

hatchR: A toolset to predict when fish hatch and emerge

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Abstract

Understanding the timing of key life history events is essential for effective fish conservation and management. Traditionally, predicting hatch and emergence timing in wild fish populations was challenging due to the reliance on average incubation temperature as a primary model parameter, which is often difficult to obtain in natural settings. Recent advancements have refined these models, enabling their application in wild environments using spawning dates and daily water temperature records. However, their broader use remains constrained by a lack of parameterizations for many species, with most applications focused on Salmonids. Here we introduce **hatchR**, a software ecosystem designed to predict hatch and emergence timing for a wide range of wild fishes, requiring only spawning date and water temperature data. **hatchR** offers users access to established phenological models and the flexibility to incorporate custom parameterizations using external datasets. The software is available in two formats: an open-source R package for advanced customization and an HTML-based graphical user interface for those unfamiliar with scripting. To illustrate its utility, we present two case studies demonstrating its application in research and management. By expanding access to predictive modeling tools, **hatchR** has the potential to advance studies of fish early life history and support conservation efforts across diverse species.

Introduction

Fish, as poikilothermic organisms, have developmental rates that are strongly influenced by ambient environmental conditions. This close relationship has allowed researchers to develop statistical models that predict developmental phenology with high accuracy. Historically, these models were created in aquaculture settings under the assumption of constant temperatures throughout development (Alderdice & Velsen, 1978; Beacham & Murray, 1990; McPhail & Murray, 1979). As a result, accumulated thermal unit thresholds (ATU models) have become widely used to predict development in wild populations. However, this approach can be inaccurate, particularly at the cold and warm extremes of temperature ranges, due to the non-linear relationship between average incubation temperature and the thermal units required for development (Quinn, 2018).

In colder environments, fish require fewer accumulated thermal units to develop than those in warmer conditions, yet still experience longer overall developmental periods. Recognizing this non-linear relationship, Sparks et al. (2019) introduced the Effective Value model, which improves prediction accuracy by accounting for daily average temperatures following spawning. This model predicts hatch or emergence timing based on the cumulative effective values reaching a threshold of one, offering a more precise framework for forecasting developmental phenology in wild fish populations.

The effective value approach has since been widely applied to Salmonids, for which aquaculture-derived parameterizations are readily available. For example, Pacific Salmon (*Oncorhynchus* spp.) models developed by Beacham & Murray (1990) have been applied across various species and populations (Adelfio et al., 2019, 2024; Kaylor et al., 2021), while Bull Trout (*Salvelinus confluentus*) models from McPhail & Murray (1979) were extended by Austin et al. (2019). Despite its growing adoption, applications of the effective value model remain largely confined to Salmonids, likely due to the availability of existing parameterizations and the commercial, recreational, and cultural importance of these species.

To extend these modeling capabilities beyond Salmonids and facilitate broader applications, we developed **hatchR**, a software ecosystem designed to predict hatch and emergence timing for wild fish populations. **hatchR** enables users to input standard raw or summarized water temperature datasets commonly collected in field settings, conduct basic data validation, and apply built-in parameterizations such as those from Beacham & Murray (1990) or Sparks et al. (2017). Users can also develop custom models using their own or published temperature and phenological data within the effective value framework for application to unparameterized species or populations.

To maximize accessibility, **hatchR** is available in two formats. The first is an R package, **hatchR**, distributed

via The Comprehensive R Archive Network (CRAN), providing advanced customization and automation for analyzing multiple variables, such as phenology type, spawn timing, or thermal regimes. Comprehensive documentation is available on the **hatchR** website (<https://bmait101.github.io/hatchR/>). The second format is a Shiny-based web application (Chang et al., 2024), offering a graphical user interface for those unfamiliar with R, balancing ease of use with much of the R package’s core functionality (https://elifelts.shinyapps.io/hatchR_demo/). Below, we provide an overview of **hatchR** and present case studies demonstrating its application in research and management.

hatchR Overview

hatchR is designed primarily as a tool for predicting early life history phenology in wild fishes. To maintain focus on this core function, we provide minimal built-in data validation and visualization tools, as users are expected to understand and check their own data. Given the diversity of potential data types, it is impractical to implement comprehensive validation checks. However, we include basic data-checking and summarization functions (`check_continuous()`, `summarize_temp()`) and limited built-in visualization capabilities (`plot_check_temp()`, `plot_phenology()`), the process for which is highlighted in **Box XXX**. Intuitive functions are provided for users to apply models—either existing models from the literature using the `model_select()` function or fitting custom functions from data using the `fit_model()` function. Users can then apply these models to water temperature data (*e.g.*, from an in-stream temperature logger) to predict when hatching phenology will occur. This is accomplished with the `predict_phenology()` function. The R package provides user-friendly example articles that build in complexity for teaching users the functionality and flexibility of each aspect of the work flow as well as customization for plotting model outputs. In contrast, the Shiny application includes a default output plot and an option to download results for external visualizations. For a high-level overview of **hatchR**’s applications, see Figure 1. Additional details on key functions and workflows, particularly for automating phenology predictions across multiple variables, are available in articles hosted on the software’s webpage.

Effective value models

Effective value models were introduced by Sparks et al. (2019) to predict developmental timing in wild populations, initially for Sockeye Salmon (*O. nerka*). Their development was necessitated by limitations in traditional models, such as those in Beacham & Murray (1990), which relied on average incubation temperature over the full developmental period. In wild settings, estimating this average temperature was

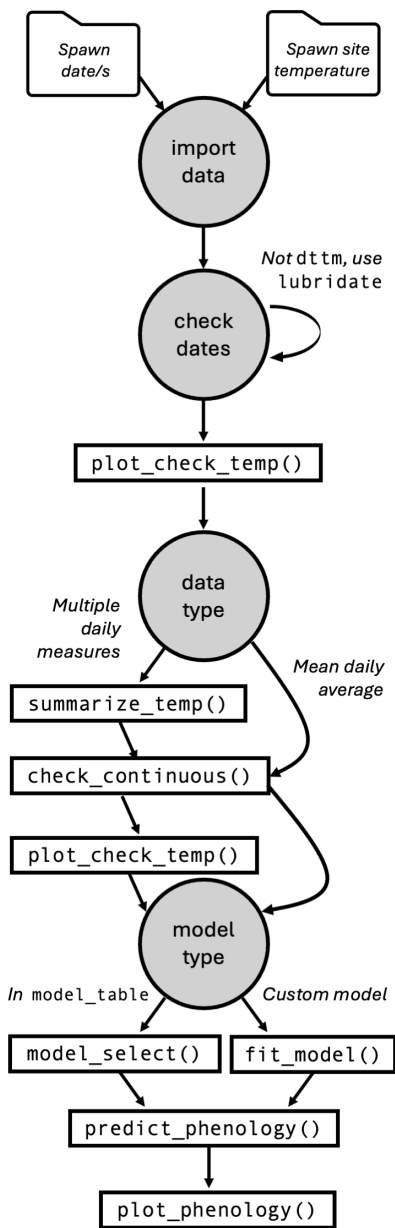


Figure 1: hatchR workflow. Data inputs are represented by folders, data processes by filled circles, hatchR functions as plain text rectangles, and decision choices as italicized text.

impracticable since hatch timing was unknown, even when spawning dates were recorded. To overcome this challenge, Sparks et al. (2019) reformulated model 2 from Beacham & Murray (1990) by taking its reciprocal and assigning an *effective value* to each day of development based on the daily average temperature. This approach allowed for cumulative tracking of developmental progress, enabling hatch and emergence predictions without requiring prior knowledge of incubation temperatures.

