Estivation-responsive microRNAs in a hypometabolic terrestrial snail

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ABSTRACT

When faced with extreme environmental conditions, the milk snail (Otala lactea) enters a state of dormancy known as estivation. This is characterized by a strong reduction in metabolic rate to <30% of normal resting rate that is facilitated by various behavioural, physiological, and molecular mechanisms. Herein, we investigated the regulation of microRNA in the induction of estivation. Changes in the expression levels of 75 highly conserved microRNAs were analysed in snail foot muscle, of which 26 were significantly upregulated during estivation compared with controls. These estivation-responsive microRNAs were linked to cell functions that are crucial for long-term survival in a hypometabolic state including anti-apoptosis, cell-cycle arrest, and maintenance of muscle functionality. Several of the microRNA responses by snail foot muscle also characterize hypometabolism in other species and support the existence of a conserved suite of miRNA responses that regulate environmental stress responsive metabolic rate depression across phylogeny.

INTRODUCTION

Native to seasonally arid regions of southern Europe, the milk snail, Otala lactea, experiences seasonal exposures to high temperatures and limited food and water. To withstand these harsh environmental conditions, the snail spends months in a dormant state known as estivation (Ramnanan & Storey, 2006a; Ramnanan & Storey, 2006b). Estivation is a survival strategy characterized by a strong reduction of basal metabolic rates to <30% of normal resting rates. Such metabolic rate depression (MRD) is a widespread survival strategy used by both vertebrates and invertebrates as a means of reprioritizing energy usage towards vital cellular processes, and preserving/stabilizing macromolecules to maximize survival when faced with unfavourable environmental conditions (Storey, 2002; Storey & Storey, 2004; Storey & Storey, 2010). However, retreating into an estivating state exposes the snail to cellular stresses including dehydration, hypoxia (oxygen limitation), hypercapnia (elevated CO2 levels), and acidosis (Churchill & Storey, 1989; Storey, 2002; Storey & Storey, 2012). As such, the milk snail has developed various adaptations to overcome estivation-associated stresses and to facilitate survival over prolonged periods of dormancy (Storey, 2002; Storey & Storey, 2012).

Although dormancy is a life-saving strategy for many animals, this phenomenon is far from being a simple feat. Indeed, animals need to undergo profound biochemical changes...
(alterations to gene and protein expression and enzyme regulation) that modulate cellular metabolism to allow for safe transitions into dormancy while stabilizing long-term viability. As such, these regulatory mechanisms must be rapid, reversible, easily inducible, and have a low energy demand (Storey & Storey, 2007). MicroRNAs (miRNAs) possess all of these characteristics, making them excellent candidate regulators of hypometabolism (Biggar & Storey, 2011). Indeed, miRNAs have been shown to be master regulators of virtually all cellular processes, exerting widespread control over activities including signal transduction pathways, cell cycle, apoptosis, atrophy, energy metabolism, and many others. These small non-coding RNA molecules act post-transcriptionally to modulate gene expression by permanently repressing translation by directing mRNA transcripts to degradation, or temporarily targeting mRNAs into storage in stress granules or P-bodies (Nilsen, 2007). In addition, miRNAs provide remarkable regulatory potential and flexibility since an individual miRNA type can target multiple different mRNA transcripts, and equally, a single type of mRNA transcript can be targeted by diverse miRNAs (Mazière & Enright, 2007; Biggar & Storey, 2011).

MiRNA-mediated regulation of gene expression has been well established as part of a typical stress response (e.g., during ischemic injury, cardiac hypertrophy, etc.) and recent studies have highlighted their role in multiple forms of environmental stress-induced hypometabolism (Biggar & Storey, 2018) including mammalian hibernation (Hadj-Moussa et al., 2016), hypoxia/anoxia tolerance (Biggar et al., 2012; Hadj-Moussa et al., 2018), estivation (Chen et al., 2013; Wu, Biggar & Storey, 2013; Chen & Storey, 2014; Luu & Storey, 2015), insect cold-hardiness/diapause (Lyons et al., 2015), and freeze tolerance (Biggar et al., 2012; Lyons et al., 2013; Hadj-Moussa & Storey, 2018). These studies linked differential miRNA expression to protective actions in the hypometabolic state including minimizing apoptosis and muscle wasting, limiting non-essential energy-expensive processes, promoting cell cycle arrest, and differentially regulating selected signalling cascades (Biggar & Storey, 2011). Additionally, miRNAs have also been shown to mediate survival responses to desiccation and osmotic and heat stress in marine molluscs (Zhou et al., 2014; Zhao et al., 2016; Chen et al., 2017). Consequently, we predicted that miRNAs are also likely to be involved in facilitating the successful transition into estivation in O. lactea. In this study, we examined the expression levels of 75 conserved miRNAs in snail foot muscle, comparing active snails to those in prolonged estivation, to better understand the molecular processes involved in mediating estivation and stress survival. This represents the first report of differential miRNA expression in an estivating land snail.

