

Basic multivariate analysis

A useful analysis pathway (including the Example Analysis Pathway done above, with its five steps), can be accomplished in one fell swoop using the **Basic multivariate analysis** wizard. This will perform a suite of multivariate analyses commonly performed for either biotic or environmental data types, with options available that match the typical choices made for handling these different types of data.

Run the 'Basic multivariate analysis' wizard

Let's suppose you wanted to repeat the step-by-step analyses we did before, but this time using a different pre-treatment option. For example, you may decide you'd like to analyse the same data using presence/absence information only, so as to emphasise only the turnover in species identities (and not differences in abundance values) across sample units. It would be great to run this whole set of analyses quickly, rather than going through them again one at a time.

Click on the original 'Fal nematode abundance' datasheet in the Explorer tree and then click **Wizards > Basic multivariate analysis....**

PRIMER 7

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Basic multivariate analysis...
Matrix display...
Coherence plots...

Fal_Workspace
Fal nematode abundance
Overall Transform1
4th-root
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BC_4th-root
CLUSTER1
Graph1
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Graph2
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Graph4
Graph5
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Graph6

Fal nematode abundance
Fal estuary nematodes
Abundance

	Samples						
	R1	R2	R3	R4	R5	R6	R7
Anoplostoma vivip	0	0	0	0	0	0	0
Halalaimus gradis	0	0	0	0	0	0	0
Halalaimus longica	0	0	0	0	0	0	0
Oxystomina elonga	0	0	0	0	0	0	0
Viscosia viscosa	0	0	0	0	0	0	0
Tripyloides gradis	149	181	385	289	170	614	
Atrochromadora mi	0	0	0	0	0	0	0
Chromadora macro	0	4	29	63	263	123	
Chromadora nudica	0	0	0	0	0	0	7
Chromadorella ?du	0	0	0	0	0	0	0
Chromadorita nana	0	0	0	0	0	0	0
Chromadorita tenta	0	0	0	0	0	0	0
Dichromadora geop	5	0	0	0	0	0	7
Hypodontolaimus b	40	44	25	18	5	14	
Ptycholaimellus po	174	424	178	99	107	123	
Neochromadora po	0	0	0	0	0	0	0

Row 1 Col 1

In the dialog box that follows, we can see that PRIMER is offering to perform a suite of basic multivariate analyses that are commonly performed for 'Biotic Data' (shown in bold blue font at the