The model follows the general format of:

$$E_i = \frac{1}{\exp(\log_e(a) - \log_e(T_i - b))}$$

where E_i is the effective value and T_i the temperature for day i , and a and b are model parameterization estimates (i.e. species- or model-specific constants). A fish hatches or emerges when the cumulative sum of effective values reaches one:

$$\sum_{i=1}^n E_i = 1$$

To demonstrate this approach, we use European Grayling (*Thymallus thymallus*) that were raised in five experimental average temperatures ranging from 2-14 °C (Quinn, 2018). We fit a developmental model to those experimental data (Figure 2A), reparameterize that model into an effective value model, and predict effective values for daily average temperatures ranging from 2-14 °C (Figure 2B).

This framework is the foundation for phenological models in **hatchR**. The package includes a predefined `model_table` containing established parameterizations, including those from Beacham & Murray (1990), Sparks et al. (2017), and Austin et al. (2019) (who extended McPhail & Murray (1979)). While `model_table` incorporated more complex models from Beacham & Murray (1990), users can also fit custom models using the `fit_model()` function. This flexibility allows for the incorporation of new parameterizations as they are developed, expanding the utility of **hatchR** beyond Salmonids.

Data format

Water temperature datasets collected in the field typically fall into two categories: 1) summarized daily data, where mean daily temperatures are pre-computed, or 2) raw high-frequency data, such as those recorded by HOBO TidbiT loggers, which require summary into mean daily temperatures before use. Additionally, new statistical models that predict daily stream temperatures, such as those developed by Siegel et al. (2023),

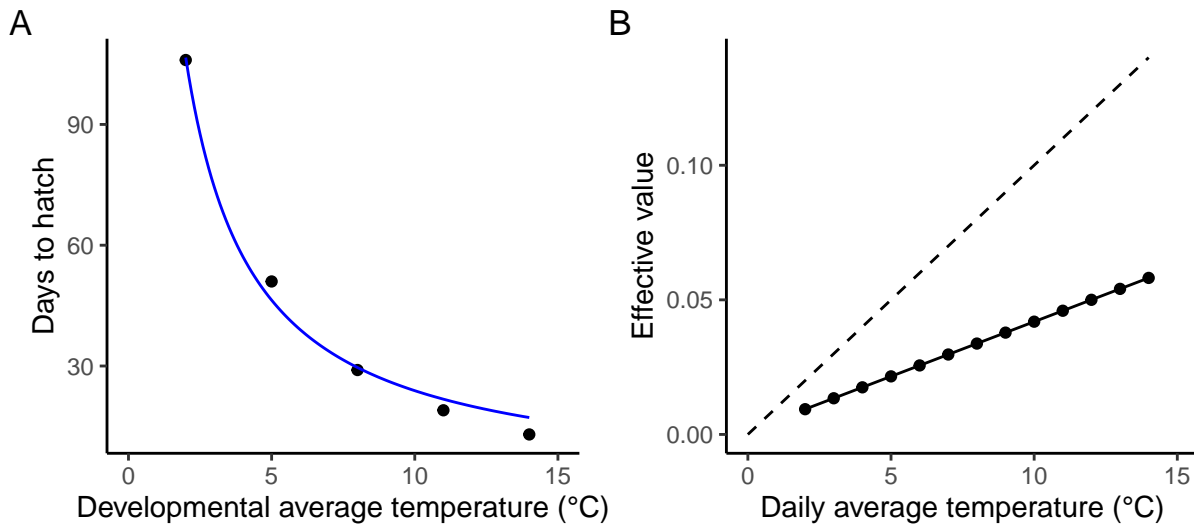


Figure 2: Effective value model fit and application for European Grayling. Panel A shows developmental rates (days to hatch) at five different average rearing temperatures and the corresponding model fit (blue line). Panel B uses the effective value model derived from the blue line in panel A to predict the effective value of daily average temperatures ranging from 2 to 14 °C. The dashed line is a 0.01 increase in effective value for every 1 °C increase as a point of reference.

could also be implemented into this framework.

hatchR assumes input data consists of at least two required columns: a date column indicating the date (and optionally time in the case of fiedl) of each temperature measurement, and a temperature column providing the corresponding temperature measurement (in °C). Other columns may be present, but column names should not include spaces. Data should follow the format outlined in Table 1.

Table 1: Example temperature data for use in hatchR.

date	temperature
2000-01-01	2.51
...	...
2000-07-01	16.32
...	...
2000-12-31	3.13

hatchR will function with missing values, but gaps in the dataset may affect predictions. Since the package does not automatically handle missing data, users must check for gaps or errors before running analyses with

the provided check functions (see section *Checking Data*). **hatchR** supports temperatures as low as 0 °C, though such values yield extremely small effective values, potentially extending hatch or emergence timing to a year or more. Users should critically assess whether such data align with biological expectations.

For R users, **hatchR** can import data in any format, provided it is converted into a `data.frame` or `tibble`, where each row represents a single temperature record. The Shiny application requires data to be uploaded as a .csv (comma separated values) file, which can easily be exported from spreadsheet software such as Microsoft Excel or Google Sheets.

Checking Data

hatchR is designed to analyze daily average temperatures. While high-frequency data (*e.g.*, from HOBO loggers) can be used, it must be summarized into daily averages. **hatchR** provides built-in functionality for this summarization in R but requires pre-summarized data for use in the Shiny app.

To help users identify potential issues, **hatchR** includes basic data checking functions that highlight outliers or missing values both visually and programatically. These checks ensure data integrity before model application.

