences is becoming a standard technique for studying gene regulation. However, only a limited number of tools are currently available for the analysis of multiple genomic sequences. An extensive data set for the testing and training of such tools is provided by the *SCL* gene locus. Here we have expanded the data set to eight vertebrate species by sequencing the dog *SCL* locus and by annotating the dog and rat *SCL* loci. To provide a resource for the bioinformatics community, all *SCL* sequences and functional annotations, comprising a collation of the extensive experimental evidence pertaining to *SCL* regulation, have been made available via a Web server. A Web interface to new tools specifically designed for the display and analysis of multiple sequence alignments was also implemented. The unique *SCL* data set and new sequence comparison tools allowed us to perform a rigorous examination of the true benefits of multiple sequence comparisons. We demonstrate that multiple sequence alignments are, overall, superior to pairwise alignments for identification of mammalian regulatory regions. In the search for individual transcription factor binding sites, multiple alignments markedly increase the signal-to-noise ratio compared to pairwise alignments.

[Supplemental data is available at www.genome.org and http://hscl.cimr.cam.ac.uk/supplementary_data.html. DNA sequence as described in the paper has been deposited in the GenBank database under accession no. AL731652. The following individuals kindly supplied reagents, samples, or unpublished information as indicated in the paper: R. Li, P. de Jong, and R. Huss.]

Among the vertebrates alone, whole-genome sequences have been determined for human, mouse, rat, and two pufferfish species. Sequencing has started for chimpanzee, chicken, frog (*Xenopus tropicalis*), and zebrafish, and projects to sequence the genomes of dog and cow are due to commence in the near future. Faced with the task of identifying putative regulatory elements of a gene, it is already a simple matter to obtain homologous human and mouse genomic sequences for the locus of interest, perform an alignment with one of a variety of algorithms, and examine in more detail those regions that are conserved between the two species. Within these regions, phylogenetic footprinting (Tagle et al. 1988; Gumucio et al. 1993; Shelton et al. 1997) can be used to determine potential transcription factor binding sites. Sequence-specific transcription factor binding sites are much more likely to occur within blocks of complete identity than outside such blocks (Wasserman et al. 2000).

Alignments of more than two species (multiple alignments), separated by similar evolutionary distances, may offer advantages over pairwise alignments. Provided that the evolutionary distances between the chosen species are sufficiently small, problems of detecting only a subset of regulatory elements with distant comparisons (Miles et al. 1998; Göttgens et al. 2000, 2002a; Cioffi et al. 2001; Flint et al. 2001; Gilligan et al. 2002) are mini-

mized. Moreover, having more than two species adds substantial resolving power, because each lineage diverges independently after separation from a common ancestor, resulting in evolutionary distances that are effectively additive (Stojanovic et al. 1999). Thus, the likelihood of residual similarities in nonselected regions is markedly reduced. This additive effect was recently demonstrated with identification of gene regulatory elements by multiple sequence comparisons among primates (Boffelli et al. 2003). However, despite the significance of the latter study, multiple alignments involving eutherian mammals other than primates are likely to prove important. If the approach is to be universally applicable, one is limited to sequence comparisons involving those species for which whole-genome sequences are available. Furthermore, murine models are likely to remain the prime in vivo experimental system for studying mammalian gene expression. Finally, statistical modeling has suggested that comparisons across a wide range of mammalian sequences are likely to provide the most useful resolution at the single-nucleotide level (Cooper et al. 2003).

Given the promise of multiple sequence comparisons, there is a relative paucity of tools for the analysis of multiple genomic sequences (Miller 2001). Essential to their design are data sets that can be used to train and evaluate algorithms. One prime model for such a data set is the locus of the basic helix-loop-helix (bHLH) stem cell leukemia (*SCL*) gene. There are several reasons for this. Firstly, *SCL* is a pivotal regulator of hematopoiesis and vascular development (for review, see Begley and Green 1999), with a pattern of expression that is highly conserved throughout

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vertebrate evolution, from mammals to teleost fish. Second, the *SCL* locus is not too large—there is evidence that all elements necessary for its transcriptional pattern are contained in a region spanning approximately 54 kb (Barton et al. 2001; Sinclair et al. 2002). Third, there is a wealth of experimental data on the transcriptional regulation of human and mouse *SCL* loci (Begley and Green 1999, and references therein; Fordham et al. 1999; Sanchez et al. 1999, 2001; Sinclair et al. 1999, 2002; Göttgens et al. 2000, 2001, 2002a,b; Barton et al. 2001). Fourth, *SCL* sequences are currently available for six species: zebrafish, pufferfish (*Fugu* and *Tetraodon*), chicken, mouse, and human (Göttgens et al. 2000, 2002a; Barton et al. 2001). Finally, *SCL* is not flanked by paralogous genes, unlike, for example, the *hox*, *interleukin*, and *globin* loci. Therefore, characterized functional elements do not regulate multiple genes. Thus, inferring the functional significance of conserved elements may be more straightforward.

