Insights from the Den: How Hibernating Bears May Help Us Understand and Treat Human Disease

Maria Berg von Linde, M.S., Lilith Arevström, M.S., and Ole Fröbert, M.D., Ph.D.

Abstract
Hibernating brown bears (Ursus arctos) and black bears (Ursus americanus) spend half of the year in a physically inactive state inside their winter dens without food intake and defecating and no or little urination. Under similar extreme conditions, humans would suffer from loss of lean body mass, heart failure, thrombosis, azotemia, osteoporosis, and more. However, bears exit the den in the spring strong without organ injuries. Translational animal models are used in human medicine but traditional experimental animals have several shortcomings; thus, we believe that it is time to systematically explore new models. In this review paper, we describe physiological adaptations of hibernating bears and how similar adaptations in humans could theoretically alleviate medical conditions. The bear has solved most of the health challenges faced by humans, including heart and kidney disease, atherosclerosis and thrombosis, and muscle wasting and osteoporosis. Understanding and applying this library of information could lead to a number of major discoveries that could have implications for the understanding and treatment of human disease. Clin Trans Sci 2015; Volume 8: 601–605

Keywords: arteriosclerosis, heart rate, bone loss

Introduction
The use of experimental models in medical research typically adheres to the following chain of actions: (1) select a model organism, (2) induce disease in this model organism, (3) interpret the results, and (4) develop a treatment. The most frequently used model animals in research laboratories are rodents, especially mice and rats. However, these models have a number of shortcomings. Experimental mice and rats are inbred, hypertensive, glucose-intolerant, prone to cancer and kidney failure, and on a trajectory to premature death. Most often these animals are kept at room temperature, which is well below their thermoneutral zone (30–32°C for mice) that in combination with light and absence of hiding space cause constant physiological stress. Focusing on a few animal species confines the providence to those particular organisms, which could partly explain the decline in medical research breakthroughs that lead to major disease elimination.

In this paper, we use brown bears (Ursus arctos) and black bears (Ursus americanus) as experimental animal models as inspiration for understanding human medicine. The pursuit of learning from Nature’s solution is termed biomimicry—a discipline successfully exploited within engineering, architecture, and medicine.

Brown and black bears remain physically inactive inside their winter dens for half a year without eating, defecating, and with no or intermittent urination. Under similar conditions, humans would develop cardiovascular disease, kidney failure, muscle loss (sarcopenia), osteoporosis, and other deleterious conditions; however, bears readily exit their dens when spring arrives and show no signs of organ damage. In this review, we systematically discuss a number of key areas in which bear physiology might lead us to discover new ways to understand and treat human disease.

Methods
We executed the literature searches using medical subjects headings terms in the PubMed database and free text searches in Google Scholar. To frame the search systematically, we identified Population, Intervention, Comparative intervention, and Outcomes (PICO) in the examined articles. These PICO components defined the inclusion for our bibliographic search strategy, identifying research articles of relevance. We included research papers involving markers for heart failure, atherosclerosis, kidney failure, sarcopenia, and osteoporosis, which also showed relevance to the effects of hibernation in bears with comparisons to active bears or other mammals (including humans). Table 1 summarizes our literature search results.

Results
Down-regulated heart rate with respiratory sinus arrhythmia
Extreme heart rate reductions with profound respiratory sinus arrhythmia have been documented in hibernating bears (Figure 1). Respiratory sinus arrhythmia has dramatic manifestations, where the duration of the cardiac cycle rises up to three times longer than the cycle length in active state (cycle length variations of 20% would be normal in humans) and up to 13- to 14-second episodes of asystole have been reported in studies of free-ranging black bears (n = 37; n = 15). These effects are accompanied by extremely low respiratory rates. The change in heart rhythm is probably primarily mediated by cardiac vagal tone. The biological function of respiratory sinus arrhythmia is to maintain interplay between cardiovascular and respiratory systems in order to meet metabolic demands over highly variable conditions.

