$1\,\,\,\,$ E. coli Glucose data with alpha $=0.01\,\,\,$

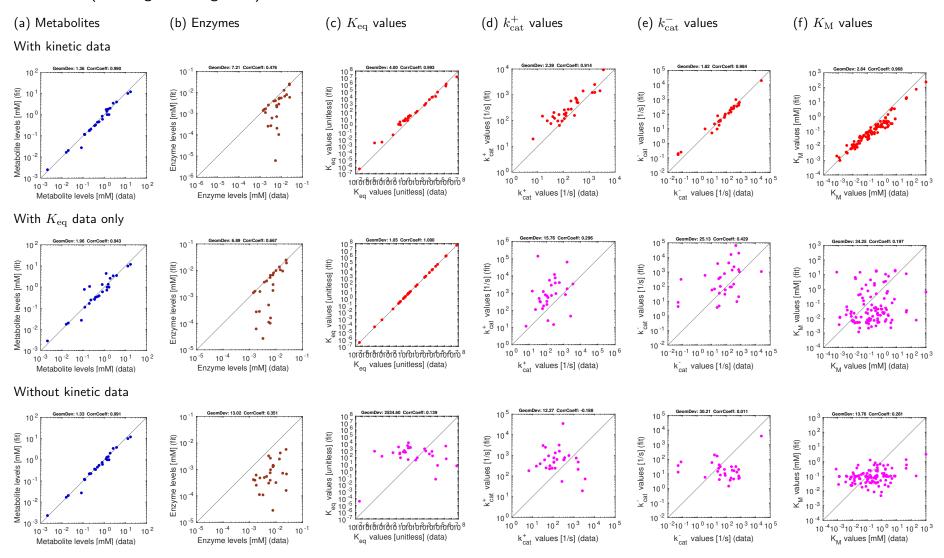


Figure 3: Results for *E. coli* central metabolism with experimental data (aerobic growth on glucose). The kinetic data stem from previous parameter balancing based on *in-vitro* data. Top: estimation using kinetic data. Centre: estimation using equilibrium constants as the only kinetic data. Centre: estimation without usage of kinetic data. The same metabolite, enzyme, and kinetic data were used in [?].

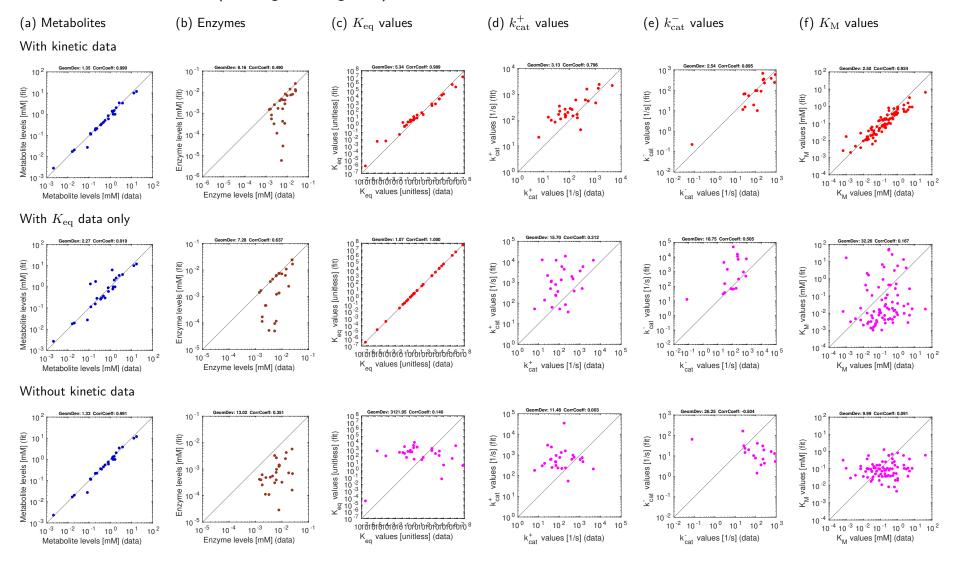


Figure 4: Results for *E. coli* central metabolism with experimental data (aerobic growth on glucose). Same as Figure 23, but based on original kinetic *in-vitro* data instead of balanced kinetic data.

