

# Blixem User Manual

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*17 January 2011*

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## Introduction

This manual explains how to configure, run and use Blixem. Blixem is an interactive browser of pairwise matches displayed as multiple alignments. It is not strictly a multiple alignment tool, rather a 'one-to-many' alignment. It is used to check the alignments of nucleotide and amino acid sequences against a reference sequence.

Blixem is maintained by the Wellcome Trust Sanger Institute and is available as part of the SeqTools package. The software can be downloaded from the Sanger Institute's website: <http://www.sanger.ac.uk>.

### An aside about the name “Blixem”

“BLIXEM” was originally an acronym for “BLast matches In an X-windows Embedded Multiple alignment”, although this is a bit of a misnomer now because Blixem can handle any kind of alignment, not just BLAST matches. We have dropped the acronym, and the capital letters, so the correct name is just “Blixem”.

## Getting Started

### Running Blixem

As a minimum, Blixem takes the following required arguments:

```
Blixem --display-mode N|P <features_file>
```

Where <features\_file> is the path name of a GFF version 3 file containing the alignments and any other features.

The '`--display-mode`' argument is the only mandatory option. It defines the display mode: 'N' for nucleotide or 'P' for protein. .

Run 'blixem' without any arguments to see further usage information.

### Input files

Blixem takes one or two files as input: a mandatory GFF version 3 file containing the features and, optionally, a separate file containing the reference sequence in FASTA format.

```
Blixem -m N|P [<reference_sequence_file>] <features_file>
```

If the reference sequence file is not provided, the reference sequence must be supplied in FASTA format at the end of the GFF file, following a comment line that reads '##FASTA'.

*Note that the reference sequence must always be a nucleotide sequence and match sequences must be the correct type for the mode, i.e. nucleotide sequences for nucleotide mode or protein sequences for protein mode.*

### GFF file:

Blixem uses the GFF version 3 file format. In this section we give a very brief description of this file format; see <http://www.sequenceontology.org/gff3.shtml> for a full description.

The GFF file should start with the following two comment lines. (Additional comments can be included but may be ignored.)

```
##gff-version 3
##sequence-region chr4-04_210623-364887 44144 154265
```

Each subsequent line defines a feature. A feature line must have the following 8 tab-separated columns:

reference_sequence_name	source	type	start	end	score	strand	phase
-------------------------	--------	------	-------	-----	-------	--------	-------

An optional 9<sup>th</sup> column defines any tags (separated by semi-colons). Blixem supports the following GFF tags. (Additional tags can be supplied but may be ignored.)

- Target** (required for alignments)
- Gap** (required for gapped alignments)
- ID** (required for parent features)
- Name** (required for transcripts and SNPs)
- Parent** (required for child features)

In addition, Blixem supports the following custom tags.

- percentId** (only applicable to alignments; populates the %ID column)
- sequence** (only applicable to alignments; supplies the sequence data)
- variant\_sequence** (only applicable to variations; supplies the variation data)
- url** (only used by variations; GFF3 special characters must be escaped)

### Transcripts

Note that exons should have a Parent transcript defined, and the Name tag should be set in the parent rather than the child exons. Note that Blixem *will* recognise exons that do not have a Parent tag if they have a Name tag instead, but they may not get grouped correctly with other exons from the same transcript.

Typically, one defines the parent transcript, the exons, and the CDS regions; Blixem will then calculate the missing components (in this case, the UTR regions and the introns). Blixem will recognise other combinations of inputs, and will always calculate the missing components as long as enough information is provided.

### Variations

SNPs, insertions and deletions are supported, as well as combined variations. One may use the generic 'sequence\_alteration' type for these but it is good practice to use more specific types such as 'SNP' or 'deletion' where applicable.

### Sample GFF file

A sample GFF file may look like this ('...' denotes that text has been omitted).

```
##gff-version 3
##sequence-region chr4-04_210623-364887 44144 154265
chr4-04_210623-364887 EST_Human nucleotide_match 79195 79311 95.000000
- . Target=DA692754.1 287 403 +;percentID=90.6;sequence=GATCTGGC...
chr4-04_210623-364887 EST_Human nucleotide_match 79195 79323 121.000000
+ . Target=AI095103.1 326 454 +;percentID=96.9;sequence=TTTAAATT...
chr4-04_210623-364887 ensembl_variation deletion 80798 80799 . +
. Name=rs60725655;url=http%3A%2F%2Fwww.ensembl.org%2FHomo_sapiens
%2Fvariation%2Fsummary%3Fv%3Drs60725655;variant_sequence=AA/-;
chr4-04_210623-364887 ensembl_variation sequence_alteration 80799 80799 .
```

```

+ . Name=rs57681246;url=http%3A%2F%2Fwww.ensembl.org%2FHomo_sapiens
%2FVariation%2FSummary%3Fv%3Drs57681246;variant_sequence=A/-/C;
chr4-04_210623-364887 ensembl_variation SNP 81040 81040 . + .
Name=rs2352935;url=http%3A%2F%2Fwww.ensembl.org%2FHomo_sapiens%2FVariation
%2FSummary%3Fv%3Drs2352935;variant_sequence=T/C;
chr4-04_210623-364887 ensembl_variation insertion 82229 82230 . +
. Name=rs35105663;url=http%3A%2F%2Fwww.ensembl.org%2FHomo_sapiens
%2FVariation%2FSummary%3Fv%3Drs35105663;variant_sequence=-/G;
chr4-04_210623-364887 Augustus mRNA 119534 119941 . - .
ID=transcript21;Name=AUGUSTUS000000051712
chr4-04_210623-364887 Augustus exon 119534 119941 . - .
Parent=transcript21
chr4-04_210623-364887 Augustus CDS 119534 119941 . - 0
Parent=transcript21

```

### FASTA file:

A FASTA file has a header line that starts with '>' and contains the sequence name. The next line contains the start of the sequence data. The sequence data can be on a single line or separated by newlines; it is usually separated by newlines every 50 characters to aid readability.

```

>chr4-04_210623-364887
tcttgtttctgtaggagaggccatctccatcagctataaccccccccccc
accccccaactcctctttttgacaagtttgtaaagcctgtccatctgggtc
tataataatcctccaggccctatgccactcctctttattcagccagttca
...

```

### Combined GFF and FASTA file:

```

##gff-version 3
##sequence-region chr4-04_210623-364887 44144 154265
chr4-04_210623-364887 EST_Human nucleotide_match 79195 79311 95.000000
- . Target=DA692754.1 287 403 +;percentID=90.6
chr4-04_210623-364887 EST_Human nucleotide_match 79195 79323 121.000000
+ . Target=AI095103.1 326 454 +;percentID=96.9
...
##FASTA
>chr4-04_210623-364887
tcttgtttctgtaggagaggccatctccatcagctataaccccccccccc
accccccaactcctctttttgacaagtttgtaaagcctgtccatctgggtc
tataataatcctccaggccctatgccactcctctttattcagccagttca
...

```

### Configuration file

Note that if the sequence data for the match sequences is not supplied via the 'sequence' tag in the GFF file then Blixem will try to fetch the data from a server using a program called 'pfetch'. Currently this is only supported for internal users at the Sanger Institute. Details of the server are supplied via a .ini-style configuration file using the '-c' argument.

## The Blixem Window

The Blixem window consists of two main sections: an overview section called the “big picture”, and a detail section showing the actual sequence data. These sections are separated by a splitter bar, so you can maximise the space for the area you are interested in. You can also hide sections of the window using the ‘View’ menu.

Blixem can show sequences in nucleotide or protein mode.

