

Video Article

Using SCOPE to Identify Potential Regulatory Motifs in Coregulated Genes

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Abstract

SCOPE is an ensemble motif finder that uses three component algorithms in parallel to identify potential regulatory motifs by over-representation and motif position preference¹. Each component algorithm is optimized to find a different kind of motif. By taking the best of these three approaches, SCOPE performs better than any single algorithm, even in the presence of noisy data¹. In this article, we utilize a web version of SCOPE² to examine genes that are involved in telomere maintenance. SCOPE has been incorporated into at least two other motif finding programs^{3,4} and has been used in other studies⁵⁻⁸.

The three algorithms that comprise SCOPE are BEAM⁹, which finds non-degenerate motifs (ACCGGT), PRISM¹⁰, which finds degenerate motifs (ASCGWT), and SPACER¹¹, which finds longer bipartite motifs (ACCnnnnnnnGGT). These three algorithms have been optimized to find their corresponding type of motif. Together, they allow SCOPE to perform extremely well.

Once a gene set has been analyzed and candidate motifs identified, SCOPE can look for other genes that contain the motif which, when added to the original set, will improve the motif score. This can occur through over-representation or motif position preference. Working with partial gene sets that have biologically verified transcription factor binding sites, SCOPE was able to identify most of the rest of the genes also regulated by the given transcription factor.

Output from SCOPE shows candidate motifs, their significance, and other information both as a table and as a graphical motif map. FAQs and video tutorials are available at the SCOPE web site which also includes a "Sample Search" button that allows the user to perform a trial run.

SCOPE has a very friendly user interface that enables novice users to access the algorithm's full power without having to become an expert in the bioinformatics of motif finding. As input, SCOPE can take a list of genes, or FASTA sequences. These can be entered in browser text fields, or read from a file. The output from SCOPE contains a list of all identified motifs with their scores, number of occurrences, fraction of genes containing the motif, and the algorithm used to identify the motif. For each motif, result details include a consensus representation of the motif, a sequence logo, a position weight matrix, and a list of instances for every motif occurrence (with exact positions and "strand" indicated). Results are returned in a browser window and also optionally by email. Previous papers describe the SCOPE algorithms in detail^{1,2,9-11}.

Video Link

The video component of this article can be found at <http://www.jove.com/video/2703/>

Protocol

1. Prepare a list of names for genes that you believe are co-regulated for analysis by SCOPE.

Save the list as a text file or copy it to the clipboard to paste into SCOPE in step 3. The file should contain one gene name per line with no additional information. Alternatively, you can prepare the list as a FASTA file containing the actual sequences to be analyzed.

2. Start your web browser and connect to the URL: <http://genie.dartmouth.edu/SCOPE/>

3. Enter the information that SCOPE needs to perform the analysis.

The initial SCOPE page is shown in Figure 1. Different sections are addressed in this step.

1. Use the 'Species' popup menu to choose the species you will be examining. It is important to choose the correct species because SCOPE refers to the genome to calculate background frequencies of occurrence for any candidate motif it is examining.

2. Use the 'upstream sequence' radio buttons to choose either intergenic or fixed length. Intergenic will analyze all the sequence between of the gene you are looking at and the previous (upstream) gene. This will mean that different upstream lengths will be used for each gene. Choosing fixed length will look at exactly that number of nucleotides upstream from the start of the current gene. In this case, SCOPE will examine the same length of upstream sequence for each gene, even if that extends into the previous gene (or not). Typically, 800 nts is the best length to choose, but this can vary with species.
3. Next tell SCOPE what gene set to analyze either by pasting in your gene list into the gene list text box, or by pressing the 'choose file' button to select the file containing the list of genes that you created earlier. You may, alternatively, paste in a FASTA sequence file into the same text box.
4. The next section of the page contains a checkbox for 'Examine genome for other genes containing found motif(s)?' This option can add considerable analysis time since SCOPE has to evaluate every other gene in the genome. However, this can be very useful in identifying other genes that are good candidates for being co-regulated with the genes in the starting gene set. Since SCOPE analyses are relatively quick, it is suggested that you leave this off in your initial analysis. It can always be turned on from the results page to rerun the analysis, as explained in the results section.
5. The 'Results must include' section can be used to enter a motif that you want SCOPE to include in its analysis. You might want to do this if you are looking for a specific motif.
6. The last section on the page can be used to enter your email address and a comment to be saved with the analysis. If this is filled in, SCOPE will send an email with a link back to the web page containing results, and it will also include two attachments. One is a plain text file that has all the analysis results in human readable format. The second attachment contains an XML file that has every result that SCOPE has found in a computer readable format. If you want to do some additional analysis on the results, the XML file is very useful. Both files are "zipped" before being sent with the email.
7. For this demo, we will start with the same information. This can be easily achieved by pressing the 'Sample Search' button which will fill in the necessary information. Press this button now. Three genes will be entered for you and appropriate choices made for the other fields. Leave these as they are set. The three genes are involved in telomere maintenance in *Saccharomyces cerevisiae*. The filled in form is shown in Figure 2. Press the 'Run SCOPE' button at the bottom of the page to start the analysis.

