

₁ Chapter 1

₂ The Pella-Tomlinson Model

Abstract

Stock assessments often assume a two-parameter functional form (e.g., Beverton-Holt or Ricker) for the expected recruitment produced by a given level of spawning output. Mangel et al. (2013) and others have shown that biological reference points such as $\frac{F^*}{M}$ and $\frac{B^*}{B(0)}$ are largely determined by a single parameter (steepness) when using two-parameter relationships. These functions introduce strong correlations between reference points that are pre-determined by the functional form, rather than a biological characteristic of the stock. Mangel et al. note that use of a three-parameter stock-recruitment relationship allows for independent estimation of these reference points. This research seeks to understand the nature of biases in reference points resulting from fitting a two-parameter functional form when the true relationship follows a three-parameter stock-recruitment relationship. This work demonstrates the useful limits of misspecified two-parameter models, and suggests the mechanisms of model failure which arise from mapping a three-dimensional parameter space into two dimensions.

1 Introduction

The most fundamental model in modern fisheries management is the surplus-production model. These models focus on modeling population growth via nonlinear parametric ordinary differential equations (ODE). Key management quantities called reference points (RPs) are commonly derived from the ODE equilibrium equations and depend upon the parameterization of biomass production. Two-parameter forms of the production function have been shown to limit the theoretical domain of RPs (Mangel et al., 2013). The limited RP-space of two parameter models are a major source of model misspecification for RPs and thus induce bias in RP estimation. The behavior of RP estimation bias is not well understood and as a result often underappreciated. A metamodeling approach is developed here to describe RP biases and explore mechanisms of model failure under the most common two parameter models.

Data for a typical surplus-production model comes in the form of an index of abundance through time which is assumed to be proportional to the reproducing biomass for the population of interest. The index is often observed alongside a variety of other known quantities, but at a minimum, each observed index will be observed in the presence of some known catch for the period. Figure (1.1) shows the classic Namibian Hake dataset exemplifying the form.

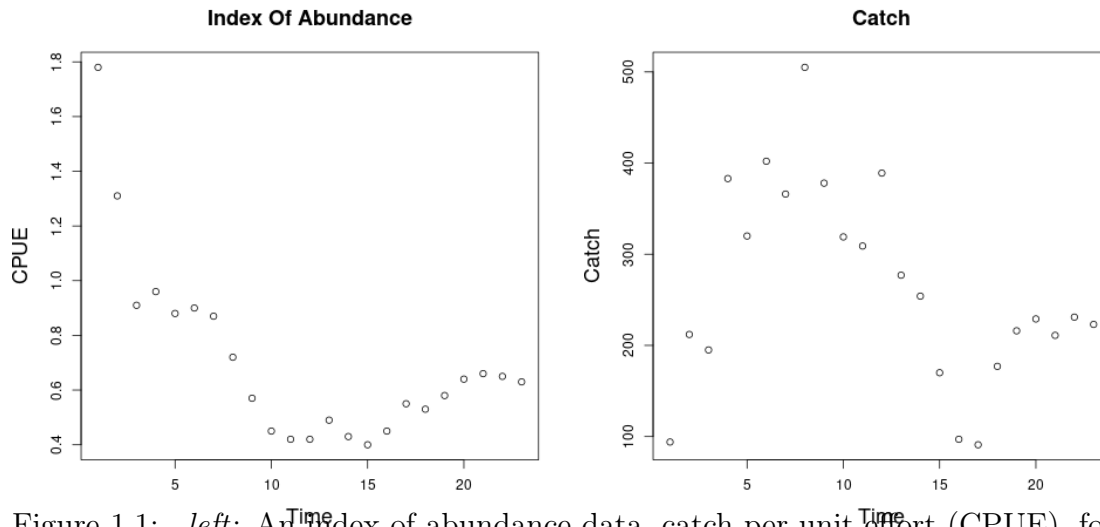


Figure 1.1: *left*: An index of abundance data, catch per unit effort (CPUE), for Namibian Hake from 1965 to 1987 (Hilborn & Mangel, 1997). *right*: The associated catch data for Namibian Hake over the same time period.

Indices are assumed to have multiplicative log-normal errors, and thus the following observation model arises naturally,

$$I_t = qB_te^\epsilon \quad \epsilon \sim N(0, \sigma^2). \quad (1.1)$$

Above q is often referred to as the “catchability parameter”; it serves as the proportionality constant mapping between the observed index of abundance and biomass. σ^2 models residual variation. Biologically speaking q and σ^2 are often treated as nuisance parameters with the “biological parameters” entering the model through a process model on biomass.