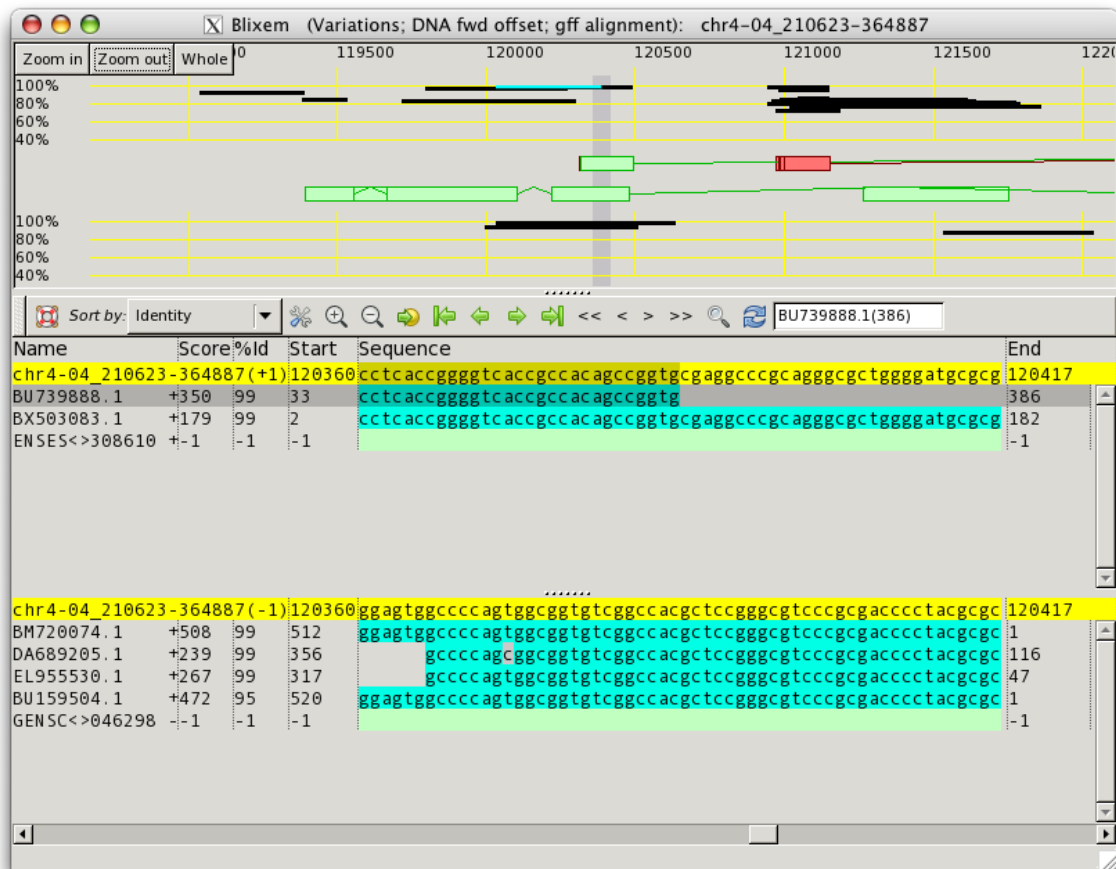


Figure 1: Nucleotide mode. There are two panes in the detail-view, one for each strand. The active strand is shown at the top. The active strand can be changed by hitting the 'Toggle' button or the 't' shortcut key.



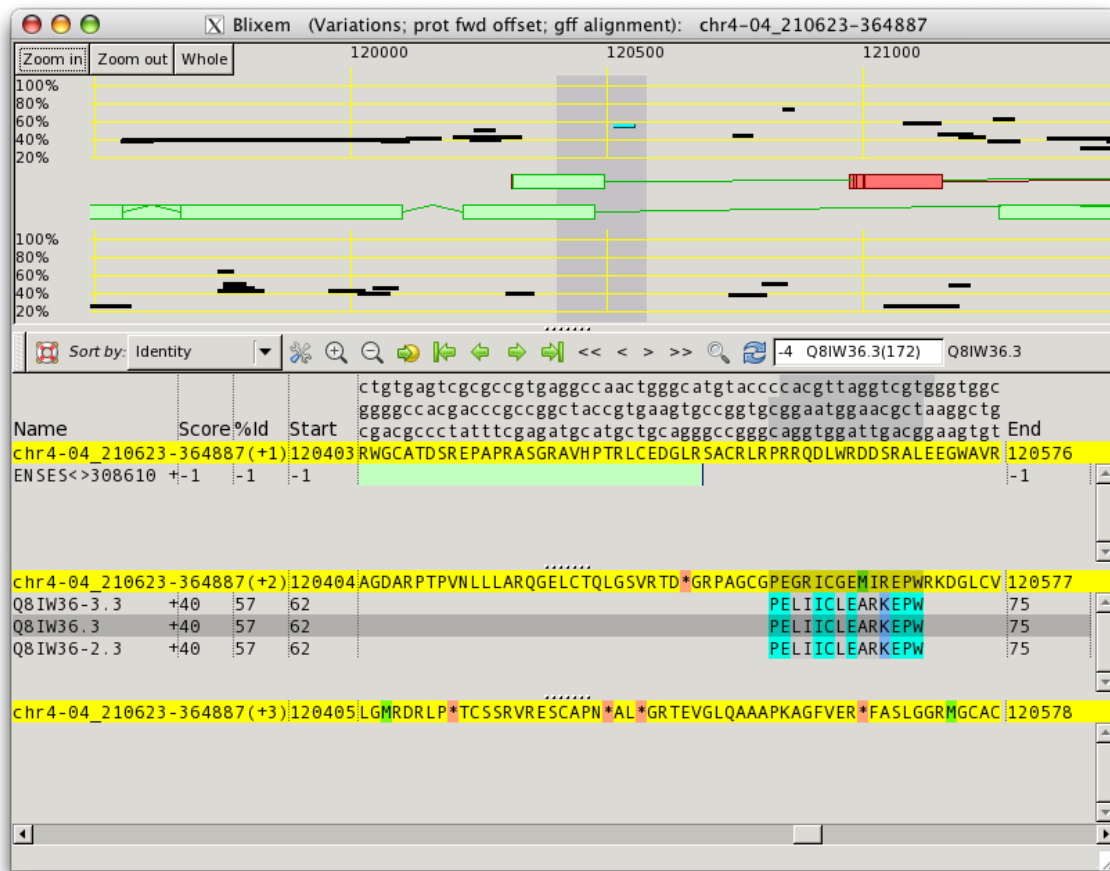



Figure 2: Protein mode. There are three panes in the detail-view; one for each reading frame of the active strand. The other strand can be activated by hitting the 'Toggle' button or the 't' shortcut key.

## Active Strand

The "active" reference sequence strand in Blixem controls the orientation of the display – coordinates are shown increasing from left-to-right for the forward strand and decreasing for the reverse strand. The active strand is always shown at the top – i.e. the top grid and top transcript view in the big picture and the top pane in the detail view.

In protein mode, only the active strand is shown in the detail view. One must toggle the strand to view the other strand.

Toggle which strand is active by:

- pressing the 'Toggle' button  on the toolbar; or
- pressing the 't' key.

By default, Blixem assumes that the reference sequence passed to it is the forward strand, unless otherwise specified by the '--reverse-strand' command line argument.

## Big Picture

The 'Big Picture' section shows an overview of the reference sequence. The reference sequence coordinates are shown along the top. You can zoom in to view a shorter range by using the 'Zoom in' button at the top left of the screen. Use 'Zoom out' or 'Whole' to zoom out – 'Whole' zooms out to view the full length of the reference sequence.

The big picture consists of two grids showing the alignments for each strand, and two sections between these grids showing the transcripts for each strand. The grids have a scale on the left-hand side showing the percent-ID, and alignments are plotted against this scale. The scale and extents of the grids can both be edited - see the Grid properties section in the Settings dialog.

The active strand alignments and transcripts are shown at the top and the other strand at the bottom. The direction of the coordinates is determined by the active strand. The active strand can be toggled using the 't' shortcut key or the 'Toggle strand' button on the toolbar.