4. Representative Results:

The main results of the analysis are shown in Figure 3. The top of the page contains a table of information about the motifs that were found by SCOPE. The first column contains a list of motifs that were found and small colored squares serve as a legend for the graphical motif map shown below. The display of any given motif may be toggled on or off by clicking in the colored box (or where the colored box would be). This can be very useful to hide the display of highly repeated motifs that might make it difficult to see the less prevalent motif patterns.

Other columns of data are Count (the number of occurrences of that motif in the entire gene set), Sig value (an indication of the significance of that motif), Coverage (the percentage of the submitted genes that contain at least one instance of that motif), and Algorithm (which of the three component algorithms was used to detect the motif).

Clicking on any of the listed motifs will take the user to a page containing detailed information for that motif. The results details are shown for the cyan motif (atgnnnttg) in Figure 4. On this page, the motif is represented in three ways: a sequence logo, a position weight matrix, and a list of all motif instances with their positions, strands and genes.

A little further down the page are some additional details about the results of looking for other genes containing this motif. As can be seen, in this case there were 1344 other genes containing the motif, all of which actually improved the Sig value when added to the original gene set. Pressing 'Add checked genes to search' will return to the SCOPE setup page with these genes added to the original gene set and the parameters set as they were previously. In this case, 10 extra genes are added to the original three.

Figure 5 shows the results of the analysis containing the extra genes for this motif. The original three genes are on the bottom of the results (in lower case). Looking at the pattern of motifs in the upstream region of these extra genes clearly shows that they are similar. In fact, many of these genes are involved in telomere maintenance as were the original three genes. Note also that the original motif is now the highest scoring motif in this set.

Another set of SCOPE results is shown in Figure 6. In this case, the set of genes are those that are involved in ribosome biogenesis in *Saccharomyces cerevisiae*. These genes are not actually part of the ribosome but are responsible for assembling ribosomes and include a number of modification enzymes. What is clear in the figure is that the red and green motifs form a reliable pattern that is likely to be involved in regulation of the genes in this set. We are investigating this pattern of "modules" in more detail and will report on it in a later publication.

Welcome to SCOPE (Suite for Computational Identification Of Promoter Elements), an ensemble of programs aimed at identifying novel *cis*-regulatory elements from groups of upstream sequences.

The SCOPE motif finder is designed to identify candidate regulatory DNA motifs from sets of genes that are coordinately regulated. SCOPE motif finder uses an ensemble of three programs behind the scenes to identify different kinds of motifs - BEAM identifies nondegenerate motifs (e.g. ACGTGC), PRISM identifies degenerate motifs (e.g. AWCGRYH), and SPACER identifies bipartite motifs (e.g. ACCNNNNNNNGTT). All parameters are automatically set to find the optimal length motif and degree of degeneracy in the reported motifs. To get started, choose a species and an upstream length, then enter a list of gene names (or FASTA sequences) and press the "Run SCOPE" button. For more details, see FAQs and Publications links at the top of the page.

Species: Upstream sequence: Intergenic Fixed [Help](#)

Enter gene list or FASTA DNA sequences: [Help](#)

- OR -

Upload file with that info: no file selected

Examine genome for other genes containing found motif(s)? [Help](#)
(note: this can take considerable extra time)

Results must include the following motifs:

Search: Plus and minus strands Plus strand only

Only search for these motifs?

If you would like the results (also) returned to you by email, please fill in the fields below.

Email address:

Email subject:

Figure 1. Main SCOPE input page. This page is used to enter the genes to be analyzed and to define the species and the length of upstream region to be examined. Optionally, the user can request the results by email or restrict the search to any specified motif. Video help is also available.