Here, we demonstrate the utility of three functions—`summarize_temp()`, `plot_check_temp()`, and `check_continuous()`—using a simulated year-long dataset (`year_sim`). This dataset contains temperature readings taken every thirty minutes, and its structure (dimensions and first six rows) is shown below:

```
# simulated object (year_sim) dimensions (rows and columns)
dim(year_sim)
```

```
## [1] 17568      2
```

```
# first 6 rows of year_sim
head(year_sim)
```

```
##           date      temp
## 1 2000-07-01 00:00:00  8.318573
## 2 2000-07-01 00:29:55  9.309468
## 3 2000-07-01 00:59:50 14.676125
## 4 2000-07-01 01:29:45 10.211525
## 5 2000-07-01 01:59:40 10.387863
## 6 2000-07-01 02:29:35 15.145195
```


We recommend using `plot_check_temp()` to visually inspect imported data for outliers or unusual values (Figure 3). This function plots the temperature data against time, allowing users to set custom thresholds for minimum and maximum temperatures. In this example, we set the minimum and maximum thresholds to 0 and 25 °C, respectively.

```
# visual check of temperature data  
plot_check_temp(data = year_sim,  
                dates = date,  
                temperature = temp,  
                temp_min = 0,  
                temp_max = 25)
```

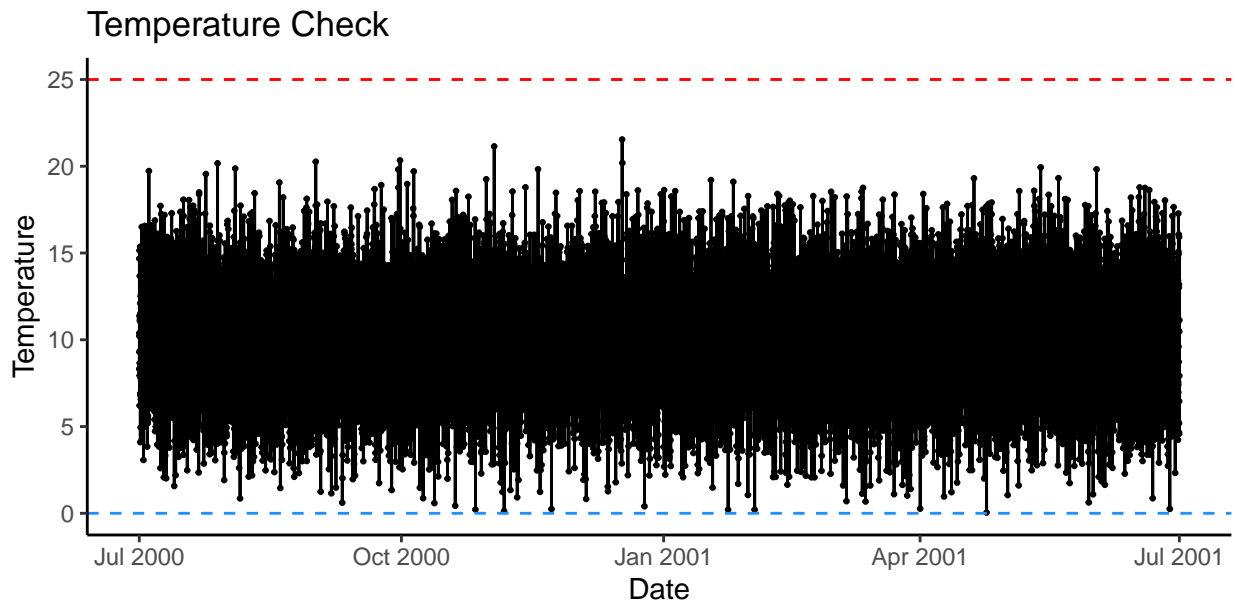


Figure 3: Output of hatchR function `plot_check_temp()`, which is used as a visual data check on the raw `year_sim` dataset. Users can set custom thresholds for minimum and maximum temperatures (dashed lines).

In this case, no obvious outliers are present. However, each day contains 48 records, so the data must be summarized to daily mean temperature using `summarize_temp()`. After summarization, `check_continuous()` should be used to identify any missing days. We also suggest running `plot_check_temp()` again on the summarized data to verify its integrity, though we omit the resulting plot here for space efficiency.

```
# summarize temperature data
year_sim_summ <- summarize_temp(
  data = year_sim,
  dates = date,
  temperature = temp
)

# check dimensions of summarized data
dim(year_sim_summ)
```

```
148 ## [1] 365  2
```

```
# check if continuous
check_continuous(
  data = year_sim_summ,
  dates = date
)
```

```
149 ## i No breaks were found. All clear!
```

```
# we can demonstrate an error by removing Oct. 8 (100th day)
check_continuous(
  data = year_sim_summ[-100, ],
  dates = date
)
```

150 hatchR Workflow

151 Users can select from published model parameterizations contained in `model_table`, or generate custom
 152 model parameterizations using `fit_model()`, in both the R and Shiny deployments of **hatchR**. The models
 153 in `model_table` are included because their parameterizations are well-documented in the literature, though
 154 they are currently limited to Pacific Salmon and Bull Trout (see Quinn (2018) pg. 183, for additional
 155 Salmonid models). To ensure reliability, we restrict `model_table` to well vetted models with experimental

156 ranges spanning 2-17 °C, instead requiring users to carefully asses whether their own parameterized models
157 are appropriate for the temperature ranges in their datasets.

158 To illustrate parameter selection and phenology prediction, we will replicate a portion of the analysis from
159 Sparks et al. (2019) using the `woody_island` dataset included with **hatchR**. Specifically, we predict both
160 hatch and emergence timing for Sockeye Salmon at Woody Island in 1990.

161 First, we obtain the model expression for hatching using `model_select()`, which retrieves the appropriate
162 parameterizations from `model_table`:

```
# use model_select() to get sockeye hatch model
sockeye_hatch_mod <- model_select(
  author = "Beacham and Murray 1990",
  species = "sockeye",
  model = 2,
  development_type = "hatch"
)
```

163 The selected model expressions are then implemented using `predict_phenology()` to estimate the days to
164 hatch (or emergence):

```
# use predict_phenology() to predict sockeye hatch
WI_hatch <- predict_phenology(
  data = woody_island,      # temperature data
  dates = date,             # date column
  temperature = temp_c,     # temperature column
  spawn.date = "1990-08-18", # spawning date
  model = sockeye_hatch_mod # model expression
)
```

165 The returned object provides several outputs, including days to hatch and the full development period,
166 allowing us to assess phenological patterns under the recorded thermal conditions:

```
WI_hatch$days_to_develop; WI_hatch$dev_period
```

167 **## [1] 74**

```

168 ##           start           stop
169 ## 1 1990-08-18 1990-10-30

```

170 Understanding your results

171 The output from `predict_phenology()` (in this example, our named object `WI_hatch`) contains multiple
 172 elements in a list, which can be accessed using the `$` operator. Each component provides different insight into
 173 the predicted phenology:

```
summary(WI_hatch)
```

```

174 ##           Length Class      Mode
175 ## days_to_develop 1      -none-   numeric
176 ## dev_period      2      data.frame list
177 ## ef_table        5      tbl_df   list
178 ## model_specs     5      spec_tbl_df list

```

179 `WI_hatch$days_to_develop` – Returns the predicted number of days required for development.