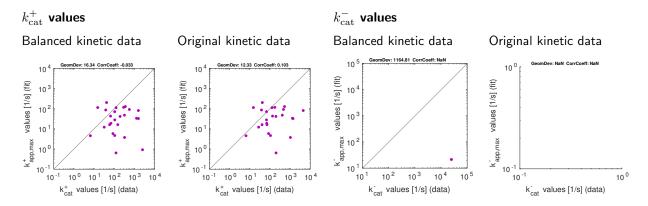
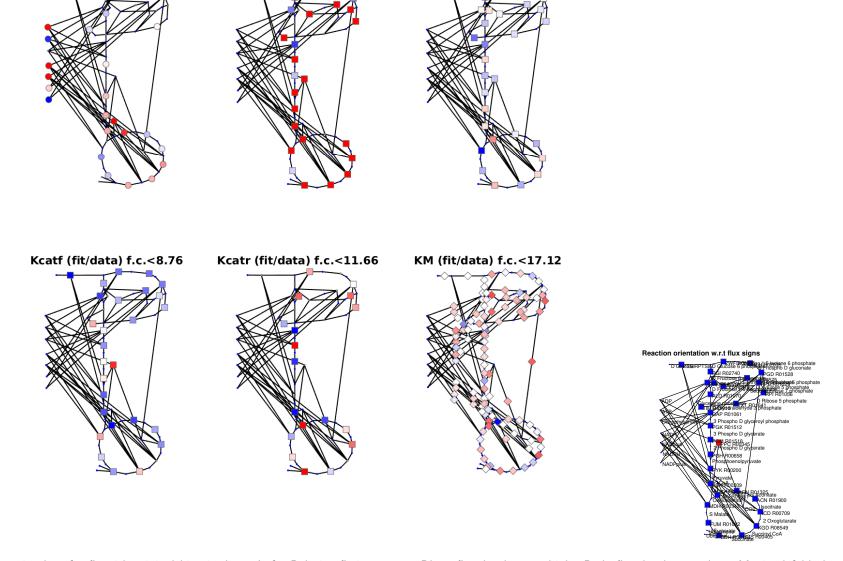


Figure 5: Catalytic constants in E. coli central metabolism model (aerobic growth on glucose), estimated by kinetic profiling [?].





met (fit/data) smpl 1 f.c.<3.02enz (fit/data) smpl 1 f.c.<888.33 Keq (fit/data) f.c.<252.77

Figure 6: Diagnostic plots for fit with original kinetic data. Left: Relative fitting errors. Blue: fitted value too high. Red: fitted value too low. Maximal fold changes (whether up or down) are given by numbers. Right: Reaction orientatation (defining "forward" and "reverse"). Blue: in flux direction; Red: against flux direction.