MATERIALS AND METHODS

Animal care and treatment

Otala lactea snails, imported from Morocco, were purchased from a local seafood retailer. In the laboratory, snails were treated as described by Ramnanan, Groom & Storey (2007). All animals were held at ~22 °C in plastic boxes lined with damp paper towels and fed shredded carrots and cabbage (sprinkled with crushed chalk) every 2–3 days. After 4 weeks, half of the snails were moved to a container with dry paper towels and no food where they
quickly entered estivation. Other snails were maintained under active conditions. After 10 days, active (control) and estivating snails were euthanized, and foot muscle was dissected out, flash frozen in liquid nitrogen, and stored at −80 °C.

**Total RNA isolation**

Total RNA was isolated from foot muscle as previously described by Hadj-Moussa *et al.* (2018). In brief, samples were homogenized in Trizol (Invitrogen), RNA was isolated using chloroform, and precipitated with isopropanol. RNA pellets were resuspended in RNase-free water, and samples were stored at −20 °C until use.

**Polyadenylation and stem-loop reverse transcription**

RNA samples were prepared for miRNA analysis as described by Biggar, Wu & Storey (2014). Briefly, RNA samples were polyadenylated using a PolyA tailing kit (Cat # .PAP5104H; Epicenter, Madison, WI, USA). Reverse transcription was performed to generate cDNA from polyadenylated RNA using a stem-loop adapter primer (Table S1). Serial dilutions of the cDNA were prepared and stored at −20 °C until use.

**Primer design**

Snail miRNA-specific forward primers were designed using the annotated gastropod (*Lottia gigantea*) miRNA sequences and synthesized by Integrated DNA Technologies (Biggar, Wu & Storey, 2014). The primers were validated by assessing the miRNA sequence conservation of *L. gigantea* against other molluscs including gastropods (*Aplysia californica, Lymnaea stagnalis*), a bivalve (*Crassostrea gigas*) and a cephalopod (*Octopus bimaculoides*) using NCBI BLASTn and CLUSTAL OMEGA. All miRNA-specific forward primers were designed with a modified short adapter at the 5′ end of the mature miRNA sequence, with the following general sequence 5′-ACA CTC CAG CTG GGN NNN NNN NNN NNN NN-3′, where N denotes miRNA-specific sequence binding regions (Biggar, Wu & Storey, 2014) (Table S1).

**Relative miRNA quantification**

qRT-PCR reactions were performed as previously described (Pellissier *et al.*, 2006) following MIQE guidelines (Bustin *et al.*, 2009) using a BioRad MyIQ Detection System. Target-specific forward primers were used in conjunction with a universal reverse primer to amplify each miRNA target (Table S1). A post-run melt-curve was performed after all PCR reactions to ensure the amplification of a single product. Reactions that amplified multiple products were rejected.

Relative miRNA expression levels were calculated using the comparative ΔΔCq method as previously described (Luu & Storey, 2015). In brief, raw Cq values were converted to the $2^{-\Delta Cq}$ form and normalized to the 5S rRNA reference gene. The 5S rRNA gene displayed abundant and stable expression in *O. lactea* foot muscle under control and estivation conditions making it a suitable reference gene (Schmittgen & Livak, 2008). Standardized values were expressed as mean relative expression (mean ± SEM, with $n = 3−4$ independent biological replicates). Statistical analysis used a two sample Student’s *t*-test (Table S2).
RESULTS AND DISCUSSION

The roles of transcriptional, translational, and post-translational modifications in supporting MRD have been extensively analyzed in multiple animal systems over the years (including in hibernation, daily torpor, anaerobiosis, and estivation), whereas post-transcriptional miRNA-mediated metabolic regulation has only recently become evident (Storey & Storey, 2004; Biggar & Storey, 2011; Storey, 2015). Indeed, various post-translational regulatory controls on O. lactea signal transduction and metabolic enzymes have been reported (Brooks & Storey, 1990; MacLean et al., 2016; Ramnanan et al., 2009; Ramnanan et al., 2010; Ramnanan & Storey, 2006a; Ramnanan & Storey, 2006b; Whitwam & Storey, 1990) but, to date, the role of miRNA had not been explored. This study evaluates this level of post-transcriptional control. Studies have shown that O. lactea can arouse from dormancy in as little as 5–10 min and be immediately capable of extending its foot from the shell (Storey, 2002). As such, the readily inducible, rapid, and reversible nature of miRNAs makes them excellent candidates for quickly mediating metabolic changes needed to support entry into or exit from dormancy in the snail. Although the genome of O. lactea has not been sequenced, the high degree of evolutionary conservation of miRNAs across species, including both vertebrates and invertebrates, allowed for this miRNA analysis (Friedman et al., 2009).