180 `WI_hatch$dev_period` – A 1x2 dataframe containing the spawning date (as input via `predict_phenology(spawn.date`
 181 `= ...)`) and predicted development completion date.

182 `WI_hatch$ef_table` – An $n \times 5$ tibble (n = number of days to hatch or emerge), containing a row index, the
 183 date, each day's temperature and effective value, and the cumulative sum of the effective values. This table
 184 serves as a foundation for users to create custom visualizations beyond the built-in functionality discussed
 185 below.

186 `WI_hatch$model_specs` – Provides details about the model used for prediction, including whether it was
 187 retrieved from `model_select()` or generated using `fit_model()`. Most importantly, it contains the model
 188 expression (*i.e.*, the formula) used for phenology predictions.

189 Plotting phenology

190 **hatchR** includes a built in function, `plot_phenology()`, for visualizing phenology predictions (Figure 4).
 191 This function generates plots with three specific components: 1) the temperature regime over the prediction
 192 period, 2) the cumulative sum of effective values, and 3) the effective value for each day within the prediction

span. By default, `plot_phenology()` produces a comprehensive figure that includes all three elements with corresponding labels and titles. However, users can customize the output to focus on specific aspects of interest, allowing for tailored visual representations of their results. This function provides a quick and effective way to interpret model outputs, facilitating comparisons between temperature regimes or species-specific phenological responses.

```
plot_phenology(WI_hatch)
```

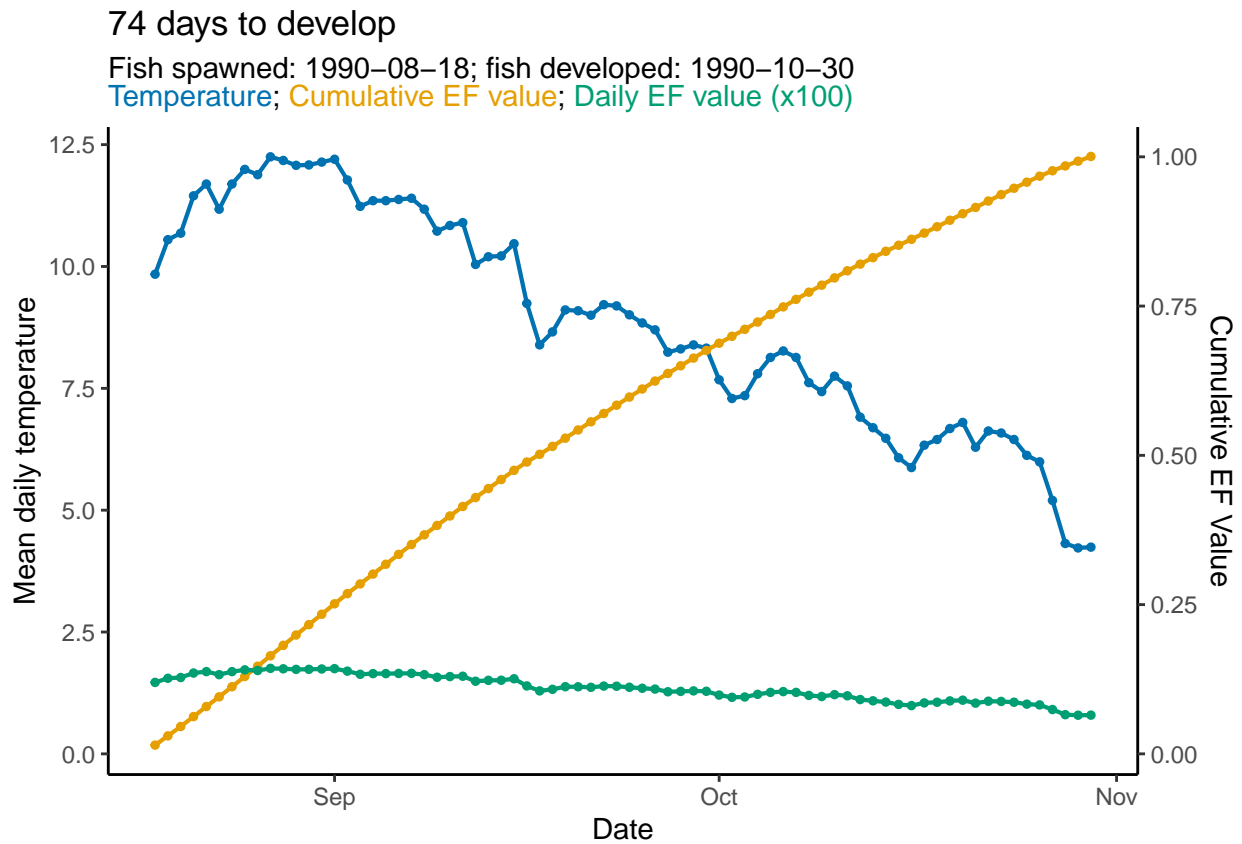


Figure 4: Output of `plot_phenology()` function using predicted hatch time from `woody_island` dataset.

Fitting models for other fishes

To expand the applicability of the effective value approach beyond Salmonids, **hatchR** includes a `fit_model()` function, which only requires species' development follows a power law relationship with temperature. The function takes two input vectors: average incubation temperature (°C) and the number of days to a given phenological event. A model is then fit to the data using `stats::nls()`, which performs nonlinear least squares regression to estimate the parameters $\log_e a$ and b . Because the optimization process in `nls()` is

sensitive to initial parameter values, `fit_model()` first fits a linear model to the log-transformed data to provide initial parameter estimates, which are then used to

This approach allows users to generate models tailored to any species of fish, or even different populations within species, provided they have experimental or field data linking development to temperature. However, users should be mindful of several factors, such as extrapolation risks (models may not generalize beyond the temperature range for which they were parameterized) and species-specific variation (genetic differences among populations may affect developmental responses). Future expansions of **hatchR** could incorporate additional vetted parameterizations for other taxa, such as non-Salmonid fishes, amphibians, or invertebrates, provided sufficient validation in the literature.