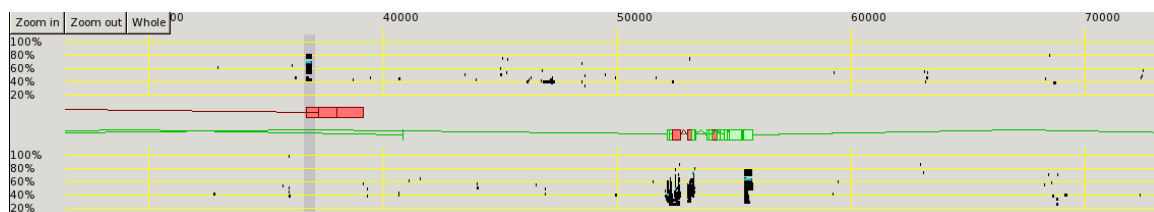


Figure 3: The Big Picture section

## Bumping the transcript view

By default, exons and introns for the same strand are drawn overlapping each other. They can be expanded (or 'bumped') by pressing the 'b' shortcut key or by enabling the relevant option in the View dialog (see Hiding sections of the window).

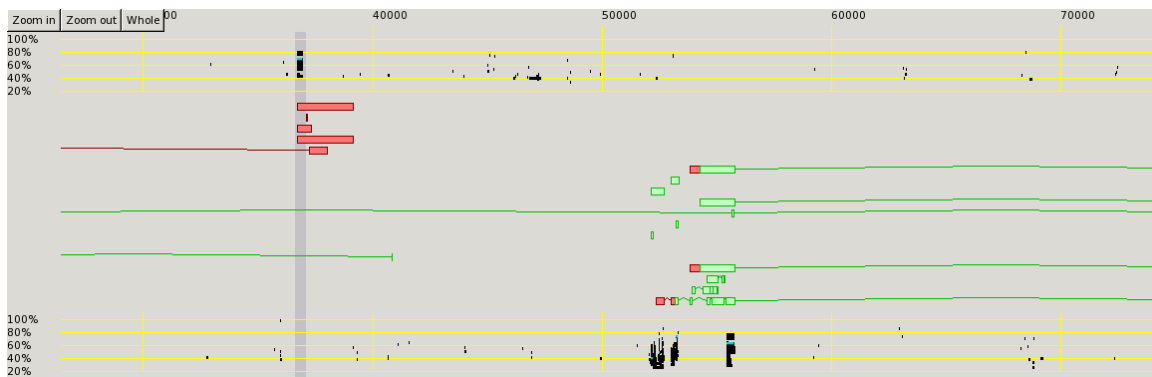


Figure 4: Expanded transcript view

## Detail View

The 'Detail View' shows the actual sequence data for the match sequences. Match sequences are lined up underneath the relevant section of reference sequence, and individual bases are highlighted in different colours to indicate how well they match.

## Match colours

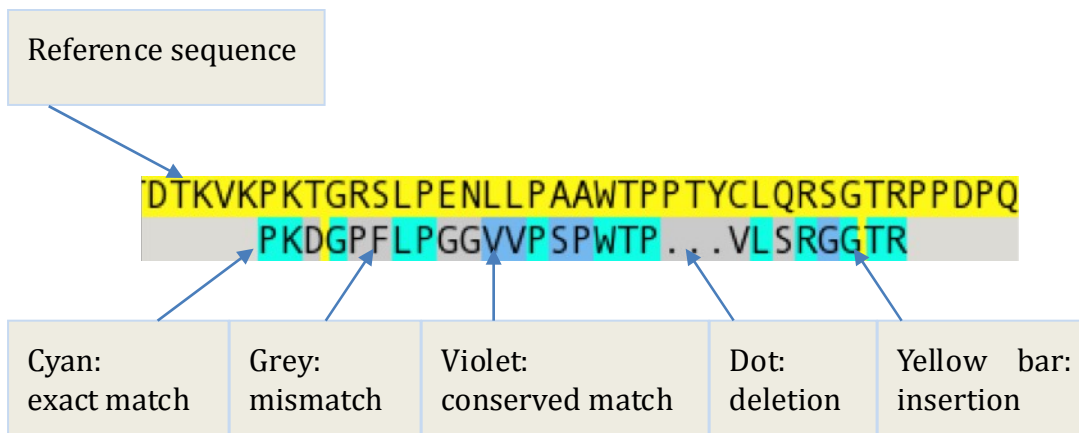


Figure 5: Alignment colour key

## Alignment lists

There are separate lists of alignments for each strand and reading frame of the reference sequence. Each list has a yellow header bar containing the reference sequence. At the left, the yellow bar shows the reference sequence name and which strand/frame it is, e.g. (+1) means forward strand, reading frame 1; (-2) means reverse strand, reading frame 2.

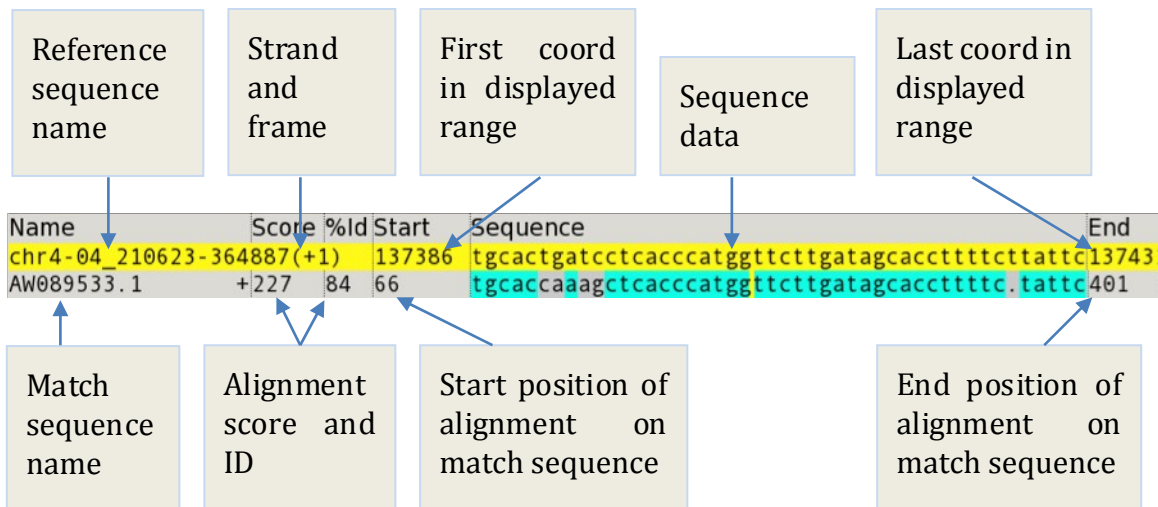


Figure 6: Alignment list details

### Nucleotide mode

There are two sections to the detail view in nucleotide mode: one for each strand. The active strand is shown at the top and defines the coordinate direction (increasing if the forward strand is active, decreasing if the reverse is active).

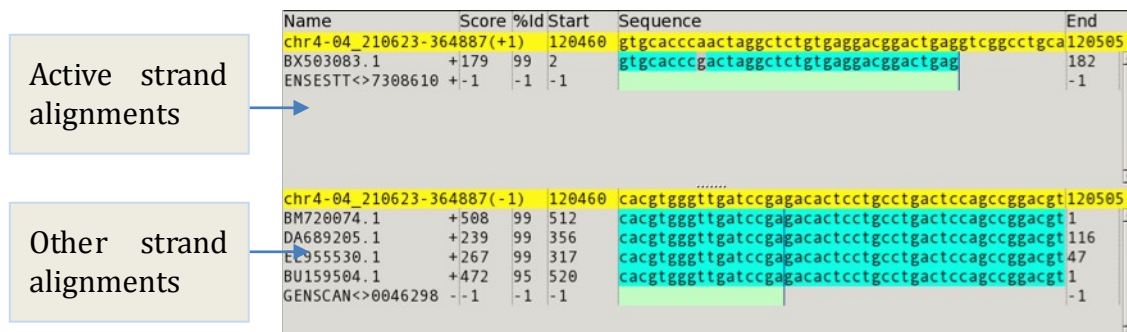


Figure 7: Alignment lists: nucleotide mode

### Protein mode

There are three sections in the detail view in protein mode: one for each of the three reading frames for the active strand. Only the active strand is shown. To view the other strand, toggle the display using the 'Toggle strand' button or the 't' shortcut key.