Biomass is assumed to evolve as an ODE; in this case I focus on the following form

$$\frac{dB}{dt} = P(B(t); \boldsymbol{\theta}) - Z(t)B(t). \quad (1.2)$$

Here biomass is assumed to change in time by two processes, net production of biomass into the population, $P(B)$, and various sources of biomass removal, Z , from the population.

Firstly, the population grows through a production function, $P(B)$. Production in this setting is defined as the net biomass increase due to all reproduction and maturation processes. The production function is assumed to be a parametric (generally non-linear) function relating the current biomass of the population to an aggregate production of biomass.

Secondly, the population decreases as biomass is removed by various sources that are assumed to remove biomass linearly with biomass. Above, $Z(t)$, is an aggregate rate of removal. When the fishing rate, $F(t)$, is the only source of removal $Z(t) = F(t)$, however often models will also included other linear terms in $Z(t)$. Commonly the rate of “natural mortality”, M , is also included as an additional term so that $Z(t) = M + F(t)$.

From a management perspective a major goal of modeling is to accurately infer a quantity known as *maximum sustainable yield* (MSY). One could maximize simple yield at a particular moment in time (and only for that moment) by fishing all available biomass in that moment. This strategy is penny-wise but pound-foolish (not to mention ecologically devastating) since it doesn’t leave biomass in the population to reproduce in the future. We seek to fish in a way that allows (or even encourages) future productivity in the population. This is accomplished by maximizing the equilibrium level of catch over time. Equilibrium yield is considered by

replacing the steady state biomass (\bar{B}) in the assumed form for catch, so that $\bar{Y} = F\bar{B}(F)$, where $\bar{\cdot}$ indicates a value at steady state. MSY is found by maximizing $\bar{Y}(F)$ with respect to F , and F^* is the fishing rate at MSY. Going forward let $*$ decorate any value derived under the condition of MSY.

Fisheries are very often managed based upon reference points which serve as simplified heuristic measures of population behavior. The mathematical form of RPs depends upon the model assumptions through the production function. While a number of different RPs exist which describe the population in different (but related) ways, the most common RPs revolve around the concept of MSY (or robust ways of measuring MSY (Hilborn, 2010; Punt et al., 2016)). Here the focus is primarily on the RPs $\frac{B^*}{B(0)}$ and F^* ($\frac{F^*}{M}$ when appropriate) for their pervasive use in modern fisheries (Punt & Cope, 2019).

F^* is the afore mentioned fishing rate which results in MSY. $\frac{B^*}{B(0)}$ is the depletion of the stock at MSY. That is to say $\frac{B^*}{B(0)}$ describes the fraction of the unfished population biomass that will remain in the equilibrium at MSY. In general $F^* \in \mathbb{R}^+$ and $\frac{B^*}{B(0)} \in (0, 1)$, however under the under the assumption of a two parameter production function production models will be structurally unable to capture the full theoretical range of RPs.

Many of the most commonly used production functions depend only on two parameters. For example, the Schaefer model depends only on the biological parameters r and K , and limits RP inference so that under the Schaefer model $(F^*, \frac{B^*}{B(0)}) \in (\mathbb{R}^+, \frac{1}{2})$. The two parameter Fox model (Fox Jr., 1970) limits $(F^*, \frac{B^*}{B(0)}) \in (\mathbb{R}^+, \frac{1}{e})$. Similarly the two parameter Cushing (Cushing, 1971), Beverton-Holt (Beverton & Holt, 1957, BH) and Ricker (Ricker, 1954) production functions do not model the full theoretical space of RPs (Mangel et al., 2013; Yeakel & Mangel, 2015).

The bias-variance trade-off (Ramasubramanian & Singh, 2017) makes it clear that the addition of a third parameter in the production function will necessarily reduce estimation bias. However the utility of this bias reduction is still under debate because the particular mechanisms and behavior (direction and magnitude) of these biases for key management quantities are not fully understood or described. Lee et al. (2012) provides some evidence that estimation of productivity parameters are dependent on biomass contrast as well as model specification. Conn et al. (2010) comes to similar conclusions via calibration modeling

techniques. These studies indicate important factors that contribute to inferential failure. However they do not offer mechanisms of model failure, nor do their experimental designs allow for the control of different types of model misspecification.