As a response to their low respiratory rate, bears have optimized oxygen transport during hibernation. Erythrocyte count and hemoglobin are increased during hibernation in captive brown bears (n = 7) and free-ranging black bears (n = 48). We recently documented that bears have higher oxygen affinity when hibernating compared to active state. This increased affinity is associated with a significant decrease in red cell hemoglobin-cofactor 2,3-diphosphoglycerate, which binds to the low-affinity conformation of mammalian hemoglobin and halves
hemoglobin-oxygen affinity compared to active state. These changes may be crucial for maintaining adequate tissue oxygen tension during hibernation.14

Protective cardiac gene expression
Humans suffering from bradycardia may develop cardiac chamber dilatation due to volume overload during the long diastolic pauses. Studies of captive grizzly bears (n = 4 [9]; n = 8 [8]) have revealed no significant reduction in ejection fraction during hibernation (although the left ventricular mass decreases), indicating sustained heart function.

One way to investigate how bear myocardium tolerates bradycardia is to study myocardial gene expression during hibernation compared to the active state. Such examinations have revealed significant alternations in the expressions of several heart-related genes. Several groups have investigated the genes titin (involved in muscle passive elasticity) and RNA-binding protein motif 3 (enhances protein synthesis at mildly hypothermic temperatures). Expression of the stiffer N2B isoform of titin is significantly elevated during hibernation, indicating increased ventricular passive rigidity,9 which could reduce the risk of chamber overload. This finding is supported by faster ventricular filling deceleration. A collaborative study has shown that phosphorylating titin in animal models with diastolic heart failure could restore diastolic function by reducing ventricular stiffness.9 RNA-binding protein motif 3 is expressed at a sevenfold higher level during hibernation.10 Elevated levels of this gene were also seen in hibernating ground squirrels.7 It has been suggested that RNA-binding protein motif 3 facilitates global protein synthesis under hypothermia by binding to 60S ribosomal subunits and reducing levels of microRNAs. Therefore, the protein biosynthesis of RNA-binding protein motif 3 may prevent myocardial atrophy and atrophy of skeletal muscle over an extended period of bradycardia and immobility as during hibernation.10

In hibernating bears, atrial strain imaging have showed decreased atrial contractility and increased atrial stiffness. The reportedly increased alpha-myosin heavy-chain protein expression in the atria (n = 14) may influence cardiac contraction characteristics. Atrium-specific expression of creatine kinase, consistent with lowered energy requirements, was reduced by 50% (n = 17), indicating depressed atrial emptying. These reported molecular changes of the atria would reduce ventricular overload and seem to be a protective response.9

Human heart failure is a clinical challenge that requires effective interventions. Therefore, the genes which seem to contribute to endured heart function in bears during hibernation could be suggested for therapeutic targets in experimental studies.

Resistance to atherosclerosis and thrombosis
Metabolism is mainly based on lipids from stored fat during hibernation leading to increases in plasma lipid levels. Most important, groups have noted increased plasma levels of low-density lipoprotein (“bad cholesterol”) and triglycerides in hibernating bears. In humans, long periods of bed rest and high plasma lipid concentrations are associated with atherosclerosis.22 However, no atherosclerotic histological changes—such as fatty streaks, cell infiltration, or inflammation—were detected in the coronary arteries and aortic arches of bears aged 1.5–12 years,21 thus suggesting atherosclerotic resistance (although

Table 1. Hibernating bear physiology and human health challenges. The human diseases of chamber dilatation, atherosclerosis, thrombosis, azotemia, muscle atrophy, osteoporosis are all disabilities that hibernating bears manage to avoid. The right column summarizes hypotheses that explain these scenarios.

| Hypotheses for why hibernating bears do not develop these diseases |
|---------------------------------|-----------------|---------------------------------|-----------------|---------------------------------|
| Bradycardia                     | Chamber dilatation |
| Changes in cardiac gene expression: upregulated isoform of titin (passive elasticity), RNA-binding protein motif 3 (protein synthesis) and myosin heavy-chain protein (contraction) |
| Physical inactivity and high cholesterol |
| Atherosclerosis and thrombosis |
| Positive selection for apolipoprotein B high-density lipoprotein (“good cholesterol”) |
| Lower platelet activity |
| Anuria |
| Azotemia |
| Urea recycled into amino acids |
| Unloaded muscles |
| Sarcopenia |
| Decreased expression of genes involved in the urea cycle |
| Increased expression of amino acid transporters in muscle Alterations in muscle catabolic pathway |
| Unloaded bones |
| Osteoporosis |
| Protective levels of serum markers |
| Black bear PTH increases bone volume fraction |

Table 1.

