

A quantitative, theoretical framework for understanding mammalian sleep

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Sleep is one of the most noticeable and widespread phenomena occurring in multicellular animals. Nevertheless, no consensus for a theory of its origins has emerged. In particular, no explicit, quantitative theory exists that elucidates or distinguishes between the myriad hypotheses proposed for sleep. Here, we develop a general, quantitative theory for mammalian sleep that relates many of its fundamental parameters to metabolic rate and body size. Several mechanisms suggested for the function of sleep can be placed in this framework, e.g., cellular repair of damage caused by metabolic processes as well as cortical reorganization to process sensory input. Our theory leads to predictions for sleep time, sleep cycle time, and rapid eye movement time as functions of body and brain mass, and it explains, for example, why mice sleep ≈ 14 hours per day relative to the 3.5 hours per day that elephants sleep. Data for 96 species of mammals, spanning six orders of magnitude in body size, are consistent with these predictions and provide strong evidence that time scales for sleep are set by the brain's, not the whole-body, metabolic rate.

allometric scaling | brain | cellular repair | metabolic rate | sleep times

In contrast to other obvious physiological phenomena such as eating, breathing, and walking, neither the function nor the mechanism by which sleep occurs is well understood, despite its ubiquity (1). Indeed, the quest for a fundamental theory of sleep is considered one of the most important, unsolved problems in science (2–7). Recent neurobiological studies have made great advances in understanding the mechanisms involved in sleep. Hormones, cells, and enzymes whose levels of activity and expression vary during sleep and between sleeping and waking states have been identified (2, 3, 8–10). Although these studies have been unable to determine the purpose of sleep, such investigations will play an increasingly important role in determining what processes are specific to sleep.

Among the most studied and best known hypotheses for the purpose of sleep are the following: (i) rest for the body or brain and prevention from overheating (11–13); (ii) cortical reorganization and processing associated with memory and learning (14–17); and (iii) cellular repair in the body or brain (3, 18–21, 54). Some hypotheses for sleep are easy to eliminate. For example, hypothesis *i* has been rejected because the energy saved during sleep is minimal, amounting to approximately one frankfurter bun worth of calories per night for humans (11, 12). Moreover, because rates of heating and cooling are set by mass-specific metabolic rate and body size, which should scale identically, equating the heat increase that leads to overheating with the heat lost to return to normal temperatures predicts that the ratio of sleep time to awake time is invariant with respect to body size. This is clearly counter to the observation that sleep times decrease with body size.

Distinguishing among other theories of sleep (e.g., hypotheses *i* and *ii*), however, has been much more difficult. Here, we develop a quantitative theory whose general structure can, in principle, be used to test any of these hypotheses for the function of sleep, particularly when combined with interspecific analyses of mammalian sleep data. Recently, Siegel (21) noted that such data could help illuminate and distinguish between different theories of sleep,

and he specifically discussed empirical data in relation to sleep as a process of neuronal repair (hypothesis *iii*).

Our starting point is the observation that all of these suggested underlying processes are related to metabolic rate. By specifying how each of these processes (hypotheses *i–iii*) is related to metabolic rate, we can derive allometric scaling relationships for several important sleep rates and times, thereby predicting, for example, how total sleep time, sleep cycle time, and the fraction of rapid eye movement (REM) sleep scale with body mass, M . These scaling relationships are usually some combination of simple power laws, $\approx aM^p$, where a is the scaling coefficient and p is the scaling exponent. As we demonstrate, the scaling exponent, p , alone is sufficient to distinguish whether sleep times are set by the whole body's or the brain's metabolic rate and to discern whether hypotheses *ii* and *iii* are consistent with empirical data and, thus, are viable theories. Our quantitative theory makes explicit, quantitative predictions regarding scaling exponents, p , which are tested by using empirical data for organisms that span six orders of magnitude in body size. Our analyses show sleep times are set by the brain's, not the whole body's, metabolic rate and that hypotheses *ii* and *iii* are both consistent with empirical data.

To distinguish between hypothesis *ii* (reorganization) and hypothesis *iii* (repair) requires knowledge regarding the scaling coefficient, a . In the theory, this is related to specific physiological, most likely cellular, processes that are independent of body mass. Unfortunately, little or nothing is known about these, thereby limiting our current ability to make quantitative predictions regarding a . Consequently, experiments that measure the values, or possibly even the magnitudes, of the relevant physiological variables would allow us to further distinguish between reorganization and repair. In *Discussion*, we address some experiments that would need to be performed to achieve this.