This article describes the addition of dog and rat sequences to the *SCL* data set and the development of a Web server that provides a highly annotated testing and training set as a resource for the bioinformatics community. To examine the practical utility of multiple alignments, all combinations of mammalian *SCL* alignment—six pairwise, four three-way, and one four-way alignment—were generated. These alignments were compared using the annotations and with new tools designed to analyze multiple alignments, made available to run via the server. Multiple alignments, and particularly the four-way alignment, markedly narrowed the search for putative transcription factor binding sites in the locus, with minimal loss of sensitivity.

RESULTS

Development of the *SCL* Data Set as a Genomics Research Resource

The first goal was to extend the *SCL* data set. A single clone of 169,460 bp in length, containing the entire dog *SIL*, *SCL*, and *MAP17* genes, was therefore isolated and fully sequenced (see Methods). The locations of noncoding exons were defined by RT-PCR using RNA from two dog cell lines. Exons 1a and 1b were spliced directly to exon 3, suggesting that there are no dog homologs to human exons 2a and 2b. The rat *SCL* sequence was obtained from the Berkeley Genome Pipeline (<http://pipeline.lbl.gov/rat>) and fully annotated. The extracted region extended from within the *SIL* gene to approximately 2 kb downstream of *MAP17*. The choice of upstream and downstream boundaries for the contig was based on the minimum region of conserved synteny previously reported (Barton et al. 2001) and the observation that a human YAC containing this region could rescue *SCL* $-/-$ mouse embryos from the normally lethal phenotype (Sinclair et al. 2002).

Algorithms for predicting regulatory regions benefit considerably from training and testing data sets with experimentally confirmed functional annotations (Miller 2001; Elnitski et al. 2003). The *SCL* data set now includes eight vertebrate species with a wealth of experimental information on regulatory elements within the locus. A Web server that provides access to the *SCL* sequences together with annotations of functional noncoding regions has therefore been implemented (http://hscl.cimr.cam.ac.uk/genomic_datasets.html). From here, there are three main links. One is to the *SCL* sequences themselves. A second is to annotations of these sequences in Gen-

eral Feature Format (GFF) files (see <http://www.sanger.ac.uk/Software/formats/GFF/>). Each GFF file details the locations of coding and noncoding exons, which have been determined experimentally in all species except rat, and repeat regions. The known *SCL* regulatory regions have been defined experimentally in the murine locus. These have therefore been annotated in the mouse GFF file, and the equivalent regions are defined in the GFF files for the other mammals. The final link is to a table (Table 1), which serves as an index to detailed descriptions of these known functional regions. The nomenclature of the regions refers to their distance from promoter 1a in the murine *SCL* locus. The extent of all regions except the +23 kb enhancer has been experimentally confirmed to be in an open chromatin configuration. The core region with full activity is described, together with the positions of transcription factor binding sites where these have been experimentally determined.

Within the Web site, Table 1 contains hyperlinks for each regulatory region, each of which leads to a more detailed review of underlying experimental evidence. These functional data include (where determined) methylation status, activity in transient and stable transfection assays, activity in transgenic mice, and the effect of mutating known transcription factor binding sites in the various experimental systems. All transcription factor binding sites known to be involved in the regulation of *SCL* are preserved in alignments of the four mammalian species (see below). These binding sites thus represent rigorous sensitivity criteria for training or testing algorithms concerned with genomic sequence alignment and binding site prediction. Therefore, for each region, known binding sites are indicated on an alignment of the mammalian sequences in that region. Finally, references to the original publications are given. The Web site will continue to be updated as more data on *SCL* regulation emerge.

