

Universal scaling of production rates across mammalian lineages

Marcus J. Hamilton, Ana D. Davidson, Richard M. Sibly and James H. Brown

Proc. R. Soc. B published online 26 August 2010
doi: 10.1098/rspb.2010.1056

Supplementary data

["Data Supplement"](#)

<http://rsbp.royalsocietypublishing.org/content/suppl/2010/08/24/rspb.2010.1056.DC1.html>

References

[This article cites 30 articles, 6 of which can be accessed free](#)

<http://rsbp.royalsocietypublishing.org/content/early/2010/08/24/rspb.2010.1056.full.html#ref-list-1>

P<P

Published online 26 August 2010 in advance of the print journal.

Subject collections

Articles on similar topics can be found in the following collections

[ecology](#) (1690 articles)

Email alerting service

Receive free email alerts when new articles cite this article - sign up in the box at the top right-hand corner of the article or click [here](#)

Advance online articles have been peer reviewed and accepted for publication but have not yet appeared in the paper journal (edited, typeset versions may be posted when available prior to final publication). Advance online articles are citable and establish publication priority; they are indexed by PubMed from initial publication. Citations to Advance online articles must include the digital object identifier (DOIs) and date of initial publication.

To subscribe to *Proc. R. Soc. B* go to: <http://rsbp.royalsocietypublishing.org/subscriptions>

Universal scaling of production rates across mammalian lineages

Marcus J. Hamilton^{1,2,3,*}, Ana D. Davidson^{1,4}, Richard M. Sibly⁵
and James H. Brown^{1,3}

¹Department of Biology, and ²Department of Anthropology, University of New Mexico, Albuquerque, NM 87131, USA

³Santa Fe Institute, 1399 Hyde Park Road, Santa Fe, NM 87501, USA

⁴Instituto de Ecología, Universidad Nacional Autónoma de México, México, DF, México

⁵School of Biological Sciences, University of Reading, Reading, UK

Over many millions of years of independent evolution, placental, marsupial and monotreme mammals have diverged conspicuously in physiology, life history and reproductive ecology. The differences in life histories are particularly striking. Compared with placentals, marsupials exhibit shorter pregnancy, smaller size of offspring at birth and longer period of lactation in the pouch. Monotremes also exhibit short pregnancy, but incubate embryos in eggs, followed by a long period of post-hatching lactation. Using a large sample of mammalian species, we show that, remarkably, despite their very different life histories, the scaling of production rates is statistically indistinguishable across mammalian lineages. Apparently all mammals are subject to the same fundamental metabolic constraints on productivity, because they share similar body designs, vascular systems and costs of producing new tissue.

Keywords: production; life history; metabolic ecology; placental; marsupial; monotreme

1. INTRODUCTION

The three lineages of mammals, placentals (eutherians), marsupials (metatherians) and monotremes (prototherians), diverged more than 100 Ma ago [1–3]. They diversified rapidly on different landmasses as the supercontinent Pangaea fragmented and drifted apart [4–9]. As a consequence of their independent evolution, the extant members of these lineages differ conspicuously in physiology, life history and reproductive ecology [10–12]. Metabolic rates of marsupials and monotremes are about 33 and 50 per cent lower, respectively, than placentals [13], and the differences in life history are dramatic (figure 1). Monotremes lay eggs; the stored energy reserves support embryonic development, and then the females supply milk to fuel growth from hatching to independence. Marsupials have a very short period of intrauterine embryonic development, are born at a small size and undeveloped state and then are protected and lactated in a pouch for a long period before they become independent [14]. By contrast, placentals have a more prolonged period of intrauterine embryonic development, are born at larger size and are nourished by lactation until they become independent.

A common feature of these seemingly divergent life histories is that the mother supplies all of the nutrition to rear offspring to independence [15]. The metabolic rate of the mother fuels the production of biomass, and this limits the ways in which life-history traits can trade off to meet energy and material requirements of offspring at different stages of development. Because the metabolic

rate of the mother is highly constrained by body size (cf. [16–20]), many life-history traits correlate closely with adult body size, scaling allometrically following the form

$$R = R_1 M^\beta, \quad (1.1)$$

where R is the trait of interest, R_1 is a taxon- and mass-specific normalization constant, M is adult body size and β is a scaling exponent. Metabolic scaling theory predicts $\beta \simeq 3/4$ for whole-organism metabolic rates, $\beta \simeq 1/4$ for biological times such as duration of pregnancy and lactation and lifespan and $\beta \simeq -1/4$ for most biological rates such as ontogenetic and population growth rates and the rate of offspring production [16].

Mechanistically, rate of offspring production depends on the rate at which biomass can be produced by the mother. Productivity is constrained by the logistical problems of transporting key materials such as oxygen and nutrients around the body [21]. Because all mammals have similar body plans and energetic costs of producing biomass [22], we expect them to be subject to similar constraints on production. Mass-specific production rate, p , can be estimated as the mean mass of offspring produced per year, normalized by adult body size (or the annual amount of biomass produced per unit of adult body size), so

$$p = \frac{m_b}{m} \cdot l \cdot n, \quad (1.2)$$

where m is adult body mass, m_b is mass at birth, l is litter size and n is the number of litters per year [23]. In placental mammals, mass-specific production rate scales with an exponent between $-1/4$ and $-1/3$, similar to mass-specific metabolic rate [23]. There is considerable variation in production rate across different

* Author for correspondence (marcusj@unm.edu).