In constructing our theory, we speculate that periods of cortical reorganization and/or repair of neuronal damage are induced primarily by activity during alert wakefulness. Because neurons typically are regenerated slowly or not at all during the lifetime of the organism, faithful repair of cellular damage is critical for maintaining the long-term integrity and functioning of the brain. Moreover, this extremely low turnover of neurons suggests that maintenance of the brain and processes such as learning involve changes within, or connections between, existing neurons, i.e., reorganization.

Cellular repair is especially crucial for the brain because, unlike most other organs and tissues, its cells are not continually being replaced. Dissipative energy is necessarily produced as a by-product

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Abbreviations: CI, confidence interval; OLS, ordinary least squares; REM, rapid eye movement.

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of metabolic processes and, inevitably, leads to cellular damage. Dissipation occurs both at the basic biochemical level, as, for example, in the production of free radicals, and at the capillary level in the circulatory system due to hemodynamic analogs of viscous drag forces. Experimental evidence that brain cell damage is caused by sleep deprivation (22, 23) has been reported, although other studies did not corroborate these findings (24, 25). Furthermore, the recent findings of Cirelli *et al.* (26) demonstrate that *Drosophila* Shaker mutants with reduced sleep time experience a concurrent and identical decrease in lifespan. Assuming all other factors such as metabolic rate are held constant (consistent with measures of activity; ref. 26), these reduced sleep times result in increased periods of damage and reduced periods of repair. Thus, lifespan for these *Drosophila* Shaker mutants should be reduced, just as observed, suggesting that sleep indeed is linked to repair (26).

Studies have shown that sleep time is positively correlated to metabolic rate, body weight, brain weight, and other physiological variables (11, 13, 27). Because cellular damage is directly tied to metabolism and its repair is a leading hypothesis for sleep, we develop our theory with this scenario in mind, similar to ideas recently expressed by Siegel (21). However, as we explain, the framework's general structure can accommodate other mechanisms, most notably structural and functional reorganization.

According to these ideas, because smaller animals have proportionately higher metabolic rates per unit mass than larger animals, they require more sleep over a fixed period. Such qualitative predictions that agree with empirical observations are clearly useful, but much more detailed analyses and tests of hypotheses are possible once a quantitative theory has been developed. Below, explicit relationships among sleep, metabolic rate, and body size are derived. These provide a way to distinguish whether cellular repair during sleep occurs solely in the brain or throughout the entire body and whether cellular damage occurs at the same rate during sleep and wakefulness.

Theory

Mass Dependence of the Ratio of Sleep Time to Awake Time. We begin by assuming that metabolic processes during wakefulness cause most cellular damage and dictate the rate at which sensory input is processed. We also assume that cellular repair and/or neural reorganization occur primarily during sleep. These assumptions will be relaxed below. Dissipative metabolic energy, e.g., viscous-type forces in metabolite transportation networks, including the exchange of oxygen from hemoglobin to tissue, occurs primarily in network terminal units (capillaries and mitochondria), which are approximately constant in size and do not vary appreciably with body mass (28). This invariance implies that both the total rate of metabolic energy dissipation and the metabolic rate itself, B , scale linearly with the number of such terminal units. Consequently, the total rate of cellular damage caused by energy dissipation is a mass-independent fraction, f , of metabolic rate, B . This invariant fraction is calculable at the capillary level (28). At the mitochondrial level, it also holds because mammals share a common biochemistry of metabolism, and the stoichiometry generates a predictable quantity of energy with free radicals as byproducts.

The rate at which metabolic energy is dissipated in an average cell therefore is fB_c , where B_c is the average *in vivo* cellular metabolic rate. The total metabolic rate of the organ, tissue, or whole body being considered is $B = N_c B_c$, where N_c is the corresponding total number of cells. During sleep, the total amount of energy causing damage that needs to be repaired therefore is given by fBt_A , where t_A is total time awake (alert wakefulness) per day, and the number of damaged units is given by $N_d = fBt_A/\Delta E_d$, where ΔE_d is the average energy per damage event. Further, we assume that the process of cellular repair is similar across mammals and occurs at a subcellular level, so that the power density (per unit volume) required for repair, P_R , is independent of body size (see *Appendix* for a more detailed discussion). Thus, the number of repairs is