Table 1. Functional Annotation of the Mouse *SCL* Locus

Region	Open chromatin (approx. location)	Minimally defined active element	Transcription factor binding site
– 8/– 9 kb	20265–20665 20765–21165	18884–22017	Unknown
– 4 kb	24965–25445	25095–25488	Ets family 25213 Ets family 25262 Ets family 25361 Ets family 25369 C/EBP 29119
Promoter 1a	28965–29265	29044–29257	Ap1 29130 Gata1 29162 Sp1 29168 Gata1 29194
Promoter 1b	29425–29585	29440–29558	Sp1/3 29472 Ets family 29494 Ets family 29498 Ets family 29502 C/EBP 29520
+1 kb	29865–30165	29558–30376	Unknown
+3 kb	31865–32065	30377–36071	Unknown
+7 kb	35625–36065	34510–37251	Unknown
+18 kb	47265–47595	47299–47728	Unknown
+19 kb	48045–48345	47728–48369	AATAA 48210 Ets family 48217 Ets family 48237 Gata2 48262
+23 kb	Unknown	50934–52609	Unknown

The names of the regulatory regions, approximate regions of open chromatin, the minimally defined active element, and known transcription factor binding sites are indicated. On the Web server, entries in the first column serve as hypertext links to detailed information on the individual regions.

Use of Multiple Mammalian Alignments to Identify *SCL* Regulatory Regions

It has been proposed that multiple mammalian sequences represent the best comparison for detecting functional regulatory regions (Frazer et al. 2003). Statistical modeling has suggested that they also provide a more useful resolution at the nucleotide level than pairwise alignments (Cooper et al. 2003). The new data set provided the opportunity to address these issues by comparing the abilities of multiple and pairwise alignments of mammalian *SCL* sequences to detect regulatory elements. To this end, the SynPlot module (Göttgens et al. 2000, 2001) was modified to process multiple alignments and to run via the Web server (http://hscl.cimr.cam.ac.uk/syn_plot.html). SynPlot displays a plot of alignment score within a sliding window, which moves across the alignment. To accommodate multiple alignments, the alignment score is generated by the sum of pairs method (see Methods) and expressed as the proportion of the maximum attainable score in the window. All possible two-, three-, and four-way alignments of human, dog, mouse, and rat *SCL* were generated and a SynPlot produced. In addition to exons and repeats, all known regions of open chromatin were shaded. All such sites in the *SCL* locus have transcriptional regulatory activity, or map to larger regions with such activity, in transfection and/or transgenic assays. In addition, the +23 kb enhancer (outside mapped regions of open chromatin) was shaded.

Figure 1 is the SynPlot for the four-way alignment (SynPlots for all the other alignments are available as Supplemental data at www.genome.org and hscl.cimr.cam.ac.uk/supplementary_data.html). As expected, coding exons are highly conserved, each represented by a peak with 77% of the maximum possible alignment score or greater. The regions of open chromatin and the +23 kb region—known functional regions, as depicted in Table 1—are shaded in green. They are also highly conserved between the four species, the only exceptions being the –9 kb and –8 kb regions (71% and 64% of maximum score, respectively). In addition, there are a few other noncoding regions with a high degree of sequence conservation, most notably the –10 kb, +26 kb, and +30 kb regions indicated (92%, 80%, and 83% of maximum score, respectively). The pattern of peak distribution was similar in the SynPlots of the other mammalian *SCL* alignments (Supplemental data).

Noncoding peaks were divided into two classes: those corresponding to regions of open chromatin and the +23 kb region (known regulatory regions), and those representing all other noncoding sequence conservation. A number of analyses were then performed to compare the abilities of the various two-, three-, and four-way alignments to selectively identify *SCL* regulatory elements and transcription factor binding sites. The methods and a detailed discussion of the results are available from http://hscl.cimr.cam.ac.uk/additional_info.html. The tools used in the analysis were designed to run via a user-friendly Web interface (http://hscl.cimr.cam.ac.uk/genomic_tools.html), and form an additional resource for biologists working in the field of gene regulation. The results of the analyses are summarized below.

In terms of identifying regulatory regions of the order of a few hundred base pairs in length, multiple alignments were, on the whole, more selective than pairwise alignments, with no loss of sensitivity. The four-way alignment was the most specific. Some pairwise alignments performed well, but this could not be predicted from the evolutionary distances separating the species. In terms of individual transcription factor binding sites known to regulate *SCL*, all were conserved in all alignments. When the alignments are searched for other putative conserved transcription factor binding sites, fewer are found in three-way than pair-

wise alignments, and the fewest are found in the four-way alignment, as would be expected. However, putative sites become concentrated within the known regulatory regions as the number of species is increased, suggesting a genuine increase in the signal-to-noise ratio of multiple over pairwise mammalian alignments in the search for transcription factor binding sites.