Electronic supplementary material is available at <http://dx.doi.org/10.1098/rsob.2010.1056> or via <http://rsob.royalsocietypublishing.org>.

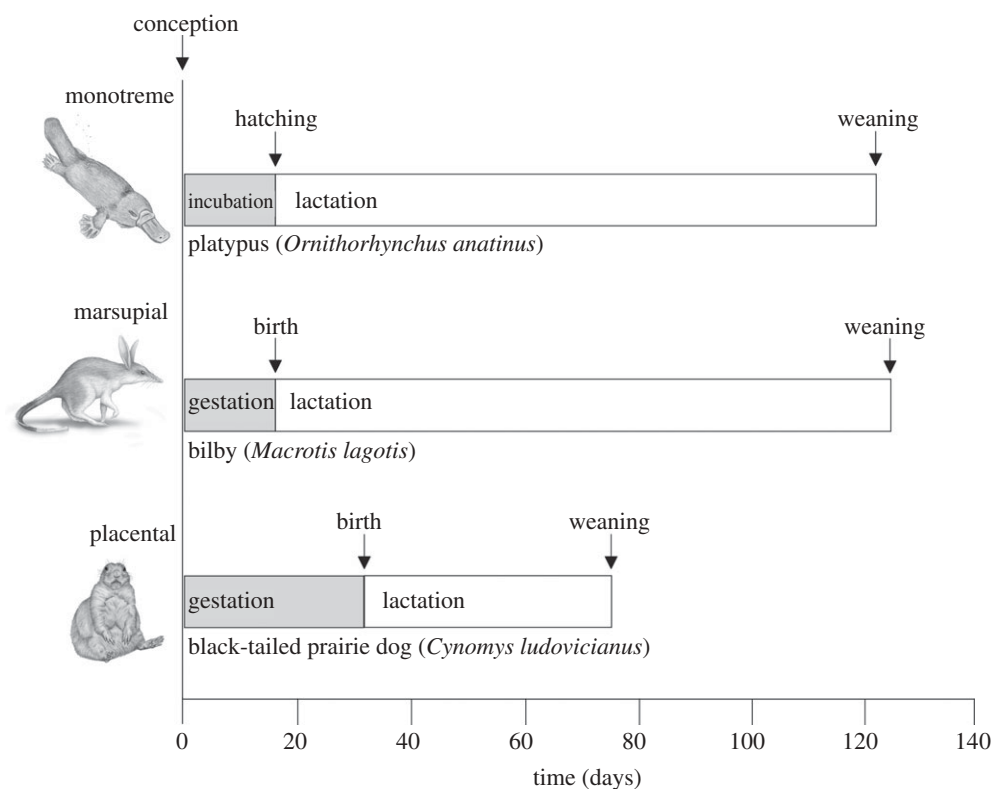


Figure 1. Schematic of differences in timing of the life history in similar-sized monotremes, marsupials and placentals. Female body mass *ca* 1500 g.

functional and taxonomic groups of eutherian mammals, but most of this variation is in the normalization constant, R_1 ; the exponents are relatively constant across these groups because production rates are highly constrained by body size according to equation (1.1) [23,24]. However, because placentals, marsupials and monotremes give birth to offspring of very different sizes and developmental stages, a better and more standardized measure of production substitutes mass at weaning, m_w , the stage at which mothers cease direct investment of energy to offspring growth, for mass at birth. Indeed, figure 2*a* shows that while size of offspring at birth scales predictably with adult body size in all lineages, newborn placentals are both absolutely larger and scale more steeply with adult body size than marsupials or monotremes. However, figure 2*b* shows that weaning mass is remarkably predictable across all lineages of mammals. So, mass-specific production measured at weaning is

$$p = \frac{m_w}{m} \cdot l \cdot n. \quad (1.3)$$

We address the effects of pre-weaning mortality on this measure of mass-specific production later in the discussion. Because all mammals have similar body plans, vascular systems and costs of producing new tissue, we hypothesize that all mammals should produce biomass at an equivalent mass-specific rate despite the differences between lineages in the size and timing of the life-history stages.

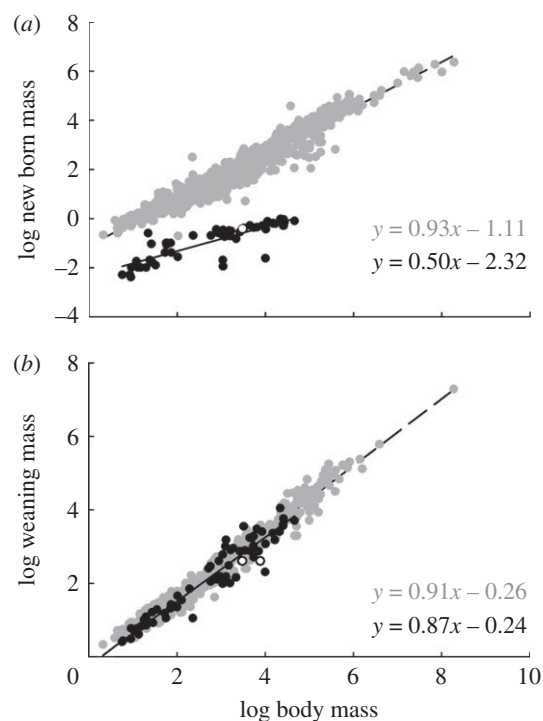


Figure 2. Bivariate log-log plots of offspring body mass (a) at birth and (b) newly weaned as a function of adult mass in grams. All logs are log-base 10, all mass units are grams and the figure legend is the same for all figures. Filled grey circles, placental; dashed lines, placental OLS; filled black circles, marsupial; solid lines, marsupial OLS; open circles, monotreme.

