

Table S1. Furukawa et al.

Table S1. Primers used for quantitative PCR

gene name		primer sequence (5' – 3')	amplicon size (bp)	accession number
<i>g6pca.1</i>	F	TTGTTTCAGGATTGGATGTATTG	131	NM_001003512
	R	GAAGCAGCAAAGCCACAG		
<i>g6pca.2</i>	F	CACACGGCTGCTCTCTTCT	89	NM_001163806
	R	GATAAGCAGTACGGGATGATG		
<i>pck1</i>	F	CAGTAAACACGGCTGAAGACAC	122	NM_214751
	R	CGGTTTTGATGCACTTGAGA		
<i>pck2</i>	F	TCTGGCAGAAGGAAACACA	116	NM_213192
	R	TCAATCCCTCACTCTCTCCTC		
<i>glud1a</i>	F	CACTTCTGATTTTCAAGAAAGGA	125	NM_212576
	R	CAAGGTATATACCGGTTGGCT		
<i>glud1b</i>	F	TGAGTGGCTGAAGAACCTGA	84	NM_199545
	R	ATCAGCAGGTGGTAGTTGGAG		
<i>glsa</i>	F	ATCAGCAGGTGGTAGTTGGAG	114	NM_001045044
	R	GGGCTGTAGGTGTTGTGATAG		
<i>glsb</i>	F	CTATAGGATGGAGACTGTTGGAGA	117	XM_688079
	R	TGACCCGAATGAGAACCAG		
<i>gl2a</i>	F	TTTCTGGGTTGGAGGACTTG	100	BX927144, ENSDARG00000069095
	R	TGCAAACCTGTGCTTTTGAG		
<i>gl2b</i>	F	ACAGAAACAAGTCGGTGGTG	110	NM_001083825, ENSDARG00000002917
	R	GTCATAGTCCCTCAACTCCATATTC		
<i>gl2-like</i>	F	GAGCTGGTTTCAACCTTCCA	142	ENSDARG000000062781
	R	CCAGGTCCCCTCTATAAGCAG		
<i>rpl13a</i>	F	TCTGGAGGACTGTAAGAGGTATGC	150	NM_212784.1
	R	TCAGACGCACAATCTTGAGAGCAG		

F, forward; R, reverse.

Table S2. Primers used to generate templates for RNA probe synthesis

gene name		primer sequence (5' – 3')	amplicon size (bp)
<i>g6pca.1</i>	F	CCGTCCACTTACAGATCAAATAC	957
	R	GAAGCAGCAAAGCCACAG	
<i>g6pca.2</i>	F	CTCATTACAAAGATGCCCAAG	970
	R	GATAAGCAGTACGGGATGATG	
<i>pck1</i>	F	CGCTCATCATGCCTCCTC	2007
	R	CGGTTTTGATGCACTTGAGA	
<i>pck2</i>	F	CAATTTACCTGCACCAAAG	1931
	R	TCAATCCCTCACTCTCTCCTC	
<i>glud1a</i>	F	CAGGACCCCGTGTAAGGA	1166
	R	CAAGGTTATACCGTTGGCT	
<i>glud1b</i>	F	ATGGTGGAGGGCTTCTTTG	1199
	R	ATCAGCAGGTGGTAGTTGGAG	
<i>glsa</i>	F	TTCGAGGGCGTTAAAGGAG	1695
	R	GGGCTGTAGGTGTTGTGATAG	
<i>glsb</i>	F	GCTTCAATTGTGCTGAAGGAG	1763
	R	TGACCCGAATGAGAACCAG	
<i>gls2a</i>	F	TGCTGGAAAATGGGAAAGAG	1327
	R	TGCAAACCTGTGCTTTTGAG	
<i>gls2b</i>	F	TCCAGTGTCTCATTTCACTGCT	1294
	R	GTCATAGTCCCTCAACTCCATATTC	
<i>gls-like</i>	F	ATCTCTCAGACGTAAATGGAGAAAG	1619
	R	CCAGGTCCCCTCTATAAGCAG	

F, forward; R, reverse.

Table S3. List of mRNA localization detected by *in situ* hybridization

gene name	localization	putative function
<i>g6pca.1</i>	yolk syncytial layer, liver (ER)	glucose production
<i>g6pca.2</i>	yolk syncytial layer, liver (ER)	glucose production
<i>pck1</i>	liver (cytosolic)	PEP synthesis, cataplerosis
<i>pck2</i>	liver, intestinal bulb (mitochondrial)	PEP synthesis, cataplerosis
<i>glud1a</i>	liver, intestinal bulb (mitochondrial)	glutamate catabolism
<i>glud1b</i>	liver, intestinal bulb (mitochondrial)	glutamate catabolism
<i>glsa</i>	brain (cytosolic)	glutamine catabolism
<i>glsb</i>	pectoral fin (cytosolic)	glutamine catabolism
<i>gls2a</i>	liver (mitochondrial)	glutamine catabolism
<i>gls2b</i>	liver, intestinal bulb (mitochondrial)	glutamine catabolism
<i>gls-like</i>	swim bladder (no data)	(glutamine catabolism?)

Putative intracellular localization of the protein product was written in parentheses. ER, endoplasmic reticulum; PEP, phosphoenolpyruvate.