DISCUSSION

Visual Representation of Multiple Alignments

Visual interfaces to alignments are required to gain an appreciation of the quality of the alignment and to identify regions of sequence conservation, which may represent functional elements. Effective representation of pairwise alignments can be achieved by using a sliding window to calculate the percentage identities across the alignment, which can then be plotted, for example, PipMaker (<http://bio.cse.psu.edu/pipmaker>; Schwartz et al. 2000), Vista (http://www.gsd.lbl.gov/vista/vista_cite.html; Loots et al. 2000), and the original version of SynPlot (Göttgens et al. 2000). However, percentage identity cannot be calculated for more than two species simultaneously. Therefore, to deal with multiple alignments, Vista and MultiPipMaker (Schwartz et al. 2003) construct a series of successive pairwise plots comparing various species in an alignment with a reference sequence. This may be the most efficient way of visualizing very large-scale alignments, for example, whole genomes. However, there is necessarily a loss of information: Differences in genomic organization cannot be seen; misalignments are often not clear; and prioritizing peaks of sequence conservation can be difficult. Thus, we felt that for examining multiple alignments of single gene loci, an alternative approach could be useful.

SynPlot was modified to process multiple alignments, generating an alignment score, to replace percentage identity, via a sliding window. The alignment score is calculated by the sum of pairs method (see Methods). This allows all the information from a multiple alignment to be considered simultaneously, rather than requiring a reference sequence. It may therefore better delineate the very highest peaks of sequence conservation. It does not yet take into account the evolutionary relationships between the species involved, but this approach may be implemented in future versions. Another advantage of the SynPlot approach is that all features of all the sequences can be visualized above the plot, which is important when the genomic organization is not identical.

SCL Data Set

One of the many uses of comparative genomic sequence analysis is the identification of gene regulatory elements. Central to its success are algorithms that are capable of selectively identifying biologically meaningful conserved regions. In particular, with the increasing availability of whole-genome sequences, tools for analyzing multiple sequences are likely to gain prominence. However, such algorithms are in their infancy. Their success will depend on robust data sets (Miller 2001).

The *SCL* locus is a prime candidate for such a testing or training data set, and its use has precedents. It was used in the training of the LAGAN algorithm and in the testing of the Chaos-Dialign combination (Brudno et al. 2003, Brudno et al. 2004). The data set has now been extended to eight species by the complete sequencing and annotation of dog *SCL* and annotation of rat *SCL*. By implementing a Web server, convenient access to the full data set is now provided. The sequences can be easily obtained from a single source. Also supplied are complete functional annotations. There are other multiple mammalian genomic sequence data sets available (e.g., Stojanovic et al. 1999; Thomas et al. 2003). However, these loci have not been subject to

Figure 1 A long-range four-way mammalian alignment is capable of detecting all known *SCL* regulatory regions. A SynPlot representation of a human/dog/mouse/rat alignment of the *SCL* locus. Numbers on the horizontal axis do not refer to any one of the sequences in the alignment, but instead represent distance in nucleotides from the beginning of the aligned file (i.e., gaps in the alignment are counted). Numbers on the vertical axis represent an alignment score in a 100-bp window moved by 25-bp increments across the entire alignment. The horizontal black lines above the profile represent the human (H), dog (D), mouse (M), and rat (R) sequences and illustrate the position of gaps introduced to permit optimum alignment. Coding and noncoding exons are demonstrated by red and brown boxes, respectively, and repeats are illustrated by the smaller blue boxes. Regions of open chromatin and the +23 kb enhancer are shaded in green over the plot. Peaks of conserved sequence can be seen within each known regulatory region. Three well conserved peaks in regions of unknown function (−10 kb, +26 kb, and +30 kb) are also indicated. Pr1a, promoter 1a; Pr1b, promoter 1b.

the same degree of in-depth analysis (including extensive transgenic experiments) as the *SCL* locus. In the case of the *CFTR* data set, the length of the locus means that there are large stretches of

sequence that have not been subjected to any functional studies. As a result, at 63% sensitivity, 49 times as many conserved non-coding regions without known function than those with known

function are identified in the CFTR locus (Thomas et al. 2003). In the *SCL* data set, at 63% sensitivity, three times fewer conserved noncoding regions without known function than those with known function are identified. The β -globin data set (Stojanovic et al. 1999) is weakened by incomplete genomic coverage and the complexities of interpreting comparative sequence analyses in a locus containing clusters of paralogous genes. Neither the β -globin data set nor the CFTR data set offers a single resource with a comprehensive review of the underlying experimental data to support the definition of minimally defined regulatory elements and the transcription factor binding sites contained therein.