A total of 75 miRNAs were successfully detected (Fig. 1), of which 26 were significantly upregulated, the increase during estivation ranging from 1.39–4.95 fold relative to expression in control foot muscle (Fig. 2; Table S2). Only five of the assessed miRNAs were downregulated but none of these showed a significant change compared with controls. Among the subset of upregulated miRNAs, miRNAs ola-miR-2a-3p, ola-miR-12-5p, ola-miR-190-5p, ola-miR-281-5p, ola-miR-723-5p, ola-miR-745b-3p, ola-miR-1989-5p, ola-miR-2001-5p showed strong upregulation by over 2.5 fold. The 26 miRNAs significantly elevated in the estivating state, suggest that their target genes were translationally suppressed during dormancy. Overall, these significantly upregulated miRNAs were implicated in regulating cell survival mechanisms that constituted three main functional groups: (1) anti-apoptosis, (2) tumour suppression, and (3) muscle maintenance responses. The present study showed elevated levels of five anti-apoptosis miRNAs (miR-2a-3p, miR-2c-3p, miRNA-124c, miRNA-153, and miRNA-190) in foot muscle (Gennarino et al., 2012; Wu et al., 2013; Dwivedi, 2011; Jia et al., 2016). The miR-2 family is an invertebrate-specific group of miRNAs involved in neural development and maintenance (Marco, Hooks & Griffiths-Jones, 2012). However, they have also been shown to specifically target the pro-apoptotic genes reaper, grim, and sickle, suggesting that miR-2 prevents apoptosis under stress and contributes to long-term life extension (Gennarino et al., 2012). Indeed, this miRNA was also upregulated in the intertidal snail Littorina littorea under freezing and anoxia stresses, as well as in the Humboldt squid Dosidicus gigas in response to hypoxia (Biggar et al., 2012; Hadj-Moussa et al., 2018). This suggests that miR-2 upregulation is a conserved invertebrate response to cellular stresses imposed by harsh environmental conditions.
Another anti-apoptotic miRNA identified was miR-153, a miRNA that is known to target the PTEN protein phosphatase to relieve Akt suppression (Wu et al., 2013). Akt signal transduction regulates central cell functions including metabolism, cell growth, transcriptional regulation, and cell survival (Zhang et al., 2011). Akt directly regulates apoptosis and promotes survival by targeting pro-apoptotic Bcl-2 related proteins, Forkhead factors (FOX), CREB, and p53 activity (Song, Ouyang & Bao, 2005; Zhang et al., 2011). As such, Akt has a key regulatory function to suppress apoptosis during dormancy. Indeed, a previous study of O. lactea reported elevated levels of Akt that in turn led to reduced levels of BAD and FOXO which acted to promote anti-apoptotic processes during estivation (Ramnanan, Groom & Storey, 2007). The role of miR-153 in stress survival has also been documented in the sea cucumber in response to hypoxia, indicating that miR-153 may also exert conserved functions in invertebrates (Zhao et al., 2014). Furthermore, miR-124 is known to relieve the suppression of CREB, a transcription factor responsible for mediating Akt-induced expression of anti-apoptotic factors (i.e., Bcl-2 and Mcl-1) (Du & Montminy, 1998; Song, Ouyang & Bao, 2005). As such, the upregulation of miR-124 in O. lactea may further promote survival by enhancing anti-apoptotic responses. Members of the miR-124 family were also upregulated in hibernating bats and thirteen-lined ground squirrels, suggesting a conserved function of these miRNAs in facilitating hypometabolism (Yuan et al., 2015; Wu et al., 2016). Lastly, although the exact function of miR-190, another miRNA targeting Akt members, is not fully understood, it has been shown to suppress TGFβ and FOXP2, two pro-apoptotic factors (Myatt & Lam, 2007; Ramesh, Wildey & Hoyeck et al. (2019), PeerJ, DOI 10.7717/peerj.6515, 5/13
Figure 2  Differentially expressed miRNAs during estivation. Changes in the relative abundance of 26 miRNAs in the foot muscle of estivating snails, *O. lactea*, as determined by RT-qPCR. MiRNA abundance was standardized against the 5S rRNA reference gene. Data are means ± SEM, *n* = 3 – 4 biological replicates. Statistical analysis was performed using a Student *t*-test and expression levels significantly different from control (*p* < 0.05) are indicated by *.