$N_R = P_R v_c N_c t_S / \Delta E_R$, where v_c is the average cell volume, t_S is sleep time per day, and ΔE_R is the average energy required for a single repair. For faithful repair, all damage must be repaired, so $N_d \approx N_R$. Combining this with our previous relationships gives $fBt_A \approx \varepsilon P_R v_c N_c t_S$, where $\varepsilon = \Delta E_d / \Delta E_R$. Because ΔE_d and ΔE_R are energies (not rates) that presumably occur at subcellular levels, ε is independent of body size. (Note that our conclusions hold so long as $\varepsilon \propto P_R^{-1}$.) The total volume (of the organ, tissue, or whole body) considered is $V = N_c v_c$, leading to

$$\frac{t_S}{t_A} \approx \left(\frac{\rho f}{\varepsilon P_R} \right) \frac{B}{M} \quad [1]$$

where M is the corresponding total mass, and $\rho \equiv M/V$, the tissue density, which is a constant. Eq. 1 can be expressed in terms of purely cellular quantities as $t_S/t_A \approx (\rho f/\varepsilon P_R) B_c/m_c$, where m_c is average cell mass. If neural reorganization is the origin of sleep, Eq. 1 still holds but with P_R now interpreted as the power density required for performing such an activity, f the fraction of metabolic energy used for processing information, and ε the measure of efficiency in neural reorganization. In either case, the quantity related to metabolic rate is neither sleep nor awake time but rather, a ratio of the two.

The scaling of whole-body metabolic rate with body size, $B = B_0 M^{3/4}$, where B_0 is the normalization constant, has been shown to reflect general properties of resource distribution networks. These networks are assumed to be space filling and have invariant terminal units. Minimizing energy expenditure in the circulatory system leads to quarter-power allometric scaling for many physiological rates and times (28). Organs such as the brain, which are supplied by major arteries, behave as nearly autonomous subunits and can effectively be treated as independent systems subject to the constraints of the theory. Their metabolic rate, B_i , therefore is predicted to scale approximately as $M_i^{3/4}$, where M_i is the organ mass. If organ mass itself scales with total body mass as $M_i \propto M^{a_i}$, then the metabolic rate of the organ scales with body mass as $B_i \propto M^{(3/4)a_i}$. From this, it immediately follows that the mass-specific metabolic rate of the organ scales with body mass as $B_i/M_i \propto M_i^{-1/4} \propto M^{-p_i}$, where $p_i \equiv (1/4)a_i$. The mass of most organs scales approximately linearly with body mass $a_i \approx 1$, so mass-specific metabolic rate scales as a typical physiological rate, $B_i/M_i \propto M^{-1/4}$. Brains, however, are exceptional. Reported empirical values for a_b (b denotes brain) range from 0.65 ($\approx 2/3$) to 0.76 ($\approx 3/4$) (29–31). Consequently, $B_b/M_b \propto M_b^{-p_b}$ with p_b ranging from $\approx 1/6$ to $3/16$. Unfortunately, no *in vivo* data are available to test this. However, the limited available data for rates of oxygen consumption in brain tissue *in vitro* are not inconsistent with these predictions (see *Appendix*).

Thus, if the primary purpose of sleep is to repair damage done to brain cells, then

$$\frac{t_S}{t_A} \propto M_b^{-1/4} \propto M^{-p_b}, \quad [2]$$

with p_b in the range 0.16–0.19. This equation explicitly predicts the exponent for how sleep time scales allometrically with body and brain size. Note, however, that the absolute value of t_S/t_A could be predicted from Eq. 1 if repair and dissipative rates (P_R and f) were known. The differences in interpretation between repair and reorganization correspond to different values for the mass-independent constants in Eq. 1. Thus, if the values, or possibly even the magnitudes of P_R , f , and ε are known, they can be used to distinguish between reorganization (hypothesis *ii*) and repair (hypothesis *iii*) as the function of sleep. It is important to realize that both processes may equally contribute to the function of sleep and that also would be revealed by such a quantitative analysis. Moreover, if the primary purpose were to repair damage to organs other than the brain, then the exponent in Eq. 2 would be $-1/4$ rather than $-1/6$ or $-3/16$.