To demonstrate how `fit_model()` may be used to create custom parameterizations for species beyond the Salmonids included in `model_table`, we provide parameterizations for three warm-water species: Smallmouth Bass (*Micropterus dolomieu*) (Webster, 1948), Channel Catfish (*Ictalurus punctatus*) (Small & Bates, 2001), and Lake Sturgeon (*Acipenser fulvescens*) (Smith & King, 2005). These species were selected due to their common use in aquaculture and sport fisheries. For concision, we present parameterization for Smallmouth Bass below, while the full implementation details for all species are available in the `paper.Rmd` on the GitHub project repository (<https://github.com/bmait101/hatchR>).

```
# Smallmouth Bass data (Webster 1948)

smallmouth <- matrix(NA, 10, 2) |> data.frame()
colnames(smallmouth) <- c("hours", "temp_F")
smallmouth$hours <- c(52, 54, 70, 78, 90, 98, 150, 167, 238, 234)
smallmouth$temp_F <- c(77, 75, 71, 70, 67, 65, 60, 59, 55, 55)

# change °F to °C and hours to days
smallmouth <- smallmouth |>
  mutate(
    days = ceiling(hours / 24),
    temp_C = (temp_F - 32) * (5 / 9)
  )

# estimate parameters for Smallmouth Bass using fit_model()
smb_mod <- fit_model(
```

```

temp = smallmouth$temp_C,
days = smallmouth$days,
species = "smb",
development_type = "hatch"
)

```

220 Note the R^2 fit from the models below. You can see they generally perform well and are close to values from
 221 model 2 of Beacham & Murray (1990), which fall between 0.95 and 0.99.

```

# R^2 values from model fits
c(smb_mod$r_squared, cat_mod$r_squared, sturgeon_mod$r_squared)

```

```

222 ## [1] 0.9868067 0.9433598 0.9217358

```

223 Next, we generated a simulated thermal regime featuring an ascending thermograph with a mean temperature
 224 of 16 °C (available in `paper.Rmd`). Using this dataset, we apply the custom models for each species using
 225 `predict_phenology()` (only Smallmouth Bass shown below):

```

# Smallmouth Bass
smb_res <- predict_phenology(
  data = data_sim,           # simulated data
  dates = date_sim,         # simulated dates
  temperature = temp_sim,    # simulated temperatures
  spawn.date = "2000-07-01", # spawn date
  model = smb_mod$expression # model expression from fit_model()
)

```

226 Finally, we summarize the outputs across species to visualize hatch timing and total developmental duration
 227 (Figure 5). This example highlights the flexibility of **hatchR** for accommodating diverse fish species and
 228 environmental conditions, making it a valuable tool for researchers and managers working outside of Salmonid
 229 systems.

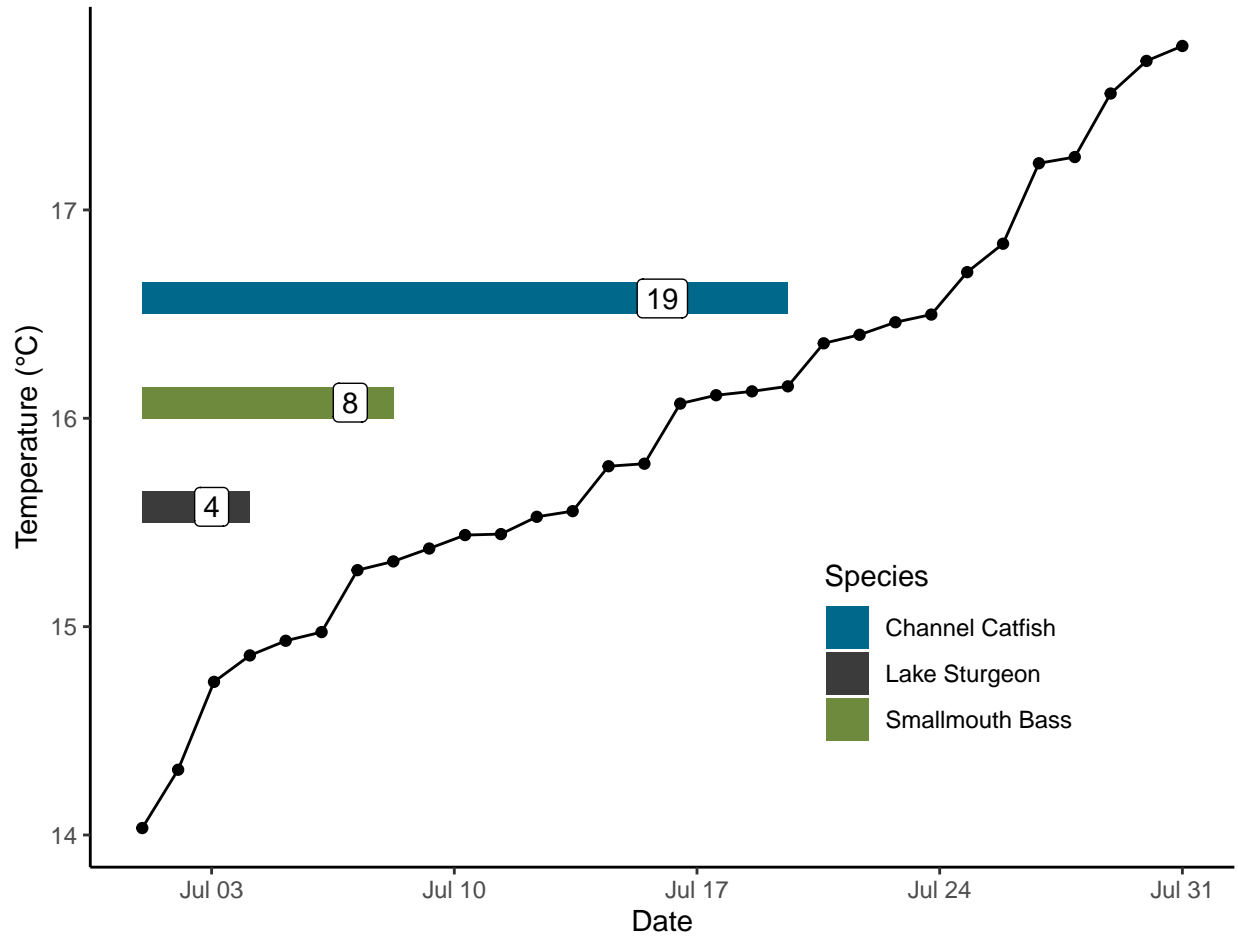


Figure 5: Predicted days to hatch for three warmwater species with custom parameterizations using a random thermal regime with an ascending thermograph with a mean temperature of 16 °C over 30 days.