In protein mode, the yellow header bars show the translated reference sequence for that reading frame. STOP and MET codons in the reference sequence are highlighted in red and green. There is also an additional header section at the top showing the nucleotide sequence.

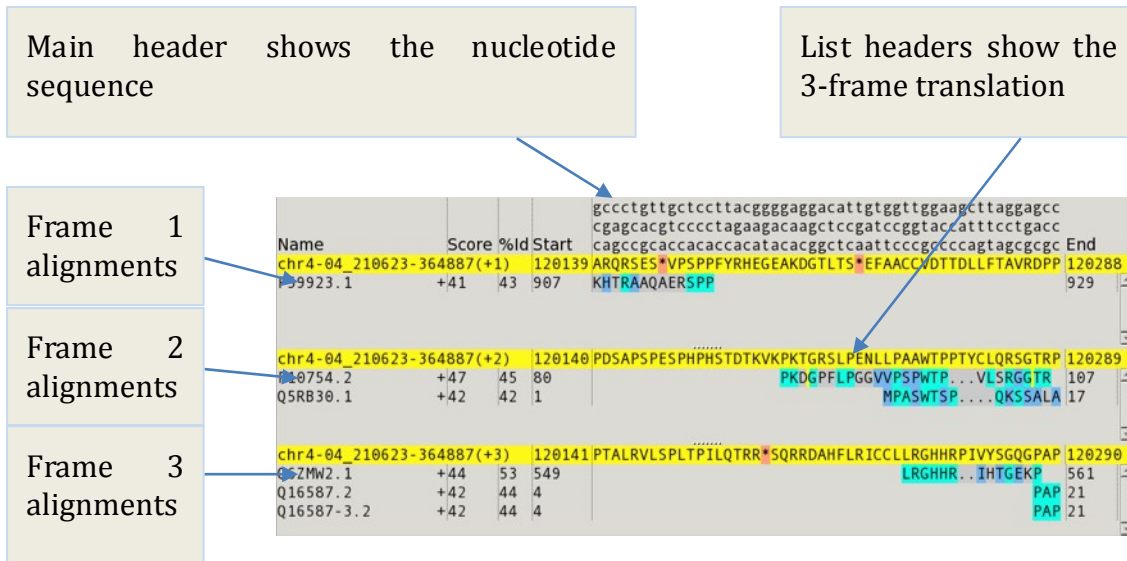


Figure 8: Alignment lists: protein mode

In the nucleotide-sequence header, codons are read from top-to-bottom and then left-to-right, starting at row 1 for frame 1, row 2 for frame 2 etc. Middle-clicking on a coordinate will highlight the three nucleotides for the selected codon and the currently-active reading frame (by default, frame 1). Left-clicking in an alignment list sets the active reading frame.

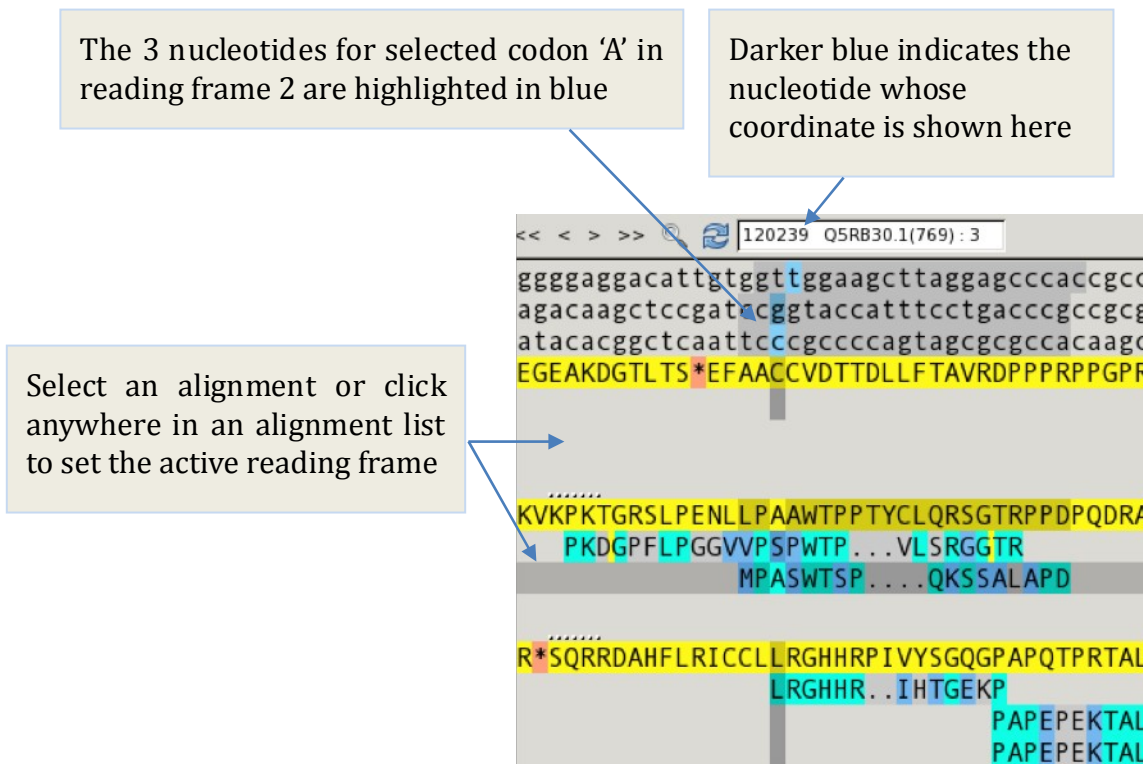















Figure 9: Selected reading frame and codon

## The toolbar

The detail-view toolbar contains the following functions. Note that the Help and Settings buttons are included in the detail-view toolbar even though they apply to Blixem as a whole.



Figure 10: Detail-view toolbar

	<b>Help:</b>	Show help about how to use Blixem
	<b>Sort-by:</b>	Select which column to sort the match sequences by
	<b>Settings:</b>	Show the Settings dialog.
	<b>Zoom in:</b>	Increase the font size in the detail-view
	<b>Zoom out:</b>	Decrease the font size in the detail-view
	<b>Go to:</b>	Go to a particular coordinate
	<b>First match:</b>	Go to the first coordinate of the first alignment <sup>1</sup>
	<b>Previous match:</b>	Go to the start of the current alignment or the end of the previous alignment <sup>1</sup>
	<b>Next match:</b>	Go to the end of the current alignment or the start of the next alignment <sup>1</sup>
	<b>Last match:</b>	Go to the end of the last alignment <sup>1</sup>
	<b>Back one page:</b>	Scroll the detail-view range to the left by one page
	<b>Back one index:</b>	Scroll the detail-view range to the left by one base
	<b>Forward one index:</b>	Scroll the detail-view range to the right by one base

<sup>1</sup> Acts only on selected sequences, if there is currently a selection; if no sequences are currently selected, then this operation acts on all sequences.



**Forward one page:**

Scroll the detail-view range to the right by one page



**Find:**

Scrolls to the start of the first alignment from that sequence if any are found.



**Toggle strand:**

Toggle which strand is the active strand

### Feedback box

The feedback box contains information about the currently selected sequence and/or coordinate, if either is selected. Click on a row in the detail-view to select a sequence. Middle-click on a base in the detail-view to select that coordinate. Text in the feedback box can be selected and copied.