Howe, 2011; Gennarino et al., 2012; Jia et al., 2016). Dormancy is associated with multiple potential intracellular stresses (e.g., oxidative stress, accumulation of waste products, protein damage, etc.) that can lead to metabolic injury and even cell death (Storey & Storey, 2007). As such, stress-tolerant animals rely on anti-apoptotic processes to prolong survival, as seen in freeze-tolerant wood frogs, and hibernating ground squirrels and bats (Chen et al., 2008; Rouble et al., 2013; Gerber, Wijenayake & Storey, 2016). Results from the present study suggest that the milk snail induces the expression of anti-apoptotic miRNAs during estivation to prolong survival.

One of the most energy-expensive activities of an organism is the proliferation of cells to support growth and development, or replacement of damaged cells (Mazia, 1961; Storey & Storey, 2004). As such, cell cycle arrest is a central energy-conserving strategy employed by most species entering a hypometabolic state including freeze-tolerant frogs, hibernating squirrels, and anoxia-tolerant turtles (Biggar & Storey, 2012; Wu & Storey, 2012; Zhang & Storey, 2012). Herein, two tumour-suppressing miRNAs, miR-7 and miR-219, were upregulated in *O. lactea* foot muscle during estivation. MiR-7 is an evolutionarily conserved
miRNA that suppresses cell growth and proliferation via numerous mechanisms such as targeting Akt and Erk1/2 (Kalinowski et al., 2014). Likewise, miR-219 inhibits Erk1/2 signaling by potentially targeting the receptor tyrosine kinases EBB3 (epidermal growth factor receptor family), although this finding requires further analysis (Lei et al., 2013). The Erk signal transduction cascade regulates various growth and proliferation cell processes including cell cycle progression, cell survival, differentiation, metabolism, transcription, amongst others (Roskoski, 2012). As such, the upregulation of miR-7 and miR-219 during estivation suggests that O. lactea enhances long-term survival via energy reprioritization by suppressing cell cycle progression and cell proliferation.

Despite long periods of inactivity during dormancy, the milk snail displays no signs of muscle disuse atrophy and has been shown to regain muscle activity within minutes of arousal (Storey, 2002). The snail’s reliance on its foot muscle to attach itself to its surroundings suggests that it has adaptive strategies in place to overcome the onset of foot muscle atrophy that would be detrimental to survival. In the present study, miR-133 was upregulated during estivation. MiR-133 is a muscle-specific miRNA that has been shown to regulate myosin genes (i.e., MEF and MyoD) and muscle signaling pathways (i.e., MAPK and Erk1/2), thereby protecting against atrophy and promoting muscle differentiation (Biggar & Storey, 2011; Feng et al., 2013). Both MEF2 and MyoD are well-characterized muscle transcription factors that regulate muscle plasticity, differentiation, proliferation, cell survival, and apoptosis (Tessier & Storey, 2016), and together these factors have been implicated in stress survival in numerous hypometabolic species (Black & Olson, 1998; Hadj-Moussa et al., 2016; Tessier & Storey, 2016; Hoyeck, Hadj-Moussa & Storey, 2017). As such, the estivation-induced upregulation of miR-133 alludes to its role in preserving muscle function, suppressing cell cycle progression, and promoting survival in O. lactea. These findings are supported by previous discoveries of miR-133 upregulation in response to hypoxia in the jumbo squid and following freezing and anoxia in L. littorea, suggesting that miR-133 may provide a conserved mechanism for maintaining muscle function during hypometabolism (Biggar et al., 2012; Hadj-Moussa et al., 2018).

