# The $E.\ coli$ molecular phenotype under different growth conditions

#### Supplementary material

Mehmet U. Caglar, John R. Houser, Craig S. Barnhart,
Daniel R. Boutz, Sean M. Carroll, Aurko Dasgupta, Walter F. Lenoir,
Bartram L. Smith, Viswanadham Sridhara, Dariya K. Sydykova,
Drew Vander Wood, Christopher J. Marx,
Edward M. Marcotte\*, Jeffrey E. Barrick\*, Claus O. Wilke\*

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	structural constituent of ribosome     structural molecule activity		lowMg
			highMg
		structural constituent of ribosome     structural molecule activity	highNa
	structural constituent of ribosome     structural molecule activity		glycerol
			gluconate
	structural constituent of ribosome     structural molecule activity		lactate
В	mRNA	Protein	lowMg
В	mRNA	Protein	lowMg highMg
В	mRNA	Protein  1. structural constituent of ribosome 2. structural molecule activity	
В	mRNA	structural constituent of ribosome	highMg
В	mRNA	structural constituent of ribosome	highMg highNa

Figure S1: Significantly differentially expressed Molecular Functions generated by GO annotations. For each condition, we show the top-5 differentially expressed MF as determined by either mRNA or protein abundances. (A) exponential phase. (B) stationary phase.

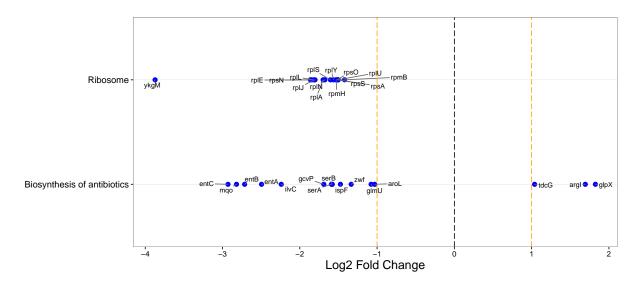


Figure S2: Significantly differentially expressed KEGG pathways and associated genes with glycerol as carbon source in exponential phase, as determined by mRNA abundances. The top differentially expressed KEGG pathways are shown along the y axis, and the relative fold change of the corresponding genes is shown along the x axis. In figure we show up to 10 most significantly changed pathways and for each pathway we show up to 15 of the most significantly changing genes.

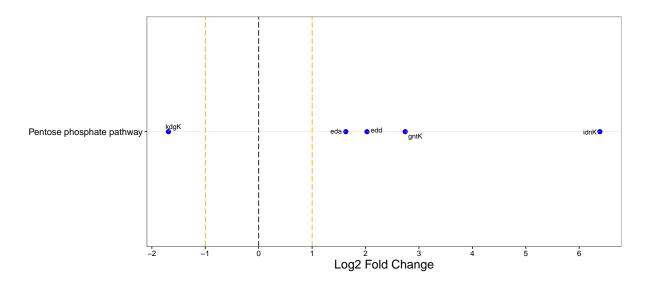


Figure S3: Significantly differentially expressed KEGG pathways and associated genes with gluconate as carbon source in exponential phase, as determined by mRNA abundances. The top differentially expressed KEGG pathways are shown along the y axis, and the relative fold change of the corresponding genes is shown along the x axis. In figure we show up to 10 most significantly changed pathways and for each pathway we show up to 15 of the most significantly changing genes.

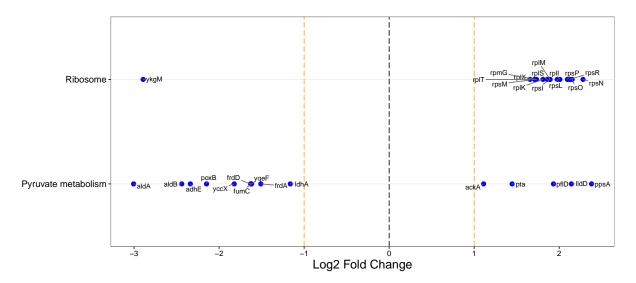


Figure S4: Significantly differentially expressed KEGG pathways and associated genes with lactate as carbon source in exponential phase, as determined by mRNA abundances. The top differentially expressed KEGG pathways are shown along the y axis, and the relative fold change of the corresponding genes is shown along the x axis. In figure we show up to 10 most significantly changed pathways and for each pathway we show up to 15 of the most significantly changing genes.