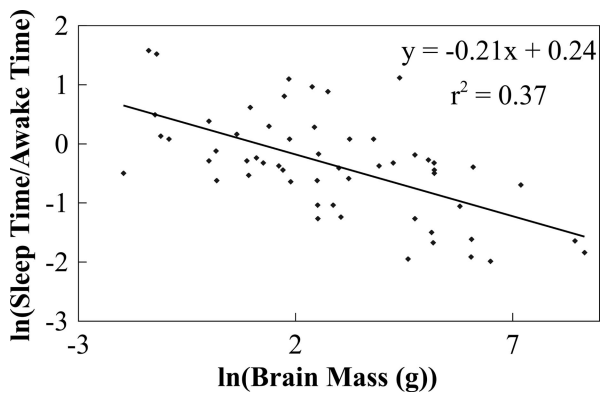


Fig. 2. Plot of the logarithm of the ratio of total sleep time, t_s , to total awake time per day, t_a , versus the logarithm of brain mass, $\ln(M_b)$, in grams. The slope computed by using OLS regression is -0.21 ($P < 0.0001$; $n = 56$, 95% CI: -0.28 , -0.14). Note that the 95% CI includes the predicted value of $-1/4$, providing more evidence that sleep is driven primarily by the brain's metabolic rate.

time is set directly by the brain's metabolic rate. Finally, in Fig. 4, we show $\ln(R)$ versus $\ln(M)$, for 61 species of mammals, spanning five orders of magnitude. The mean value of R is 0.17, with a slope of -6×10^{-3} ($P = 0.74$; $n = 61$, 95% CI: -0.04 , 0.03), consistent with zero and REM sleep being a constant fraction of total sleep time, as predicted. See Elgar *et al.* (13) for additional evidence.

Discussion

Based on these and others results, we speculate that the process identified as sleep is a special state of the brain that is devoted primarily to the critical activities of repair and reorganization. This leads to the conclusion that other organs or tissues do not require an analogous special state because they can be repaired or reorganized during waking or resting periods without significantly interfering with "normal" functionality (5).

Because lifespan scales like a typical physiological time ($\approx M^{1/4}$ or slightly shallower) (30), whereas sleep times scale according to Eqs. 1–3 [$\approx M^{-1/6}/(1 + aM^{-1/6})$], smaller mammals spend less time asleep on average during a lifetime than larger ones. Conversely, larger animals spend a much smaller fraction of their lives asleep than smaller ones. These observations raise interesting questions regarding the possible ecological and evolutionary consequences of sleep (e.g., predator–prey effects) and the corresponding advantages and/or disadvantages it confers on different size mammals (27,

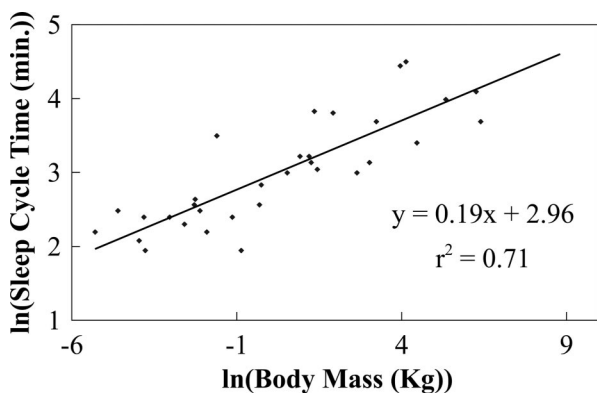


Fig. 3. Plot of the logarithm of sleep cycle time in minutes, the period between REM and non-REM sleep, versus the logarithm of body size, $\ln(M)$, in kilograms. The slope computed by using OLS regression is 0.19 ($P < 0.0001$; $n = 32$, 95% CI: 0.14 , 0.23). Note that the 95% CI includes the predicted values of $3/16$ and $1/6$ but excludes $1/4$ [Reproduced with permission from ref. 53 (Copyright 2006, Springer).].