2 E. coli Glucose data with alpha = 0.1

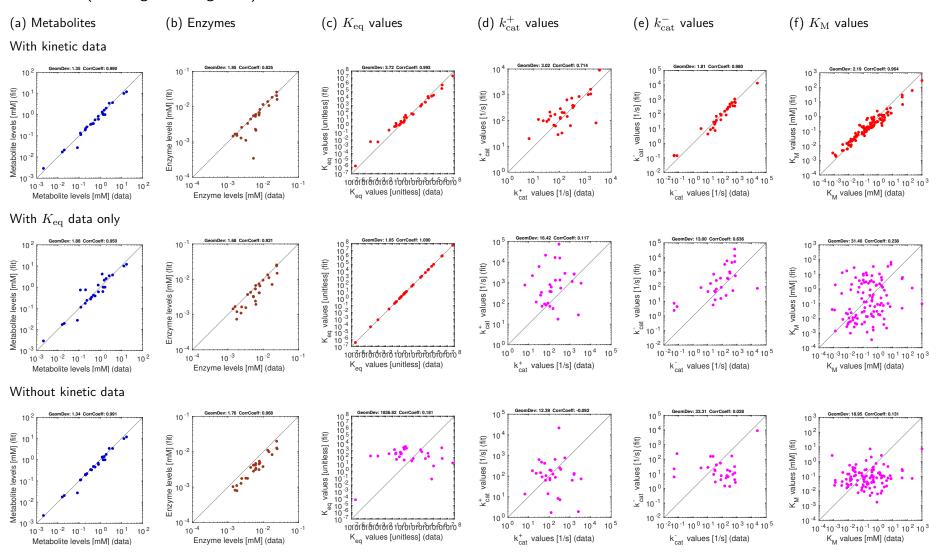


Figure 7: Results for *E. coli* central metabolism with experimental data (aerobic growth on glucose). The kinetic data stem from previous parameter balancing based on *in-vitro* data. Top: estimation using kinetic data. Centre: estimation using equilibrium constants as the only kinetic data. Centre: estimation without usage of kinetic data. The same metabolite, enzyme, and kinetic data were used in [?].

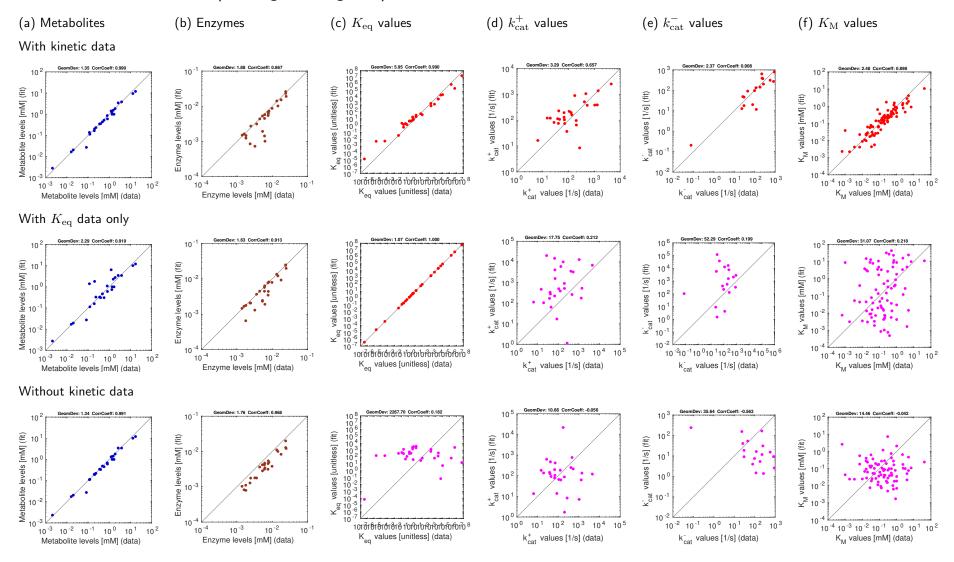


Figure 8: Results for *E. coli* central metabolism with experimental data (aerobic growth on glucose). Same as Figure 23, but based on original kinetic *in-vitro* data instead of balanced kinetic data.

