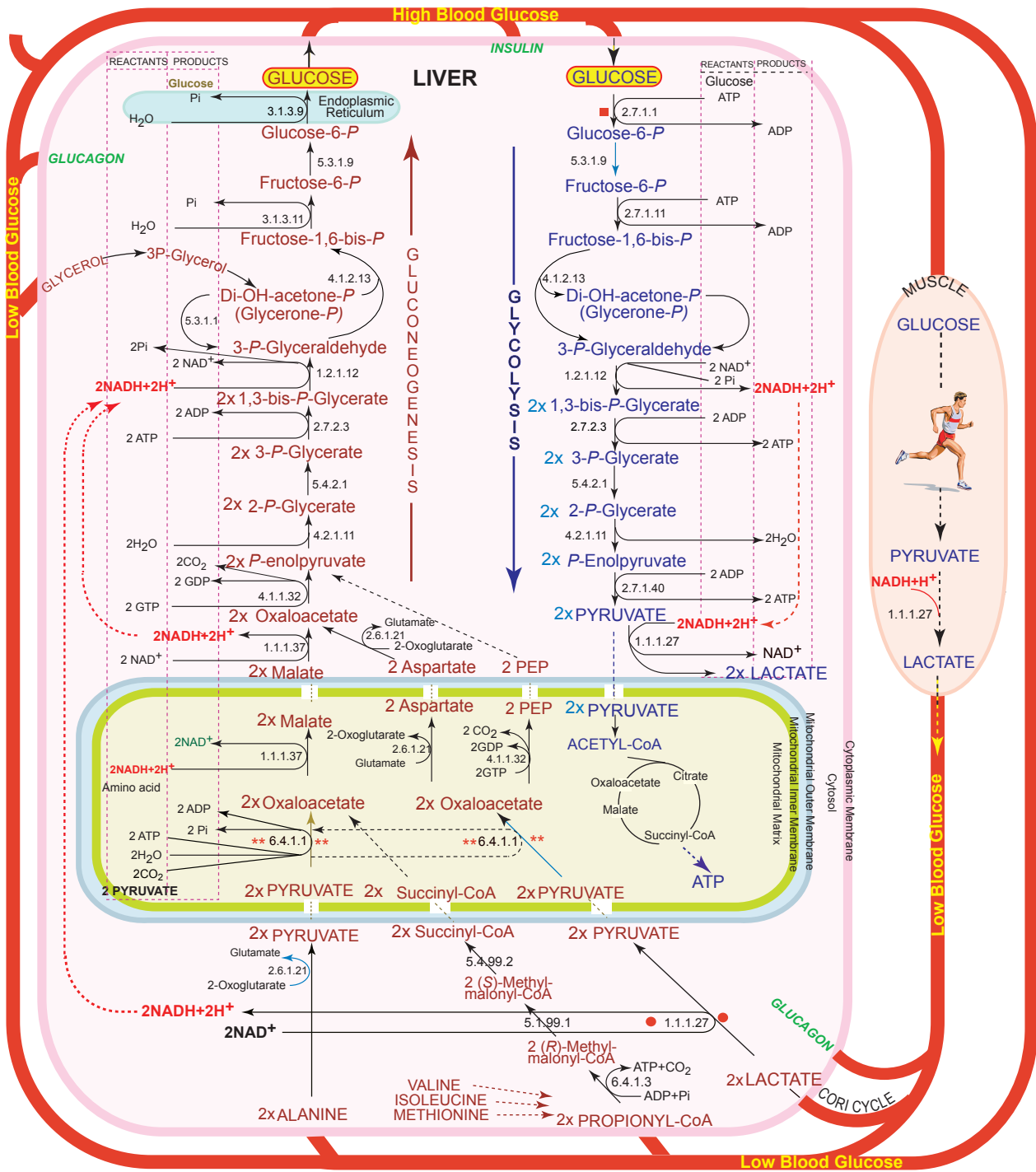


# GLUCONEOGENESIS & GLYCOLYSIS



Comparison of the two equations shows that:

**OXIDATION** of 1 **GLUCOSE** to 2 **PYRUVATE** produces 2**ATP** whereas **REDUCTION** of 2 **PYRUVATE** to 1 **GLUCOSE** requires 4**ATP** + 2**GTP**  
 To make **GLUCONEOGENESIS** thermodynamically possible requires the hydrolysis of the equivalent of 6**ATP**

Gluconeogenesis needs NADH for the reduction of 1,3-bis-P-glycerate to triose-P. For most precursors this is formed in the mitochondria and transported across the mitochondrial membrane via malate. Lactate is unique in that lactate dehydrogenase only occurs in the cytosol so that NADH is directly available in the cytosol for gluconeogenesis. However, the resulting pyruvate can only be converted into oxaloacetate within mitochondria since this is the only site of pyruvate carboxylase. The oxaloacetate so formed can then be transported into the cytosol by direct carboxylation to PEP or by transamination to aspartate in the mitochondria and then reverse transamination to oxaloacetate in the cytosol

ENZYMES		
● 1.1.1.27 Lactate dehydrogenase (only in cytosol)	2.7.1.40 Pyruvate kinase	5.1.99.1 Methylmalonyl-CoA epimerase
1.1.1.37 Malate dehydrogenase	2.7.2.3 Phosphoglycerate kinase	5.3.1.1 Triosephosphate isomerase
1.2.1.12 Glyceraldehyde-3-P dehydrogenase	3.1.3.9 Glucose-6-phosphatase	5.4.2.1 Phosphoglycerate mutase
■ 2.7.1.1 Hexokinase	3.1.3.11 Fructose-bisphosphatase	5.4.99.2 Methylmalonyl-Co mutase
2.7.1.30 Glycerolkinase	4.1.1.32 Phosphoenolpyruvate carboxykinase	6.2.1.17 Propionate-CoA ligase
2.7.1.11 6-Phosphofructokinase	4.1.2.13 Fructose-bisphosphatealdolase	** 6.4.1.1 Pyruvate carboxylase (only in matrix)
■ Liver Hexokinase is Hexokinase D (or IV) (Often called Glucokinase)	4.2.1.11 Phosphopyruvate hydratase	6.4.1.3 Propionyl-CoA carboxylase