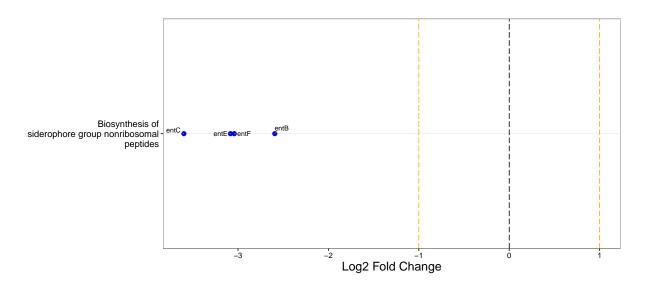


Figure S5: Significantly differentially expressed KEGG pathways and associated genes with gluconate as carbon source in exponential phase, as determined by protein abundances. The top differentially expressed KEGG pathways are shown along the y axis, and the relative fold change of the corresponding genes is shown along the x axis. In figure we show up to 10 most significantly changed pathways and for each pathway we show up to 15 of the most significantly changing genes.

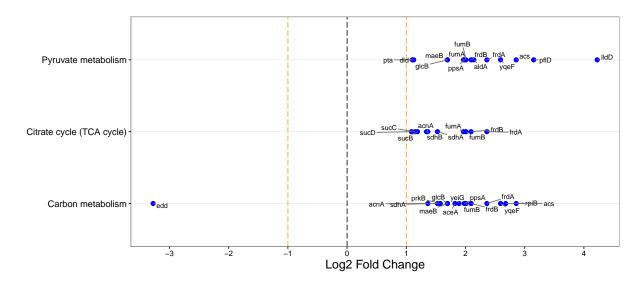


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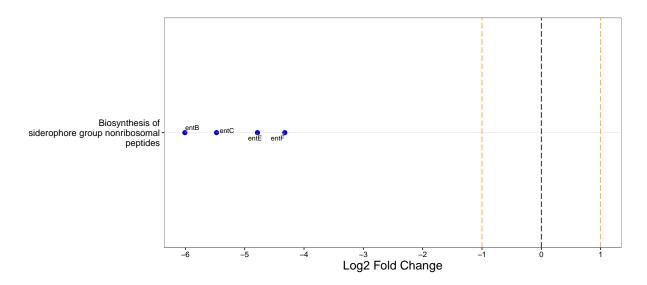


Figure S7: Significantly differentially expressed KEGG pathways and associated genes with glycerol as carbon source in stationary phase, as determined by protein abundances. The top differentially expressed KEGG pathways are shown along the y axis, and the relative fold change of the corresponding genes is shown along the x axis. In figure we show up to 10 most significantly changed pathways and for each pathway, we show up to 15 of the most significantly changing genes.

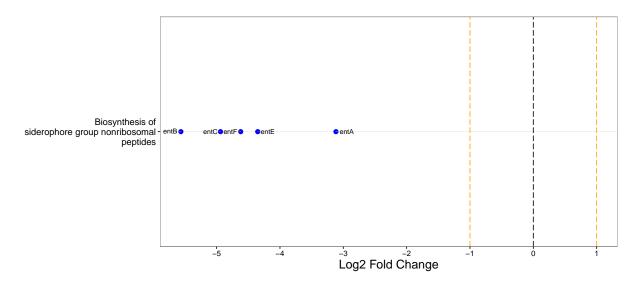


Figure S8: Significantly differentially expressed KEGG pathways and associated genes with gluconate as carbon source in stationary phase, as determined by protein abundances. The top differentially expressed KEGG pathways are shown along the y axis, and the relative fold change of the corresponding genes is shown along the x axis. We show up to 10 most significantly changed pathways and for each pathway, we show up to 15 of the most significantly changing genes.