Case Study 1

Predicting Emergence Timing for Management Actions

A common management application of developmental phenology is assessing whether fish will be free-moving before a scheduled management action, such as stream section access for grazing or road work. For instance, will fish have emerged from redds before construction begins, reducing potential disturbance? This type of predictive modeling helps managers make informed, science-based decisions that balance conservation priorities with land-use activities.

In this scenario, we consider road work near the upper portion of Crooked River in the Boise River watershed in Idaho, USA, home to a key Bull Trout (*S. confluentus*) population. Bull Trout, a federally threatened species under the Endangered Species Act (Nolfi et al., 2024), are particularly sensitive to sediment disturbance. The Forest Service Fisheries Biologist overseeing the project wants to determine whether Bull Trout fry will likely be out of the gravel and free-swimming by June 1st. In this system, Bull Trout typically complete spawning by the end of September, so we consider the latest possible spawn date: September 30th.

To demonstrate this case study, we use the **hatchR** graphical user interface. Users begin by uploading their temperature dataset through the **Import Data** window, selecting their file, specifying the appropriate temperature and date columns, and providing the date format (e.g., year-month-day or day-month-year). For this example, we use the **crooked_river** dataset, which is included in **hatchR**.

Once uploaded, **hatchR** automatically generates a visual data check using `plot_check_temp()`. After confirming data integrity, users navigate to the **Model Phenology** window. For this case study, we use the pre-loaded Bull Trout parameterization from Austin et al. (2019), selecting the **Existing** model option via dropdown menus. The user then may chooses multiple spawn dates using an interactive calendar. Here, we focus on September 30th (in the 2014 spawn year) as outlined in our management scenario.

Following model selection, **hatchR** outputs results in two key locations: the **Phenology Summaries** tab, which provides a table with predictions for each spawn date, and the **Timeline Plot** tab, which shows the corresponding visualization of development timing. Both the prediction table and plot can be downloaded directly from their respective tabs. The full process is demonstrated through a detailed walkthrough in the supplementary video file (Supplementary Materials).

In this example, the model predicts that the last Bull Trout will emerge on April 21, before the road work target date (June 1). This suggests that the Fisheries Biologist can confidently approve the work in the area without concern for sediment disturbance impacting fish developing in the gravel.

Case Study 2

Large Scale Predictions of Bull Trout Development Timing

For the second case study, we demonstrate a more complex, large-scale application of **hatchR**, highlighting its full flexibility when applied programmatically in R. This example also focuses on Bull Trout, but extends beyond a single site to a broad spatial analysis across 226 locations in the greater Snake River watershed in Idaho, USA.

We use the `idaho` dataset from Isaak et al. (2018), which includes four years of temperature data (2011-2014) for these sites. To identify putative Bull Trout Spawning locations, we apply a filtering criterion based on mean August temperature, as outlined in Isaak et al. (2015), selecting only sites with mean August temperature at or less than 13 °C, a known thermal threshold for Bull Trout spawning suitability. The filtering process reduced the dataset to 139 potential spawning sites.

To predict hatch timing across these sites, we first set up the necessary models and data, using the built-in Bull Trout parameterization (for concision, we omit this setup here, but full details are available in the `paper.Rmd` file in the GitHub repository, or in the *Predict fish phenology: Nested* article on **hatchR**'s website).

We then apply `predict_phenology()` across all 139 sites, running predictions for three representative spawn dates each year (Early: September 1st, Peak: September 15th, Late: September 30th).

```
# apply predict_phenology() over multiple sites and spawn dates
hatch_res <- isaak_summ_bt |>
  mutate(
    dev_period = map2(
      summ_obj,          # summarized temperature data object
      spawn_dates,      # list of spawn dates
      predict_phenology, # function to apply
      temperature = daily_temp, # column of temperature data in `summ_obj`
      dates = date,      # column of date in `summ_obj`
      model = bt_hatch   # model to use (from model_select())
    ) |>
    map_df("dev_period") |> # pull out just dev_period results
    list()
  ) |>
```

```

select(site, dev_period) |>           # just select the columns we want
unnest(cols = c(dev_period)) |>      # un-nest everything
mutate(days_to_hatch = stop - start) # make a new column of days to hatch

```

By mapping predictions across this broad spatial extent, we generate a large-scale assessment of Bull Trout phenology, illustrating how hatch timing varies across different spawning habitats. The results of this analysis are presented in Figure 6, providing insights into how hatch timing might vary under different thermal regimes and across the species' geographic range and across years. This case study underscores the power of **hatchR** for large-scale ecological applications, particularly in conservation planning and habitat suitability assessments.

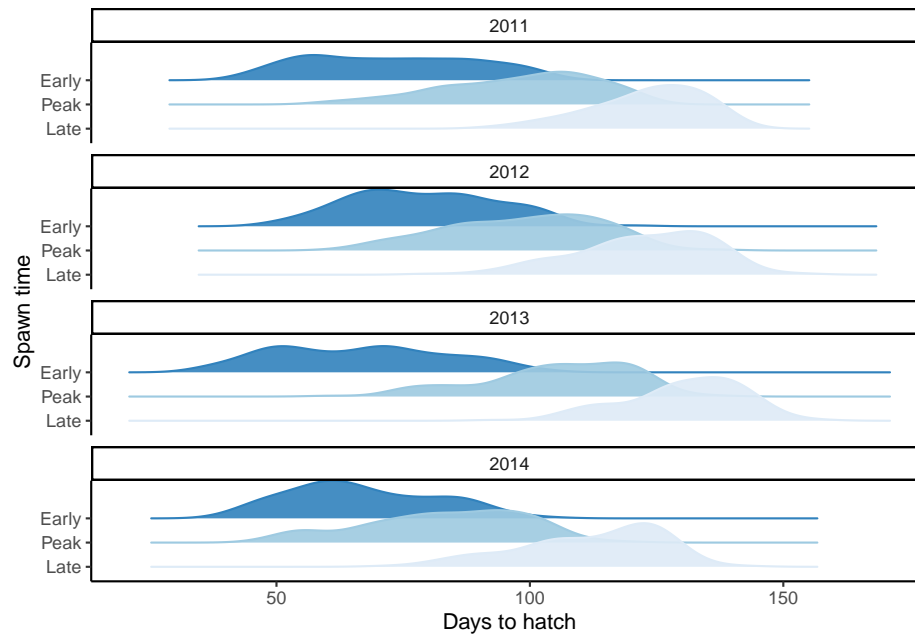


Figure 6: Predicted days to hatch for 139 putative bull trout populations over three spawning periods (Early = September 1, Peak = September 15, Late = September 30) and four years of temperature data.