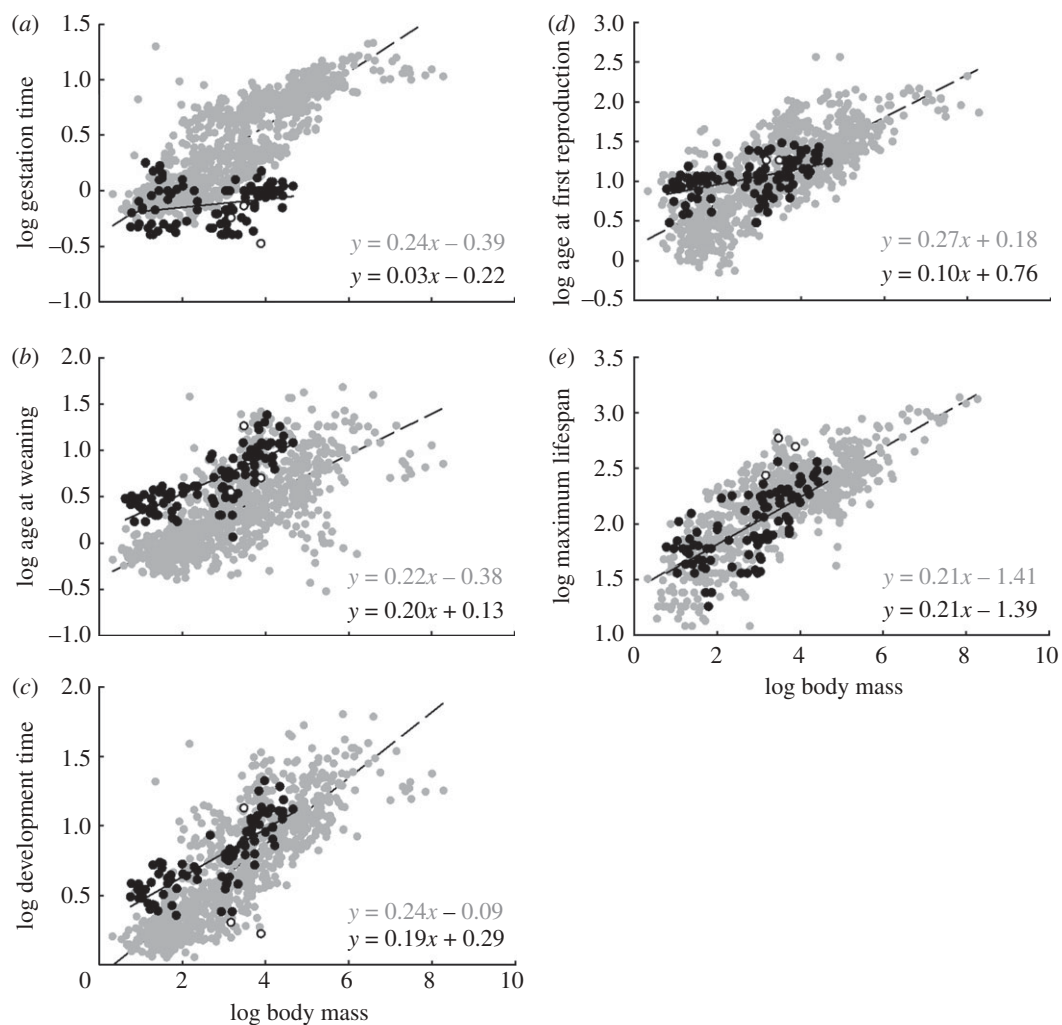


Figure 3. Bivariate log–log plots of life-history timings in months as a function of adult body mass: (a) gestation time; (b) age at weaning; (c) development time; (d) age at first reproduction; (e) maximum lifespan.

2. METHODS

To evaluate these predictions and explore variation in allocation to reproduction across placentals, marsupials and monotremes, we compiled a life-history database for more than 4000 non-volant mammalian species from both primary and secondary published sources (for more details, see electronic supplementary material, table S1). For each species, we recorded neonate mass (g), mass at weaning (g), adult body size (g), gestation time (months), time to weaning (months), age at first reproduction (months), maximum lifespan (months), litter size and litter frequency (per year), where data were available. For the placentals and marsupials, we used ordinary least squares (OLS) regression, fitted to bivariate log–log plots, to quantify allometric scaling of these life-history traits in terms of equation (1.1). For the mass-specific production analyses, we first used OLS regressions to estimate slopes within lineages, and then we used a general linear model (GLM) to test for significant differences between the lineages. For the GLMs between lineages, we controlled for taxonomy by nesting order within lineage because body size is confounded by phylogeny. Within the placental lineage, there are 20 orders represented in our dataset, and in the marsupial lineage, there are 7 orders represented. There were only three monotreme

species, and so we show their data in figures, but do not include them in statistical analyses. All statistical analyses were performed in MINITAB 15.0.