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Species: Upstream sequence: Intergenic Fixed [Help](#)

Comment: November 21, 2006 (ftp://ftp.ncbi.nih.gov/)

[Sample Search](#)

Enter gene list or FASTA DNA sequences: [Help](#)

- OR -

Upload file with that info: no file selected

Examine genome for other genes containing found motif(s)? [Help](#)
(note: this can take considerable extra time)

Results must include the following motifs:

Search: Plus and minus strands Plus strand only
 Only search for these motifs?

If you would like the results (also) returned to you by email, please fill in the fields below.

Email address:

Email subject:

Figure 2. Main SCOPE input page with values filled in for performing a search. These parameters are the result of pressing the 'Sample Search' button. In this case, the check box to find other genes containing the motifs found by SCOPE is checked. This option takes longer to compute (every gene in the genome has to be examined) but can provide interesting insights.

Top 10 Results from SCOPE: [View Top 19 Results](#)

Consensus Sequence (Caveat)	Count	Sig Value (What is this?)	Coverage (What is this?)	Algorithm (What is this?)
■ acadmkdhtgty	14	69.6	100.0%	SPACER
■ scwgsaw	23	40.3	100.0%	PRISM
■ swaasttw	12	30.5	100.0%	PRISM
■ ttcgnnnnnnacaa	7	29.5	100.0%	SPACER
■ atgnnnnttg	8	29.0	100.0%	SPACER
agannnnnnnnnctg	8	26.8	66.7%	SPACER
aatnnnnnnnact	10	22.0	100.0%	SPACER
atwgтта	8	21.7	100.0%	PRISM
aagnnnnnnnnnnnaca	10	20.8	100.0%	SPACER
actnnnnnnnnnttgc	5	19.7	100.0%	SPACER

Here are the results of individual programs:

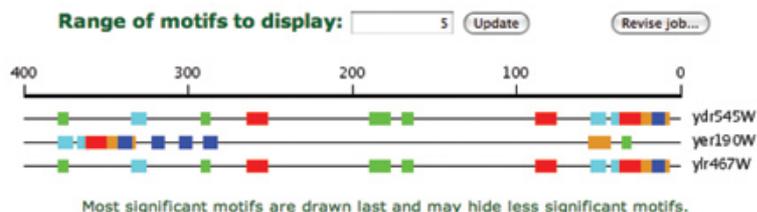


Figure 3. Main SCOPE results page. This page summarizes the results of the SCOPE search. A list of all high scoring motifs is provided and a color coded motif map shows the positioning of the identified motifs in the set of genes analyzed. Clicking on a colored box next to a motif will toggle the display of that motif on or off in the motif map. In addition to a significance score (Sig value), the fraction of genes containing the motif (coverage), and the algorithm used to find that motif are also provided.