In this study I consider the behavior of inference when index data are simulated from three parameter PT and Schnute production models, but the simulated data are fit using intentionally misspecified two parameter logistic or BH production models. The work begins with a derivation of RPs under the three parameter models. A method is then presented for generating simulation designs based on the parametric form of RPs which serves as a control on the nature of simulated model misspecification. Finally a Gaussian Process (GP) metamodel (Gramacy, 2020) is constructed for exploration and analysis of RP biases.

A key insight of this approach is that bias is considered broadly across RP-space to uncover patterns and correlations between RPs. The GP metamodel is explicit about trade-offs between RPs so as to inform the full utility of reducing bias, as well as to suggest mechanisms for understanding what causes bias. Further, the effect of contrast on estimation is considered together with model misspecification.

2 Methods

2.1 Pella-Tomlinson Model

The three parameter Pella-Tomlinson (PT) family has a convenient form that includes, among others (Fox Jr., 1970; Rankin & Lemos, 2015), the logistic production function as a special case. PT production function is parameterized so that $\boldsymbol{\theta} = [r, K, \gamma]$ and the family takes the following form,

$$P_p(B; [r, K, \gamma]) = \frac{rB}{\gamma - 1} \left(1 - \left(\frac{B}{K} \right)^{(\gamma-1)} \right). \quad (1.3)$$

γ is a parameter which breaks PT out of the restrictive symmetry of the logistic curve. In general $\gamma \in (1, \infty)$, with the logistic model appearing in the special case of $\gamma = 2$, and the Fox model appearing as a limiting case as $\gamma \rightarrow 1$. The parameter r controls the maximum reproductive rate of the population in the absence of competition for resources (i.e. the slope of production function at the origin). K is the so called "carrying capacity" of the population. In this context the carrying capacity can be formally stated as steady state biomass in the absence of fishing (i.e. $\bar{B}(0) = K$). In Figure (3.1) PT recruitment is shown for a range of parameter values so as to demonstrate the various recruitment shapes that can be achieved by PT recruitment.

While the form of the PT curve produces some limitations (Fletcher, 1978), importantly the introduction of a third parameter allows enough flexibility to fully describe the space of reference points used in management. To see this, the reference points are analytically derived for the PT model below.

PT Reference Points

With $B(t)$ representing biomass at time t , under PT production, the dynamics of biomass are defined by the following ODE,

$$\frac{dB}{dt} = \frac{rB}{\gamma - 1} \left(1 - \left(\frac{B}{K} \right)^{\gamma-1} \right) - FB. \quad (1.4)$$

An expression for the equilibrium biomass is attained by setting Eq (1.4) equal to zero, and rearranging the resulting equation to solve for B . Thinking of the result as a function

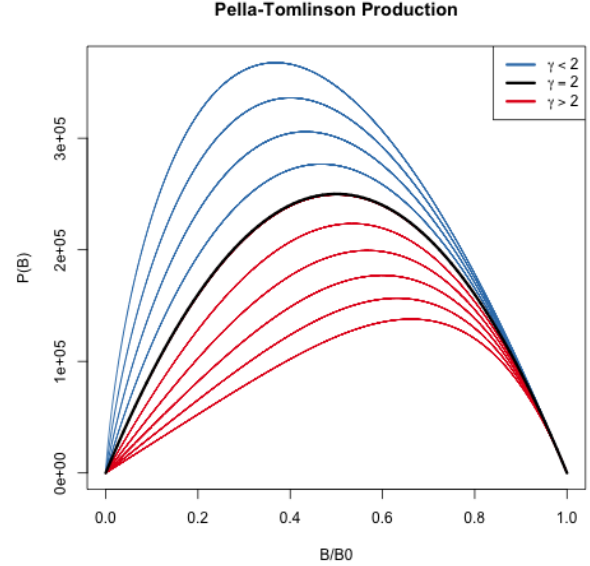


Figure 1.2: The Pella-Tomlinson production function plotted across a variety of parameter values. The special cases of Logistic production is shown in black, and the left-leaning and right-leaning regimes are shown in blue and red respectively.

of F gives,

$$\bar{B}(F) = K \left(1 - \frac{F(\gamma - 1)}{r} \right)^{\frac{1}{\gamma-1}}. \quad (1.5)$$

