

# bioSyntax Manual v0.1

April 13th, 2018

Grok your data

The objective of bioSyntax is to bring you closer to your data, giving you an intuitive & empathetic understanding of biology. To appreciate all that bioSyntax has to offer read this short manual (~10 minutes) and go explore.

1. Getting Started
2. Reading bioSyntax
3. Supported Formats
4. Developing New Formats & Themes

## Getting Started

**See: Installing bioSyntax**

bioSyntax integrates seamlessly with **vim** (*Linux / Mac / Win*), **sublime** (*Linux / Mac / Win*), **gedit** (*Linux / Win*), & **less** (*Linux / Mac*). After installing bioSyntax files will automatically be detected by file-extension.

### Reading large-data

For very large data sets, it's often slow to open them in a text editor. It's best to use the command-line program **less** which will read your file from a data-stream.

### Read your large-data set with less directly

```
# If your file is uncompressed, it can be read directly.  
# less will recognize the file extension (.XYZ)
```

```
cd ~/myData/
```

```
less dbSNP107_common.vcf
```

```
less hg19.fa
```

### Streaming your data directly into less with pipes |

```
# If your file is compressed, you can 'pipe' the data
# using the "|" operator from decompression, directly into
# less. You must prefix the file extension you want
# as file formats are not recognized within streams.
```

```
cd ~/myCompressedData/
```

```
samtools view -h NA12878_hg38.bam | sam-less
```

```
gzip -dc dbSNP107_rare.vcf.gz | vcf-less
```

```
gzip -dc hg38.fa.gz | fa-less
```

### Bypassing bioSyntax (data in plain-text)

For vim Type :syntax off in vim

For less

```
# You may want to view your data without syntax highlighting
# such as where a file is improperly formatted or very large
# files where syntax highlighting may be slow (i.e. VCF files
# with hundreds of columns).
```

```
# 1. Pipe your data through cat
cat snp_1000genomes.vcf | less -
```

```
# 2. Within less, switch to a visual editor
less snp_1000genomes.vcf
# press 'CTRL-C' to stop process
# press 'v' to switch to visual editor
```

## Reading Data

### Nucleotides

bioSyntax implements a novel, full IUPAC Nucleotide Code coloring (Figure 1). Ambiguous bases are represented by an ~additive color-mixing of the parent bases. For example, Thymine (blue) + Cytosine (red) are both pYrimidines (magenta).