The comprehensive functional annotation of the *SCL* data set and the successful alignment of all known functional elements and binding sites allow strict performance criteria to be defined for multiple species analysis tools. These criteria can be used to set parameters during algorithm training or as a gold standard for sensitivity in evaluation. The detailed experimental work underlying functional annotation of the *SCL* locus allowed us to test the performance of pairwise and multiple alignments in their identification of regulatory elements with a degree of rigor that was probably hitherto impossible. Multiple alignments were superior to pairwise alignments in the selective identification of both regulatory elements a few hundred base pairs in length and individual transcription factor binding sites. The four-way alignment performed better than any of the three-way alignments. This suggests that increasing the number of species progressively increases the signal-to-noise ratio of the alignment. It is important to note that our approach in these analyses has been empirical. Thus, although it would be interesting to predict how much more useful multiple alignments would become as the number of mammalian species is increased (above the four presented herein), the lack of available sequence data for the *SCL* locus in other mammalian species precludes this. We did not use the chicken or fish *SCL* loci in our analysis, as most of the mammalian regulatory elements cannot be found in these sequences (Göttgens et al. 2000, 2002a). However, they are included in the data set, as comparisons of more divergent sequences may be useful in addressing other important questions, such as how similar patterns of gene expression can be achieved with a different organization of regulatory elements.

Functional annotation of the noncoding portion of the human genome represents the next major milestone of the Human Genome Project (Collins et al. 2003). The ENCODE program, to be launched later this year, has been set up with the objective of developing the necessary functional and computational resources. It is anticipated that the wealth of functional information, combined with the comparative sequence data, will make the *SCL* locus a valuable model for this next phase of genomics research.

METHODS

Production of Probes for Library Screening

The human *SCL* bHLH probe was amplified by PCR from genomic DNA, using standard procedures. Primers were CCATTCTCCTAACTCTTGTCCTC and CCAGCAGCCTAAGAACGCC, and the annealing temperature was 55°C. *SIL* and *MAP17* whole cDNA (available from the authors on request) were used as probes for these genes.

Sequencing, Annotation, and Alignments of *SCL* Loci

To isolate a clone covering the *SCL* locus, the RPCI canine BAC library 81 (Li et al. 1999) was screened using the *SCL* bHLH probe as described (Chapman et al. 2003). DNA was sequenced from a single dog clone, RP81-7016, containing *SIL*, *SCL*, and *MAP17* as described elsewhere (Lander et al. 2001). Briefly, 2–4 kb of sheared fragments were subcloned into pUC19. The sequences

were assembled with 1500 sequence reads. During the finishing process, difficult gaps were subjected to transposon libraries. The locations of coding exons were predicted with BLAST searches (<http://www.ncbi.nlm.nih.gov/BLAST/>; Altschul et al. 1990). The coding frame was maintained in each, and there was no ambiguity in the start or stop site of the various exons. RT-PCR from two dog cell lines, D064 (bone marrow) and IIIG7 (peripheral blood), was used to locate noncoding exons. Rat *SCL* sequence was extracted from the Berkeley Genome Pipeline (<http://pipeline.lbl.gov/rat/>). Pairwise BLAST analysis with mouse sequence was used to predict both noncoding and coding exon locations. Human, dog, mouse, and rat *SCL* sequences were trimmed from about 2 kb upstream of the fifth last exon of *SIL* to about 2 kb downstream of the last exon of *MAP17*. Repeats were masked using RepeatMasker (<http://repeatmasker.genome.washington.edu/cgi-bin/RepeatMasker>) on slow setting. The LAGAN server (<http://lagan.stanford.edu>) was used to create all possible two-, three-, and four-way alignments using these sequences.