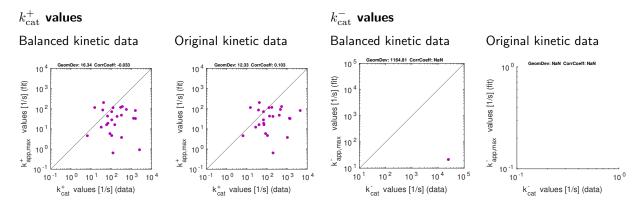
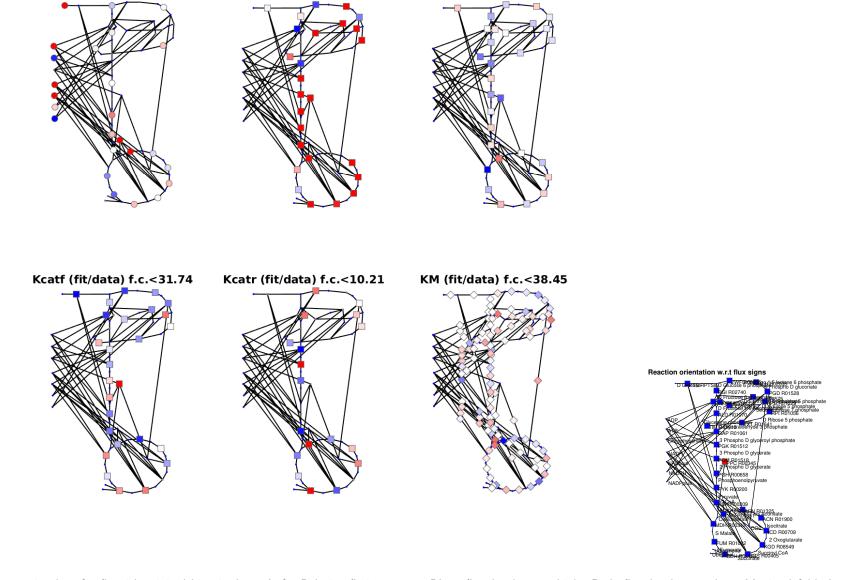


Figure 9: Catalytic constants in E. coli central metabolism model (aerobic growth on glucose), estimated by kinetic profiling [?].



met (fit/data) smpl 1 f.c.<3.03 enz (fit/data) smpl 1 f.c.<6.11 Keq (fit/data) f.c.<259.22

Figure 10: Diagnostic plots for fit with original kinetic data. Left: Relative fitting errors. Blue: fitted value too high. Red: fitted value too low. Maximal fold changes (whether up or down) are given by numbers. Right: Reaction orientatation (defining "forward" and "reverse"). Blue: in flux direction; Red: against flux direction.

3 E. coli Glucose data with alpha = 0.5

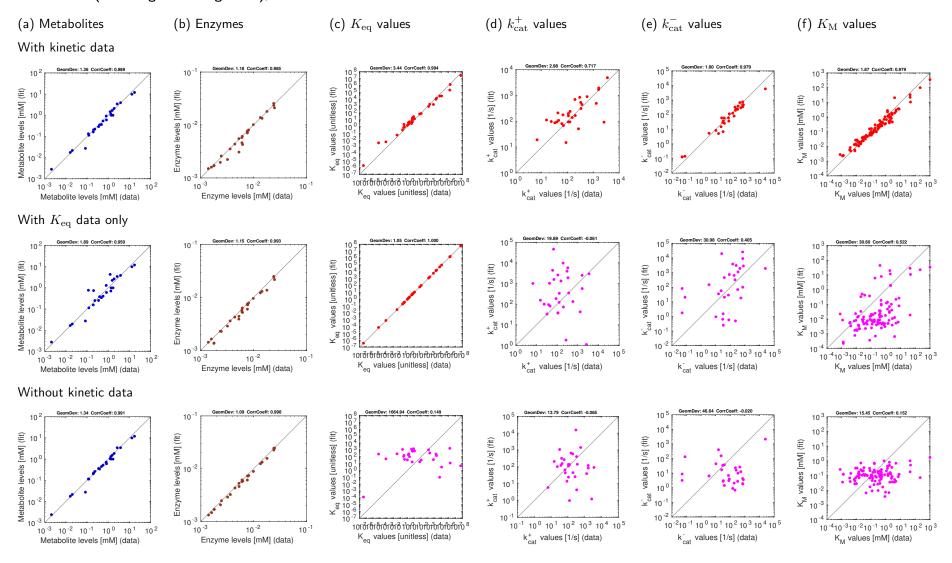


Figure 11: Results for *E. coli* central metabolism with experimental data (aerobic growth on glucose). The kinetic data stem from previous parameter balancing based on *in-vitro* data. Top: estimation using kinetic data. Centre: estimation using equilibrium constants as the only kinetic data. Bottom: estimation without usage of kinetic data. The same metabolite, enzyme, and kinetic data were used in [?].