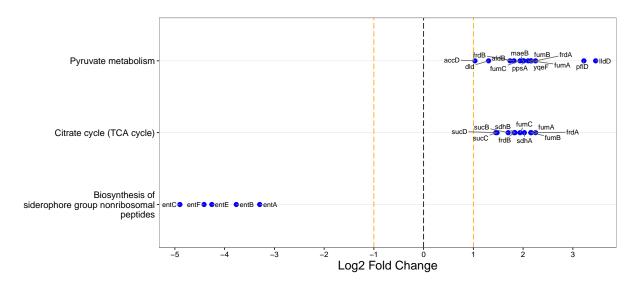


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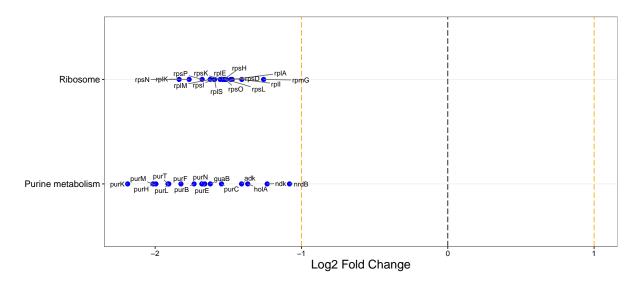


Figure S10: Significantly differentially expressed KEGG pathways and associated genes with low  $\mathrm{Mg^{+2}}$  levels in exponential phase, as determined by mRNA abundances. The top differentially expressed KEGG pathways are shown along the y axis, and the relative fold change of the corresponding genes is shown along the x axis. We show up to 10 most significantly changed pathways and for each pathway, we show up to 15 of the most significantly changing genes.

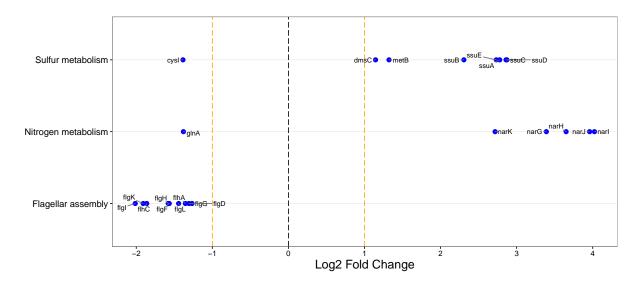


Figure S11: Significantly differentially expressed KEGG pathways and associated genes with high  $\mathrm{Mg^{+2}}$  levels in exponential phase, as determined by mRNA abundances. The top differentially expressed KEGG pathways are shown along the y axis, and the relative fold change of the corresponding genes is shown along the x axis. We show up to 10 most significantly changed pathways and for each pathway, we show up to 15 of the most significantly changing genes.

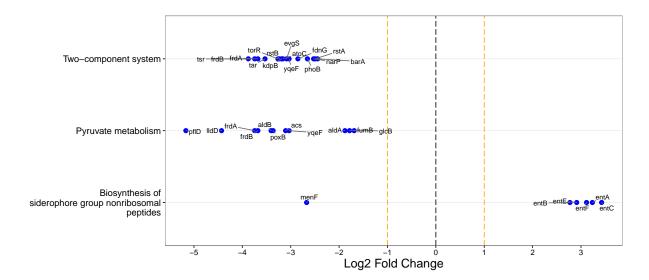


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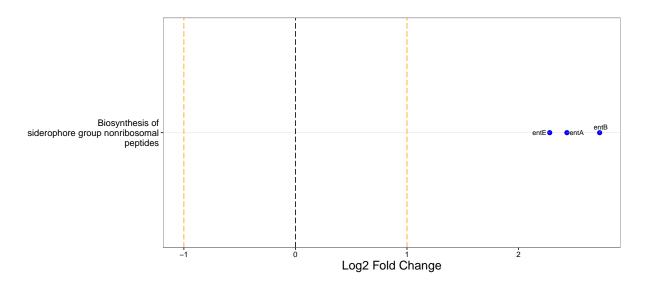


Figure S13: Significantly differentially expressed KEGG pathways and associated genes with high  $\mathrm{Mg^{+2}}$  levels in stationary phase, as determined by mRNA abundances. The top differentially expressed KEGG pathways are shown along the y axis, and the relative fold change of the corresponding genes is shown along the x axis. We show up to 10 most significantly changed pathways and for each pathway, we show up to 15 of the most significantly changing genes.

