

Correction: Integration of flux measurements to resolve changes in anabolic and catabolic metabolism in cardiac myocytes

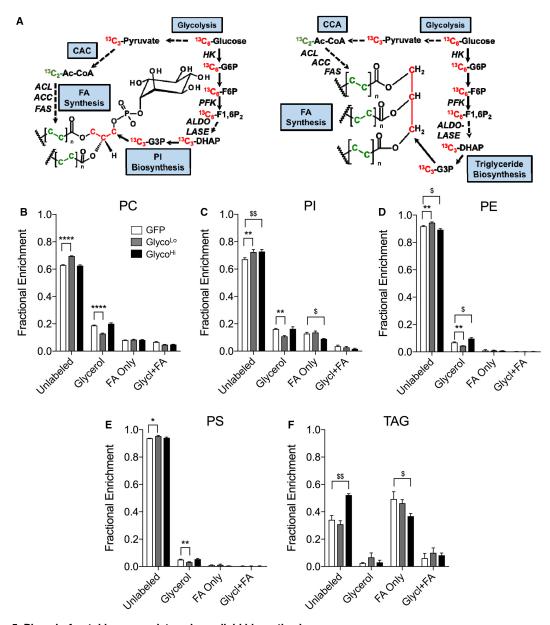
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In the isotopologue analysis of glycero(phospho)lipids (Figure 5), the authors mistakenly included glycerolipids with only the glycerol moiety labeled (i.e., m+3) in the Glycerol + Fatty Acid labeling groups (Glycl+FA). Fractional enrichment of 13 C into Glycl+FA of panels B–F should include only those glycerolipids having both the glycerol and fatty acyl chains labeled (e.g. this would include m+5, m+7, m+9, m+11, m+13, m+15, etc.). The corrected figure is presented here. These changes do not change the conclusions drawn from the study and do not require modifications to the text of the manuscript.





 $\label{lem:figure 5.} \textbf{Phosphofructokinase} \ \textbf{regulates} \ \textbf{glycerolipid} \ \textbf{biosynthesis}.$

Stable isotope tracing of phospholipids and triacylglycerols in cardiomyocytes incubated with media containing 13 C₆-glucose for 18 h: (**A**) atom-resolved map illustrating the biological and biochemical history of 13 C incorporation into glycerolipids; fractional enrichment values of 13 C into: (**B**) PC; (**C**) PI; (**D**) PE; (**E**) PS; and (**F**) TAG. Graph represents three replicates per group from one isolation. *,* $^{\$}P$ < 0.05, **,* $^{\$}P$ < 0.01, **** $^{**}P$ < 0.0001.