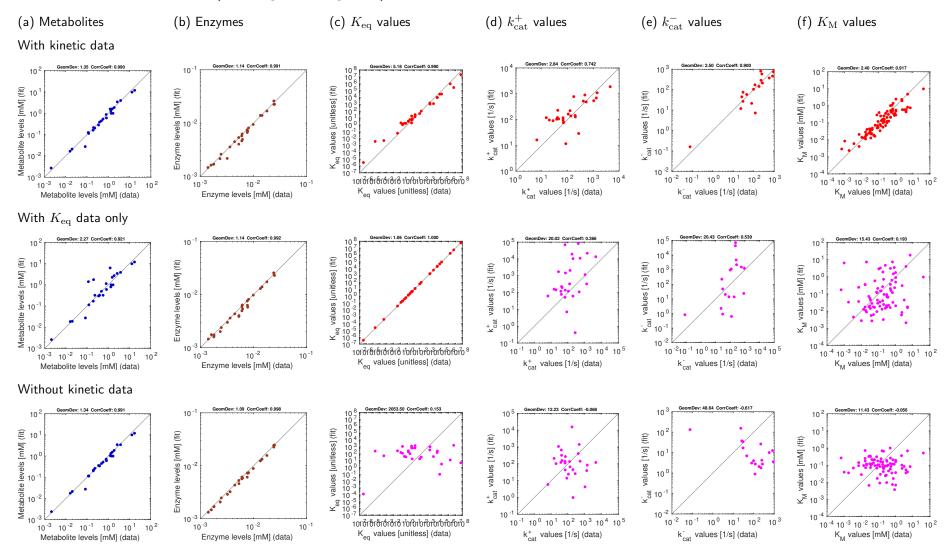


Figure 12: Results for *E. coli* central metabolism with experimental data (aerobic growth on glucose). Same as Figure 23, but based on original kinetic *in-vitro* data instead of balanced kinetic data.

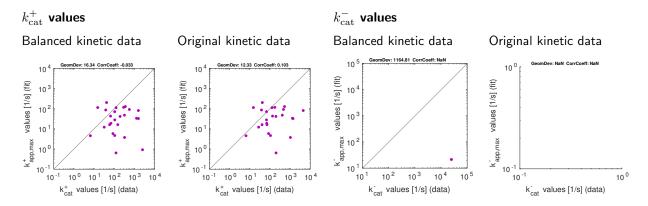
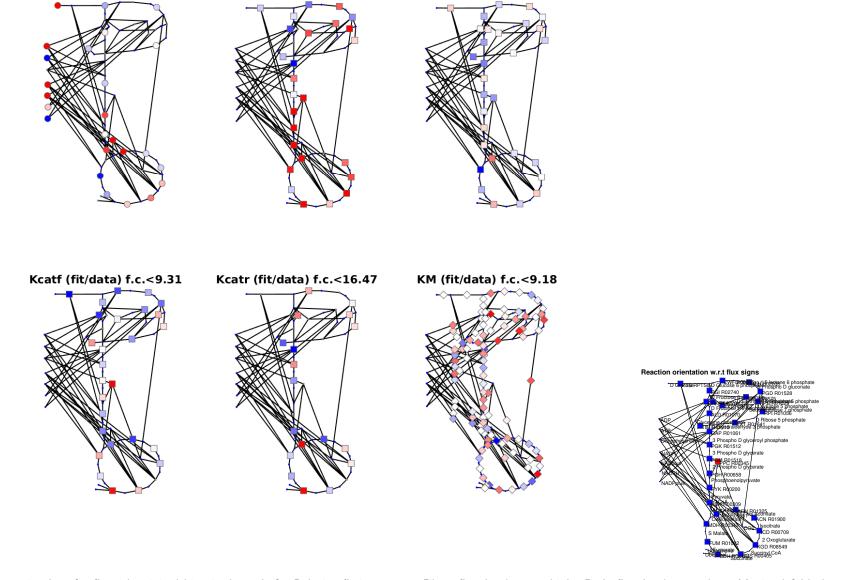