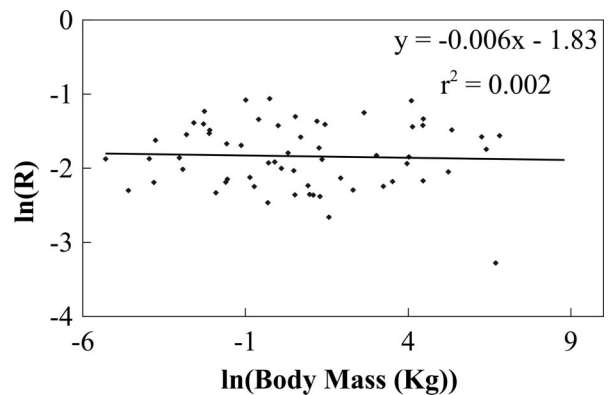


Fig. 4. Plot of the logarithm of the ratio of REM sleep time to total sleep time per day, $\ln(R)$, versus the logarithm of body size, $\ln(M)$, in kilograms. The slope computed by using OLS regression is $-6e-3$ ($P = 0.74$; $n = 61$, 95% CI: -0.04 , 0.03). The average value for R is 0.17. Note that the 95% CI for the slope is narrow and includes zero. The large outlier on the right side of the plot corresponds to the pilot whale (*G. scammoni*), a marine mammal.

34, 35). Indeed, the fraction of time spent asleep must eventually become limiting for small animals because some fraction of time is necessary just to forage and to reproduce (13, 34, 35).

For example, separating carnivores, herbivores, and omnivores suggests differences. Performing regressions of the sleep time to awake time ratio on body size, as in Fig. 1, for each of these groups gives the allometric exponents: -0.12 for carnivores ($P = 0.02$; $n = 27$, 95% CI: -0.22 , -0.03), -0.20 for herbivores ($P < 0.005$; $n = 26$, 95% CI: -0.20 , -0.13), and -0.10 for omnivores ($P = 0.03$; $n = 30$, 95% CI: -0.18 , -0.01). [Note that our results for carnivores differ from Siegel (21) because he excludes marine mammals.] Importantly, for all of these cases, the 95% CIs include our prediction. The variability, however, is quite large because the separation of these groups decreases both the number of data points and the mass range covered. Notably, the variability for herbivores is significantly less than that for carnivores or omnivores. This difference may have arisen because carnivores and omnivores experience less selective pressure, because of lower risks of predation and a higher calorie diet, to maximize awake time and, thus, to optimize the sleep time (21). Including such effects could enhance this theory.

Another natural extension of our analyses is to mammals during ontogenetic growth and to other taxa. This is particularly interesting for birds because their brain allometry and partial use of unihemispheric sleep distinguish them from most mammals (1, 2, 30), and for ectotherms because the temperature dependence of metabolic rate as well as the variation in cell and genome size adds other parameters to the theory (32). Unfortunately, in both cases there are, at present, insufficient data for definitive analyses.

We have presented a theory to address sleep-related questions in a well defined, general framework whose quantitative predictions can be compared directly with existing data. Our results demonstrate that there is substantial evidence that the function of sleep is related to metabolic processes in the brain and, in particular, to the repair of neuronal damage and/or reorganization (3, 18–21). The theory is able to explain sleep times that differ by a factor of seven across organisms that differ in mass by six orders of magnitude, revealing why a mouse sleeps four times longer per day than an elephant. Further work needs to be done to understand the detailed mechanisms underlying damage and repair and their relationship to metabolic rate and sleep. Our work suggests several avenues of related experimental investigation. These include (i) tests of whether the power density of cellular repair and/or neural organization is body mass independent, (ii) measurements of the fraction of metabolic energy given to damage or neural reorganization, and (iii) determination of the scaling of brain metabolic rate with body

size. Our results strongly suggest that metabolic processes in the brain control the underlying mechanisms for the function of sleep and are a major determinant of why sleep cycle time increases with body size and why mammals need the amounts of sleep they do.

Methods

All of the sleep data are from Zepelin (33), and methods are detailed therein. Data for body masses were not given by Zepelin (33), so most values for body mass were taken from Meddis (27), which is an earlier compilation that Zepelin (33) draws from heavily. When sleep measurements listed in Zepelin (33) were not given or did not match sleep data in Meddis (27), we used the masses given in Smith *et al.* (36), and when masses could not be found there, we used the average of the range of values given in Nowak (37). All of the data and the sources for the body mass values are listed in SI Table 1. Some of the original sources given in Zepelin (33) were consulted to determine which species were used. In a few cases, the logarithmic averages of body masses were calculated for groups of species (e.g., four species of *Microtus* and five species of *Peromyscus*). This was done to be consistent with the original sleep data in Zepelin (33). Data for brain mass were taken from Meddis (27), but two values were excluded because the sleep values given did not match those given in Zepelin (33).