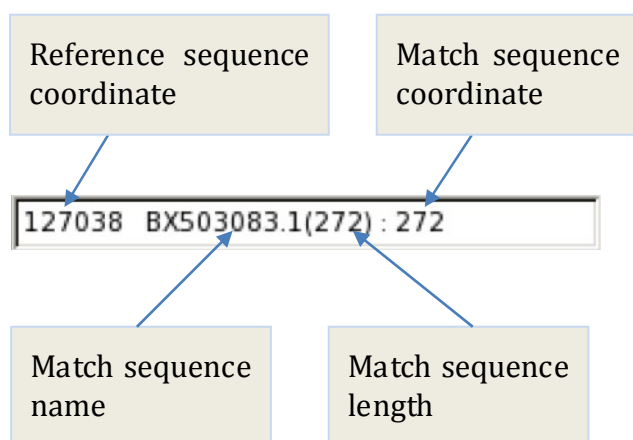


Figure 11: Feedback box

### Moused-over item feedback area

The area to the right of the toolbar contains information about the currently moused-over item (e.g. a match sequence in the alignment list or a variation in the variations track). For a match sequence, this information includes the sequence name and optional data such as organism and tissue type that can be parsed from EMBL files (currently only available to authorised users). To load optional data, see the Settings dialog. Note that the optional data may be incomplete due to the inconsistent information available from the EMBL files.

BX503083.1; Hs Homo sapiens (human); liver

Figure 12: Moused-over item feedback area

## The main menu

Right-click anywhere in the Blixem window to pop up the main menu.

<u>Q</u> uit	Ctrl-Q
<u>H</u> elp	Ctrl-H
<u>P</u> rint	Ctrl-P
<u>S</u> ettings	Ctrl-S
<hr/>	
<u>V</u> iew	v
Create Group	Shift-Ctrl-G
Edit <u>G</u> roups	Ctrl-G
Toggle <u>m</u> atch set group	g
Deselect <u>a</u> ll	Shift-Ctrl-A
<hr/>	
<u>D</u> otter	Ctrl-D

The options are:

<b>Quit</b>	<i>Ctrl-Q</i>	Close Blixem and any spawned processes
<b>Help</b>	<i>Ctrl-H</i>	Display the user help
<b>Print</b>	<i>Ctrl-P</i>	Printing options
<b>Settings</b>	<i>Ctrl-S</i>	Edit settings
<b>View</b>	<i>v</i>	Show/hide parts of the display
<b>Create Group</b>	<i>Shift-Ctrl-G</i>	Create a group of sequences
<b>Edit Groups</b>	<i>Ctrl-G</i>	Edit properties for groups
<b>Deselect all</b>	<i>Shift-Ctrl-A</i>	Deselect all sequences
<b>Dotter</b>	<i>Ctrl-D</i>	Run Dotter on the currently selected sequence

## Hiding sections of the window

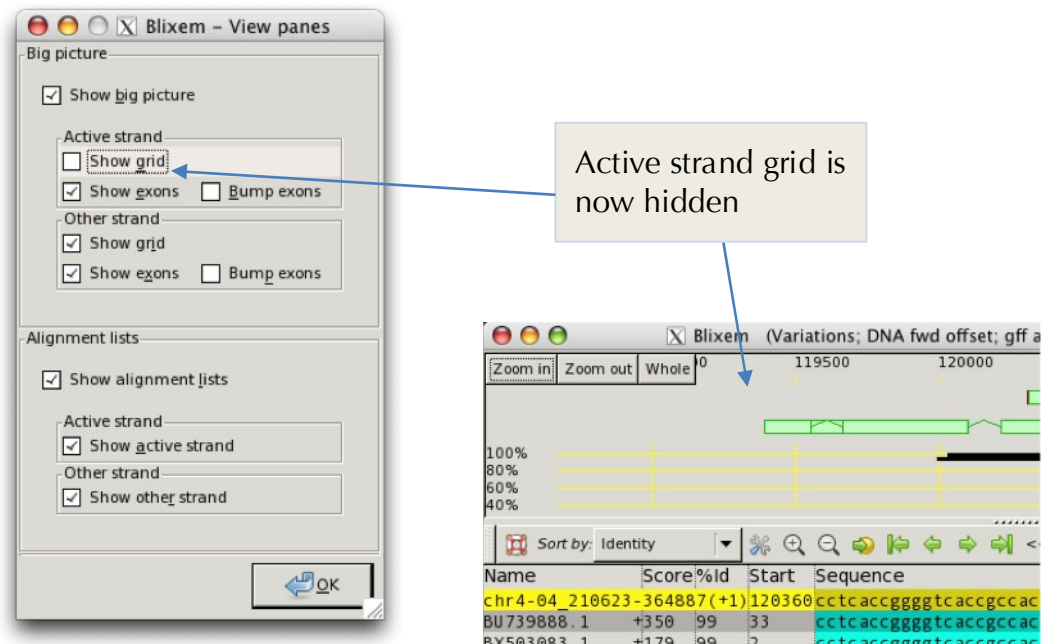
Use to 'View' dialog to show/hide sections of the window.

1. Right-click and select the View option, or hit the 'v' shortcut key.

<u>Q</u> uit	Ctrl-Q
<u>H</u> elp	Ctrl-H
<u>P</u> rint	Ctrl-P
<u>S</u> ettings	Ctrl-S
<hr/>	
<u>V</u> iew	v
Create Group	Shift-Ctrl-G
Edit <u>G</u> roups	Ctrl-G
Toggle <u>m</u> atch set group	g
Deselect <u>a</u> ll	Shift-Ctrl-A
<hr/>	
<u>D</u> otter	Ctrl-D

2. Toggle check marks on or off to show/hide sections.





Alternatively, use the following keyboard shortcuts to toggle visibility of a component:

- 1** Hide top pane in detail view
- 2** Hide second pane in detail view
- 3** Hide third pane in detail view (protein mode only)
- Ctrl-1** Hide top grid in big picture (active strand)
- Ctrl-2** Hide bottom grid in big picture (other strand)
- Shift-Ctrl-1** Hide top exon view (active strand)
- Shift-Ctrl-2** Hide bottom exon view (other strand)


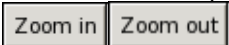
## Operation

### Navigation

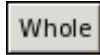
#### Scrolling

<b>Middle-click/drag in big picture</b>	Select a region to jump to.
<b>Middle-click/drag in detail view</b>	Select and centre on a base.
<b>Horizontal scrollbar</b>	Scroll the detail-view range.
<b>Vertical scrollbars</b>	Scroll up/down an alignment list.
<b>Horizontal mouse-wheel</b>	Scroll the detail-view range (if your mouse has a horizontal scroll-wheel).
<b>Vertical mouse-wheel</b>	Scroll up/down the currently moused-over alignment list
<b>Ctrl-left Ctrl-right</b>	Scroll to the start/end of the previous/next match (limited to currently-selected sequences, if any are selected; includes all sequences otherwise).
<b>Home End</b>	Scroll to the start/end of the display.
<b>Ctrl-Home Ctrl-End</b>	Scroll to the start/end of the currently-selected alignments (or to the first/last alignment if none are selected).
<b>',' (comma) '.' (full-stop)</b>	Scroll the detail-view range one nucleotide to the left/right.
<b>Ctrl-, Ctrl-.</b>	Scroll the detail-view range one page to the left/right.
<b>Go-to button or 'p' key</b>	Scroll to a specific coordinate position.

#### Zooming

<b>= - keys and</b> 	Zoom in/out of the detail-view
<b>Ctrl-= or Ctrl-- keys and</b> 	Zoom in/out of the big-picture

Shift-Ctrl-- and



Zoom the big picture out to view the full length of the reference sequence.