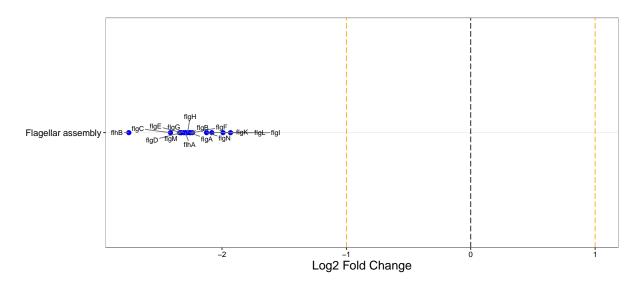


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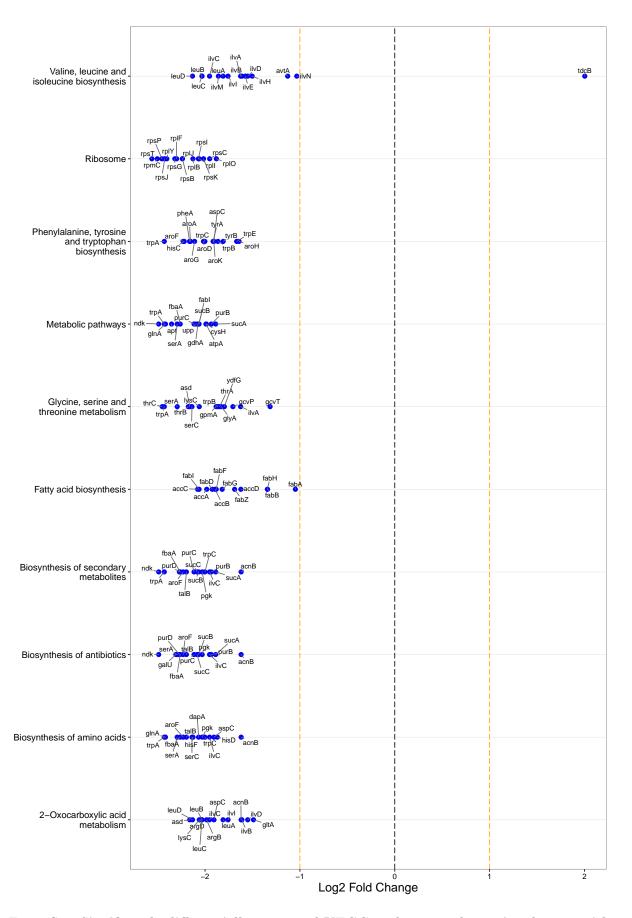


Figure S15: Significantly differentially expressed KEGG pathways and associated genes with high Na<sup>+1</sup> levels in exponential phase, as determined by protein abundances. The top differentially expressed KEGG pathways are shown along the y axis, and the relative fold change of the corresponding genes is shown along the x axis. We show up to 10 most significantly changed pathways and for each pathway, we show up to 15 of the most significantly changing genes.

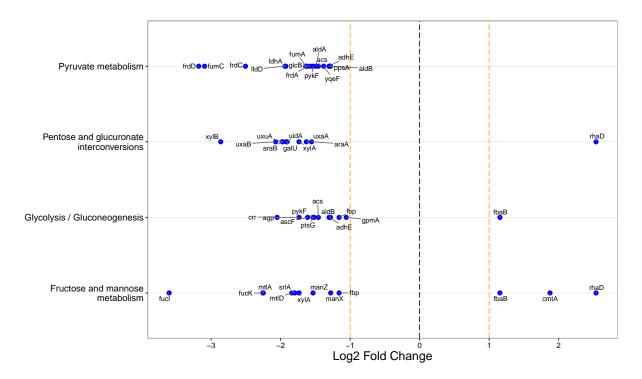


Figure S16: Significantly differentially expressed KEGG pathways and associated genes with high  $\mathrm{Na^{+1}}$  levels in stationary phase, as determined by mRNA abundances. The top differentially expressed KEGG pathways are shown along the y axis, and the relative fold change of the corresponding genes is shown along the x axis. We show up to 10 most significantly changed pathways and for each pathway, we show up to 15 of the most significantly changing genes.