Our knowledge of miRNA functions in non-model species is far from complete. Consequently, the function of several miRNAs that showed upregulation in this study remains unknown. To our knowledge, this is the first report suggesting that miR-87, -281, -723, -745, -1542, -1989, -2176, -6833 and -7428 may be implicated in stress survival. Only one report on osteoarthritis suggests a role for miR-6833 in cell proliferation, metabolism, homeostasis, and response stimuli (Balaskas et al., 2017). Likewise, the function of miR-2001 has not been elucidated, although it has been associated with the stress response in marine lifeforms, including the squid D. gigas (Zhou et al., 2015; Hadj-Moussa et al., 2018). Lastly, very limited information is available on miR-12, -1175, or -1985, with only one reported finding of differential expression of these miRNAs in response to hypoxia in D. gigas, suggesting a potential role in invertebrate (or possibly only mollusc) hypometabolic regulation (Hadj-Moussa et al., 2018).
CONCLUSION

This study is the first to explore the stress-induced regulation of miRNAs in an estivating land snail. All estivation-responsive miRNAs identified in this study were upregulated, suggesting a role for miRNAs in energy reprioritization for vital processes by reducing mRNA translation in the hypometabolic state. Our findings indicated that estivation-induced miRNAs are implicated in anti-apoptotic, tumour suppressive, and muscle maintenance responses. Many of our findings follow similar expression patterns reported in other hypometabolic models, suggesting that miRNAs serve a conserved role necessary for survival during prolonged dormancy. Our present study suggests that miRNAs regulate these survival mechanisms by targeting the Akt and Erk1/2 signaling pathways, as well as myosin genes. Overall, this study deepens our understating of the molecular mechanisms involved in transitioning to and from a hypometabolic state and highlights conserved miRNA-mediated invertebrate and stress-responsive adaptations.

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ADDITIONAL INFORMATION AND DECLARATIONS

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Competing Interests

Kenneth B. Storey is an Academic Editor for PeerJ.

Author Contributions

- Myriam P. Hoyeck performed the experiments, analyzed the data, prepared figures and/or tables, authored or reviewed drafts of the paper, approved the final draft.
- Hanane Hadj-Moussa conceived and designed the experiments, performed the experiments, analyzed the data, authored or reviewed drafts of the paper, approved the final draft.
• Kenneth B. Storey conceived and designed the experiments, contributed reagents/materials/analysis tools, approved the final draft.

Data Availability
The following information was supplied regarding data availability:

The raw data are available in a Supplemental File.

Supplemental Information
Supplemental information for this article can be found online at http://dx.doi.org/10.7717/peerj.6515#supplemental-information.

REFERENCES

Balaskas P, Goljanek-Whysall K, Clegg P, Fang Y, Cremers A, Emans P, Welting T, Peffers M. 2017. MicroRNA profiling in cartilage ageing. International Journal of Genomics 2017(2):1–11 DOI 10.1155/2017/2713725.

Biggar KK, Kornfeld SF, Maistrovski Y, Storey KB. 2012. MicroRNA regulation in extreme environments: differential expression of microRNAs in the intertidal snail Littorina littorea during extended periods of freezing and anoxia. Genomics, Proteomics & Bioinformatics 10:302–309 DOI 10.1016/j.gpb.2012.09.002.

Biggar KK, Storey KB. 2011. The emerging roles of microRNAs in the molecular responses of metabolic rate depression. Journal of Molecular Cell Biology 3:167–175 DOI 10.1093/jmcb/mjq045.

Biggar KK, Storey KB. 2012. Evidence for cell cycle suppression and microRNA regulation of cyclin D1 during anoxia exposure in turtles. Cell Cycle 11:1705–1713 DOI 10.4161/cc.19790.

Biggar KK, Storey KB. 2018. Functional impact of microRNA regulation in models of extreme stress adaptation. Journal of Molecular Cell Biology 10(2):93–101 DOI 10.1093/jmcb/mjx053.

Biggar KK, Wu CW, Storey KB. 2014. High-throughput amplification of mature microRNAs in uncharacterized animal models using polyadenylated RNA and stem-loop reverse transcription polymerase chain reaction. Analytical Biochemistry 462:32–34 DOI 10.1016/j.jab.2014.05.032.

Black BL, Olson EN. 1998. Transcriptional control of muscle development by myocyte enhancer factor-2 (MEF2) proteins. Annual Review of Cell and Developmental Biology 14:167–196 DOI 10.1146/annurev.cellbio.14.1.167.

Brooks SPJ, Storey KB. 1990. Glycolytic enzyme binding and metabolic control in estivation and anoxia in the land snail Otala lactea. Journal of Experimental Biology 151:193–204.

Bustin SA, Benes V, Garson JA, Hellemans J, Huggett J, Kubista M, Mueller R, Nolan T, Pfafli MW, Shipley GL, Vandesompele J, Wittwer CT. 2009. The MIQE guidelines: minimum information for publication of quantitative real-time PCR experiments. Clinical Chemistry 55:611–622 DOI 10.1373/clinchem.2008.112797.
Chen H, Xin L, Song X, Wang L, Wang W, Liu Z, Zhang H, Wang L, Zhou Z, Qiu L, Song L. 2017. A norepinephrine-responsive miRNA directly promotes CgHSP90AA1 expression in oyster haemocytes during desiccation. *Fish and Shellfish Immunology* **64**:297–307 DOI 10.1016/j.fsi.2017.03.020.