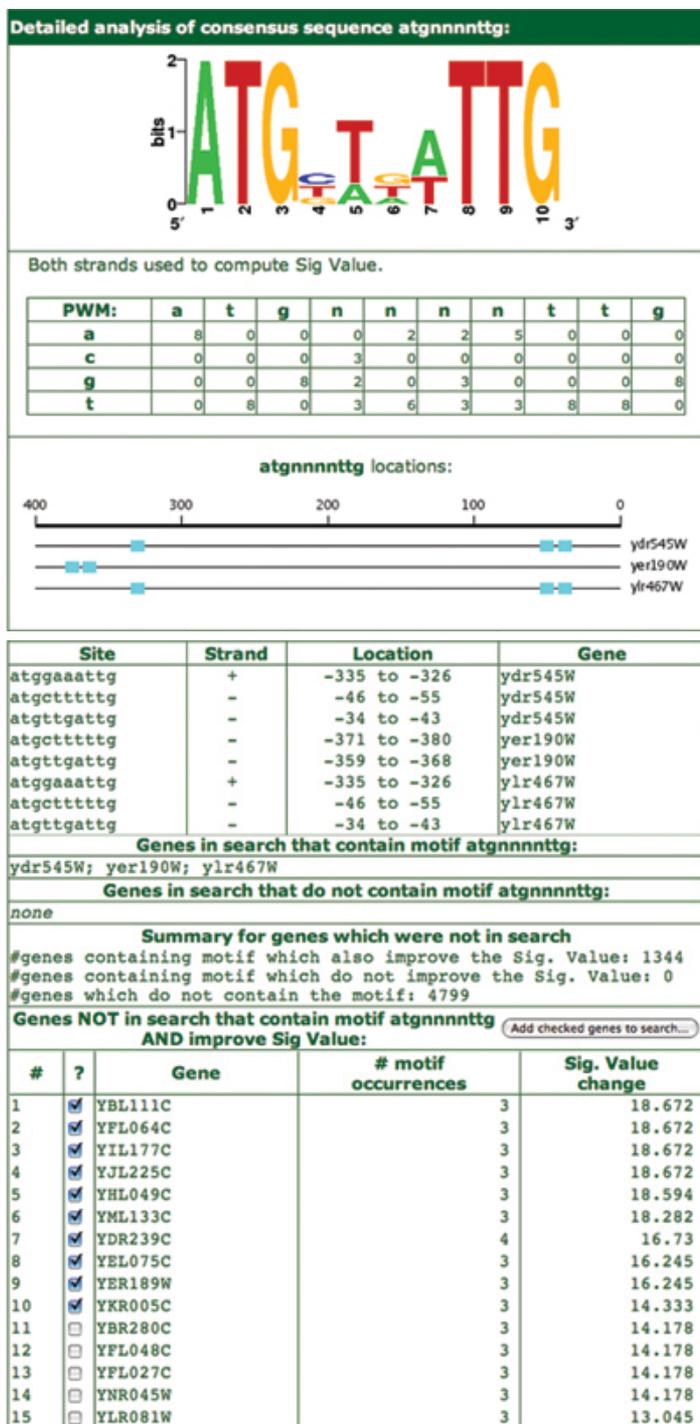


Figure 4. This results detail page is brought up when a specific motif is clicked in the main results page. It shows details of the individual motif. The sequence logo, the position weight matrix, and the consensus sequence each represent a different kind of summary of the list of motif instances also on the page. Since 'find extra genes' was checked in the original search setup, there is also information on this page about any other genes in the genome that contain this motif. From this page it is also possible to start another SCOPE run including the extra genes identified on this page.

Top 10 Results from SCOPE: View Top 21 Results

Consensus Sequence <small>(Caveat)</small>	Count	Sig Value <small>(What is this?)</small>	Coverage <small>(What is this?)</small>	Algorithm <small>(What is this?)</small>
atgnnnttg	39	171.7	100.0%	LOOKUP
acadhmkdhtgty	60	363.9	84.6%	SPACER
agahddnbabwdwctg	38	231.8	76.9%	SPACER
kvwgmw	314	208.7	100.0%	PRISM
atgnnnttg	39	168.3	100.0%	SPACER
ttgtnnnttcgaaa	22	138.7	84.6%	SPACER
aacaatnnnnnnnnntgta	20	134.5	76.9%	SPACER
tcannnnggaaat	20	127.7	76.9%	SPACER
cagnnnnncttatac	20	125.0	76.9%	SPACER
cagannctgc	20	121.5	76.9%	SPACER

Here are the results of individual programs: BEAM
PRISM
SPACER

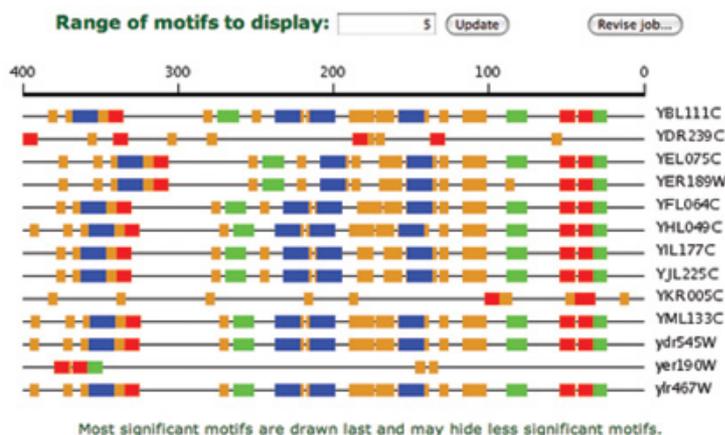


Figure 5. This figure shows the results of looking for extra genes for the motif 'atgnnnttg' shown in Figure 4. The original three genes are in lower case at the bottom of the motif map. The additional genes are shown in upper case. There is a clear pattern to the motifs in the upstream regions of these genes. Notice also that the specified motif shows an algorithm as 'LOOKUP' because that is how it was identified. It actually matches the 5th motif found by SPACER in this analysis.