3. RESULTS

(a) Life-history times

Figure 3a–e shows variation in life-history times as a function of body mass, all plotted on logarithmic axes. Comparing marsupials and placentals: (i) gestation time is short and nearly invariant in marsupials, whereas it is substantially longer and increases significantly with body size in placentals, (ii) time to weaning is substantially longer in marsupials than in placentals, but scales similarly with size, (iii) total development time (the sum of gestation time and time to weaning) is longer in marsupials than in placentals, (iv) age at first reproduction is widely variable and overlaps greatly, but tends to be greater in small marsupials and to scale less steeply with body size than in placentals, and (v) maximum lifespan is similar across lineages and scales very similarly with mass (see table 1 for statistical results). Slopes generally are close to the predicted 1/4, except for gestation time and age of first reproduction in marsupials, which are shallower.

Table 1. Results of regressions of life-history variables and adult body mass.

trait	lineage	slope	s.e.	intercept	s.e.	d.f.	r^2	p
newborn mass	placentals	0.93	0.01	-1.11	0.03	888	0.95	0.00
	marsupials	0.50	0.04	-2.32	0.14	54	0.70	0.00
weaning mass	placentals	0.91	0.01	-0.26	0.03	391	0.97	0.00
	marsupials	0.87	0.03	-0.24	0.10	64	0.92	0.00
gestation time	placentals	0.24	0.00	-0.39	0.02	1098	0.73	0.00
	marsupials	0.04	0.02	-0.22	0.05	94	0.06	0.02
age at weaning	placentals	0.22	0.01	-0.38	0.02	925	0.52	0.00
	marsupials	0.20	0.01	0.13	0.04	109	0.63	0.00
development time	placentals	0.24	0.01	-0.08	0.02	853	0.70	0.00
	marsupials	0.17	0.01	0.29	0.04	83	0.66	0.00
age at first reproduction	placentals	0.27	0.01	0.18	0.03	886	0.57	0.00
max. lifespan	marsupials	0.10	0.02	0.76	0.06	101	0.22	0.00
litter size	placentals	0.21	0.01	1.41	0.02	737	0.65	0.00
	marsupials	0.21	0.02	1.39	0.07	93	0.51	0.00
litters per year	placentals	-0.13	0.00	0.74	0.01	1432	0.41	0.00
	marsupials	-0.27	0.02	1.09	0.05	167	0.57	0.00
annual fecundity	placentals	-0.12	0.01	0.52	0.02	869	0.39	0.00
	marsupials	-0.02	0.02	0.23	0.06	111	0.01	0.40
lifetime fecundity	placentals	-0.26	0.01	1.34	0.03	847	0.59	0.00
	marsupials	-0.28	0.03	1.33	0.08	110	0.47	0.00
	placentals	-0.02	0.01	1.24	0.03	530	0.01	0.10
	marsupials	-0.02	0.04	1.43	0.13	93	0.00	0.57

(b) Fecundity

Figure 4*a,b* shows variation in the different components of fecundity. Litter size (figure 4*a*) is highly variable and scales more steeply in marsupials than in placentals, and the number of litters per year (figure 4*b*) overlaps greatly between marsupials and placentals of small body size, but is approximately invariant with size in marsupials and scales negatively in placentals. We combine litter size and the number of litters per year to estimate annual fertility, F_Y , by rearranging equation (1.2) so that $F_Y = l \cdot n = p \cdot (m/m_w)$. Figure 4*c* shows that for both placentals and marsupials, annual fecundity scales similarly, and close to $-1/4$ power. So, despite the very different allocations to litter size and frequency, the number of offspring produced per year is identical across lineages, but the relationship is not tight in either group. Further, lifetime fertility, F_L , is annual fertility $\times$ adult lifespan, a , or $F_L = a \cdot p \cdot (m/m_w)$ (where adult lifespan = maximum lifespan - age at sexual maturity). Figure 4*d* shows that lifetime fertility is statistically invariant with adult body size across the lineages, consistent with Charnov *et al.* [25]. So, even though the allocations to the components of fertility differ conspicuously between the lineages in terms of litter size and frequency, the overall scaling of both annual fecundity and lifetime fertility is predictable though with limited precision from adult body size.

For all variables, values for monotremes tend to overlap those of similar-sized marsupials.

(c) Mass-specific production

Mass-specific production rate, calculated using equation (1.3), is very similar between marsupials and placentals. This is evidenced both by the extensive overlap in values for the two groups in figure 5 and by the parameters of the fitted regression models (table 2): marsupials $\beta = -0.40 \pm 0.08$ (95% CI) and placentals $\beta = -0.37 \pm 0.02$. A GLM testing for differences between the slopes and intercepts, while controlling for

lineage and order, shows that the groups collapse onto a single scaling function, where $\beta = -0.37 \pm 0.04$. To control for the effects of phylogeny at a finer grained taxonomic level, we also ran a model that nested families within order, but the slopes and intercepts were not significantly different from the model controlling solely for order.