SynPlot

SynPlot is a modification of the SynPlot Perl module (Göttgens et al. 2000), and is now implemented on a worldwide Web server (http://hscl.cimr.cam.ac.uk/syn_plot.html). It runs as described previously, with the exception that more than two sequences can be compared simultaneously to generate a single identity plot. To enable this, the scoring mechanism employs the sum of pairs method. Each base position within a user-defined window is examined sequentially, and the base in each species is compared with those with which it is aligned in the other species. For every matching base, 1 is scored. Therefore, the maximum score at any base position is 1 in a two-way alignment, 3 in a three-way alignment, 6 in a four-way alignment, and so on. The scores are summed for the entire window, and the alignment score is given as a fraction of maximum total possible score (window length multiplied by maximum possible score at each base position). Scores will thus range from 0 to 1.0 for each window. The window is then moved by user-defined increments along the alignment. Gaps introduced into individual sequences by the alignment are considered in the scoring; that is, they contribute to the length of the window, but not to the score. Features can be designed by the user, described in a configuration file, and annotated in GFF files. These are then drawn on an individual line for each sequence above the plot. Gaps introduced into individual sequences by the alignment will also be shown above the plot. SynPlot was designed to process global rather than local alignments; that is, it assumes that all the bases of each sequence have been used in the alignment. For this reason, gaps in the alignment will mean that the numbering on the x-axis refers to the position within the global alignment rather than in any one of the sequences. PeakExtractor is therefore provided on the Web server to enable the alignment in regions of interest to be examined (http://hscl.cimr.cam.ac.uk/peak_extractor.html; see supplementary online material at http://hscl.cimr.cam.ac.uk/additional_info.html for details). The output of the stand-alone SynPlot module is a Postscript file. The Web server uses the ps2pdf package to perform a conversion to PDF if required. For the purposes of this paper, SynPlots were generated for all the two-, three-, and four-way mammalian alignments using a 100-bp window moving at 25-bp increments. Three features in the configuration files were enabled: exons, coding sequence, and repeat regions.