## Selections

### Selecting sequences

- You can select a sequence by clicking on its row in the alignment list. Selected sequences are highlighted in cyan in the big picture.
- You can select a sequence by clicking on it in the big picture.
- The name of the sequence you selected is displayed in the feedback box on the toolbar. If there are multiple alignments for the same sequence, all of them will be selected.
- You can select multiple sequences by holding down the Ctrl or Shift keys while selecting rows.
  - You can deselect a single sequence by Ctrl-clicking on its row.
- You can deselect all sequences by right-clicking and selecting 'Deselect all', or with the Shift-Ctrl-A keyboard shortcut.
- You can move the selection up/down a row using the up/down arrow keys.

### Selecting coordinates

- You can select a nucleotide/peptide by middle-clicking on it in the detail view. This selects the entire column at that index, and the coordinate number on the reference sequence is shown in the feedback box. (The coordinate on the match sequence is also shown if a match sequence is selected.)
- By default the display will centre on the selected base when you middle click. To select a base without scrolling, hold down Ctrl when you middle click.
- For protein matches, when a peptide is selected, the three nucleotides for that peptide (for the active reading frame) are highlighted in the header in blue. (The active reading frame is whichever alignment list currently has the focus - click in a different list to change the reading frame.) Darker blue highlighting indicates the specific nucleotide that is currently selected (i.e. whose coordinate is displayed in the feedback box).

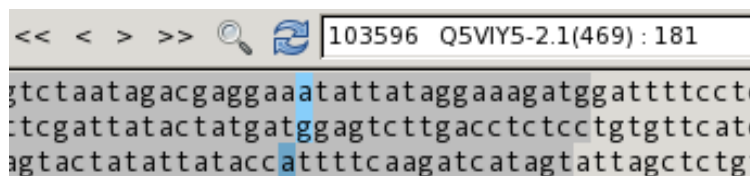



Figure 14: The 3 nucleotides for the currently-selected amino acid in reading-frame 3. Selected nucleotide 103596 is shaded in darker blue.

- You can move the selection to the previous/next index using the left and

- right arrow keys.
- In protein mode, you can move the selected nucleotide by a single base (rather than an entire codon) holding Shift while using the left and right arrow keys.
- You can move the selection to the start/end of the previous/next match by holding Ctrl while using the left and right arrow keys (limited to just the selected sequences if any are selected; includes all sequences otherwise).

### Finding sequences

The Find dialog allows the user to search for sequences by name. Press the Find  button on the toolbar or hit the 'Ctrl-F' shortcut key to open the Find dialog.

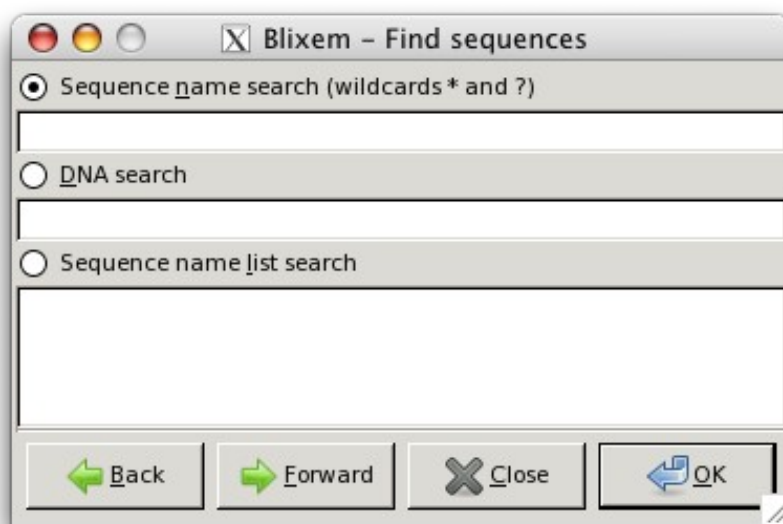


Figure 15: Find dialog

There are three search modes:

- Sequence name search: Search for match sequences by name. The wildcard '\*' means any number (or zero) of any character and '?' means 1 character (which can be any character). Any sequences whose names match the search string will be selected and the display will scroll to the start of the selection.
- DNA search: This searches for a given sub-sequence of nucleotides in the reference sequence. If the sub-sequence is found, the display will scroll to the start of the sub-sequence and the first base in the sub-sequence will be selected.
- Sequence name list search: the same as 'Sequence name search', but for multiple sequences. Each sequence names should be on a separate line.

Enter your search text in the appropriate box and click the OK button to perform the search. By default, Blixem will start the search at the beginning of the reference sequence range. To start the search from the current position, click the Forward or Back button instead of OK. This will start searching from the currently-selected base, if there is one selected; if not, it will start from the beginning of the current detail-view display range when searching forwards or from the end of the display range if searching backwards.

### **Repeat a Find**

After clicking OK on the Find dialog, press F3 to repeat the search in a forwards direction or Shift-F3 to repeat in a backwards direction. Alternatively, if you had selected the Forward or Back button in the Find dialog then click the Forward or Back buttons again to jump to the next result in that direction.

### **Copy and paste**

- When sequence(s) are selected, their names are copied to the *selection buffer* and can be pasted to another program by middle-clicking in that program.
- Sequence names can be pasted from the selection buffer into Blixem by hitting the 'f' keyboard shortcut. If the selection buffer contains valid sequence names, those sequences will be selected and the display will jump to the start of the selection.
- Sequence names can also be pasted from the selection buffer into text boxes in dialog boxes such as the Groups dialog or Find dialog.
- To copy sequence name(s) to the *default clipboard*, select the sequence(s) and hit Ctrl-C. Sequence names can then be pasted into other applications using Ctrl-V.
- The default clipboard can be pasted into Blixem using Ctrl-V. If the clipboard contains valid sequence names, those sequences will be selected and the display will jump to the start of the selection.
- Note that text from the feedback box and some text labels (e.g. the reference sequence start/end coords) can be copied to the selection buffer by selecting the required text with the mouse (or copied to the default clipboard by selecting it and then hitting 'Ctrl-C').
- Text can be pasted from the default clipboard into text entry boxes on dialogs such as the Groups or Find dialog by using Ctrl-V.

### **Sorting alignments**

- Alignments can be sorted by selecting the column you wish to sort by from the drop-down box on the toolbar.

Name
Organism
Gene name
Tissue type
Strain
Group
Score
Identity
Position

Figure 16: Sort-by list

- The default sort order may be ascending or descending depending on what makes most sense for the selected column: e.g. sorting by position is ascending by default but sorting by score or ID is descending.
- To get the inverse of the default sort order, select the 'Invert sort order' option in the Settings dialog.
- Alignments can also be sorted by group. Alignments that are part of a group will then be listed first (before any that are not in a group), and ordered according to the group's order number. See the Groups section for more details.

chr4-04_210623-364887(+3)			103527	VVYFPDCIALKHFKCKSCKCRQHKL	LFHINGRYHNCNVCS	FMVLSPFAYVNLVNPQELVISVF
Q9UC07.2	+Group1	44	43	220	YKCKFCGKAFHCLSLYLIERIH	
Q5VIY5-2.1	+Group1	51	30	163	ECISFKSFNCSSL	LKKHQI..IHLEEKQCKCDVC
Q86YE8-3.3	+Group2	50	70	381	MKHFECKECK	
Q9UII5.1	+	47	87	102	KHFKCKEC	
Q8TB69.1	+	47	87	430	KHFKCKEC	
Q8TF20.2	+	42	75	95	KHFKCNEC	
Q86YE8.3	+	50	70	419	MKHFECKECK	

Figure 17: Alignment list sorted by group

## Fetching sequences

Currently only available to authorised users at the Sanger Institute.

- Double-click a row to fetch a match sequence's EMBL file.

## Grouping sequences

Alignments can be grouped together so that they can be sorted/highlighted/hidden etc.

### Creating a group from a selection:

- Select the sequences you wish to include in the group by left-clicking their rows in the detail view. Multiple rows can be selected by holding the Ctrl or Shift keys while clicking.
- Right-click and select 'Create Group', or use the Shift-Ctrl-G shortcut key. (Note that Ctrl-G will also shortcut to here if no groups currently exist.)