4. DISCUSSION AND CONCLUSIONS

Our data analysis not only quantifies well-known differences in life histories among the lineages but also offers new insights into the variation in life-history strategies within and among these groups. The differences in absolute values and allometric scalings of the various life-history traits among placentals, marsupials and monotremes reflect different ways that these three lineages have adapted to allocate time and energy to growth and development in order to maximize reproductive output. These differences reflect a complex combination of phylogenetic constraints owing to ancient divergence and selection owing to environmental conditions during subsequent evolution. Despite these differences in reproductive strategies, however, female placentals, marsupials and monotremes of the same body size all allocate energy to reproduction at essentially the same rate.

An important life-history parameter not considered in equation (1.3) is pre-weaning survival, which would adjust production values by accounting for offspring that received energy investment from the mother, but died before weaning [25]. However, accurate measures of pre-weaning survival are extremely rare, especially in wild populations. To estimate the effect of pre-weaning survival on mass-specific production, we used a method similar to Charnov *et al.* [25]. Using a small sample of placental mammals ($n = 13$), Charnov *et al.* [25] show that pre-weaning survival rates scale with litter size as $s_w = 0.71^{-0.35}$ ($r^2 = 0.69$, $p < 0.01$). We collected

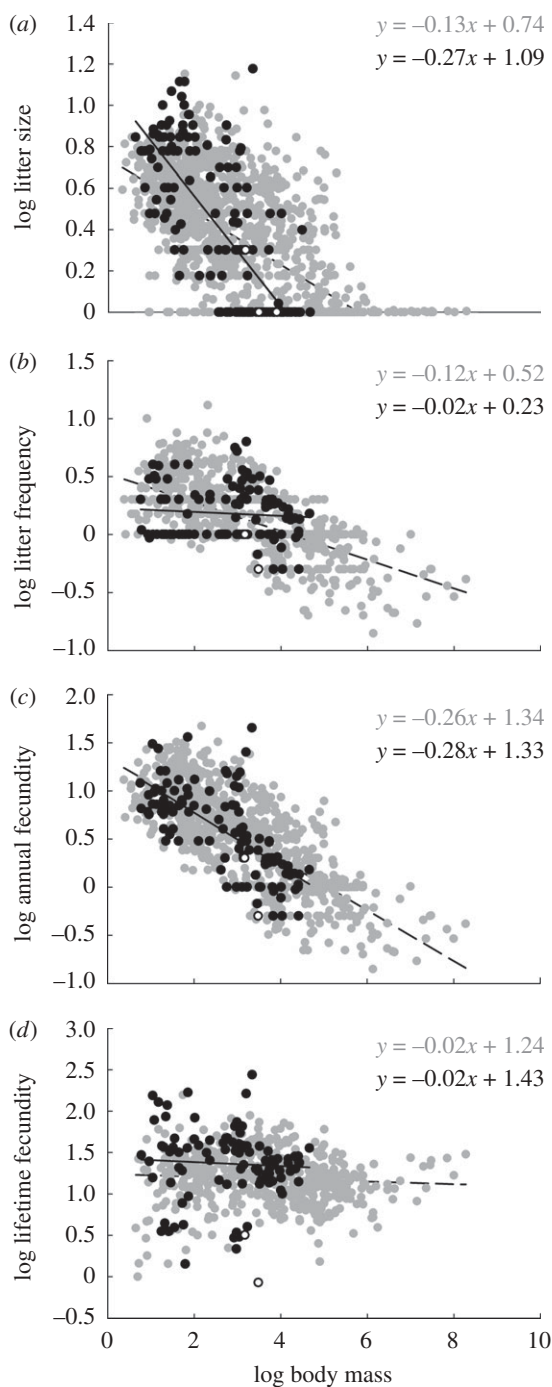


Figure 4. Bivariate log–log plots of components of fecundity as a function of adult body mass: (a) litter size, (b) litter frequency, (c) annual fecundity, and (d) lifetime fecundity.

available data on marsupial pre-weaning survival ($n = 9$), and found that pre-weaning survival is approximately invariant with litter size as $s_w = 0.6l^{0.03}$ ($r^2 = 0.02$, $p > 0.05$), but scales positively with body size, $s_w = 0.2 m^{0.1}$ ($r^2 = 0.32$, $p < 0.05$) (see appendix A for table of data and sources). The differences in the independent variables in offspring survival here probably follow from the differences in reproductive strategies between placentals and marsupials. For example, pouch-rearing in marsupials probably decreases litter size but increases pre-weaning survival as pre-weaned offspring effectively take on the mothers' mortality, whereas in placentals, pre-weaned offspring are exposed to many additional sources of mortality, resulting in lower survival and so

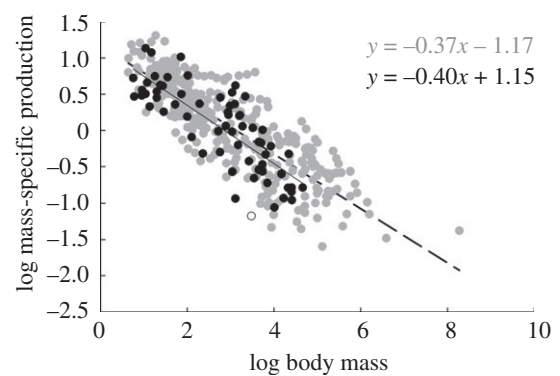


Figure 5. Bivariate log–log plot of mass-specific production as a function of adult body mass. A GLM testing for differences in scaling among lineages finds no significant difference, and gives an overall slope of -0.37 . See table 2 for further details.