Figure 1. Heart rates in the hibernating and active states. Average heart rate values in brown and black bears during the hibernation and active states. The study of Laske TG included free-ranging black bears (n = 8), and that of Tøien Ø examined captive black bears (n = 4). The papers of Nelson OL (n = 4, n = 8) and Folk GE (n = 2) are based on samples from captive brown bears. The heart rates reported from the group of Tøien Ø were unexpectedly low compared to those of other studies. This could be related to the captive environment combined with the small sample size, since heart rate is strongly affected by activity.
atherosclerosis in bears of high age has been described in one report.23

Genome analyses of polar bears and brown bears have revealed positive selection for mutated apolipoprotein B (which encodes the primary lipoprotein component of low-density lipoprotein). This may explain how polar bears can cope with enhanced cholesterol without a build-up of arterial plaque, despite their hyper-lipid diet.21 However, brown bears have higher cholesterol levels than polar bears,21 which challenge this statement.21 The absence of atherosclerosis could alternatively be explained by the high levels of high-density lipoprotein (“good cholesterol”).

Immobilization is also associated with thrombus formation, for which atrial chamber hypokinesia is an additional risk factor. Compared to human controls, hibernating bears exhibit lower platelet activity (less than half), using either adenosine diphosphate, arachidonic acid, or thrombin receptor-activating peptide as agonists.26 This may explain how the bears endure hibernation without obvious thrombus building.

It is also possible that bears benefit from the immunosuppressive effect of reduced body temperature27 or the extreme consumptions of berries (Vaccinium), containing antioxidants such as vitamin C and polyphenols,28 which have shown favorable effects on high-density lipoprotein, blood pressure, and platelet function. A berry-rich diet may reduce cardiovascular risk factors for patients suffering from, for example, hyperlipidemia and hypertension.

### Suppressed glomerular filtration rate

The glomerular filtration rate (GFR; the flow rate of plasma filtered through the kidney) decreases by about 70% in hibernating bears, resulting in anuria with a very low urine production, which is reabsorbed across the urine bladder.29 Despite GFR suppression no clinical signs of azotemia are seen in hibernating bears. The serum urea levels remain low in free-ranging bears (n = 71),31 which could be explained by urea being recycled back into amino acids. Please see recent review on kidney function in bears for a detailed overview.32

### Limited muscle atrophy

Despite months of inactivity and starvation during hibernation, skeletal muscle mass54,54 and strength44 are not reduced to the same extents as observed in humans during bed rest. Free-ranging black bears (n = 7) and captive brown bears (n = 6) do not exhibit changes in muscle-fiber number and cross-sectional area between seasons.30,31 Studies of muscle biopsies and fiber type proportions of the myosin isoforms and fast- and slow-twitch muscle fibers between hibernating and active bears,34 which is associated with suppressed disuse atrophy. Limited muscle loss during hibernation has been explained by findings suggesting periodic muscle activity during dormancy among free-ranging black bears (n = 5).37 However, transection of nerves in hibernating captive brown bears (n = 8) revealed minimal change of muscle mass,34 which challenges the importance of muscle activity.

Measurements of protein content and nitrogen stable isotope enrichment show that protein catabolism in free-ranging black bears (n = 12) is lower in winter than in summer.35 Plasma from hibernating bears incubated in the presence of rat skeletal muscle showed down-regulation of the proteolytic rate by 40% compared to control plasma. Therefore, it is conceivable that a proteolytic inhibitor could influence pathways depressing muscle wasting during immobilization.40 Moreover, expressions of genes involved in the urea cycle are depressed in the hibernating state in parallel with muscle overexpression of amino acid transporters, suggesting facilitation of protein synthesis in captive black bears (n = 11). These results suggest that amino acids are redirected from catabolic pathways to protein biosynthesis.40

Further investigation of muscle responses to hibernation could be helpful for treating people suffering from immobilization or neuromuscular diseases. In order to influence the proteolytic pathway as during hibernation, peroxisome proliferator-activated receptor-γ coactivator (PGC-1α; stimulates mitochondrial biogenesis and regulates muscle fiber type formation) could be a potential target. In a study made on ground squirrels in torpor, the animals exhibited a shift to slow-twitch muscle fibers accompanied by up-regulated activation of the endurance exercise pathway mediated by PGC-1α.41

