DEBBIES to compare life history strategies across ectotherms

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Abstract

Demographic models are used to explore how life history traits structure life history strategies across species. This study presents the DEBBIES dataset that contains estimates of eight life history traits (length at birth, puberty and maximum length, maximum reproduction rate, fraction energy allocated to respiration versus reproduction, von Bertalanffy growth rate, mortality rates) for 185 ectotherm species. The dataset can be used to parameterise dynamic energy budget integral projection models (DEB-IPMs) to calculate key demographic quantities like population growth rate and demographic resilience, but also link to conservation status or biogeographical characteristics. Our technical validation shows a satisfactory agreement between observed and predicted longevity, generation time, age at maturity across all species. Compared to existing datasets, DEBBIES accommodates (i) easy cross-taxonomical comparisons, (ii) many data-deficient species, and (iii) population forecasts to novel conditions because DEB-IPMs include a mechanistic description of the tradeoff between growth and reproduction. This dataset has the potential for biologists to unlock general predictions on ectotherm population responses from only a few key life history traits.

Background & Summary

Matrix population models (MPMs) and integral projection models (IPMs) provide the basis for exploring how variation in the demographic rates of survival, growth and reproduction fuels variation across species in life history traits (like the timing, intensity, frequency and duration of key demographic processes, such as longevity, generation time or degree of iteroparity) and in combinations of life history traits that form life history strategies (including pace of life and reproductive strategy) [1-3]. Life history traits and strategies calculated from these structured population models predict not only key demographic properties such as population growth rate and demographic resilience, but also have important connections to other disciplines like biogeography, evolutionary biology and conservation biology [4-6].

Important datasets exist that collate MPMs (COMPADRE [7]; COMADRE [8]) and IPMs (PADRINO [9]) for plants and animals, and even algae, fungi, bacteria, and viruses. Recent efforts have furthermore linked the latter datasets in a centralised meta-database of trait data (MOSAIC [10]) so they can be interrogated at the same time. While these datasets are valuable in improving both data access and data usability [11] and are used globally in networks like the Open Trait Network (https://opentraits.org/), the parameterisation of the structured population models that are within these datasets requires long-term individual-level data that are scored from birth till death. Yet, there are many organisms for which such data are not available, for example, because it is difficult to track individuals over their lifetime (e.g. micro-organisms, small (soil-dwelling) animals). Because such MPM and IPM datasets can form part of pipelines to develop, for example, essential biodiversity variables to observe and report global biodiversity change [12], it is pertinent to avoid any unwilling species bias. We thus critically need to include datasets of structured population models that can also accommodate more data-deficient species to have a taxonomically most balanced representation as possible.

One type of structured population model that does not require long-term individual-level life history data is the dynamic energy budget (DEB) IPM [13]. To parameterise a DEB-IPM for a species, one only requires eight life history traits to be estimated (traits include length measures, rates of growth, reproduction and mortality). So far, only two small datasets have been generated, comprising estimates of the eight life history traits for 13 species of marine megafauna [14] and 13 microorganisms [15]. Here, we introduce a much larger and taxonomically diverse dataset that we refer to as DEBBIES in which we compiled estimates of the eight life history traits for 185 ectotherm species. Ectotherms are taxonomically diverse and their growth and reproduction can be captured in simple energy budget models, like the one incorporated into the DEB-IPM. Also, more than 99% of species are ectotherms [16]; consequently, no biological prediction can be considered universal if it is not supported by these organisms. We find in our technical validation that our model outputs exhibit good agreement with observations on key life history traits (age at maturity, longevity, generation time). The dataset can be used for a variety of different applications of eco-evolutionary studies (e.g., [15,17]) (Fig. 1). Because DEB-IPMs (unlike MPMs and other IPMs) across species are built from the same life history traits, they can readily be used for comparative studies of life histories and population dynamics across a wide range of species, for which DEBBIES provides all the necessary input data. Finally, because DEB-IPMs include a mechanistic description of the tradeoff between growth and reproduction, they are particularly suited to create population forecasts to novel conditions (like those created by climate change) [13] for the species currently listed in the dataset.