larger litter sizes. Therefore, we estimated pre-weaning survival rates from adult body size data for marsupials and from litter size data for placentals. For each species, we calculated the survival-corrected mass-specific production, $p \cdot s_w$, and plotted this as a function of log body size (figure 6). A GLM showed no significant difference between marsupial and placental production rates, and resulted in an overall scaling slope of -0.33 ± 0.07 , which is not significantly different from the uncorrected analyses. Thus, despite the different effects of pre-weaning survival on the ecologies of marsupials and placentals, its effect on production is consistent across lineages, reducing the overall scaling exponent slightly from between -0.37 and -0.40 to -0.33 .

The relatively tight clustering of data about the regression lines in figures 5 and 6 is predicted by metabolic scaling theory on the grounds that all types of mammal are subject to essentially the same constraint on production rate. However, scaling theory would predict a slope of $-1/4$, whereas the observed slope is closer to $-1/3$. This discrepancy occurs because weaning mass is not a constant function of adult mass, but scales sublinearly with an exponent of -0.87 for marsupials and -0.91 for placentals (figure 2b). Why does relative weaning mass scale negatively? We offer three hypotheses for future consideration: (i) Larger mammals differentially allocate resources to adult survivorship at the expense of weaning mass. This should be particularly advantageous in seasonal environments, where it helps adults to survive the winter. (ii) Seasonality should place additional demands on small mammals that breed more than once a year. By producing relatively large offspring, mothers increase the chance their offspring survive a stressful period. (iii) Capital breeding (i.e. fueling reproduction from stored reserves) is increasingly common in larger mammals. Fitness tradeoffs associated with acquisition, storage and mobilization of stored reserves may favour reducing size of offspring at weaning in larger mammals.

The interactions and replacements that occurred when the previously isolated lineages came into contact, such as occurred when the Panamanian land bridge allowed interchange between North and South America approximately 3 Ma ago or when modern humans introduced placental mammals into Australia, have often been attributed to

Table 2. Results of regressions of mass-specific production and adult body mass. The ‘combined’ rows report the result of GLMs testing for the significance between intercepts and slopes between the lineages. In both models, neither the intercepts nor the slopes were significantly different between the lineages.

trait	lineage	slope	s.e.	intercept	s.e.	d.f.	r^2	p
mass-specific production	placentals	−0.37	0.01	1.17	0.05	315	0.69	0.00
	marsupials	−0.40	0.04	1.15	0.11	62	0.66	0.00
	combined	−0.37	0.02	1.16	0.08	378	0.79	0.00
corrected mass-specific production	placentals	−0.32	0.01	0.71	0.04	315	0.67	0.00
	marsupials	−0.33	0.04	0.63	0.13	46	0.59	0.00
	combined	−0.33	0.04	0.74	0.11	362	0.76	0.00

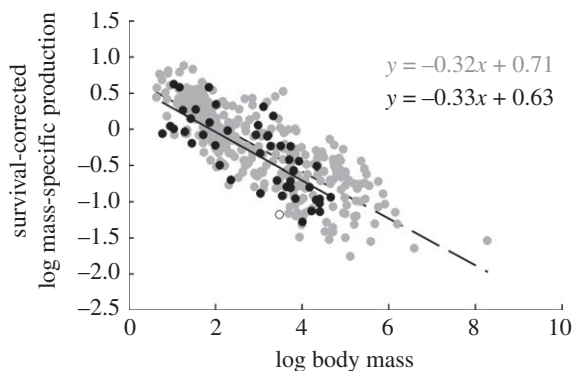


Figure 6. Bivariate log–log plot of survival-corrected mass-specific production as a function of adult body mass. A GLM testing for differences in scaling among lineages finds no significant difference, and gives an overall slope of −0.33, which is not significantly different from the uncorrected result in figure 5. See table 2 for further details.

direct superiority of one phylogenetic lineage over another [26,27]. The fact that all lineages produce biomass at equivalent rates suggests that this interpretation may be too simplistic. Instead, the different reproductive physiologies and ecologies of marsupials and placentals may reflect adaptations to different environments. For example, pouch-rearing may be particularly beneficial in the seasonal, arid and generally nutrient-poor environments of much of Australia [28,30].

Despite many millions of years of independent evolution and widely differing allocations to the components of growth and reproduction, some remarkable convergences across the lineages emphasize the central role of metabolic scaling in mammalian life-history evolution. For example, while marsupial (and monotreme) gestation and weaning times differ considerably from placentals (figure 3*a,b*), the size of offspring at weaning is strikingly consistent across lineages (figure 2*b*), suggesting that while the underlying allocations of energy to the components of pre-weaning development may be free to tradeoff in various ways, the resulting mass at weaning is under strong stabilizing selection. Similarly, despite slower development times, and ages at first reproduction in marsupials and monotremes than in placentals, the maximum lifespan of all mammals scales similarly with body size. The components of fertility also differ conspicuously among the lineages. In marsupials, litter size is approximately invariant with adult body size, and the number of litters per year decreases as approximately the $-1/4$ power of adult body size, whereas in placentals, both litter size and frequency decrease at about $-1/8$

power of body size (figure 4*a,b*), with the result that annual fecundity scales similarly across the lineages, and lifetime fertility is invariant (figure 4*c,d*).