Chen J, Yuan L, Sun M, Zhang L, Zhang S. 2008. Screening of hibernation-related genes in the brain of *Rhinolophus ferrumequinum* during hibernation. *Comparative Biochemistry and Physiology B: Biochemistry and Molecular Biology* **149**:388–393 DOI 10.1016/j.cbpb.2007.10.011.

Chen M, Storey KB. 2014. Large-scale identification and comparative analysis of miRNA expression profile in the respiratory tree of the sea cucumber *Apostichopus japonicus* during aestivation. *Marine Genomics* **13**:39–44 DOI 10.1016/j.margen.2014.01.002.

Chen M, Zhang X, Liu J, Storey KB. 2013. High-throughput sequencing reveals differential expression of miRNAs in intestine from sea cucumber during aestivation. *PLOS ONE* **8**:e76120 DOI 10.1371/journal.pone.0076120.

Churchill TA, Storey KB. 1989. Intermediary energy metabolism during dormancy and anoxia in the land snail *Ota latalecta*. *Physiological Zoology* **62**(5):1015–1030.

Du K, Montminy M. 1998. CREB is a regulatory target for the protein kinase Akt/PKB. *Journal of Biological Chemistry* **273**:32377–32379 DOI 10.1074/jbc.273.49.32377.

Dwivedi Y. 2011. Evidence demonstrating role of microRNAs in the etiopathology of major depression. *Journal of Chemical Neuroanatomy* **42**:142–156 DOI 10.1016/j.jchemneu.2011.04.002.

Feng Y, Niu LL, Wei W, Zhang WY, Li XY, Cao JH, Zhao SH. 2013. A feedback circuit between miR-133 and the ERK1/2 pathway involving an exquisite mechanism for regulating myoblast proliferation and differentiation. *Cell Death and Disease* **4**:e934 DOI 10.1038/cddis.2013.462.

Friedman RC, Farh KKH, Burge CB, Bartel DP. 2009. Most mammalian mRNAs are conserved targets of microRNAs. *Genome Research* **19**:92–105 DOI 10.1101/gr.082701.108.

Gennarino VA, D’Angelo G, Dharmalingam G, Fernandez S, Russolillo G, Sanges R, Mutarelli M, Belcastro V, Ballabio A, Verde P, Sardiello M, Banfi S. 2012. Identification of microRNA-regulated gene networks by expression analysis of target genes. *Genome Research* **22**:1163–1172 DOI 10.1101/gr.130435.111.

Gerber VEM, Wijenayake S, Storey KB. 2016. Anti-apoptotic response during anoxia and recovery in a freeze-tolerant wood frog (*Rana sylvatica*). *PeerJ* **4**:e1834 DOI 10.7717/peerj.1834.

Hadj-Moussa H, Logan SM, Seibel BA, Storey KB. 2018. Potential role for microRNA in regulating hypoxia-induced metabolic suppression in jumbo squids. *Biochimica et Biophysica Acta* **1861**:586–593 DOI 10.1016/j.bbagen.2018.04.007.

Hadj-Moussa H, Moggridge JA, Luu BE, Quintero-Galvis JF, Gaitán-Espitia JD, Nespolo RF, Storey KB. 2016. The hibernating South American marsupial, *Dromiciops gliroides*, displays torpor-sensitive microRNA expression patterns. *Scientific Reports* **6**:24627 DOI 10.1038/srep24627.
Hadj-Moussa H, Storey KB. 2018. Micromanaging freeze tolerance: the biogenesis and regulation of neuroprotective microRNAs in frozen brains. *Cellular and Molecular Life Sciences* 75:3635–3647 DOI 10.1007/s00018-018-2821-0.

Hoyeck MP, Hadj-Moussa H, Storey KB. 2017. The role of MEF2 transcription factors in dehydration and anoxia survival in *Rana sylvatica* skeletal muscle. *PeerJ* 5:e4014 DOI 10.7717/peerj.4014.

Jia W-Z, Yu T, An Q, Yang H, Zhang Z, Liu X, Xiao G. 2016. MicroRNA-190 regulates FOXP2 genes in human gastric cancer. *Onco Targets and Therapy* 6:3643–3651 DOI 10.2147/OTT.S103682.