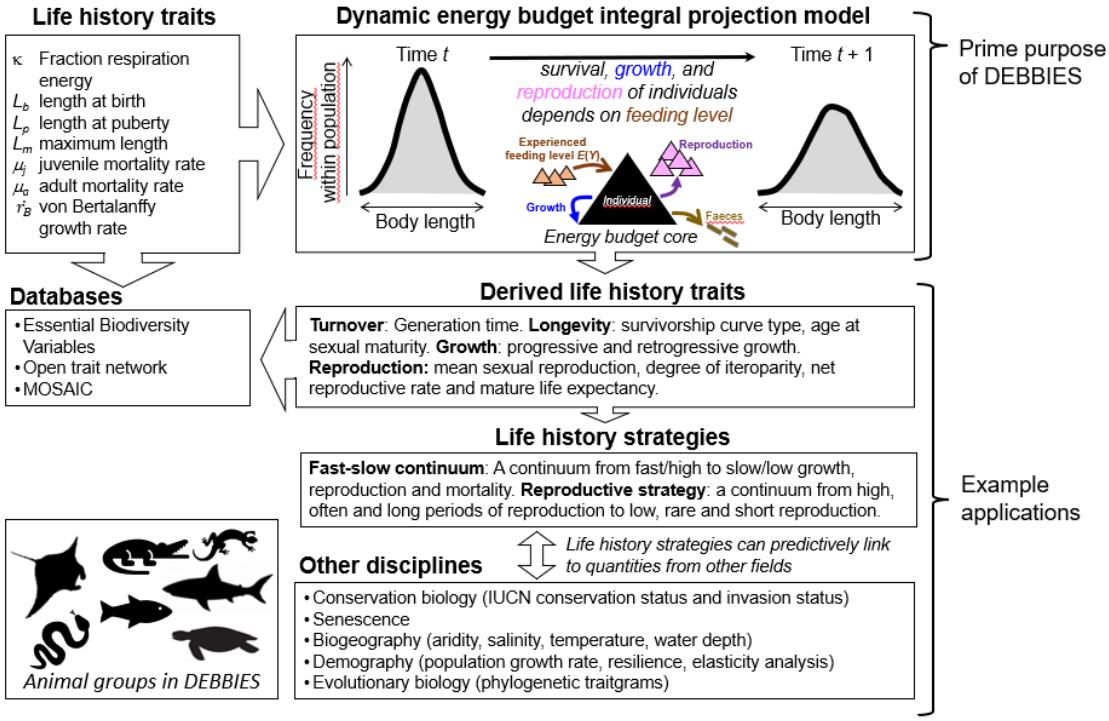

Figure 1. Workflow of parameterising a DEB-IPM and example applications, including what databases DEBBIES can feed into (Essential Biodiversity Variables [Perera et al. 2013]; MOSAIC [Bernard et al. 2022]). DEBBIES currently contains 185 ectotherms of 18 different orders. Eight life history trait values are required to parameterise a DEB-IPM. Once parameterised, it can be used to calculate further, derived life history traits that can in turn be summarised into life history strategies [1]. The resulting 'fast–slow continuum and reproductive strategy framework' can be linked to quantities from other disciplines [4].

Methods

Because the main purpose of DEBBIES is to parameterise DEB-IPMs for further analysis (Fig. 1), we first give a brief description of what a DEB-IPM is, and how a species' eight life history traits feed into a DEB-IPM, before explaining how we collated the life history data presented in DEBBIES.

Brief description of a DEB-IPM

The demographic functions that describe growth and reproduction in the DEB-IPM are derived from the Kooijman-Metz model [18], which is a simple version of the standard model of Kooijman's DEB theory, but still fulfils the criteria for general explanatory models for the energetics of individuals [19]. The Kooijman-Metz model assumes that individual organisms are isomorphic (body surface area and volume are proportional to squared and cubed length, respectively). The rate at which an individual ingests food, *I*, is assumed to be proportional to the maximum ingestion rate I_{max}, the current feeding level *Y* and body surface area, and hence to the squared length of an organism: $I = I_{max}YL^2$. Ingested food is assimilated with a constant efficiency ε. A constant fraction κ of assimilated energy is allocated to respiration; this respiration energy equals $\kappa\varepsilon I_{max}YL^2$ and is used to first cover maintenance costs, which are proportional to body volume following ξL^3 (ξ is the proportionality constant relating maintenance energy requirements to cubed length), while the remainder is allocated to somatic growth. The remaining fraction $1-\kappa$ of assimilated energy, the reproduction energy, is allocated to reproduction in case of adults and to the development of reproductive organs in case of juveniles, and equals $(1-\kappa)\varepsilon I_{max}YL^2$. This means that, if an individual survives from time *t* to time *t*+1, it grows from length *L* to length

L' following a von Bertalanffy growth curve, $\frac{dL}{dt} = \dot{r}_B(L_m \cdot Y - L)$, where $\dot{r}_B$ is the von Bertalanffy growth rate and $L_m = \kappa\varepsilon I_{max}/\xi$ is the maximum length under conditions of unlimited resource. Both κ and I_{max} are assumed to be constant across experienced feeding levels, and therefore L_m is also assumed constant. If a surviving female is an adult, she also produces offspring. According to the Kooijman-Metz model, reproduction, i.e. the number of offspring produced by an individual of length L between time t and $t + 1$, equals $Y \cdot R_m \cdot L^2/L_m^2$. The parameter R_m is the maximum reproduction rate of an individual of maximum length L_m. Note that R_m is proportional to $(1 - \kappa)$ [18], whereas L_m is proportional to κ, which controls energy conservation. However, the role of κ in the DEB-IPM is mostly implicit, as κ is used as input parameter only in the starvation condition (see below), whereas R_m and L_m are measured directly from data. Like L_m, R_m is also proportional to I_{max}; since both κ and I_{max} are assumed to be constant across experienced feeding levels, R_m is also assumed constant.

The above individual life history events are captured in the DEB-IPM by four fundamental functions to describe the dynamics of a population comprising cohorts of females of different sizes [13]: (1) the survival function, $S(L(t))$, describing the probability of surviving from time t to time t+1; (2) the growth function, $G(L', L(t))$, describing the probability that an individual of body length L at time t grows to length L' at $t + 1$, conditional on survival; (3) the reproduction function, $R\big(L(t)\big)$, giving the number of offspring produced between time t and $t + 1$ by an individual of length L at time t; and (4) the parent-offspring function, $D(L', L(t))$, the latter which describes the association between the body length of the parent L and offspring length L' (i.e. to what extent does offspring size depend on parental size). The DEB-IPM assumes no effect of temperature on fundamental functions. Denoting the number of females at time t by $N(L, t)$ means that the dynamics of the body length number distribution from time t to t+1 can be written as:

