Table S4. Enriched KEGG pathways identified with DAVID (p<0.05).

<u>Category</u>	‡ <u>Term</u>	≑ RT	Genes	Count	♦ <u>%</u> :	P-Value	Benjamini 💠
KEGG_PATHWAY	<u>Circadian rhythm</u>	<u>RT</u>	i	9	0.6	4.0E-7	6.8E-5
KEGG_PATHWAY	Pathways in cancer	<u>RT</u>	=	39	2.7	5.0E-4	4.2E-2
KEGG_PATHWAY	Insulin signaling pathway	RT		18	1.2	1.1E-2	4.8E-1
KEGG_PATHWAY	Small cell lung cancer	<u>RT</u>	i .	13	0.9	1.1E-2	3.9E-1
KEGG_PATHWAY	TGF-beta signaling pathway	<u>RT</u>	i i	13	0.9	1.4E-2	3.8E-1
KEGG_PATHWAY	Alanine, aspartate and glutamate metabolism	<u>RT</u>	i i	7	0.5	1.4E-2	3.3E-1
KEGG_PATHWAY	Renal cell carcinoma	RT	i i	11	0.8	1.9E-2	3.7E-1
KEGG_PATHWAY	Glycine, serine and threonine metabolism	<u>RT</u>	i	7	0.5	1.9E-2	3.3E-1
KEGG_PATHWAY	Cysteine and methionine metabolism	<u>RT</u>	i	7	0.5	2.2E-2	3.4E-1
KEGG_PATHWAY	Nicotinate and nicotinamide metabolism	<u>RT</u>	i i	6	0.4	2.4E-2	3.4E-1
KEGG_PATHWAY	Adipocytokine signaling pathway	<u>RT</u>	i i	10	0.7	3.6E-2	4.3E-1
KEGG_PATHWAY	PPAR signaling pathway	<u>RT</u>	i i	11	0.8	4.0E-2	4.4E-1
KEGG_PATHWAY	Prostate cancer	<u>RT</u>	i i	12	0.8	4.0E-2	4.2E-1
KEGG_PATHWAY	Oocyte meiosis	<u>RT</u>	i i	14	1.0	4.6E-2	4.4E-1

The BMAL1 putative targets in the three most significant pathways are shown in the Figure S5.