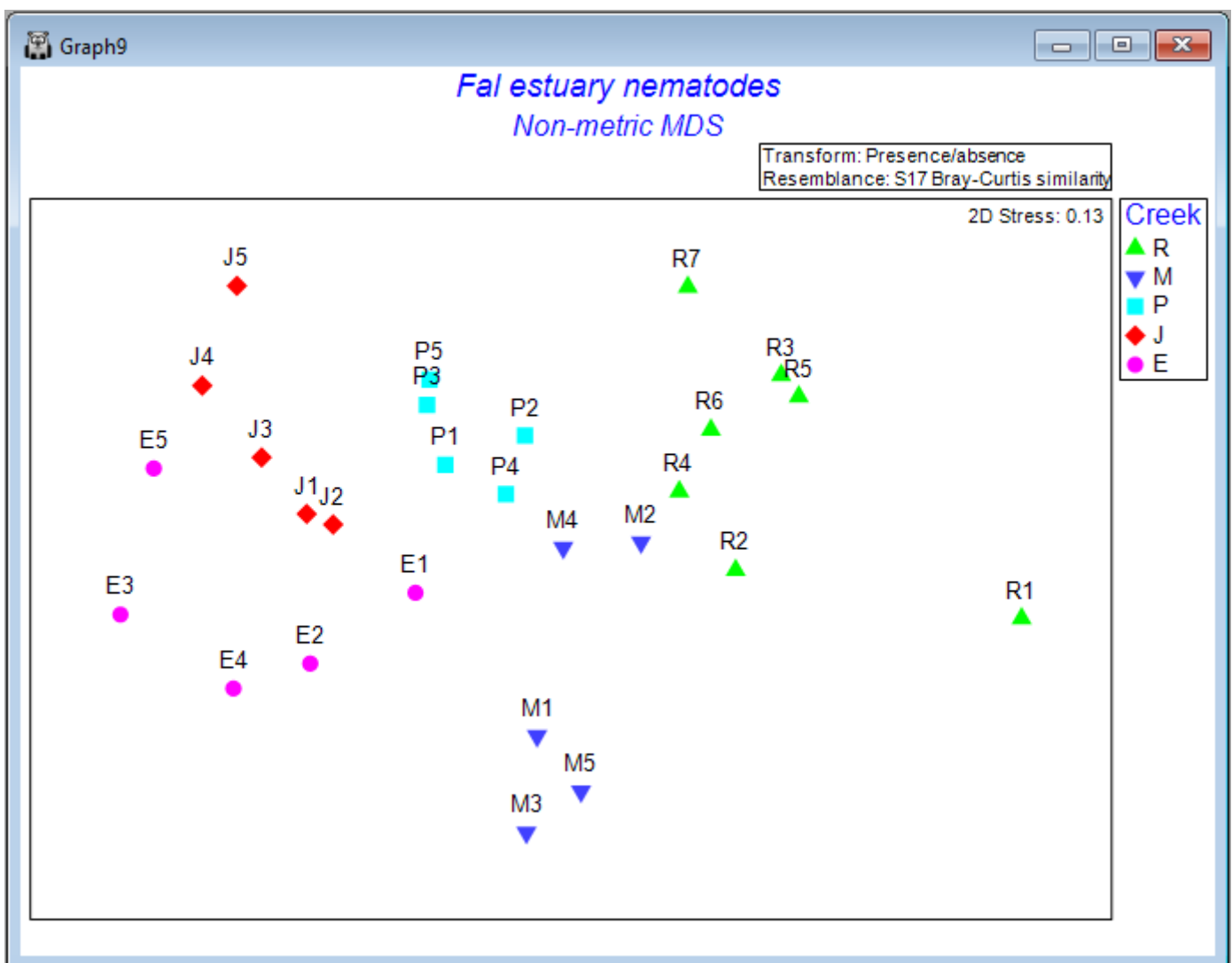
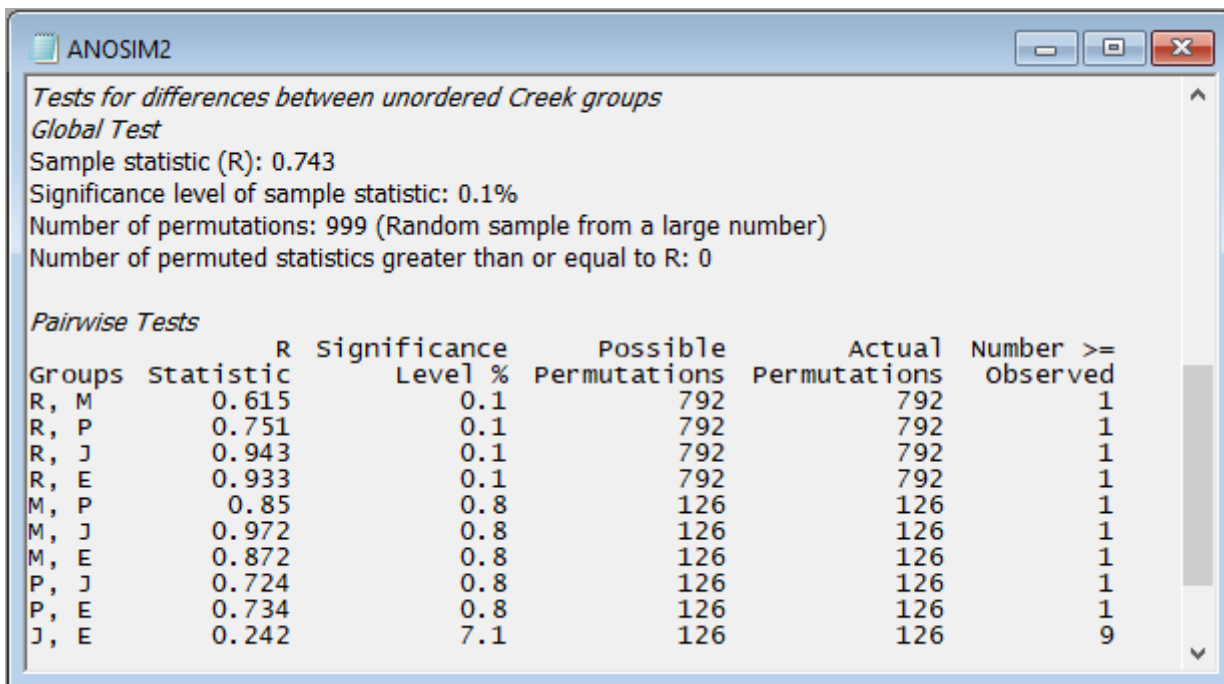
top). You can choose from a couple of (common) pre-treatment options, and then it will perform the analyses you choose (*via* the relevant checkboxes under 'Analyse'), using the default options for each routine. Note that different options would be shown for data of a different type (e.g., for environmental data). Recall that the 'type' of data is specified by you when you import the data into PRIMER. This can be changed for a given data sheet by clicking on **Edit > Properties** at any time.

Under 'Pre-treatment', choose 'Transformation: **Presence/absence**' and under 'Analyse', leave all of the default options, except you can *untick* the checkbox next to the 'SIMPER' routine (for now), then click **Finish**.