Kalinowski FC, Brown RAM, Ganda KM, Epis MR, Horsham J, Leedman PJ. 2014. MicroRNA-7: a tumor suppressor miRNA with therapeutic potential. *International Journal of Biochemistry and Cell Biology* 54:312–317 DOI 10.1016/j.biocel.2014.05.040.

Lei H, Zou D, Li Z, Luo M, Dong L, Wang B, Yin H, Ma Y, Liu C, Wang F, Zhang J, Yu J, Li Y. 2013. MicroRNA-219-2-3p functions as a tumor suppressor in gastric cancer and is regulated by DNA methylation. *PLOS ONE* 8:e60369 DOI 10.1371/journal.pone.0060369.

Luu BE, Storey KB. 2015. Dehydration triggers differential microRNA expression in *Xenopus laevis* brain. *Gene* 573:64–69 DOI 10.1016/j.gene.2015.07.027.

Lyons PJ, Crapoulet N, Storey KB, Morin PJ. 2015. Identification and profiling of miRNAs in the freeze-avoiding gall moth Epiblema scudderiana via next-generation sequencing. *Molecular and Cellular Biochemistry* 410:155–163 DOI 10.1007/s11010-015-2547-3.

Lyons P, Poitra J, Courteau L, Storey K, Morin PJ. 2013. Identification of differentially regulated microRNAs in cold-hardy insects. *Cryo Letters* 34:83–89.

MacLean IA, Mattice AMS, Adam NJ, Storey KB. 2016. Purification and characterization of lactate dehydrogenase in the foot muscle and hepatopancreas of *Otala lactea*. *Protein Journal* 35:467–480 DOI 10.1007/s10930-016-9689-3.

Marco A, Hooks S, Griffiths-Jones S. 2012. Evolution and function of the extended miR-2 microRNA family. *RNA Biology* 9:242–248 DOI 10.4161/rna.19160.

Mazia D. 1961. Biochemistry of the dividing cell. *Annual Review of Biochemistry* 30:669–688 DOI 10.1146/annurev.bi.30.070161.003321.

Mazière P, Enright AJ. 2007. Prediction of microRNA targets. *Drug Discovery Today* 12:452–458 DOI 10.1016/j.drudis.2007.04.002.

Myatt SS, Lam EW-F. 2007. The emerging roles of forkhead box (Fox) proteins in cancer. *Nature Reviews* 7:847–859 DOI 10.1038/nrc2223.

Nilsen TW. 2007. Mechanisms of microRNA-mediated gene regulation in animal cells. *Trends Gene* 23:243–249 DOI 10.1016/j.tig.2007.02.011.

Pelissier F, Glogowski CM, Heinemann SF, Ballivet M, Ossipow V. 2006. Lab assembly of a low-cost, robust SYBR green buffer system for quantitative real-time polymerase chain reaction. *Analytical Biochemistry* 350:310–312 DOI 10.1016/j.ab.2005.12.002.
Ramesh S, Wildey GM, Howe PH. 2011. Transforming growth factor β (TGF β)-induced apoptosis. *Cell Cycle* 8:11–17 DOI 10.4161/cc.8.1.7291.

Ramnanan CJ, Allan ME, Groom AG, Storey KB. 2009. Regulation of global protein translation and protein degradation in aerobic dormancy. *Molecular and Cellular Biochemistry* 323:9–20 DOI 10.1007/s11010-008-9959-2.

Ramnanan CJ, Groom AG, Storey KB. 2007. Akt and its downstream targets play key roles in mediating dormancy in land snails. *Comparative Biochemistry and Physiology—B Biochemistry and Molecular Biology* 148:245–255 DOI 10.1016/j.cbpb.2007.06.002.

Ramnanan CJ, McMullen DC, Groom AG, Storey KB. 2010. The regulation of AMPK signaling in a natural state of profound metabolic rate depression. *Molecular and Cellular Biochemistry* 335:91–105 DOI 10.1007/s11010-009-0246-7.

Ramnanan CJ, Storey KB. 2006a. Suppression of Na+/K+-ATPase activity during estivation in the land snail *Otala lactea*. *Journal of Experimental Biology* 209:677–688 DOI 10.1242/jeb.02052.

Ramnanan CJ, Storey KB. 2006b. Glucose-6-phosphate dehydrogenase regulation during hypometabolism. *Biochemical and Biophysical Research Communications* 339:7–16 DOI 10.1016/j.bbrc.2005.10.036.

Roskoski R. 2012. ERK1/2 MAP kinases: structure, function, and regulation. *Pharmacological Research* 66:105–143 DOI 10.1016/j.phrs.2012.04.005.