Figure 13: Catalytic constants in E. coli central metabolism model (aerobic growth on glucose), estimated by kinetic profiling [?].



met (fit/data) smpl 1 f.c.<3.03 enz (fit/data) smpl 1 f.c.<1.48 Keq (fit/data) f.c.<196.14

Figure 14: Diagnostic plots for fit with original kinetic data. Left: Relative fitting errors. Blue: fitted value too high. Red: fitted value too low. Maximal fold changes (whether up or down) are given by numbers. Right: Reaction orientatation (defining "forward" and "reverse"). Blue: in flux direction; Red: against flux direction.

4 E. coli Glucose data with alpha = 1

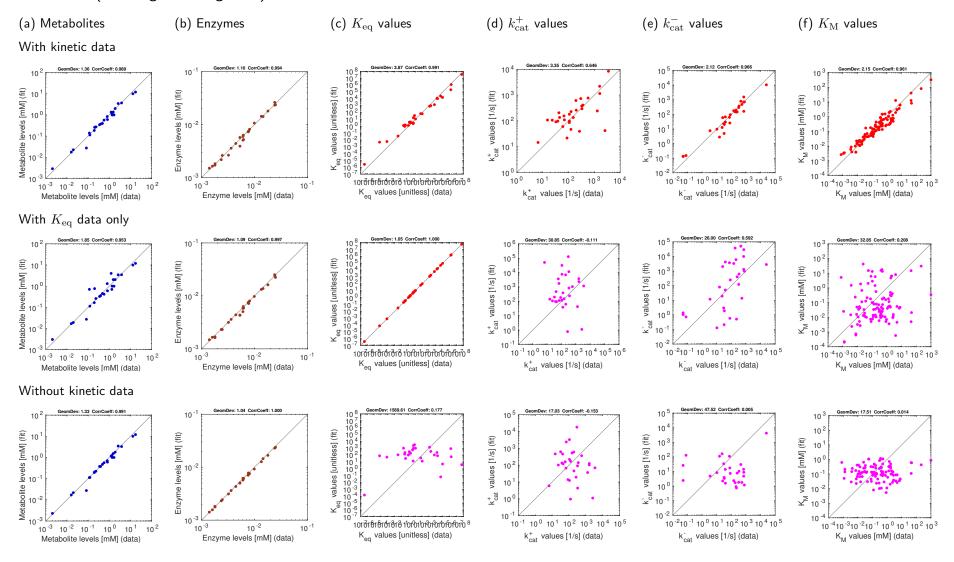


Figure 15: Results for *E. coli* central metabolism with experimental data (aerobic growth on glucose). The kinetic data stem from previous parameter balancing based on *in-vitro* data. Top: estimation using kinetic data. Centre: estimation using equilibrium constants as the only kinetic data. Centre: estimation without usage of kinetic data. The same metabolite, enzyme, and kinetic data were used in [?].