- Ensure that the 'From selection' radio button is selected, and click 'OK' or 'Apply'. If you click 'Apply', you will be shown the group you just created so that you can edit it. If you click 'OK' the group will be created with the default properties.

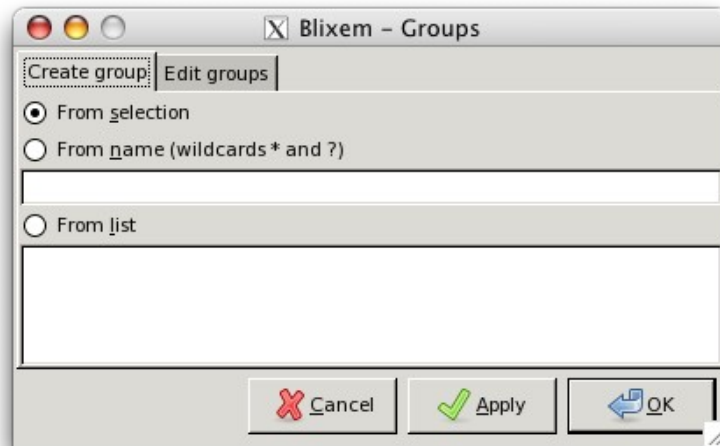


Figure 18: Groups dialog: create group

#### Creating a group from a sequence name:

- Right-click and select 'Create Group', or use the Shift-Ctrl-G shortcut key. (Or Ctrl-G if no groups currently exist.)
- Select the 'From name' radio button and enter the name of the sequence in the box below. You may use the following wildcards to search for sequences: '\*' for any number of characters; '?' for a single character.
- Click 'OK'.

#### Creating a group from sequence name(s):

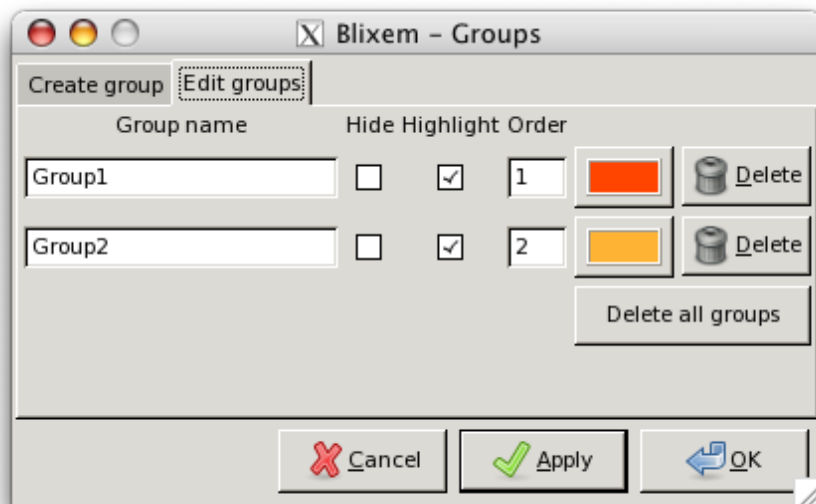
- Right-click and select 'Create Group', or use the Shift-Ctrl-G shortcut key. (Or Ctrl-G if no groups currently exist.)
- Select the 'From name(s)' radio button.
- Enter the sequence name(s) in the text box.
- You may use the following wild-cards in a sequence name: '\*' for any number of characters; '?' for a single character.
- You may search for multiple sequence names by separating them with the following delimiters: newline, comma or semi-colon.
- You may paste sequence names directly from another compatible program (e.g. ZMap): click on the feature in ZMap and then middle-click in the text box on the Groups dialog. (Grouping in Blixem works on the sequence name alone, so the feature coords output by ZMap will be ignored.)
- Click 'OK'.

### Creating a temporary 'match-set' group from the current selection:

- You can quickly create a group from a current selection (e.g. selected features in ZMap or just the current selection in Blixem) using the 'Toggle match set' option.
- To create a match-set group, select the required items and then select 'Toggle match set' from the right-click menu in Blixem, or hit the 'g' shortcut key.
- To clear the match-set group, choose the 'Toggle match set' option again, or hit the 'g' shortcut key again.
- While it is enabled (i.e. toggled on), the match-set group can be edited like any other group, via the 'Edit Groups' dialog. Any settings you change (e.g. highlight colour) will be saved even if the match-set group is toggled off and then on again.
- If you delete the match-set group using the 'Edit Groups' dialog, all of its settings will be lost; you will get the default settings again the next time you enable the match-set group. To avoid this, disable it by toggling it off using the 'Toggle match set' menu option (or 'g' shortcut key) rather than by deleting it in the Groups dialog.

## Editing groups:

To edit a group, right-click and select 'Edit Groups', or use the Ctrl-G shortcut key.



### Figure 19: Groups dialog - edit groups

You can change the following properties for a group. Click on Apply or OK to apply the changes.

**Name** You can specify a more meaningful name to help you identify the group.



<b>Hide</b>	Tick this box to hide the alignments in the alignment lists.
<b>Highlight</b>	Tick this box to highlight the alignments.
<b>Colour</b>	The colour the group will be highlighted in, if 'Highlight' is enabled. The default colour for all groups is orange, so you may wish to change this if you want different groups to be highlighted in different colours.
<b>Order</b>	When sorting by Group, alignments in a group with a lower order number will appear before those with a higher order number (or vice versa if sort order is inverted). Alignments in a group will appear before alignments that are not in a group.

To delete a group, click one of the following buttons. This will have an immediate effect (i.e. you don't have to click 'Apply').

- To delete a single group, click on the 'Delete' button next to the group you wish to delete.
- To delete all groups, click on the 'Delete all groups' button.

### Running dotter

- To start Dotter from within Blixem, or to edit the parameters for running Dotter, right-click and select 'Dotter' or use the Ctrl-D keyboard shortcut. The Dotter dialog will pop up.

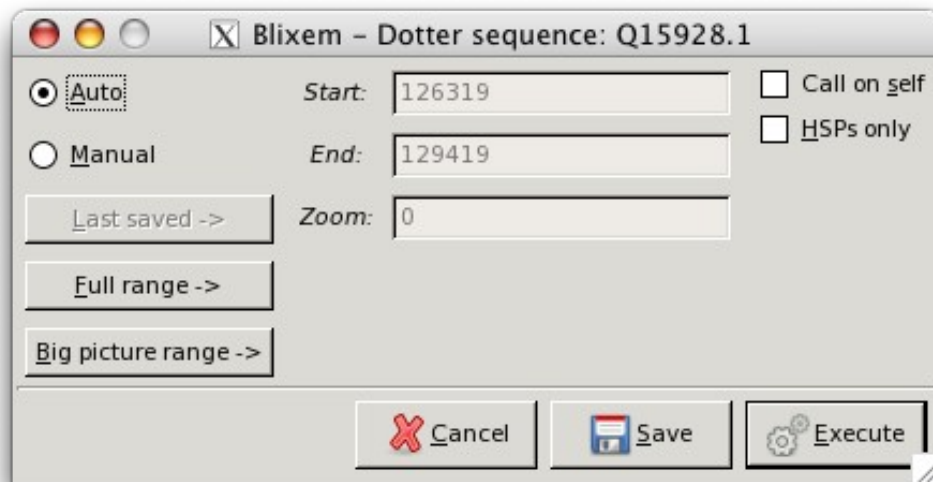


Figure 20: Dotter dialog

- Select the sequence you wish to run Dotter on before or after opening the dialog. The selected sequence name will be shown at the top of the dialog.
- Alternatively, if you just wish to edit the settings, you do not need to select a sequence.
- To run Dotter with the default (automatic) parameters, just hit RETURN, or click the 'Execute' button.
- To enter custom parameters, select the 'Manual' radio button and enter the

- values in the 'Start' and 'End' boxes.
- To save the parameters without running Dotter, click 'Save' and then 'Cancel'.
  - To save the parameters and run Dotter, click 'Execute'.
  - To revert to the last-saved manual parameters, click the 'Last saved' button.
  - To revert back to automatic parameters, click the 'Auto' radio button. The coordinates in the Start and End box will be recalculated for the currently-selected sequence.