Rouble AN, Hefler J, Mamady H, Storey KB, Tessier SN. 2013. Anti-apoptotic signaling as a cytoprotective mechanism in mammalian hibernation. *PeerJ* 1:e29 DOI 10.7717/peerj.29.

Schmittgen TD, Livak KJ. 2008. Analyzing real-time PCR data by the comparative CT method. *Nature Protocols* 3:1101–1108 DOI 10.1038/nprot.2008.73.

Song G, Ouyang G, Bao S. 2005. The activation of Akt/PKB signaling pathway and cell survival. *Journal of Cellular and Molecular Medicine* 9:59–71 DOI 10.1111/j.1582-4934.2005.tb00337.x.

Storey KB. 2002. Life in the slow lane: molecular mechanisms of estivation. *Comparative Biochemistry and Physiology Part A: Molecular & Integrative Physiology* 133:733–754 DOI 10.1016/S1095-6433(02)00206-4.

Storey KB. 2015. Regulation of hypometabolism: insights into epigenetic controls. *Journal of Experimental Biology* 218:150–159 DOI 10.1242/jeb.106369.

Storey KB, Storey JM. 2004. Metabolic rate depression in animals: transcriptional and translational controls. *Biological Reviews of the Cambridge Philosophical Society* 79:207–233 DOI 10.1017/S1464793103006195.

Storey KB, Storey JM. 2007. Tribute to P. L. Lutz: putting life on ‘pause’—molecular regulation of hypometabolism. *Journal of Experimental Biology* 210:1700–1714 DOI 10.1242/jeb.02716.

Storey KB, Storey JM. 2010. Metabolic rate depression. The biochemistry of mammalian hibernation. *Advances in Clinical Chemistry* 52:77–108 DOI 10.1016/S0065-2423(10)52003-1.
Storey KB, Storey JM. 2012. Aestivation: signaling and hypometabolism. Journal of Experimental Biology 215:1425–1433 DOI 10.1242/jeb.054403.

Tessier SN, Storey KB. 2016. Lessons from mammalian hibernators: molecular insights into striated muscle plasticity and remodeling. Biomolecular Concepts 7:69–92 DOI 10.1515/bmc-2015-0031.

Whitwam RE, Storey KB. 1990. Pyruvate kinase from the land snail Otala lactea: regulation by reversible phosphorylation during estivation and anoxia. Journal of Experimental Biology 154:321–337.

Wu CW, Biggar KK, Storey KB. 2013. Dehydration mediated microRNA response in the African clawed frog Xenopus laevis. Gene 529:269–275 DOI 10.1016/j.gene.2013.07.064.

Wu CW, Storey KB. 2012. Pattern of cellular quiescence over the hibernation cycle in liver of thirteen-lined ground squirrels. Cell Cycle 11:1714–1726 DOI 10.4161/cc.19799.

Wu C-W, Biggar KK, Luu BE, Szereszewski KE, Storey KB. 2016. Analysis of microRNA expression during the torpor-arousal cycle of a mammalian hibernator, the 13-lined ground squirrel. Physiological Genomics 48:388–396 DOI 10.1152/physiolgenomics.00005.2016.

Wu Z, He B, He J, Mao X. 2013. Upregulation of miR-153 promotes cell proliferation via downregulation of the PTEN tumor suppressor gene in human prostate cancer. The Prostate 73:596–604 DOI 10.1002/pros.22600.

Yuan L, Geiser F, Lin B, Sun H, Chen J, Zhang S. 2015. Down but not out: the role of MicroRNAs in hibernating bats. PLOS ONE 10:e0135064 DOI 10.1371/journal.pone.0135064.

Zhang J, Storey KB. 2012. Cell cycle regulation in the freeze-tolerant wood frog, Rana sylvatica. Cell Cycle 11:1727–1742 DOI 10.4161/cc.19880.

Zhao Y, Yang H, Storey KB, Chen M. 2014. Differential gene expression in the respiratory tree of the sea cucumber Apostichopus japonicus during aestivation. Marine Genomics 18:173–183 DOI 10.1016/j.margen.2014.07.001.

Zhao Y, He Y, Wang C, Zhang X. 2015. Characterization of miRNAs from hydrothermal vent shrimp Rimicaris exoculata. Marine Genomics 3:371–378 DOI 10.1016/j.margen.2015.09.004.

Zhou Z, Wang L, Song L, Liu R, Zhang H, Huang M, Chen H. 2014. The identification and characteristics of immune-related microRNAs in haemocytes of oyster Crassostrea gigas. PLOS ONE 9:e88397 DOI 10.1371/journal.pone.0088397.