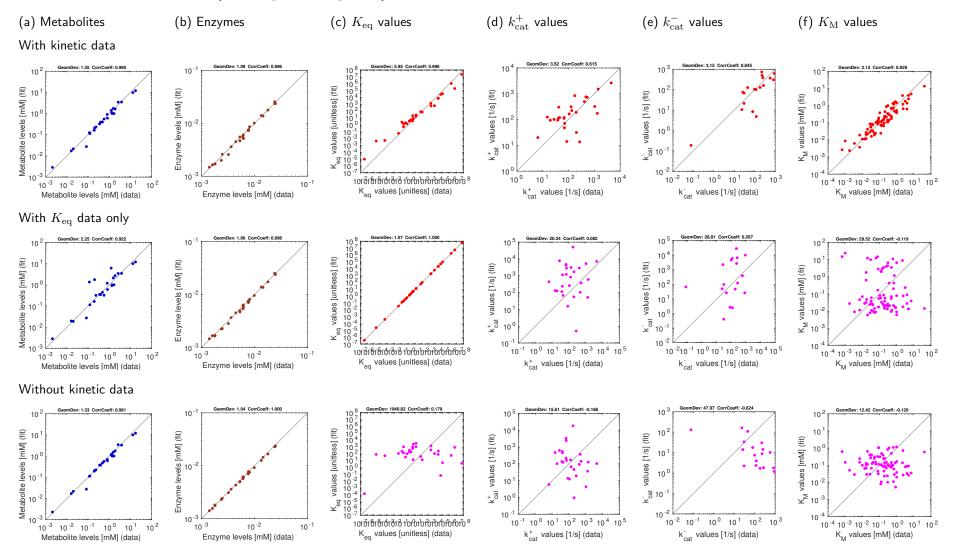


Figure 16: Results for *E. coli* central metabolism with experimental data (aerobic growth on glucose). Same as Figure 23, but based on original kinetic *in-vitro* data instead of balanced kinetic data.

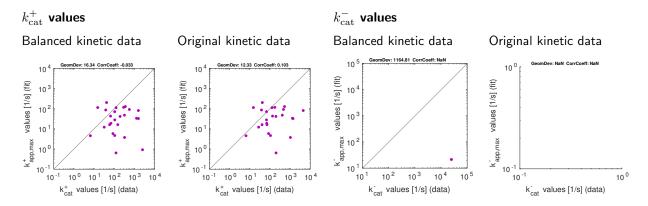


Figure 17: Catalytic constants in E. coli central metabolism model (aerobic growth on glucose), estimated by kinetic profiling [?].

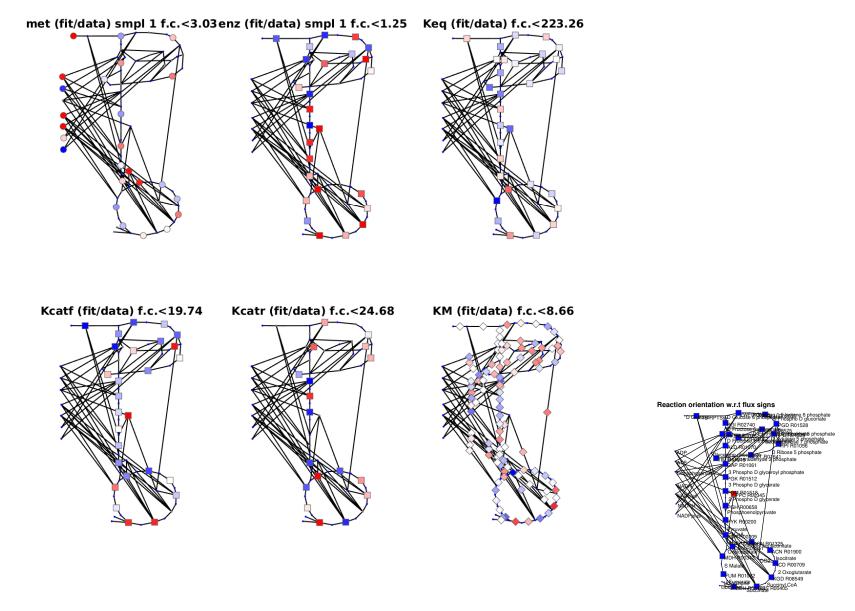


Figure 18: Diagnostic plots for fit with original kinetic data. Left: Relative fitting errors. Blue: fitted value too high. Red: fitted value too low. Maximal fold changes (whether up or down) are given by numbers. Right: Reaction orientatation (defining "forward" and "reverse"). Blue: in flux direction; Red: against flux direction.