Certain marine mammals sleep with one hemisphere of the brain at a time. Because in Eq. 1 the variable t_s is the amount of sleep time per cell or per tissue, total sleep time for these marine mammals must be divided by two to obtain the appropriate sleep time. [There is often an asymmetry between amounts of sleep for the left and right hemisphere in marine mammals, so dividing by two should be regarded as the average sleep per neuron (38).] The original sleep data from Zepelin (33) were adjusted accordingly for the three species: *Tursiops truncatus*, *Globiocephalus scammoni*, and *Phocoena phocoena*. Moreover, marine mammals are known to have very small or perhaps nonexistent amounts of REM sleep, and in accord with several other studies, the data for REM sleep for *T. truncatus* and *P. phocoena* were excluded (33). Data for the REM sleep of *G. scammoni* was included and corresponds to the large outlier in Fig. 3. Finally, REM sleep time and sleep cycle time for *Elephas maximus* were excluded because Zepelin (33) denotes those values as doubtful, and the REM sleep time of 0.0 for *Tachyglossus aculeatus* was excluded because recent studies have shown that differentiating between REM and non-REM sleep for this primitive species is especially difficult (39).

Allometric exponents were determined by using OLS regression on ln-ln plots of the data. CI (95%) and P values were computed by using Mathematica (Wolfram Research, Inc., Champaign, IL). It is most appropriate to perform regressions and calculate statistics in ln-ln space because that is the space in which the error for each data point is approximately the same, thus satisfying one of the central assumptions of linear regression. For example, within a species, the variance in body mass increases approximately linearly with the mean body mass, so that the relative error is approximately constant (≈ 0.23) (40). Consequently, in linear space, the error for each data point increases systematically with body mass (the x axis), but in ln-ln space, the error for each data point, which is the relative error of the linear values, is approximately constant and independent of the logarithm of body mass (the x axis). Similarly, the relative error of physiological rates and times, such as metabolic rates and sleep times, should be approximately constant, and, therefore, plots and regressions should be done in ln-ln space.

To test for the normality of the residuals in our plots, we used Matlab (The MathWorks, Natick, MA) to compute the Lilliefors (41) and Jarque-Bera (42) tests for our data. Both tests choose a normal distribution with unknown variance and mean as the null hypothesis. The Lilliefors test determines how much the cumulative distribution function for the residuals differs from that for a normal distribution, and the Jarque-Bera test determines how much the skew and kurtosis of the residuals differ

from that of a normal distribution (with a skew and kurtosis of zero). Because these tests depend on different characteristics of the data, we applied both tests to the residuals in our plots. Data often are considered normal if the null hypothesis of normality is not rejected for a significance level < 0.05 . For the Lilliefors test, normality was not rejected for a significance level of $\alpha_S < 0.16$ for the residuals in Fig. 1 and $\alpha_S < 0.13$ for the residuals in Fig. 2. Using Matlab, the maximum computable significance level for the Lilliefors test is 0.2, and within this bound, normality was not rejected for the residuals in Figs. 3 or 4. For the Jarque-Bera test, normality was not rejected for a significance level of $\alpha_S < 0.21$ for the residuals in Fig. 1, $\alpha_S < 0.38$ for the residuals in Fig. 2, $\alpha_S < 0.60$ for the residuals in Fig. 3, and $\alpha_S < 0.47$ for the residuals in Fig. 4. Thus, the residuals passed normality tests for all cases considered.

Appendix

Dependence of Power Density Given to Repair, P_R , on Body Size, M .

We assume that the power density (per unit volume) given to repair or reorganization, P_R , is constant. This is because we are assuming that the repair or reorganization processes are occurring locally, at a level significantly smaller than that of the cell, and because we assume that the entire cell, including cell walls, proteins, and the cytoskeleton, requires repair.