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REFERENCES

- Altschul, S.F., Gish, W., Miller, W., Myers, E.W., and Lipman, D.J. 1990. Basic local alignment search tool. *J. Mol. Biol.* **215**: 403–410.
- Barton, L.M., Göttgens, B., Gering, M., Gilbert, J.G., Grafham, D., Rogers, J., Bentley, D., Patient, R., and Green, A.R. 2001. Regulation of the stem cell leukemia (SCL) gene: A tale of two fishes. *Proc. Natl. Acad. Sci.* **98**: 6747–6752.
- Begley, C.G. and Green, A.R. 1999. The SCL gene: From case report to critical hematopoietic regulator. *Blood* **93**: 2760–2770.
- Boffelli, D., McAuliffe, J., Ovcharenko, D., Lewis, K.D., Ovcharenko, I., Pachter, L., and Rubin, E.M. 2003. Phylogenetic shadowing of primate sequences to find functional regions of the human genome. *Science* **299**: 1391–1394.
- Brudno, M., Do, C.B., Cooper, G.M., Kim, M.F., Davydov, E., Green, E.D., Sidow, A., and Batzoglu, S. 2003. LAGAN and Multi-LAGAN: Efficient tools for large-scale multiple alignment of genomic DNA. *Genome Res.* **13**: 721–731.
- Brudno, M., Chapman, M., Göttgens, B., Batzoglu, S., and Morgenstern, B. 2004. Fast and sensitive multiple alignment of large genomic sequences. *BMC Bioinformatics* (in press).
- Chapman, M.A., Charchar, F.J., Kinston, S., Bird, C.P., Grafham, D., Rogers, J., Grutzner, F., Marshall Graves, J.A., Green, A.R., and Göttgens, B. 2003. Comparative and functional analyses of LYL1 loci establish marsupial sequences as a model for phylogenetic footprinting. *Genomics* **81**: 249–259.
- Cioffi, C.C., Middleton, D.L., Wilson, M.R., Miller, N.W., Clem, L.W., and Warr, G.W. 2001. An IgH enhancer that drives transcription through basic helix-loop-helix and Oct transcription factor binding motifs. Functional analysis of the E(mu)3' enhancer of the catfish. *J. Biol. Chem.* **276**: 27825–27830.
- Collins, F.S., Green, E.D., Guttacher, A.E., and Guyer, M.S. 2003. A vision for the future of genomics research. *Nature* **422**: 835–847.
- Cooper, G.M., Brudno, M., Green, E.D., Batzoglu, S., and Sidow, A. 2003. Quantitative estimates of sequence divergence for comparative analyses of mammalian genomes. *Genome Res.* **13**: 813–820.
- Elnitski, L., Hardison, R.C., Li, J., Yang, S., Kolbe, D., Eswara, P., O'Connor, M.J., Schwartz, S., Miller, W., and Chiaromonte, F. 2003. Distinguishing regulatory DNA from neutral sites. *Genome Res.* **13**: 64–72.
- Flint, J., Tufarelli, C., Peden, J., Clark, K., Daniels, R.J., Hardison, R., Miller, W., Philipsen, S., Tan-Un, K.C., McMorrow, T., et al. 2001. Comparative genome analysis delimits a chromosomal domain and identifies key regulatory elements in the α lobin cluster. *Hum. Mol. Genet.* **10**: 371–382.
- Fordham, J.L., Göttgens, B., McLaughlin, F., and Green, A.R. 1999. Chromatin structure and transcriptional regulation of the stem cell leukaemia (SCL) gene in mast cells. *Leukemia* **13**: 750–759.
- Frazer, K.A., Elnitski, L., Church, D.M., Dubchak, I., and Hardison, R.C. 2003. Cross-species sequence comparisons: A review of methods and available resources. *Genome Res.* **13**: 1–12.
- Gilligan, P., Brenner, S., and Venkatesh, B. 2002. Fugu and human sequence comparison identifies novel human genes and conserved non-coding sequences. *Gene* **294**: 35.
- Göttgens, B., Barton, L.M., Gilbert, J.G., Bench, A.J., Sanchez, M.J., Bahn, S., Mistry, S., Grafham, D., McMurray, A., Vaudin, M., et al. 2000. Analysis of vertebrate SCL loci identifies conserved enhancers. *Nat. Biotechnol.* **18**: 181–186.
- Göttgens, B., Gilbert, J.G., Barton, L.M., Grafham, D., Rogers, J., Bentley, D.R., and Green, A.R. 2001. Long-range comparison of human and mouse SCL loci: Localized regions of sensitivity to restriction endonucleases correspond precisely with peaks of conserved noncoding sequences. *Genome Res.* **11**: 87–97.
- Göttgens, B., Barton, L.M., Chapman, M.A., Sinclair, A.M., Knudsen, B., Grafham, D., Gilbert, J.G., Rogers, J., Bentley, D.R., and Green, A.R. 2002a. Transcriptional regulation of the stem cell leukemia gene (SCL)—Comparative analysis of five vertebrate SCL loci. *Genome Res.* **12**: 749–759.
- Göttgens, B., Nastos, A., Kinston, S., Piltz, S., Delabesse, E.C., Stanley, M., Sanchez, M.J., Ciaui-Uitz, A., Patient, R., and Green, A.R. 2002b. Establishing the transcriptional programme for blood: The SCL stem cell enhancer is regulated by a multiprotein complex containing Ets and GATA factors. *EMBO J.* **21**: 3039–3050.
- Gumucio, D.L., Shelton, D.A., Bailey, W.J., Slightom, J.L., and Goodman, M. 1993. Phylogenetic footprinting reveals unexpected complexity in trans factor binding upstream from the ϵ -globin gene. *Proc. Natl. Acad. Sci.* **90**: 6018–6022.