#### **Reference sequence versus itself**

To run Dotter on the reference sequence versus itself, select the 'Call on self' tick box in the Dotter dialog and then click 'Execute'. This can be useful to analyse internal repeats etc. (see the Dotter manual for more information).

#### **Dotter HSPs only**

This starts Dotter in HSP (High-Scoring Pair) mode. See the Dotter manual for more information.

## Settings

The settings menu can be accessed by right-clicking and selecting Settings, or by the shortcut Ctrl-S.

## Features

### Highlight variations

When this option is enabled, bases in the reference sequence that have known variations (such as SNPs, insertions, deletions etc.) are highlighted in the reference sequence (nucleotide) header. If the 'Show variations track' sub-option is also enabled, then an additional line is shown above the nucleotide header showing the alternative bases for each variation. Note that the Variations track can be quickly enabled or disabled by double-clicking the nucleotide header.

### Show polyA tails

When this option is enabled, polyA tails are shown and highlighted in the alignment lists and polyA signals are highlighted in the reference sequence (nucleotide) header. If the sub-option 'Selected sequences only' is enabled, polyA features will only be shown for the currently selected sequences.

## Display options

### Show Unaligned Sequence

When this option is enabled, any additional, unaligned portions of the match sequences are displayed at the start and end of the alignments. If the 'Limit to' sub-option is also enabled, you can specify the maximum number of additional bases to display. If the 'Selected sequences only' sub-option is enabled, only the currently selected sequence(s) will display unaligned portions of sequence.

### Show Splice Sites

When this option is enabled, splice sites are highlighted in the reference sequence (nucleotide) header for the currently-selected sequence(s). The two bases from the adjacent introns are highlighted in green if they are canonical or red if they are non-canonical.

### Highlight Differences

When this option is enabled, matching bases are blanked out and mismatches are highlighted, making it easier to see where alignments differ from the reference sequence.

### Squash Matches

This groups multiple alignments from the same sequence together into the same

row in the detail view, rather than showing them on separate rows.

#### **Invert Sort Order:**

Reverse the default sort order. (Note that some columns sort ascending by default (e.g. name, start, end) and some sort descending (score and ID). This option reverses that sort order.)

### **General settings**

#### **Font**

Allows you to change the font that is used to display alignments in the detail-view. Note that you must select a monospace font; otherwise matches will not be shown aligned correctly. Blixem will warn you if the font you have selected is not monospace.

#### **Fetch mode**

Allows you to change the program used to fetch sequence EMBL entries. (Currently only available to authorised users within the Sanger Institute).

### **Columns**

#### **Load optional data**

Click this button to load optional data from EMBL entries (currently only applicable to authorised users within the Sanger Institute). Note that this operation can take a long time if there are many sequences. The button will be greyed out once optional data has been loaded.

#### **Column visibility**

Tick/un-tick the check-marks to show/hide individual columns. Adjust the column width by entering the new width in the text box in pixels. Note that if you enter a zero width then the column will be hidden, regardless of whether the check-mark is ticked or not. Greyed-out columns are optional-data columns, and will only become available once optional data has been loaded.

### **Grid properties**

#### **%ID per cell**

Use this to change the vertical scale of the grid; a smaller value means the grid will be more spaced out, a larger value means the grid will be more compact.

#### **Max %ID**

Defines the maximum cut-off value for the %ID scale.

#### **Min %ID**

Defines the minimum cut-off value for the %ID scale.

## Appearance

### Use print colours

Select this option to make Blixem use grey-scale colours, suitable for printing.

### Display colours

Change any of Blixem's custom display colours, such as the colour aligned bases are shown in or the colour stop codons are highlighted in etc. There are four colours for each item:

- Normal: this is the standard display colour;
- Normal (selected): this is the colour used when the item is selected (if applicable). Typically one would use a slightly darker or lighter shade of the Normal colour for this, so that the item does not look radically different when it is selected;
- Print: this is the standard colour used when the 'Use print colours' option is enabled;
- Print (selected): this is the colour used when 'Use print colours' is enabled and the item is selected.

## Key

In the detail view, the following colours and symbols have the following meanings:

Alignment list header	Yellow background	Reference sequence
Alignment list	Cyan background	Identical residues
Alignment list	Violet background	Conserved residues
Alignment list	Grey background	Mismatch
Alignment list	'.' with grey background	Deletion
Alignment list	Yellow vertical line	Insertion
Alignment list	Thin blue vertical line	Boundary of an exon
Nucleotide header (protein mode)	Sky-blue background	The three nucleotides for the currently-selected codon; darker blue indicates the nucleotide whose coordinate is displayed in the feedback box
Alignment list header (protein mode)	Pale red background	STOP codon
Alignment list header (protein mode)	Green background	MET codon

## Keyboard shortcuts

<b>Ctrl-Q</b>	Quit
<b>Ctrl-H</b>	Help
<b>Ctrl-P</b>	Print
<b>Ctrl-S</b>	Edit settings
<b>V</b>	Show/hide sections of the display
<b>Shift-Ctrl-G</b>	Create group
<b>Ctrl-G</b>	Edit groups (or create a group if none currently exist)
<b>Ctrl-A</b>	Select all sequences in the current list
<b>Shift-Ctrl-A</b>	Deselect all sequences
<b>Ctrl-D</b>	Dotter
<b>Left-arrow</b>	Move coordinate section one index to the left <sup>2</sup>
<b>Right-arrow</b>	Move coordinate section one index to the right <sup>2</sup>
<b>Shift-Left</b>	Same as Left, but in protein mode it scrolls by a single nucleotide
<b>Shift-Right</b>	Same as Right, but in protein mode it scrolls by a single nucleotide
<b>Ctrl-Left</b>	Scroll to the start/end of the previous alignment <sup>3</sup>
<b>Ctrl-Right</b>	Scroll to the start/end of the next alignment <sup>3</sup>
<b>Up-arrow</b>	Move row selection up
<b>Down-arrow</b>	Move row selection down
<b>Home</b>	Scroll to the start of the display
<b>End</b>	Scroll to the end of the display
<b>Ctrl-Home</b>	Scroll to the start of the first alignment <sup>3</sup>
<b>Ctrl-End</b>	Scroll to the end of the last alignment <sup>3</sup>
<b>=</b>	Zoom in detail view
<b>-</b>	Zoom out detail view
<b>Ctrl-=</b>	Zoom in big picture
<b>Ctrl--</b>	Zoom out big picture
<b>Shift-Ctrl--</b>	Zoom out big picture to view the whole reference sequence
<b>,</b>	Scroll left one coordinate
<b>.</b>	Scroll right one coordinate
<b>P</b>	Go to position
<b>T</b>	Toggle the active strand
<b>G</b>	Toggle the 'match set' Group
<b>1</b>	Toggles visibility of the 1 <sup>st</sup> alignment list
<b>2</b>	Toggles visibility of the 2 <sup>nd</sup> alignment list
<b>3</b>	Toggles visibility of the 3 <sup>rd</sup> alignment list (protein mode only)
<b>Ctrl-1</b>	Toggles visibility of the 1 <sup>st</sup> big picture grid
<b>Ctrl-2</b>	Toggles visibility of the 2 <sup>nd</sup> big picture grid
<b>Shift-Ctrl-1</b>	Toggles visibility of the 1 <sup>st</sup> exon view
<b>Shift-Ctrl-2</b>	Toggles visibility of the 2 <sup>nd</sup> exon view

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<sup>2</sup> Only applicable if a coordinate is currently selected; middle-click a coordinate to select it.

<sup>3</sup> Limited to just the selected sequences, if any are selected; otherwise, acts on all sequences.