$$N(L', t+1) = \int_{\Omega} \left[D(L', L(t))R\big(L(t)\big) + G(L', L(t))S(L(t))\right]N(L, t)dL \qquad \text{eqn 1}$$

where the closed interval Ω denotes the length domain. Implicitly underlying the population-level model of eqn 1, like in any IPM, is a stochastic, individual-based model, in which individuals follow Markovian growth trajectories that depend on an individual's current state [20]. This individual variability is in standard IPMs modelled in the functions describing growth, $G(L', L(t))$, and the parent-offspring association, $D(L', L(t))$, using a probability density distribution, typically Gaussian [20]. In the DEB-IPM, this individual variability arises from how individuals experience the environment; specifically, the experienced feeding level Y follows a Gaussian distribution with mean $E(Y)$ and standard deviation $\sigma(Y)$. It means that individuals within a cohort of length L do not necessarily experience the same feeding level due to demographic stochasticity (e.g. individuals, independently of each other, have good or bad luck in their feeding experience).

The survival function $S(L(t))$ in eqn 1 is the probability that an individual of length L survives from time t to $t + 1$:

$$S(L(t)) = \begin{cases} e^{-\mu_j} & \text{for } L_b \le L < L_p \quad \& \quad L \le L_m E(Y)/\kappa\,, \\ e^{-\mu_a} & \text{for } L_p \le L \le L_m \quad \& \quad L \le L_m E(Y)/\kappa\,, \\ 0 & \text{otherwise} \end{cases} \qquad \text{eqn 2}$$

where $E(Y)$ can range from zero (empty gut) to one (full gut). Individuals die from starvation at a body length at which maintenance requirements exceed the total amount of assimilated energy, which occurs when $L > L_m \cdot E(Y)/\kappa$ and hence, then, $S\big(L(t)\big) = 0$ (e.g., an individual of size L_m will die of starvation if $E(Y) < \kappa$). Juveniles and adults often have different mortality rates, and, thus, juveniles ($L_b \le L < L_p$) that do not die of starvation (i.e. $L \le L_m \cdot$

$E(Y)/\kappa$) have a mortality rate of μ_j, and adults ($L_p \leq L \leq L_m$) that do not die of starvation (i.e. $L \leq L_m \cdot E(Y)/\kappa$) have a mortality rate of μ_a.

The function $G(L', L(t))$ is the probability that an individual of body length L at time t grows to length L' at $t + 1$, conditional on survival, and, following common practice [20-22], follows a Gaussian distribution:

$$G(L', L(t)) = \frac{1}{\sqrt{2\pi\sigma_L^2(L(t+1))}} e^{\frac{-(L'-E(L(t+1))^2}{2\sigma_L^2(L(t+1))}} \qquad \text{eqn 3}$$

with the growth realized by a cohort of individuals with length $L(t)$ equalling

$$E\big(L(t+1)\big) = \begin{cases} L(t)e^{-\dot{r_B}} + \big(1 - e^{-\dot{r_B}}\big)L_m E(Y) & \text{for } L \leq L_m E(Y) \\ L(t) & \text{otherwise} \end{cases}, \qquad \text{eqn 4}$$

and the variance in length at time $t + 1$ for a cohort of individuals of length L as

$$\sigma^2(L(t+1)) = \begin{cases} (1 - e^{-\dot{r_B}})^2 L_m^2 \sigma^2(Y) & \text{for } L \leq L_m E(Y) \\ 0 & \text{otherwise} \end{cases} \qquad \text{eqn 5}$$

where $\sigma(Y)$ is the standard deviation of the expected feeding level.

The reproduction function $R(L(t))$ gives the number of offspring produced between time t and $t + 1$ by an individual of length L at time t:

$$R(L(t)) = \begin{cases} 0 & \text{for } L_b \leq L < L_p \\ E(Y)R_m L(t)^2/L_m^2 & \text{for } L_p \leq L \leq L_m E(Y) \\ \dfrac{R_m}{1-\kappa}\left[E(Y)L(t)^2 - \dfrac{\kappa L(t)^3}{L_m}\right] & \text{for } L_m E(Y) < L \leq L_m E(Y)/\kappa \end{cases} \qquad \text{eqn 6}$$

Individuals are mature when they reach puberty at body length L_p and only surviving adults reproduce (eqn 1); thus, only individuals within a cohort of length $L_p \leq L \leq L_m Y/\kappa$ reproduce.

The probability density function $D(L', L(t))$ gives the probability that the offspring of an individual of body length L are of length L' at time $t + 1$, and hence describes the association between parent and offspring character values:

$$D\big(L', L(t)\big) = \begin{cases} 0 & \text{for } L < L_p \\ \dfrac{1}{\sqrt{2\pi\sigma_{L_b}^2(L(t))}} e^{\frac{-(L'-E_{L_b}(L(t))^2}{2\sigma_{L_b}^2(L(t))}} & \text{otherwise} \end{cases} \qquad \text{eqn 7}$$

where $E_{L_b}\big(L(t)\big)$ is the expected size of offspring produced by a cohort of individuals with length $L(t)$, and $\sigma_{L_b}^2(L(t))$ the associated variance. For simplicity, we set $E_{L_b}\big(L(t)\big)$ constant and assumed its associated variance, $\sigma_{L_b}^2\big(L(t)\big)$, to be very small.