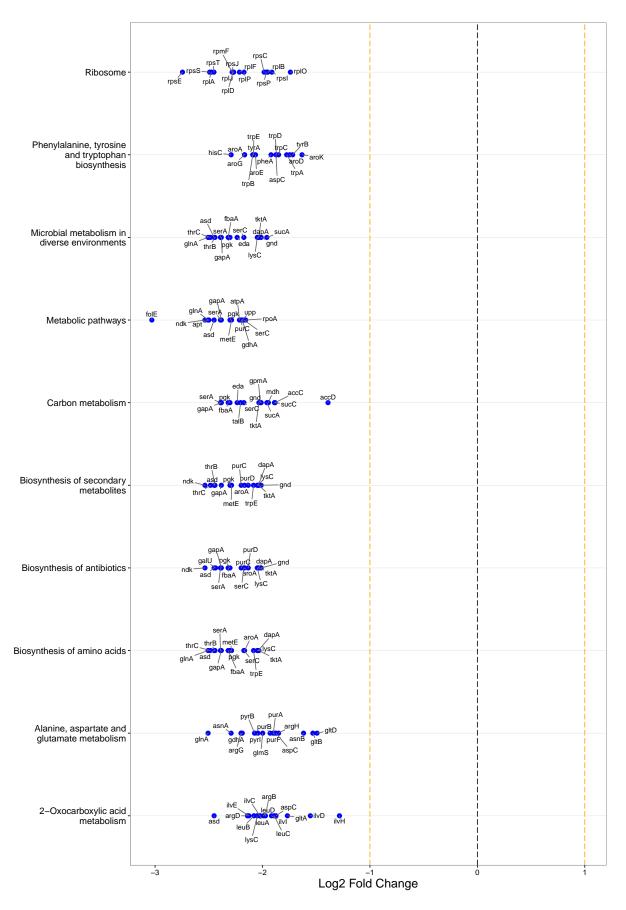


Figure S17: Significantly differentially expressed KEGG pathways and associated genes with high Na<sup>+1</sup> levels in stationary phase, as determined by protein abundances. The top differentially expressed KEGG pathways are shown along the y axis, and the relative fold change of the corresponding genes is shown along the x axis. We show up to 10 most significantly changed pathways and for each pathway, we show up to 15 of the most significantly changing genes.

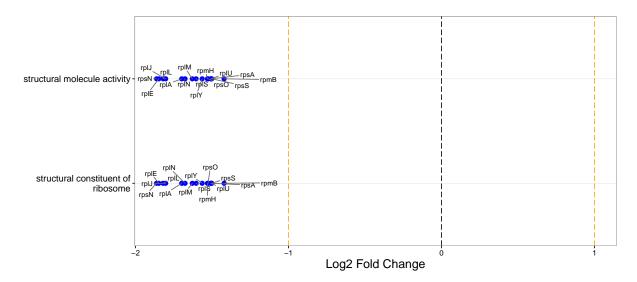


Figure S18: Significantly differentially expressed GO annotations related with molecular functions and associated genes with glycerol as carbon source in exponential phase, as determined by mRNA abundances. The top differentially expressed KEGG pathways are shown along the y axis, and the relative fold change of the corresponding genes is shown along the x axis. We show up to 10 most significantly changed pathways and for each pathway, we show up to 15 of the most significantly changing genes.

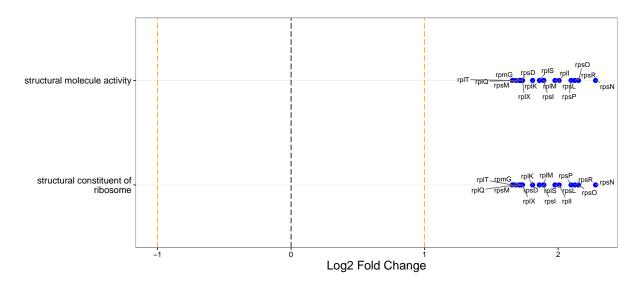


Figure S19: Significantly differentially expressed GO annotations related with molecular functions and associated genes with lactate as carbon source in exponential phase, as determined by mRNA abundances. The top differentially expressed KEGG pathways are shown along the y axis, and the relative fold change of the corresponding genes is shown along the x axis. We show up to 10 most significantly changed pathways and for each pathway, we show up to 15 of the most significantly changing genes.

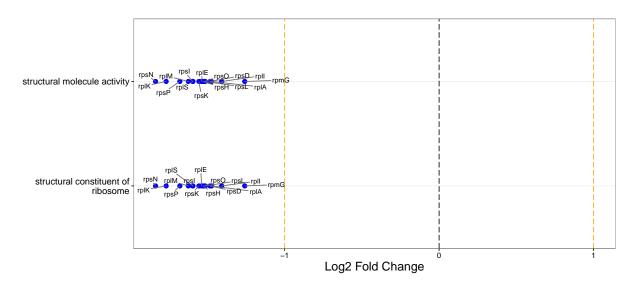


Figure S20: Significantly differentially expressed GO annotations related with molecular functions and associated genes with low  $Mg^{+2}$  levels in exponential phase, as determined by mRNA abundances. The top differentially expressed KEGG pathways are shown along the y axis, and the relative fold change of the corresponding genes is shown along the x axis. We show up to 10 most significantly changed pathways and for each pathway, we show up to 15 of the most significantly changing genes.