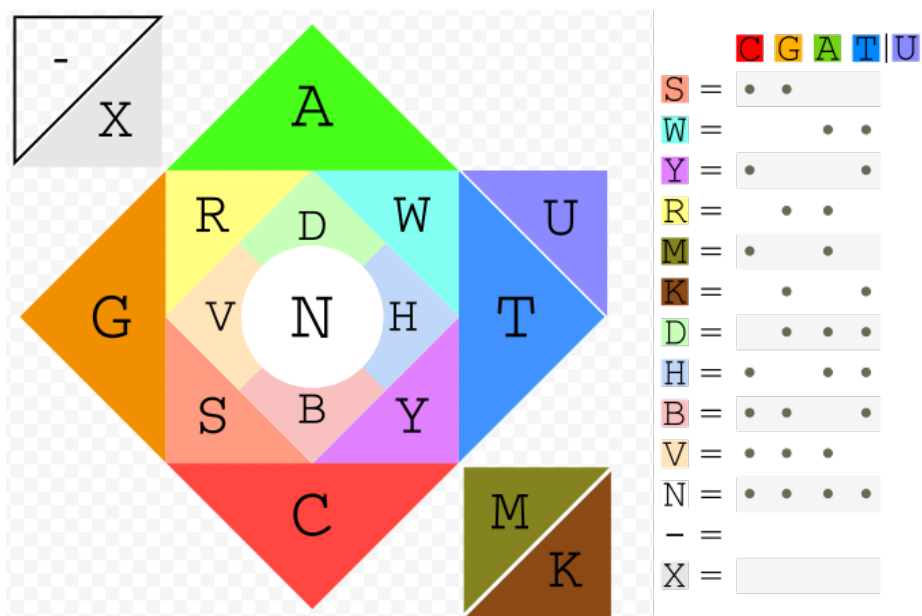


Figure 1: bioSyntax IUPAC nucleotide colouring

An intuitive feature of the bioSyntax color scheme is that the ‘GC-content’ of a sequence can be quickly approximated by how warm (high GC, red-orange) or cool (low GC, blue-green) a sequence looks (Figure 2).

### PHRED Scores

When available, bioSyntax will highlight PHRED quality scores in a step-gradient of blacks (PHRED = 0-10) to whites (PHRED = 40+) (Figure 3).

### CIGAR Strings

In `.sam` files the Query:Reference alignment is summarized efficiently but illegibly as a CIGAR String. With a little bit of highlighting these become much easier to read (Figure 4).

### Amino Acid Color Schemes

You can choose from several color-schemes for amino-acid fasta files. The **Fasta Clustal** (Default) syntax colors amino acids based on their physiochemical properties, so does **Fasta Hydrophobicity**, or you may prefer better discrimination of each amino acids with **Fasta Zappo** or **Fasta Taylor**.



101M	101M
101M	101M
2S99M	2S99M
15M3I13M2I50M2I16M	15M3I13M2I50M2I16M
101M	101M
82M1D19M	82M1D19M
10S63M1D18M2I7M	10S63M1D18M2I7M
13M1X37M50H	13M1X37M50H
101M	101M
101M	101M
101M	101M
101M	101M
101M	101M

Figure 4: CIGAR strings colouring

## Support

### Report a bug / Ask a question

The fastest way to get an answer is to:

- 1) Search / Open an issue on the bioSyntax Repo.

Please Include: - A detailed and descriptive title. - Enough information about what did for someone else to replicate the problem. - Information about the operating system / software you're using (`uname -a`) - If it's a syntax highlighting issue: a screenshot of the error and a small bit of the input file you used.

- 2) If you really don't want to make a (fake) github account. Email [info@bioSyntax.org](mailto:info@bioSyntax.org) and we'll open the issue, but it will be slower.

## Collaborating on bioSyntax

bioSyntax is a community-oriented project for scientific syntax highlighting. We encourage you to change and customize it to suit your needs.

Check out the Development page to create syntax-highlighting for custom file-formats and for other ways to help out.

## Supported File Formats

File format and software compatibility matrix for bioSyntax.

### Core bioSyntax

File Format	Description	sublime	vim	gedit	less
.fasta	Generic nt/aa sequence	X	X	X	X
.fastq	Fasta + PHRED quality	X	X	X	X
.clustal	Multiple Sequence Alignment	X	X	X	X
.bed	Genomic Ranges	X	X	X	X
.gtf	Genomic Annotation	X	X	X	X
.pdb	Protein Structure	X	X	X	X
.vcf	Variant Call Format	X	X	X	X
.sam	NGS Sequence Data	X	X	X	X

### Auxillary Syntaxes

File Format	Description	sublime	vim	gedit	less
.fasta	fasta alternative AA colors				
-	Clustal	X	-	X	-
-	Taylor	X	-	X	-
-	Zappo	X	-	X	-
-	Hydrophobicity	X	-	X	-
.fai	Fasta Index (faidx)	X	X	X	X
.flagstat	samtools flag summary	X	X	X	X
.wig	Wiggle data	-	-	X	-
.pdbx	Protein Structure (large)	-	-	-	-
.phylip	Multiple Sequence Alignment	-	-	-	-
.newick	Tree Format	-	-	-	-
.nexus	Phylogenetics data	-	X	-	-
.cwl	Common Workflow Language	-	-	-	-

See Also: [Alternative/User Syntax Definitions](#)