Collection of life history trait data required to parameterise a DEB-IPM

Running a DEB-IPM for a species requires estimates for eight life history traits: the fraction respiration energy κ, length at birth L_b, length at puberty L_p, maximum length L_m, juvenile mortality rate μ_j, adult mortality rate μ_a, von Bertalanffy growth rate $\dot{r_B}$ and maximum reproduction rate R_m (Fig. 1).

For the elasmobranchs, we obtained von Bertalanffy growth rate values using the following search, in order of priority: (i) primary literature, using female growth curve, measured empirically using data from animals, (ii) from Froese [23] supplementary material, or (iv) using the equation $\dot{r}_B = -\log(1-0.95)/t_{max}$, where $t_{\max}$ is a species' longevity (years), sourced from Fishbase [24]. If multiple values were available, only those in the highest priority group were kept. If there were still multiple values, their median value was used. Any values listed as 'Questionable' on Fishbase were not used. Body lengths for 152 species were sourced from Sharks of the World [25] or Rays of the World [26]. Total lengths were converted to fork lengths using scalar values on Fishbase [24]. Length at birth was sourced from the IUCN red list [27] for five species (*M. ambigua, M. birostris, A. parmifera, D. trachyderma* and *R. australiae*). Length at puberty for *Maculabatis ambigua* was sourced from primary literature [28]. We calculated maximum reproduction rate R_m as $R_m = (c \times n)/i$ [3], where *c* is the mean clutch size, *n* is the mean number of litters produced per year, and *i* is the remigration interval, which is the minimal number of years between reproductive seasons. Minimum and maximum pup numbers were sourced from Sharks of the World [25], Rays of the World [26], IUCN red list [27], Fishbase [24], or Barrowclift et al. [29]. The sources for maximum and minimum pup numbers were not prioritised, and therefore some maximums and minimums were obtained from different sources. Breeding intervals were found in the same way, except for *C. granulosus*, *S. californica*, *F. macki* and *R. porosus,* which data we sourced from primary literature [30-33]. If the number of pups or breeding interval were not found, we took those data from the next closest species within the same genus. Mode of reproduction was sourced from the Sharks of the World [25] and Rays of the World [26]. Adult mortality rate was calculated by taking the inverse of the mean of longevity and age at maturity (*a*) ($\mu_a = 1/[(t_{max} + a)/2]$ [34]. Juvenile mortality rate was calculated as $\mu_j = -\log\,(l_\alpha^{\frac{1}{a}})$ [35], where survival to maturity, l_α, equals $l_\alpha = e^{-\mu_a(t_{max}-a)}$ [36].

Values for all other species were taken directly from the primary literature. Finally, for most species, we were able to take values for κ from the Add My Pet database [37]. If no values for κ were available for a species, we assumed κ = 0.8 as is common practice [38].

Data Records

The DEBBIES dataset (Version 4) includes eight life history trait records for 185 ectotherm species that were sourced from the scientific literature and are stored in one Excel file on FigShare [39]. Each row in the file represents one species record. For most species, we sourced life history traits from different studies as rarely only one single study provided estimates for all traits for one species. Each data record therefore describes a species' general life history and is not specific to a particular population. The columns of the data file are described in Table 1.

Technical Validation

Two model performance tests on a subset of DEBBIES that have validated the reliability of the dataset are presented elsewhere [13,14]. Here, we conducted a similar model performance test on the full dataset by exploring the distribution of predicted population growth rates, and by comparing predicted and observed generation time (years), age at maturity (years), and longevity (the sum of age at maturity and mature life expectancy; years) for three different feeding levels. Because not all species were represented at all feeding levels (less than half of all species in the dataset can persist at low feeding levels), any effect of feeding level cannot be estimated independently. Therefore, we ran the model performance tests separately for each feeding level. Predicted values of the latter quantities

are calculated as explained below in the usage notes (see also Fig. 1: derived life history traits). For ray-finned fish, observed values were obtained from Fishbase [24] and, if unavailable, we obtained generation times from the IUCN Red List [27] and age at maturity values from the Animal Diversity Web [40]. For the cartilaginous fish, we obtained generation times (for all species) and longevities (18 species) from the IUCN Red List [27], and longevity and age at maturity values from the Sharks of the World book [25], the Rays of the World book [26] or from Barrowclift et al. [29]. For one cartilaginous fish, we obtained the longevity value directly from the scientific literature (silvertip shark *Carcharhinus albimarginatus* [41]) and for the Galapagos shark *Carcharhinus galapagensis* from Fishbase [24]. For all other species, we first consulted the Animal Diversity Web [40], then the IUCN Red List [27] for observations on generation time and AnAge [42] for observations on age at maturity and/or longevity if these were unavailable in the Animal Diversity Web [40]. If, for a particular quantity, a value range was given, we took the median; if a series of observations was given, we took the mean.

Table 1. Description of the columns in the data records file; each row in the file is a species.