### Bone remodeling

In humans, long-term bed rest leads to unbalanced bone remodeling and skeletal changes leading to disuse osteoporosis.42 Despite months of disuse, hibernating free-ranging black bears (n = 65) exhibit no loss of cortical bone geometry or mechanical properties.52,54 Bone loss can also be assessed by measuring bone metabolism markers. Studies measuring various serum biochemical markers have identified several parameters that significantly differ between hibernation and the active state (Table 2).41–50 Interestingly, hibernating bears seem to retain a consistent calcium level during hibernation,39,50 even though bone turnover marker levels suggest unbalanced bone remodeling. Serum concentrations of PTH remain consistent year-round,47,48,50 however, amino acid residue differences exist between human PTH and black bear PTH and induced cAMP activation of black bear PTH has been found in osteoblast cultures.51 These suggest that black bear PTH could be a potent osteoanabolic agent, important for maintaining bone mass during hibernation. Subcutaneous injection of black bear PTH in mice with dystrophin deficiency (n = 20) resulted in a sevenfold increase in bone volume fraction, as well as expanded osteoblast area and down-regulated osteoclast area.51

Clarification of the impact hierarchy of the altered bone serum markers during hibernation could be a step in the direction of developing new tools to prevent and treat humans with osteoporosis. Some osteoanabolic factors, such as calcitonin52 and PTH,53 are in use in osteoporosis treatment. However, there is room for more efficient treatments and a possibility would be to explore the potential of biochemical markers found in bears. The finding of multipotent stem cells with osteogenic capacity has raised interest for their role in bone regeneration. Adipose tissue-derived stem cells from free-ranging brown bears (n = 6) showed an inclination to undergo spontaneous and unstimulated osteogenesis and chondrogenesis in cultures, indicating an advantageous differentiation potential. Further research in osteoporosis could be meaningful with the aim to reestablish normal bone regenerative mechanisms.

### Discussion

#### Similarities between human conditions and hibernation

Some human conditions are associated with physiological changes quite similar to those found in hibernating bears. Human
myocardial hibernation is defined as reversible heart dysfunction during chronic ischemia, whereas myocardial stunning is characterized by prolonged reduced contractile function after reperfusion of severe ischemia. The underlying pathophysiology is controversial with only limited data available in humans. Some argue that myocardial hibernation is an adaptive response in which cardiac function and metabolism are down-regulated to cope with the decreased energy supply during hypoperfusion, possibly mediated by decreased β-adrenergocceptor density. This cardiac adaption has similarities to the extreme bradycardia during respiratory sinus arrhythmia and reduced metabolic demands in hibernating bears.

During accidental severe hypothermia (less than 28°C), humans react with reduced heart rate, respiratory rate, blood pressure, and kidney function. Humans exposed to severe cold instinctively try to reduce heat loss by seeking some kind of cover. This is probably partly an autonomous mechanism associated with primitive and burrowing-like behavior to avoid a cold environment, just like what is observed in hibernating animals.

Winter depression and hypersomnia (prolonged sleep) are characterized by hypometabolism and decreased heart rate. These conditions could be secondary to an up-regulated parasympathetic tone as a response to winter onset, just like in hibernation.

Strengths and limitations

Most of the studies included in the present review used a prospective study design, in which every bear served as its own control. This method enabled an optimized comparison by eliminating bias based on individual characteristics (genetics, weight, activity level, etc.). Several of the reviewed studies were limited by relatively small sample sizes and high variations within the population (sex, age, and pregnancy). Furthermore, we chose to include studies with both free-ranging and captive bears and thus, differences in diet and hibernation patterns—as well as bears from different geographic origins representing different climate zones and nutritional availability. This inclusion increases the variation between populations but also broadens the overall picture.

Conclusion

Hibernating bears have developed several survival strategies, which show promising implications for patients suffering a number of different diseases, including heart failure, atherosclerosis, kidney dysfunction, muscle wasting, and osteoporosis. Considering the limitations of traditional rodent animal models, a systematic experimental and clinical research approach based on bear physiology is warranted in order to investigate the possibility of biomimicry and isolating protective biochemical markers and genes for future patient treatments.

Acknowledgments

We received no specific funding for this paper.

References