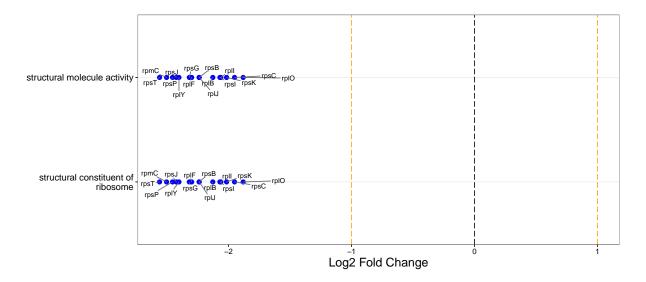


Figure S21: Significantly differentially expressed GO annotations related with molecular functions and associated genes with high Na<sup>+1</sup> levels in exponential phase, as determined by protein abundances. The top differentially expressed KEGG pathways are shown along the y axis, and the relative fold change of the corresponding genes is shown along the x axis. We show up to 10 most significantly changed pathways and for each pathway, we show up to 15 of the most significantly changing genes.

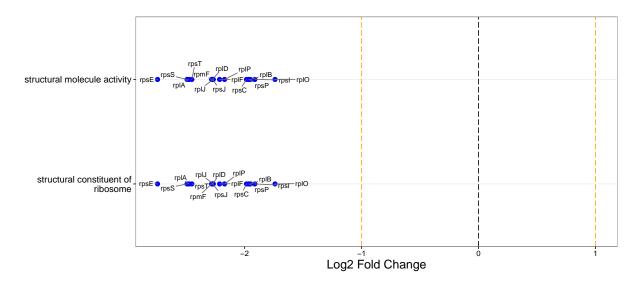


Figure S22: Significantly differentially expressed GO annotations related with molecular functions and associated genes with high Na<sup>+1</sup> levels in stationary phase, as determined by protein abundances. The top differentially expressed KEGG pathways are shown along the y axis, and the relative fold change of the corresponding genes is shown along the x axis. We show up to 10 most significantly changed pathways and for each pathway, we show up to 15 of the most significantly changing genes.

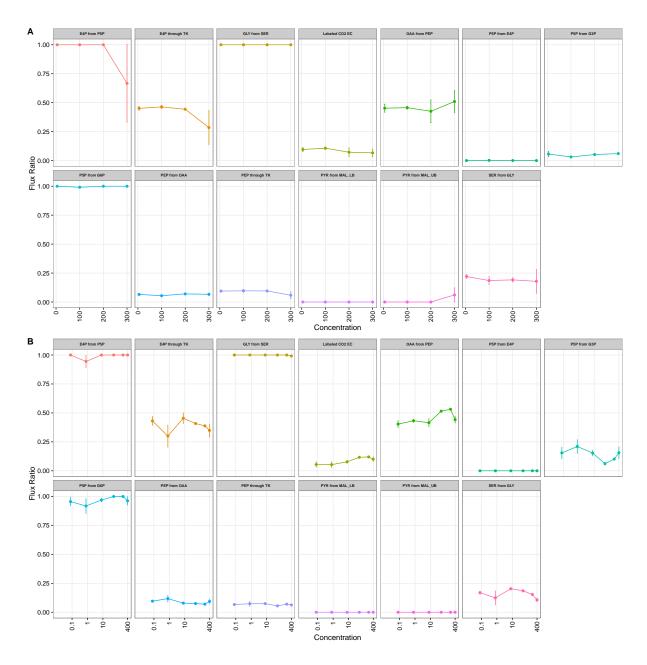


Figure S23: Flux changes with respect to salt stresses in exponential phase. 13 different flux were measured with respect to four different Na and five different Mg2+ concentrations. (A) Concentrations with respect to changing Na+ concentrations. (B) Concentrations with respect to changing Mg2+ concentrations. There is no significant trend of increase or decrease in exponential phase flux measurements with respect to either Mg2+ or Na+ concentrations after fdr correction for multiple testing. Correction. The only significant trend was for labelled CO2 EC branch at exponential phase with respect to Mg2+ concentrations before multiple testing (P=.02), but even for this branch, after multiple testing correction corresponding p value increases to .4.