Name	Location	Description
RecordID	A	Record ID
Class	B	Species class
Order	C	Species order
Family	D	Species family
Species	E	Species Latin name
Common_name	F	Species common name
kappa	G	Fraction of energy allocated to respiration (κ) (as opposed to reproduction)
L_b	H	Length at birth (cm)
L_p	I	Length at puberty (cm)
L_m	J	Maximum length (cm)
μ_j	K	Juvenile mortality rate (y^{-1})
μ_a	L	Adult mortality rate (y^{-1})
v_B	M	von Bertalanffy growth rate (y^{-1})
R_m	N	Maximum reproduction rate (y^{-1})
Contributor	O	Name of the person who collected the trait data
kappa_REF	P	Reference for κ
L_b_REF	Q	Reference for L_b
L_p_REF	R	Reference for L_p
L_m_REF	S	Reference for L_m
μ_j_REF	T	Reference for μ_j
μ_a_REF	U	Reference for μ_a
v_B_REF	V	Reference for v_B
R_m_REF	W	Reference for R_m

Our validation shows that predicted population growth rates of most species are slightly higher than $\lambda = 1$ (denoting population increase) at high experienced feeding level ($E(Y) = 0.9$), centred around $\lambda = 1$ (denoting stability) at an intermediate feeding level ($E(Y) = 0.7$), and mostly lower than $\lambda = 1$ (denoting population decline) at low feeding level ($E(Y) = 0.5$) (Fig. 2A). This is in line with the general expectation in ecology that populations increase under favourable conditions but decline when conditions deteriorate. Predicted generation times were higher than observed generation times at the lower experienced feeding levels but not significantly different from observed generation times at the highest experienced feeding level (Table 2; Fig. 2B). Similarly, predicted longevity was lower than observed longevity at the lower experienced feeding levels but not significantly different from

observed values at the highest experienced feeding level (Table 2; Fig. 2C). Finally, predicted age at maturity was higher than observed age at maturity and the best fit with observed values was at the lowest experienced feeding level (Table 2; Fig. 2D). Although age at maturity values were significantly higher than observed values, their deviation was not great and the model predicts generation times and longevity values accurately at high experienced feeding levels. Especially given the fact that predicted population growth rates are within the range that we would expect biologically, this technical validation gives us confidence in the technical quality of the dataset and its potential applications (Fig. 1).

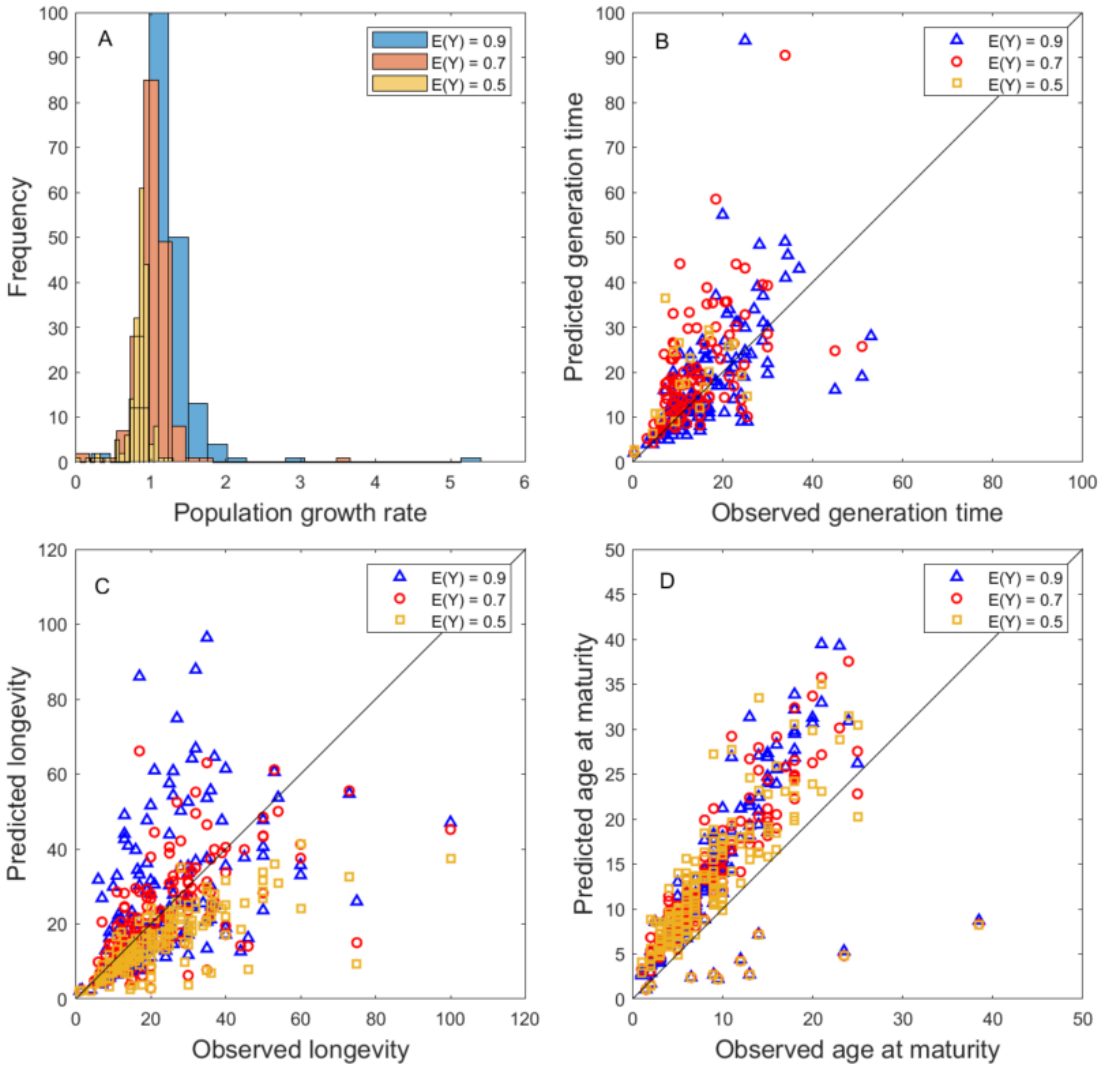

Figure 2. Technical validation. Shown are for three levels of experienced feeding level (blue: $E(Y) = 0.9$; red: $E(Y) = 0.7$; yellow: $E(Y) = 0.5$) the frequency distributions of the population growth rate λ (**A**), observed versus predicted generation time (years) (**B**); observed versus predicted longevity (years) (**C**); and observed versus predicted age at maturity (years) (**D**). The black lines in (B-D) denote the $x = y$ line of equal values.