- Lander, E.S., Linton, L.M., Birren, B., Nusbaum, C., Zody, M.C., Baldwin, J., Devon, K., Dewar, K., Doyle, M., FitzHugh, W., et al. 2001. Initial sequencing and analysis of the human genome. *Nature* **409**: 860–921.
- Li, R., Mignot, E., Faraco, J., Kadotani, H., Cantanese, J., Zhao, B., Lin, X., Hinton, L., Ostrander, E.A., Patterson, D.F., et al. 1999. Construction and characterization of an eightfold redundant dog genomic bacterial artificial chromosome library. *Genomics* **58**: 9–17.
- Loots, G.G., Locksley, R.M., Blankespoor, C.M., Wang, Z.E., Miller, W., Rubin, E.M., and Frazer, K.A. 2000. Identification of a coordinate regulator of interleukins 4, 13, and 5 by cross-species sequence comparisons. *Science* **288**: 136–140.
- Miles, C., Elgar, G., Coles, E., Kleinjan, D.J., van Heyningen, V., and Hastie, N. 1998. Complete sequencing of the Fugu WAGR region from WT1 to PAX6: Dramatic compaction and conservation of synteny with human chromosome 11p13. *Proc. Natl. Acad. Sci.* **95**: 13068–13072.
- Miller, W. 2001. Comparison of genomic DNA sequences: Solved and unsolved problems. *Bioinformatics* **17**: 391–397.
- Sanchez, M., Göttgens, B., Sinclair, A.M., Stanley, M., Begley, C.G., Hunter, S., and Green, A.R. 1999. An SCL 3' enhancer targets developing endothelium together with embryonic and adult haematopoietic progenitors. *Development* **126**: 3891–3904.
- Sanchez, M.J., Bockamp, E.O., Miller, J., Gambardella, L., and Green, A.R. 2001. Selective rescue of early haematopoietic progenitors in Scl(−/−) mice by expressing Scl under the control of a stem cell enhancer. *Development* **128**: 4815–4827.
- Schwartz, S., Zhang, Z., Frazer, K.A., Smit, A., Riemer, C., Bouck, J., Gibbs, R., Hardison, R., and Miller, W. 2000. PipMaker—A web server for aligning two genomic DNA sequences. *Genome Res.* **10**: 577–586.
- Schwartz, S., Elnitski, L., Li, M., Weirauch, M., Riemer, C., Smit, A., Green, E.D., Hardison, R.C., and Miller, W. 2003. MultiPipMaker and supporting tools: Alignments and analysis of multiple genomic DNA sequences. *Nucleic Acids Res.* **31**: 3518–3524.
- Shelton, D.A., Stegman, L., Hardison, R., Miller, W., Bock, J.H., Slightom, J.L., Goodman, M., and Gumucio, D.L. 1997. Phylogenetic footprinting of hypersensitive site 3 of the β -globin locus control region. *Blood* **89**: 3457–3469.
- Sinclair, A.M., Göttgens, B., Barton, L.M., Stanley, M.L., Pardanau, L., Klaine, M., Gering, M., Bahn, S., Sanchez, M., Bench, A.J., et al. 1999. Distinct 5' SCL enhancers direct transcription to developing brain, spinal cord, and endothelium: Neural expression is mediated by GATA factor binding sites. *Dev. Biol.* **209**: 128–142.
- Sinclair, A.M., Bench, A.J., Bloor, A.J., Li, J., Göttgens, B., Stanley, M.L., Miller, J., Piltz, S., Hunter, S., Nacheva, E.P., et al. 2002. Rescue of the lethal scl(−/−) phenotype by the human SCL locus. *Blood* **99**: 3931–3938.
- Stojanovic, N., Florea, L., Riemer, C., Gumucio, D., Slightom, J., Goodman, M., Miller, W., and Hardison, R. 1999. Comparison of five methods for finding conserved sequences in multiple alignments of gene regulatory regions. *Nucleic Acids Res.* **27**: 3899–3910.
- Tagle, D.A., Koop, B.F., Goodman, M., Slightom, J.L., Hess, D.L., and Jones, R.T. 1988. Embryonic ϵ and γ globin genes of a prosimian primate (*Galago crassicaudatus*). Nucleotide and amino acid sequences, developmental regulation and phylogenetic footprints. *J. Mol. Biol.* **203**: 439–455.
- Thomas, J.W., Touchman, J.W., Blakesley, R.W., Bouffard, G.G., Beckstrom-Sternberg, S.M., Margulies, E.H., Blanchette, M., Siepel, A.C., Thomas, P.J., McDowell, J.C., et al. 2003. Comparative analyses of multi-species sequences from targeted genomic regions. *Nature* **424**: 788–793.
- Wasserman, W.W., Palumbo, M., Thompson, W., Fickett, J.W., and Lawrence, C.E. 2000. Human-mouse genome comparisons to locate regulatory sites. *Nat. Genet.* **26**: 225–228.

WEB SITE REFERENCES

- <http://bio.cse.psu.edu/pipmaker/>; PipMaker home page.
- <http://lagan.stanford.edu/>; LAGAN home page.
- <http://pipeline.lbl.gov/rat/>; Berkeley Genome Pipeline.
- <http://repeatmasker.genome.washington.edu/cgi-bin/RepeatMasker/>; RepeatMasker server.
- <http://www.gsd.lbl.gov/vista/>; Vista home page.
- <http://www.ncbi.nlm.nih.gov/BLAST/>; BLAST home page
- <http://www.sanger.ac.uk/Software/formats/GFF/>; GFF format.
- http://hscl.cimr.cam.ac.uk/supplementary_data.html; Supplemental data for this article.
- http://hscl.cimr.cam.ac.uk/genomic_datasets.html; Data sets are described in this article.

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