Discussion

hatchR is a software ecosystem that bridges the analytical gap in predicting developmental phenology for wild fishes. It establishes a formal framework for applying effective value models from user-provided parameterizations. The software is available in two formats: 1) A fully customizable R package, ideal for complex and repetitive analyses and 2) a graphical user interface for ease of use, designed for tasks that may only need to be run once or a few times.

Both versions allow users to import data, perform basic data checks, perform basic visualizations, and apply either pre-existing Salmonid model parameterizations or generate custom models specific to other species or populations. To support users at various levels of expertise, we provide extensive documentation on the **hatchR** website (<https://bmait101.github.io/hatchR/>), covering basic and advanced applications.

Assumptions and considerations in applying effective value models

The application of **hatchR** and the effective value modeling framework relies on several key assumptions. First, environmental stressors may alter developmental timing. While effective value models predict developmental timing based on temperature, studies have shown that stressful environmental condition such as low dissolved oxygen, altered pH, high salinity, pathogen exposure, or mechanical disturbance can induce premature hatching or emergence (Cowan et al., 2024; Quinn, 2018). Users should consider how such factors may influence their predictions.

Second, developmental timing occurs as a distribution, not a fixed point. While **hatchR** provides point estimates for developmental phenology, fish spawning and development within populations occur as distributions rather than single events (Mason, 1976). We encourage users to predict phenology using early, peak, and late thresholds (*e.g.*, 5th, 50th, and 95th percentiles) or incorporate real or modeled distributions to capture variation.

Third, sensor-based temperature data may differ from actual embryonic ambient temperatures. Water temperatures recorded by environmental sensors may not fully reflect thermal conditions in spawning microhabitats, where geomorphic factors influence temperature regimes (Geist et al., 2002). Users should consider how differences between measured and actual incubation temperatures may affect predictions.

Evolutionary and population-level considerations

To date, effective value models have primarily been used to predict phenology in wild environments using species-specific parameterizations (Adelfio et al., 2024; Austin et al., 2019). However, these models fundamentally represent reaction norms, meaning that temperature-development relationships are influenced by local adaptation, gene-environment interactions, and phylogenetic differences (West-Eberhard, 2003). For example, Sparks et al. (2017) found no significant differences in developmental rates between populations in their study but did observe family-level genetic \times environment interactions under different thermal regimes. Similarly, when they reparameterized their models using western Alaskan Sockeye Salmon, they found slower developmental rates compared to populations from Canada (Beacham & Murray, 1990), consistent with

cogradient variation (Conover et al., 2009; Sparks et al., 2022). These findings highlight the importance of considering how developmental rates are keyed to specific environments but also how these underlying statistical relationships inform micro- and macro-evolutionary processes in fishes.

Expanding the utility of **hatchR**

The models within **hatchR** can be customized in multiple ways beyond the examples provided. While our current framework focuses on predicting hatch or emergence timing, it could be adapted to other key developmental milestones not reliant on exogenous feeding, such as early embryonic stages (*e.g.*, eye-up; (Velsen, 1980)), initiation or cessation of pelagic-larval dispersal, or current-mediated dispersal in riverine species, though not all cases may be as specifically tied to temperature as hatch and emergence.

Additionally, while `fit_model()` uses non-linear regression to estimate parameters, `predict_phenology()` only requires users to provide a model expression. This means that users can integrate alternative model structures, as long as they incorporate daily temperature, allowing further customization of predictions.

Finally, while **hatchR** was designed specifically for fishes, it has potential applications for other poikilothermic organisms, such as reptiles, amphibians, and invertebrates, where developmental rates similarly follow a power law relationship with temperature. Extending the effective value framework to these taxa could provide valuable insights into their life history timing under variable environmental conditions. We encourage users to submit fitted models using this framework as a pull request to **hatchR**'s Github repository provided that data used in the models are published.

Conclusion

hatchR provides a versatile and accessible tool for predicting developmental phenology in wild fish populations. It offers basic data checks and summarization tools, pre-existing and customizable model parameterization options, and scalable applications from simple site-level predictions to complex multi-site analyses.

Importantly, **hatchR** extends the effective value framework developed by Sparks et al. (2019) into a generalizable tool that can be applied to any fish species or population, provided that appropriate source data are available. We present foundational applications of **hatchR**, with additional user-friendly cases and implementation guides available on the software's website. The software is designed for both applied and fundamental research, allowing users to engage with it either through a programmatic R environment or via a user-friendly Shiny app. We expect that the examples provided here represent only a fraction of **hatchR**'s

potential applications and encourage the user community to explore and expand upon this framework for their own research and management needs.

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Conflicts of Interest

The authors declare no known conflicts of interest.

Data Availability

hatchR is fully open source and reproducible. Source code and data can be found at <https://github.com/bmait101/hatchR/>. The Rmarkdown document with all the code to reproduce the examples from this manuscript is available at <https://github.com/bmait101/hatchR/blob/master/inst/manuscript/paper.Rmd>. The latest version will be archived upon acceptance of the manuscript.

Ethics Statement

All data was derived from pre-published sources or created synthetically.