Table 2. Technical validation that fitted linear regression models without an intercept ($y \sim x$) on observed (x) and predicted (y) values of generation time (G), longevity (L) and age at maturity (A) for three experienced feeding levels $E(Y)$ = 0.5, $E(Y)$ = 0.7, and $E(Y)$ = 0.9. The 95% confidence intervals (CIs) are approximated as the regression coefficient ± twice its standard error. If a 95% CI overlaps with 1, predicted values do not significantly differ from observed values. See also Fig. 2.

E(Y)	model $y \sim x$	regression coefficient	95% CI
0.5	$G_{predicted} \sim G_{observed}$	1.24	1.10 – 1.38
0.7	$G_{predicted} \sim G_{observed}$	1.23	1.16 – 1.30
0.9	$G_{predicted} \sim G_{observed}$	1.02	0.97 – 1.07
0.5	$L_{predicted} \sim L_{observed}$	0.59	0.57 – 0.61
0.7	$L_{predicted} \sim L_{observed}$	0.84	0.81 – 0.87
0.9	$L_{predicted} \sim L_{observed}$	0.99	0.93 – 1.05
0.5	$A_{predicted} \sim A_{observed}$	1.25	1.21 – 1.29
0.7	$A_{predicted} \sim A_{observed}$	1.30	1.26 – 1.34
0.9	$A_{predicted} \sim A_{observed}$	1.45	1.40 – 1.50

Usage Notes

The main purpose of the DEBBIES dataset is to be able to parameterise a DEB-IPM for each of the species in the dataset, but the dataset on its own can also feed into other databases (Fig. 1). Once parameterised, running a DEB-IPM requires setting a value for the experienced feeding level $E(Y)$, which value can range from zero (empty gut) to one (full gut). We find that for most species, $E(Y)$ should be higher than 0.7 for the model to run.

Many different quantities can be calculated from a parameterised DEB-IPM. For example, like current MPM and IPM datasets [7-9], DEB-IPMs can be used to calculate key demographic quantities such as population growth rate and demographic resilience, but also nine key derived life history traits that inform on a species' turnover rate, longevity, growth and reproduction (Fig. 1). Additionally, because all DEB-IPMs have the same structure, one can run elasticity analyses to estimate for each species the proportional change in the population growth rate for a proportional change in each of the input life history traits (κ, L_b, L_p, L_m, μ_j, μ_a, $\dot{r}_B$, R_m). These results can be used to pinpoint those parts of an organism's life history that should be the focus of management effort, or those that contribute most to fitness. We provide MatLab code to calculate the aforementioned quantities (see Code Availability below). Some users, however, might be more interested in exploring the dynamics of single species, rather than performing cross-taxonomical analyses. For those, we have included code to explore how the aforementioned quantities vary with experienced feeding level $E(Y)$ for a particular species (see Code Availability below).

The nine derived life history traits can be used for more in-depth cross-taxonomical analyses (Fig. 1). For example, statistical analyses (e.g., a phylogenetically corrected principal component analysis [43] can be used to explore how life history variation can be structured and summarised into a few life history strategies [44]. Previous studies using MPM databases have revealed that plant and animal life history variation is predominantly structured along a fast-slow life history speed axis (rapid growth, high reproduction and short lifespan versus slow growth, low reproduction but long lifespan) and a reproductive strategy axis (concentrated versus more dispersed reproduction events across the adult lifespan) [1,2]. An interesting exercise would be to see if the same structuring holds when individual growth and reproduction rates are described mechanistically using an energy budget model (like in DEB-IPMs) instead of being estimated from observational data (like in MPMs). What is more,

because a DEB-IPM is run for a specific feeding level set by the user, one could also investigate if the same structuring applies across a range of feeding levels. Finally, if species can be ranked along one or two life history strategy axes, their position along these axes can be linked to quantities from other disciplines (Fig. 1). This opens op possibilities for users to explore the potential for life history strategies to, for example, predict the likelihood of extinction or invasion (e.g., [4]) investigate to what extent such strategies map onto phylogenetic trees (traitgrams) to answer questions about their evolution (e.g., [6]).

Finally, users should be aware that, for the species currently included in DEBBIES, the projection interval between time t and time $t + 1$ is set at 1 year; all rates are therefore expressed per year and all durations are expressed in years. Likewise, all length measurements in DEBBIES are currently expressed in centimetres. Future updates, however, could include species which demographic rates are better projected over shorter time intervals and which lengths are better expressed in millimetres or even micrometres (e.g., microorganisms [6]).

Code Availability

MatLab code and instructions in a README text file for cross-taxonomical analysis can be downloaded from FigShare (https://doi.org/10.6084/m9.figshare.13241972; folder: 'Derived life history traits across ectotherms' [39]) to calculate (i) generation time, survivorship curve, age at sexual maturity, progressive growth, retrogressive growth, mean sexual reproduction, degree of iteroparity, net reproductive rate, mature life expectancy, population growth rate and demographic resilience (damping ratio), and (ii) run an elasticity analysis for each species listed in DEBBIES for a predefined, experienced feeding level.