The strikingly similar scaling of production in monotremes, marsupials and placentals suggests that the evolutionary pathways that led to the seemingly divergent mammalian physiologies, reproductive ecologies and life histories are metabolically equivalent ways of being a mammal.

The authors thank Eric Charnov, Felisa Smith, Stephen Dobson, Oskar Burger members of the *Integrative Macroecological Patterns and Process across Scales* (IMPPS) working group (NSF Research Collaboration Network grant DEB-0541625, F. A. Smith, S. K. Lyons and S. K. M. Ernest, PIs), members of the Smith and Brown labs and other colleagues for stimulating discussions on the ideas presented here. This paper is IMPPS RCN publication no. 12. M.J.H. was funded by the IMPPS working group and the Rockefeller Foundation, A.D.D. by NSF, R.M.S. was partially supported by IMPPS and JHB by IMPPS, the Howard Hughes Medical Institute and the National Institutes of Health. The artistic images in figure 1 were reproduced from the originals with kind permission from Sharyn Davidson (*Cynomys ludovicianus*), Joe Trumpey (*Ornithorhynchus anatinus*) and Bruce Worden (*Macrotis lagotis*).

APPENDIX A. MARSUPIAL PRE-WEANING SURVIVAL DATA

species	pre-weaning survival	source
<i>Macropus giganteus</i>	0.83	[31]
<i>Dasyurus hallucatus</i>	0.34	[32]
<i>Tarsipes rostratus</i>	0.33	[33]
<i>Dasyurus maculatus</i>	0.28	[34]
<i>Trichosurus vulpecula</i>	0.38	[35]
<i>Phasogale tapoatafa</i>	0.66	[36]
<i>Didelphis virginia</i>	0.78	[37]
<i>Phascogale cinereus</i>	0.85	[38]
<i>Macropus rufogriseus</i>	0.65	[39]

REFERENCES

- Cifelli, R. L. 2001 Early mammalian radiations. *J. Paleontol.* **75**, 1214–1226. (doi:10.1666/0022-3360(2001)075<1214:EMR>2.0.CO;2)
- Luo, Z.-X. 2007 Transformation and diversification in early mammal evolution. *Nature* **450**, 1011–1019. (doi:10.1038/nature06277)
- Kemp, T. S. 2005 *The origin and evolution of mammals*. Oxford, UK: Oxford University Press.