Depending on the exact mechanisms of cellular damage and repair, however, it may be most appropriate to focus on other quantities. For example, if DNA is the dominant site of damage and repair, it is most appropriate to consider the power given to repair per DNA content. If power given to repair per nucleotide or per codon is constant because of the limited spatial access to these domains, then it follows that power given to repair per DNA content is constant. For mammals, DNA content and glial cell size are roughly invariant, but brain neuronal cell size appears to vary as $M_b^{1/4}$ (43–47). Consequently, within this scenario, power given to repair per unit volume, P_R , would still be a constant for glial cells but would decrease as $M_b^{-1/4}$ for neuronal cells. Thus, if glial cells require the most repair, the theory is exactly the same as presented in *Theory*. However, if brain neuronal cells require the most repair during sleep, most repair at the cellular level is to DNA, and if power given to repair per DNA content is constant, then the mass dependence of P_R decreases as $M_b^{-1/4}$, so the ratio of sleep time to awake time would be predicted to be mass-independent. This prediction is in clear disagreement with data and suggests that this scenario is not correct. However, it does provide a potential means for either using our theory to support reorganization (hypothesis *ii*) as the dominant function of sleep over repair (hypothesis *iii*) or for falsifying our theory altogether. That is, if this mass dependence for P_R was supported by empirical findings at the cellular level, it would suggest that our theory for repair is incorrect and, depending on the analogous findings for neural reorganization, that the influence of metabolic rate on sleep time requires reevaluation. Alternatively, if B_b/M_b was constant, which contradicts our assumption and limited empirical data (see below), and P_R decreased as $M_b^{-1/4}$, the predictions of our theory would be exactly the same as presented in *Theory*, but the source of the mass dependence would come from the power density given to repair and not from the mass-specific metabolic rate of the brain. Measurements of P_R and p_b therefore are crucial for determining both the source of mass dependencies in our theory and for assessing the validity of our theory.

Within ectotherms, cell size and genome size vary over several orders of magnitude, and this variation may be related to mass-specific metabolic rate (48, 49). Therefore, to extend this framework to ectotherms, it will be crucial to discern between the different mechanisms given above. This problem is somewhat simplified by the fact that cell size and genome size appear to vary together linearly.

Because we do not know the exact nature of the repair occurring at the cellular level (either what is being repaired or the mechanism

by which it is repaired), we focus on the power per unit volume in *Theory* because its generality does not depend on a specific repair mechanism. As explained above, however, the framework can be modified to accommodate different and/or more complicated repair mechanisms.

In Vitro Data for Scaling Exponent of Brain Metabolic Rate with Body Mass, p_b . Qualitative support for a negative value for p_b is found in Elliott and Henderson (50), Davies (51), and Tower and Young (52). Elliott and Henderson (50) measured rates of oxygen consumption in cortical tissue for three species of mammals (rat, cat, and cattle). Their measurements were taken *in vitro* within 1 h of killing of the animals. They concluded that $p_b \approx 0.1$. Subsequent studies by Davies (51) concluded that *in vitro* measurements are usually an underestimate of *in vivo* values, but using a limited amount of *in vitro* data for rates of oxygen consumption in brain tissue, he obtained a value of $p_b \approx 0.07$. Later, Tower and Young (52) performed the same measurements as Elliott and Henderson (50) with inclusion of data for fin and sperm whale and obtained similar results. Tower and Young also used data for seven species to show that the activity of acetylcholinesterase in the dorsal cerebral cortex decreases with brain mass as $M_b^{-0.2}$. Although these findings for the scaling of oxygen consumption rates in the brain are based on very limited amounts of *in vitro* data (as opposed to the *in vivo* data needed to test directly this theory), the scaling exponents are not inconsistent with the theoretical values of $-1/6$ and $-3/16$.

Estimate of Fraction of Metabolic Rate Given to Damage During Sleep, α . We performed a linear plot for t_A/t_S versus $M^{-3/16}$ and versus $M^{-1/6}$. From Eq. 3 the intercept of these plots corresponds to α . Using OLS regression, we obtained the value $\alpha = 0.35$ ($P = 0.074$; $n = 83$, 95% CI: $-0.04, 0.75$) for $p_b = -3/16$ and $\alpha = 0.27$ ($P = 0.20$; $n = 83$, 95% CI: $-0.15, 0.70$) for $p_b = -1/6$. The 95% CI is large but suggests $\alpha < 1$. Note that OLS regression assumes the same error for each datum, which is not strictly satisfied in these plots (see *Methods*). The mass dependence for the standard deviation of each datum in this plot behaves approximately like $\sigma/M^{19/16}$ and $\sigma/M^{7/6}$, respectively, where σ is the standard deviation in mass for each species. The relative standard deviation, σ/M , is roughly constant for mammals (40), so the standard deviation for each datum in this plot slightly decreases with M . Consequently, the errors for α are even larger than the ones given here, but the data likely still suggest $\alpha < 1$.

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