The screenshot shows the 'Basic analysis wizard' dialog box for 'Biotic Data'. It is divided into two main sections: 'Pre-treatment' and 'Analyse'. In the 'Pre-treatment' section, the 'Standardise samples' checkbox is unchecked, and the 'Transformation' dropdown menu is set to 'Presence/absence'. In the 'Analyse' section, the 'Resemblance' dropdown is set to 'S17 Bray-Curtis similarity'. Under the 'ANOSIM (1-way)' section, the 'Factor' dropdown is set to 'Creek'. The 'CLUSTER' checkbox is checked, and the 'SIMPROF' sub-option is also checked. The 'MDS' checkbox is checked, and the 'SIMPER' checkbox is unchecked. At the bottom of the dialog, there are five buttons: 'Cancel', '< Previous', 'Next >', 'Finish' (which is highlighted with a mouse cursor), and 'Help'.

All of the requested analyses are done, and the resulting output files and graphics are shown in the Explorer tree window. Click on any of the items in the Explorer tree to see the steps that were taken, including specific data sheets, resemblance matrices, graphical outputs and results files. Although you will generally use PRIMER to perform individual analyses, one routine at a time, this wizard provides a quick way to achieve multiple analyses (provided you know *a priori* that you want to do them) with a single stroke.

The ANOSIM results ('**ANOSIM2**') and the 2D nMDS output (**Graph9**) obtained by this run of the wizard are shown below:



In this example, we can see that there are statistically significant differences in the identities of nematode species among the five creeks (ANOSIM $R = 0.743$, $P = 0.001$ with 999 permutations), although the pattern in the nMDS plot suggests that the difference between Restronguet Creek and

the other creeks is not as great as what was observed when abundance information was included in the Bray-Curtis calculation. (Compare 'Graph9', as shown above, with 'Graph2' that we obtained earlier.)

Expanding and collapsing the Explorer tree

Note that this new set of analyses, performed using the wizard on the original imported data, will initiate its own new branch of the Explorer tree. You can always initiate a new analysis starting from a given item in the tree (e.g., from a data sheet or a resemblance matrix, etc.), and the tree will expand, generating a new branch, to accommodate these new analyses. You can use the '+' and '-' symbols in the Explorer tree to 'roll up' or 'unpack' the items in the tree belonging to a particular branch at any time.

For example, clicking on the '-' symbol next to the item named '4th-root' will 'roll up' all of the items associated with our original analysis (based on a fourth-root transformation), so the analyses that were done by the Wizard (based on presence/absence) are now closer to the top of the Explorer tree window. For clarity, we might choose to rename the sheet called 'Data1' (the data sheet produced by the wizard after performing the presence/absence transformation) to 'Pres_abs'.

The screenshot shows the PRIMER 7 software interface. On the left is the Explorer tree, which shows a hierarchy of analyses. The '4th-root' item is expanded, showing a sub-tree with 'Overall Transform1', 'Overall Transform2', 'Pres_abs', 'Resemblance2', 'Resem1', 'ANOSIM2', 'Graph7', 'CLUSTER2', 'Graph8', 'nMDS2', 'MultiPlot2', 'Graph9', and 'Graph10'. The 'Pres_abs' item is selected. On the right is a data table titled 'Fal estuary nematodes Abundance'. The table has columns for 'Samples' (R1, R2, R3, R4, R5, R6, R7, M1) and rows for various nematode species. The data is as follows:

	R1	R2	R3	R4	R5	R6	R7	M1
Anoplostoma vivip	0	0	0	0	0	0	0	0
Halalaimus gradis	0	0	0	0	0	0	0	0
Halalaimus longica	0	0	0	0	0	0	0	0
Oxystomina elonga	0	0	0	0	0	0	0	1
Viscosia viscosa	0	0	0	0	0	0	0	0
Tripyloides gradis	1	1	1	1	1	1	1	1
Atrochromadora mi	0	0	0	0	0	0	0	0
Chromadora macro	0	1	1	1	1	1	1	1
Chromadora nudica	0	0	0	0	0	0	1	1
Chromadorella ?du	0	0	0	0	0	0	0	0
Chromadorita nana	0	0	0	0	0	0	0	0
Chromadorita tenta	0	0	0	0	0	0	0	0
Dichromadora geop	1	0	0	0	0	0	1	0
Hypodontolaimus b	1	1	1	1	1	1	1	1
Ptycholaimellus po	1	1	1	1	1	1	1	1
Neochromadora po	0	0	0	0	0	0	0	0
Xinema sp.	0	0	0	0	0	0	0	0
Paracomesoma dub	0	0	0	0	0	0	0	0
Sabatieria breviseta	0	0	0	0	0	0	0	0
Sabatieria celtica	0	0	0	0	0	0	0	0
Sabatieria praedati	0	0	0	0	0	0	0	0
Sabatieria pulchra	1	1	1	1	1	1	1	1
Comesa ?cuanensis	0	0	0	0	0	1	0	0
Paracanthorchus he	0	0	0	0	0	1	1	1
Preacanthorchus pu	0	0	0	0	0	0	0	0