For users interested in exploring the dynamics of single species, MatLab code and instructions in a README text file can be downloaded from FigShare (https://doi.org/10.6084/m9.figshare.13241972; folder: 'Derived life history traits single ecotherm' [39]) to calculate (i) generation time, survivorship curve, age at sexual maturity, progressive growth, retrogressive growth, mean sexual reproduction, degree of iteroparity, net reproductive rate, mature life expectancy, population growth rate and demographic resilience (damping ratio), and (ii) run an elasticity analysis for a single species for a range of experienced feeding levels defined by the user.

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Author contributions

Isabel Smallegange - project conception – data acquisition – data validation – writing.
Sol Lucas - data acquisition – data validation – proof reading.

Competing interests

The authors declare no competing interests.

References

1. Salguero-Gómez, R., Jones, O. R., Jongejans, E., Blomberg, S. P., Hodgson, D. J., Mbeau-Ache, C., et al.. Fast–slow continuum and reproductive strategies structure plant life-history variation worldwide. *Proc. Natl Acad. Sci. U.S.A.* **113**, 230–235 (2016).
2. Paniw, M., Ozgul, A., & Salguero-Gómez, R. Interactive life-history traits predict sensitivity of plants and animals to temporal autocorrelation. *Ecol. Lett.* **21**, 275–286 (2018).
3. Smallegange, I. M., Flotats Avilés, M., Eustache, K. Unusually paced life history strategies of marine megafauna drive atypical sensitivities to environmental variability. *Front. Mar. Sci.* **7**, 597492 (2020).
4. Salguero-Gómez, R. Applications of the fast–slow continuum and reproductive strategy framework of plant life histories. *New Phytol.* **213**, 1618–1624 (2017).
5. Capdevila, P., Beger, M., Blomberg, S. P., Hereu, B., Linares, C., and Salguero-Gómez, R. Longevity, body dimension and reproductive mode drive differences in aquatic versus terrestrial life history strategies. *Funct. Ecol.* **34**, 1613–1625 (2020).
6. Romeijn, J., Smallegange, I. M. Exploring how the fast-slow pace of life continuum and reproductive strategies structure microorganism life history variation. Preprint at https://doi.org/10.1101/2022.11.28.517963 (2022).
7. Salguero-Gómez, R., Jones, O. R., Archer, C. A., Buckley, Y. M., Che-Castaldo, J., Caswell, C., et al. The COMPADRE Plant Matrix Database: an online repository for plant population dynamics. *J. Ecol.* **103**, 202-218 (2014).
8. Salguero-Gómez, R., Jones, O. R., Archer, C. R., Bein, C., de Buhr, H., Farack, C. et al. COMADRE: a global database of animal demography. *J. Anim. Ecol.* **85**, 371-384 (2016).
9. Levin, S., Compagnoni, A., Childs, D., Evers, S., Potter, T., Salguero-Gómez, R., et al.. PADRINO. *Zenodo* v0.0.1. https://doi.org/10.5281/zenodo.6573870 (2022).
10. Bernard, C., Silva Santos, G., Deere, J. A., Rodriguez-Caro, R., Capdevila, P., Kusch, E., et al. MOSAIC: a unified trait database to complement structured population models. *Sci. Data* **10**, 335 (2023).
11. Gallagher, R. V., Falster, D. S., Maitner, B. S., Salguero-Gómez, R., Vandvik, V., Pearse, W. D. et al.. Open Science principles for accelerating trait-based science across the Tree of Life. *Nature Ecol. Evol.* **4**, 294–303 (2020).
12. Kissling, W. D., Walls, R., Bowser, A., Jones, M. O., Kattge, J., Agosti, D., et al.. Towards global data products of Essential Biodiversity Variables on species traits. *Nature Ecol. Evol.* **2**, 1531-1540 (2018).
13. Smallegange, I. M., Caswell, H., Toorians, M. E. M., & de Roos, A. M. Mechanistic description of population dynamics using dynamic energy budget theory incorporated into integral projection models. *Methods Ecol. Evol.* **8**, 146-154 (2017).
14. Smallegange, I. M., Flotats Avilés, M., & Eustache, K. Unusually paced life history strategies of marine megafauna drive atypical sensitivities to environmental variability. *Front. Mar. Sci.* **7**, 597492 (2020).
15. Romeijn, J., & Smallegange, I. M. Exploring how the fast-slow pace of life continuum and reproductive strategies structure microorganism life history variation. Preprint at https://doi.org/10.1101/2022.11.28.517963 (2022).
16. Atkinson, D., & Sibly, R. M. Why are organisms usually bigger in colder environments? Making sense of a life history puzzle. *Trends Ecol. Evol.* **12**, 235-239 (1997).
17. Rademaker, M., van Leeuwen, A., & Smallegange, I. M. Why we should not necessarily expect life history strategies to inform on sensitivity to environmental change. Preprint at https://doi.org/10.22541/au.164848872.26565315/v1 (2022).