Top 9 Results from SCOPE:

Consensus Sequence (Caveat)	Count	Sig Value (What is this?)	Coverage (What is this?)	Algorithm (What is this?)
aaawtttbbh	493	758.1	98.0%	PRISM
abctcatcd	142	347.9	91.2%	PRISM
aaaaannttt	121	137.4	61.8%	SPACER
aaaannttca	89	134.2	65.7%	SPACER
aaaanntttca	92	132.2	64.7%	SPACER
atgnnttga	142	122.0	76.5%	SPACER
gcgatganntga	28	80.0	27.5%	SPACER
atgnnatgaga	28	73.1	19.6%	SPACER
gatgagngag	24	68.0	20.6%	SPACER

Here are the results of individual programs: PRISM SPACER

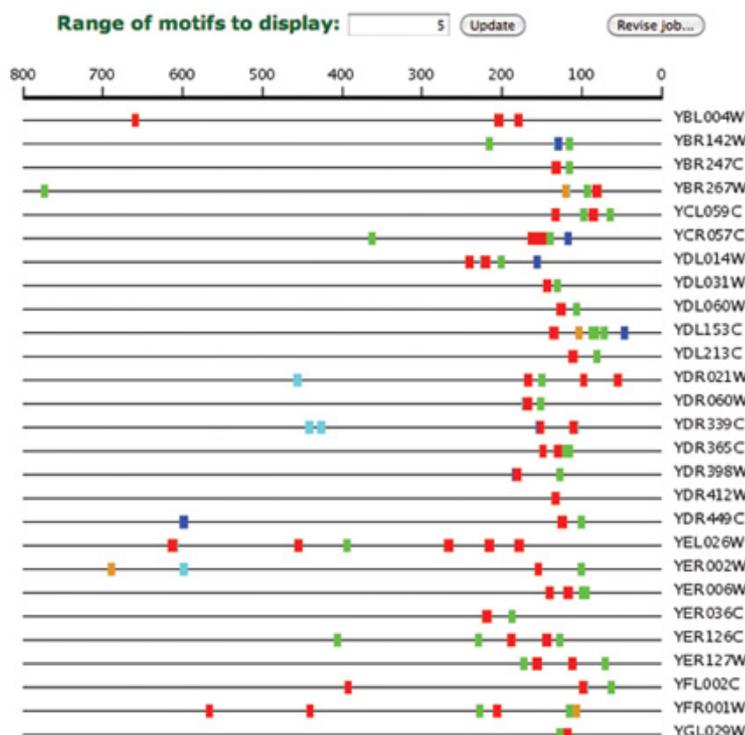


Figure 6. SCOPE output for genes involved in ribosome biogenesis in *Saccharomyces cerevisiae*. Note the conserved pattern of modules consisting of the motifs 'aaawtttbbh' (red) and 'abctcatcd' (green) separated by about 10-30 nts and present at 100-200 nucleotides upstream of transcription start for the gene.

Discussion

SCOPE provides the researcher with a powerful tool to use for the identification of potential regulatory motifs in sets of coordinately regulated genes. The user is not required to guess at the size of the motif or the number of occurrences of the motif as many other motif finding sites require. These parameters are basically unknowable until the motif is identified. The interface is very simple both for entering sequences or gene names and for viewing the output.

SCOPE output provides detailed information about all of the motifs that are identified, using three different ways of motif representation. Each instance of the motif in all of the genes is listed with position and "strand" information. Graphical results in the form of motif maps provide a visual display that is easy to understand and provides an intuitive way to see patterns in the motifs that are present.

SCOPE is very robust to the presence of noise in the data. Typically, this takes the form of extra genes being present in the starting set that might not actually be co-regulated with the rest of the genes. This often happens when starting with genes that are co-expressed in microarray experiments. Sometimes the experiment is noisy, or there may be several transcription factors activated in the experimental conditions used for the microarray experiment. These different transcription factors will likely have different target sites on the DNA. Even in the presence of 4-fold extraneous genes (noise:signal ratio is 4:1), SCOPE is still maintains 50% of its accuracy in predicting sites¹.

Although SCOPE contains over 2 million synonyms for gene names, it sometimes fails to identify some genes names. We are constantly updating our synonym lists, but sometimes find that different synonyms refer to the same gene. In those cases, we do not include the synonyms

because of the ambiguity. If you have a gene name that is not found by SCOPE, it is recommended that you refer to the genome specific site to find an alternative gene name to use in SCOPE. Examples of appropriate gene names for each species are provided by SCOPE.

SCOPE currently contains 72 species with new species being added all the time. The web site contains video help as well as FAQs. Source code is freely available to academic users by writing to RHG.

Disclosures

No conflicts of interest declared.

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