- 4 Asher, R. J., Horovitz, I. & Sánchez-Villagra, M. R. 2004 First combined cladistic analysis of marsupial mammal interrelationships. *Mol. Phylogenet. Evol.* **33**, 240–250. (doi:10.1016/j.ympev.2004.05.004)
- 5 Cifelli, R. L. & Davis, B. M. 2003 Marsupial origins. *Science* **302**, 1899–1900. (doi:10.1126/science.1092272)
- 6 Kumar, S. & Hedges, S. B. 1998 A molecular timescale for vertebrate evolution. *Nature* **392**, 917–920.
- 7 Luo, Z.-X., Ji, Q., Wible, J. R. & Yuan, X. 2003 An early Cretaceous tribosphenic mammal and metatherian evolution. *Science* **302**, 1934–1940. (doi:10.1126/science.1090718)
- 8 van Rheede, T., Bastiaans, T., Boone, D. N., Hedges, S. B., de Jong, W. W. & Madsen, O. 2006 The platypus is in its place: nuclear genes and indels confirm the sister group relation of monotremes and therians. *Mol. Biol. Evol.* **23**, 587–597. (doi:10.1093/molbev/msj064)
- 9 Woodburne, M. O., Rich, T. H. & Springer, M. S. 2003 The evolution of tribospheny and the antiquity of mammalian clades. *Mol. Phylogenet. Evol.* **28**, 360–385. (doi:10.1016/S1055-7903(03)00113-1)
- 10 Clutton-Brock, T. H. & Harvey, P. H. 1978 Mammals, resources and reproductive strategies. *Nature* **273**, 191–195. (doi:10.1038/273191a0)
- 11 Hayssen, V., Lacy, R. C. & Parker, P. J. 1985 Metatherian reproduction: transitional or transcending? *Am. Nat.* **126**, 617. (doi:10.1086/284443)
- 12 May, R. M. & Rubenstein, D. I. 1985 Reproductive strategies. In *Reproduction in mammals*, vol. 4 (eds C. R. Austin & R. V. Short), pp. 1–23. Cambridge, UK: Cambridge University Press.
- 13 Dawson, T. J., Grant, T. R. & Fanning, D. 1979 Standard metabolism of monotremes and the evolution of homeothermy. *Aust. J. Zool.* **27**, 511–515. (doi:10.1071/ZO9790511)
- 14 Tyndale-Biscoe, H. & Renfree, M. 1987 *Reproductive physiology of marsupials*. Cambridge, UK: Cambridge University Press.
- 15 Charnov, E. L. 1993 *Life history invariants: some explorations of symmetry in evolutionary ecology*. Oxford, UK: Oxford University Press.
- 16 Brown, J. H., Gillooly, J. F., Allen, A. P., Savage, V. M. & West, G. B. 2004 Toward a metabolic theory of ecology. *Ecology* **85**, 1771–1789. (doi:10.1890/03-9000)
- 17 Calder, W. A. 1984 *Size, function, and life history*. Cambridge, UK: Harvard University Press.
- 18 McNab, B. K. 2002 *The physiological ecology of vertebrates*. Ithaca, NY: Cornell University Press.
- 19 Peters, R. H. 1983 *The ecological implications of body size*. Cambridge, UK: Cambridge University Press.
- 20 Schmidt-Nielsen, K. 1984 *Scaling: why is animal size so important?* Cambridge, UK: Cambridge University Press.
- 21 West, G. B., Brown, J. H. & Enquist, B. J. 1997 A general model for the origin of allometric scaling laws in biology. *Science* **276**, 122–126. (doi:10.1126/science.276.5309.122)
- 22 Hou, C., Zuo, W., Moses, M. E., Woodruff, W. H., Brown, J. H. & West, G. B. 2008 Energy uptake and allocation during ontogeny. *Science* **322**, 736–739. (doi:10.1126/science.1162302)
- 23 Sibly, R. M. & Brown, J. H. 2007 Effects of body size and lifestyle on evolution of mammal life histories. *Proc. Natl Acad. Sci. USA* **104**, 17 707–17 712. (doi:10.1073/pnas.0707725104)
- 24 Sibly, R. M. & Brown, J. H. 2009 Mammal reproductive strategies driven by offspring mortality-size relationships. *Am. Nat.* **173**, E185–E199. (doi:10.1086/598680)
- 25 Charnov, E. L., Warne, R. & Moses, M. 2007 Lifetime reproductive effort. *Am. Nat.* **170**, E129–E142. (doi:10.1086/522840)
- 26 Bakker, R. T. 1971 Dinosaur physiology and the origin of mammals. *Evolution* **25**, 636–658. (doi:10.2307/2406945)
- 27 Simpson, G. G. 1980 *Splendid isolation: the curious history of South American mammals*. New Haven, CT: Yale University Press.
- 28 Dickman, C. & Woodford Ganf, R. 2006 *A fragile balance: the extraordinary story of Australian marsupials*. Chicago, IL: University of Chicago Press.
- 29 Low, B. S. 1978 Environmental uncertainty and the parental strategies of marsupials and placentals. *Am. Nat.* **112**, 197–213. (doi:10.1086/283260)
- 30 Russell, E. M. 1982 Parental investment and desertion of young in marsupials. *Am. Nat.* **119**, 744–748. (doi:10.1086/283950)
- 31 Poole, W. E. 1975 Reproduction in the two species of grey kangaroos, *Macropus Giganteus* (Shaw) and *M. Fuliginosus* (Desmarest). II. Gestation, parturition and pouch life. *Aust. J. Zool.* **23**, 333–353. (doi:10.1071/ZO9750333)
- 32 Oakwood, M. 2000 Reproduction and demography of the northern quoll, *Dasyurus hallucatus*, in the lowland savanna of northern Australia. *Aust. J. Zool.* **48**, 519–539. (doi:10.1071/ZO00028)
- 33 Wooller, R. D., Richardson, K. C., Garavanta, C. A. M., Saffer, V. M. & Bryant, K. A. 2000 Opportunistic breeding in the polyandrous honey possum, *Tarsipes rostratus*. *Aust. J. Zool.* **48**, 669–680. (doi:10.1071/ZO00071)
- 34 Belcher, C. A. 2003 Demographics of tiger quoll (*Dasyurus maculatus maculatus*) populations in south-eastern Australia. *Aust. J. Zool.* **51**, 611–626. (doi:10.1071/ZO02051)
- 35 Crawley, M. C. 1973 A live-trapping of Australian brush-tailed possums, *Trichosurus vulpecula* (Kerr), in the orongorongo Valley, Wellington, New Zealand. *Aust. J. Zool.* **21**, 75–90. (doi:10.1071/ZO9730075)
- 36 Soderquist, T. R. 1993 Maternal strategies of *Phascogale tapoatafa* (Marsupialia, Dasyuridae). 1. Breeding seasonality and maternal investment. *Aust. J. Zool.* **41**, 549–566. (doi:10.1071/ZO9930549)
- 37 Hossler, R. J., McAninch, J. B. & Harder, J. D. 1994 Maternal denning behavior and survival of juveniles in opossums in southeastern New York. *J. Mammal.* **75**, 60–70. (doi:10.2307/1382236)
- 38 Tobey, J. R., Andrus, C. H., Doyle, L., Thompson, V. D. & Bercovitch, F. B. 2006 Maternal effort and joey growth in koalas (*Phascolarctos cinereus*). *J. Zool.* **268**, 423–431. (doi:10.1111/j.1469-7998.2005.00041.x)
- 39 Higinbottom, K. & Johnson, C. N. 2000 Partial seasonality of breeding in red-necked wallabies (*Macropus rufogriseus banksianus*). *J. Zool.* **251**, 71–77. (doi:10.1111/j.1469-7998.2000.tb00594.x)