18. Kooijman, S. A. L. M., & Metz, J. A. J. On the dynamics of chemically stressed populations: The deduction of population consequences from effects on individuals. *Ecotoxicol. Environ. Saf.* **8**, 254–274 (1984).
19. Sousa, T., Domingos, T., Poggiale, J. C., & Kooijman, S. A. L. M. Dynamic energy budget theory restores coherence in biology. *Philos. Trans. R. Soc. Lond. B Biol. Sci.* **365**, 3413–3428 (2010).
20. Easterling, M. R., Ellner, S. P., & Dixon, P. M. Size-specific sensitivity: Applying a new structured population model. *Ecology* **81**, 694–708 (2000).
21. Coulson, T. Integral projections models, their construction and use in posing hypotheses in ecology. *Oikos* **121**, 1337–1350 (2012).
22. Merow, C., Dahlgren, J. P., Metcalf, C. J. E., Childs, D. Z., Evans, M. E. K., Jongejans, E., et al.. Advancing population ecology with integral projection models: a practical guide. *Methods Ecol. Evol*. **5**, 99-110.
23. Froese, R. Estimating somatic growth of fishes from maximum age or maturity. *Acta Ichthyol. Piscat.* **52**, 125-133 (2022).
24. Froese, R., & Pauly, D. Editors. FishBase. World Wide Web electronic publication. www.fishbase.org, (02/2023).
25. Ebert, D. A., Dando, M., & Fowler, S. *Sharks of the World. A complete guide* (Princeton University Press, 2021).
26. Last, P., White, W. T., de Carvalho, M. R., Seret, B., Stehmann, M. F. W., & Naylor, G. J. P. *Rays of the World* (CSIRO Publishing, 2016).
27. IUCN. The IUCN Red List of Threatened Species. Version 2022-2. https://www.iucnredlist.org. Accessed on [19 May 2023].
28. Temple, A. J., Stead, S. M., Jiddawi, N., Wambiji, N., Dulvy, N. K., Barrowclift, E. et al.. Life-history, exploitation and extinction risk of the data-poor Baraka's whipray (*Maculabatis ambigua*) in small-scale tropical fisheries. *J. Fish Biol.* **97**, 708-719 (2020).
29. Barrowclift, E., Gravel, S. M., Pardo, S. A., Bigman, J, S., Berggren, P., & Dulvy, N. K. Tropical rays are intrinsically more sensitive to overfishing than the temperate skates. *Biol. Cons.* **281**, 110003 (2023).
30. Guallart, J., & Vicent, J. J. Changes in composition during embryo development of the gulper shark, *Centrophorus granulosus* (Elasmobranchii, Centrophoridae): an assessment of maternal-embryonic nutritional relationships. *Environ. Biol. Fishes* **61**, 135-150 (2001).
31. Natanson, L. J., & Cailliet, G. M. Reproduction and development of the Pacific angel shark, *Squatina californica*, off Santa Barbara, California. *Copeia* **4**, 987-994 (1986).
32. Simpfendorfer, C. A., & Unsworth, P. Reproductive biology of the whiskery shark, *Furgaleus macki*, off south-western Australia. *Mar. Freshwater Res*. **49**, 687-793 (1998).
33. Mattos, S. M., Broadhurst, M., Hazin, F. H., & Jonnes, D. M. Reproductive biology of the Caribbean sharpnose shark, *Rhizoprionodon porosus*, from northern Brazil. *Mar. Freshwater Res.* **52**, 745-752 (2001).
34. Pardo, S. A., Kindsvater, H. K., Reynolds, J. D., & Dulvy, N. K. Maximum intrinsic rate of population increase in sharks, rays, and chimaeras: the importance of survival to maturity. *Can. J. Fish. Aquat. Sci.*, **73**, 1159-1163 (2016).
35. Smith, S. E., Au, D. W., & Show, C. Intrinsic rebound potentials of 26 species of Pacific sharks. *Mar. Freshwater Res.* **49**, 663-678 (1998).
36. Dulvy, N. K., Ellis, J. R., Goodwin, N. B., Grant, A., Reynolds, J. D., & Jennings, S. Methods of assessing extinction risk in marine fishes. *Fish Fish.* **5**, 255-276 (2004).
37. Add-my-pet. *Database of code, data and DEB model parameters* (www.debtheory.org) (2023)
38. Kooijman, S. A. L. M. *Dynamic energy budget theory for metabolic organization* (Cambridge, UK: Cambridge University Press 2010).

39. Smallegange, I. M. DEBBIES. A database to compare life history strategies across ectotherms. *Figshare*. Dataset. https://doi.org/10.6084/m9.figshare.13241972 (2020).
40. Myers, P., Espinosa, R., Parr, C. S., Jones, T., Hammond ,G. S., & Dewey, T. A. *The Animal Diversity Web* (online). Accessed at https://animaldiversity.org (2023).
41. Smart, J. J., Chin, A., Tobin, A. J., White, W. T., Kumasi, B., & Simpfendorfer, C. A. Stochastic demographic analyses of the silvertip shark (*Carcharhinus albimarginatus*) and the common blacktip shark (*Carcharhinus limbatus*) from the Indo-Pacific. *Fish. Res.* **191**, 95-107 (2017).
42. de Magalhaes, J. P., & Costa, J. A database of vertebrate longevity records and their relation to other life-history traits. *J. Evol. Biol*. **22**, 1770-1774 (2009).
43. Revell, L. J. phytools: An R package for phylogenetic comparative biology (and other things). *Methods Ecol. Evol*. **3**, 217–223 (2012).
44. Stearns, S. C. The influence of size and phylogeny on patterns of covariation among life-history traits in the mammals. *Oikos* **41**, 173–187 (1983).