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References

- Adelfio, L. A., Wondzell, S. M., Mantua, N. J., & Reeves, G. H. (2019). Warm winters reduce landscape-scale variability in the duration of egg incubation for coho salmon (*Oncorhynchus kisutch*) on the copper river delta, alaska. *Canadian Journal of Fisheries and Aquatic Sciences*, 76(8), 1362–1375. <https://doi.org/10.1139/cjfas-2018-0152>
- Adelfio, L. A., Wondzell, S. M., Mantua, N. J., & Reeves, G. H. (2024). Expanded, compressed, or equal? Interactions between spawning window and stream thermal regime generate three responses in modeled juvenile emergence for Pacific salmon. *Canadian Journal of Fisheries and Aquatic Sciences*. <https://doi.org/10.1139/cjfas-2023-0238>
- Alderdice, D. F., & Velsen, F. P. J. (1978). Relation between temperature and incubation yime for eggs of chinook salmon (*Oncorhynchus tshawytscha*). *Journal of the Fisheries Research Board of Canada*, 35(1), 69–75. <https://doi.org/10.1139/f78-010>
- Austin, C. S., Essington, T. E., & Quinn, T. P. (2019). Spawning and emergence phenology of bull trout *Salvelinus confluentus* under differing thermal regimes. *Journal of Fish Biology*, 94(1), 191–195. <https://doi.org/10.1111/jfb.13864>
- Beacham, T. D., & Murray, C. B. (1990). Temperature, egg size, and development of embryos and alevins of five species of Pacific salmon: A comparative analysis. *Transactions of the American Fisheries Society*, 119(6), 927–945. [https://doi.org/10.1577/1548-8659\(1990\)119%3C0927:TESADO%3E2.3.CO;2](https://doi.org/10.1577/1548-8659(1990)119%3C0927:TESADO%3E2.3.CO;2)
- Chang, W., Cheng, J., Allaire, J., Sievert, C., Schloerke, B., Xie, Y., Allen, J., McPherson, J., Dipert, A., & Borges, B. (2024). *Shiny: Web application framework for r*. <https://CRAN.R-project.org/package=shiny>
- Conover, D. O., Duffy, T. A., & Hice, L. A. (2009). The covariance between genetic and environmental influences across ecological gradients. *Annals of the New York Academy of Sciences*, 1168(1), 100–129. <https://doi.org/10.1111/j.1749-6632.2009.04575.x>
- Cowan, Z.-L., Green, L., Clark, T. D., Blewett, T. A., De Bonville, J., Gagnon, T., Hoots, E., Kuchenmüller, L., Leeuwis, R. H. J., Navajas Acedo, J., Rowsey, L. E., Scheuffele, H., Skeeles, M. R., Silva-Garay, L., Jutfelt, F., & Binning, S. A. (2024). Global change and premature hatching of aquatic embryos. *Global Change Biology*, 30(9), e17488. <https://doi.org/10.1111/gcb.17488>
- Geist, D. R., Hanrahan, T. P., Arntzen, E. V., McMichael, G. A., Murray, C. J., & Chien, Y.-J. (2002). Physicochemical characteristics of the hyporheic zone affect redd site selection by chum salmon and fall Chinook salmon in the Columbia River. *North American Journal of Fisheries Management*, 22(4), 1077–1085. [https://doi.org/10.1577/1548-8675\(2002\)022%3C1077:PCOTHZ%3E2.0.CO;2](https://doi.org/10.1577/1548-8675(2002)022%3C1077:PCOTHZ%3E2.0.CO;2)
- Isaak, D. J., Luce, C. H., Chandler, G. L., Horan, D. L., & Wollrab, S. P. (2018). Principal components of

thermal regimes in mountain river networks. *Hydrology and Earth System Sciences*, 22(12), 6225–6240.
<https://doi.org/10.5194/hess-22-6225-2018>

Isaak, D. J., Young, M. K., Nagel, D. E., Horan, D. L., & Groce, M. C. (2015). The cold-water climate shield: delineating refugia for preserving salmonid fishes through the 21st century. *Global Change Biology*, 21(7), 2540–2553. <https://doi.org/10.1111/gcb.12879>

Kaylor, M. J., Justice, C., Armstrong, J. B., Staton, B. A., Burns, L. A., Sedell, E., & White, S. M. (2021). Temperature, emergence phenology and consumption drive seasonal shifts in fish growth and production across riverscapes. *Journal of Animal Ecology*, 90(7), 1727–1741. <https://doi.org/10.1111/1365-2656.13491>

Mason, J. (1976). Some features of coho salmon, *Oncorhynchus kisutch*, fry emerging from simulated redds and concurrent changes in photobehavior. *Fish. Bull*, 74(1), 167–175.

McPhail, J., & Murray, C. (1979). *The early life-history and ecology of dolly varden (Salvelinus malma) in the Upper Arrow Lakes* [Report to B.C. Hydro and Kootenay Fish and Wildlife].

Nolfi, D., Melbiess, T., Fisher, S., & Ellis, L. (2024). *5-year status review coterminous United States population of bull trout (Salvelinus confluentus)*.

Quinn, T. P. (2018). *The Behavior and Ecology of Pacific Salmon and Trout*. University of Washington Press.

Siegel, J. E., Fullerton, A. H., FitzGerald, A. M., Holzer, D., & Jordan, C. E. (2023). Daily stream temperature predictions for free-flowing streams in the Pacific Northwest, USA. *PLOS Water*, 2(8), e0000119. <https://doi.org/10.1371/journal.pwat.0000119>

Small, B. C., & Bates, T. D. (2001). Effect of low-temperature incubation of channel catfish *Ictalurus punctatus* eggs on development, survival, and growth. *Journal of the World Aquaculture Society*, 32(2), 189–194. <https://doi.org/10.1111/j.1749-7345.2001.tb01094.x>

Smith, K. M., & King, D. K. (2005). Dynamics and extent of larval lake sturgeon *Acipenser fulvescens* drift in the upper Black River, Michigan. *Journal of Applied Ichthyology*, 21(3), 161–168. <https://doi.org/10.1111/j.1439-0426.2005.00623.x>

Sparks, M. M., Falke, J. A., Quinn, T. P., Adkison, M. D., Schindler, D. E., Bartz, K., Young, D., & Westley, P. A. H. (2019). Influences of spawning timing, water temperature, and climatic warming on early life history phenology in western Alaska sockeye salmon. *Canadian Journal of Fisheries and Aquatic Sciences*, 76(1), 123–135. <https://doi.org/10.1139/cjfas-2017-0468>

Sparks, M. M., Kraft, J. C., Blackstone, K. M. S., McNickle, G. G., & Christie, M. R. (2022). Large genetic divergence underpins cryptic local adaptation across ecological and evolutionary gradients. *Proceedings of the Royal Society B: Biological Sciences*, 289(1984), 20221472. <https://doi.org/10.1098/rspb.2022.1472>

Sparks, M. M., Westley, P. A. H., Falke, J. A., & Quinn, T. P. (2017). Thermal adaptation and phenotypic

430 plasticity in a warming world: Insights from common garden experiments on Alaskan sockeye salmon.
 431 *Global Change Biology*, 23(12), 5203–5217. <https://doi.org/10.1111/gcb.13782>
 432 Velsen, F. P. J. (1980). Embryonic development in eggs of sockeye salmon *Oncorhynchus nerka*. *Canadian*
 433 *Special Publication of Fisheries and Aquatic Sciences*, 49.
 434 Webster, D. A. (1948). Relation of temperature to survival and incubation of the eggs of smallmouth
 435 bass (*Micropterus dolomieu*). *Transactions of the American Fisheries Society*, 75(1), 43–47. [https:](https://doi.org/10.1577/1548-8659(1945)75%5B43:ROTTSA%5D2.0.CO;2)
 436 [//doi.org/10.1577/1548-8659\(1945\)75%5B43:ROTTSA%5D2.0.CO;2](https://doi.org/10.1577/1548-8659(1945)75%5B43:ROTTSA%5D2.0.CO;2)
 437 West-Eberhard, M. J. (2003). *Developmental Plasticity and Evolution*. Oxford University Press.