At this point it is convenient to notice that $\bar{B}(0) = K$. The expression for B^* is given by evaluating Eq (1.5) at F^* . To get an expression for F^* , the equilibrium yield is maximized with respect to F ,

$$F^* = \operatorname{argmax}_F F \bar{B}(F). \quad (1.6)$$

In the case of PT production this maximization can be done analytically, by differentiating the equilibrium yield with respect to F as follows,

$$\frac{d\bar{Y}}{dF} = \bar{B}(F) + F \frac{d\bar{B}}{dF} \quad (1.7)$$

$$\frac{d\bar{B}}{dF} = -\frac{K}{r} \left(1 - \frac{F(\gamma - 1)}{r} \right)^{\frac{1}{\gamma-1}-1}. \quad (1.8)$$

Setting Eq (1.7) equal to 0, substituting $\bar{B}(F)$ and $\frac{d\bar{B}}{dF}$ by Equations (1.5) and (1.8) respectively, and solving for F produces the following expression for the fishing rate required to produce MSY,

$$F^* = \frac{r}{\gamma} \quad (1.9)$$

Plugging the above expression for F^* back into Eq (1.5) gives the following expression for biomass at MSY,

$$B^* = K \left(\frac{1}{\gamma} \right)^{\frac{1}{\gamma-1}}. \quad (1.10)$$

The above derived expressions for $\bar{B}(0)$, B^* , and F^* can then be used to build a specific analytical form for the biological reference points in terms of only productivity parameters.

$$F^* = \frac{r}{\gamma} \qquad \frac{B^*}{\bar{B}(0)} = \left(\frac{1}{\gamma} \right)^{\frac{1}{\gamma-1}} \quad (1.11)$$

131 Simulation

Generating simulated indices of abundance from the PT model requires inverting the relationship between $\left(F^*, \frac{B^*}{B(0)}\right)$, and (r, γ) . It is not generally possible to analytically invert this relationship for many three parameter production functions (Punt & Cope, 2019; J. T. Schnute & Richards, 1998). Most three parameter production functions lead to RPs that require expensive numerical methods to invert; more over the numerical inversion procedure can often be unstable. That said, for the case of PT this relationship is analytically invertible, and leads to the following relationship

$$r = \gamma F^* \qquad \gamma = \frac{W\left(\frac{B^*}{B(0)} \log\left(\frac{B^*}{B(0)}\right)\right)}{\log\left(\frac{B^*}{B(0)}\right)}. \quad (1.12)$$

Above W is the Lambert product logarithm function. More details about this derivation, and the Lambert product logarithm, are given in Appendix (6).

Using Eq. (1.12) to obtain production parameters, a PT production model can be fully defined for any combination of the RPs F^* and $\frac{B^*}{B(0)}$. Since K does not enter the RP calculation its value is fixed arbitrarily at 10000.

Indices of abundance are simulated from the three parameter PT production model broadly over the space of F^* and $\frac{B^*}{B(0)}$ via a space filling design as described in Section (0.2). A small amount of residual variation, $\sigma = 0.01$, is added to the simulated index, and these data are then fit with a Schaefer model, at various degrees of misspecification, so as to observe the effect of productivity model misspecification upon RP inference.

142 PT Design

Letting \mathcal{F} and \mathcal{B} be regular grids, of size $n = 100$, on $F^* \in (0.1, 0.7)$ and $\frac{B^*}{B_0} \in (0.2, 0.6)$ respectively, a LHS design of size 100 is collected among the cells produced by $\mathcal{F} \times \mathcal{B}$.

Each of the sampled LHS design locations represent a unique PT model with the sampled RP values. Since the relationship mapping RPs analytically to productivity parameters can be found for the PT model, LHS designs the the PT model are computed directly in RP space and Eq. (1.12) is used to map the sampled RP design locations to PT productivity parameters.

2.2 Gaussian Process Metamodel

At its core, a metamodel is simply a model of some mapping of inputs to outputs (the mapping itself is typically defined by a computer model). By modeling the mapping with a statistical model (that explicitly defines the relevant features of the mapping) a metamodel defines a specific ontology for the mapping. By simulating examples of the mapping, the inferential infrastructure of the statistical model is used to empirically learn an effective emulation of the mapping within the ontology defined by the statistical model. The predictive infrastructure of the statistical model is then useful as an approximate abstraction of the system itself to better understand the system through further data collection, cheap approximation of the mapping, and/or study of the mapping itself.

In this setting, the aim of metamodeling is to study how well RPs are inferred when typical two parameter models of productivity (Logistic and BH) are misspecified for populations that are actually driven by more complicated dynamics. The simulation design, \mathbf{X} , provides a sample of different population dynamics that are driven by three parameter production functions broadly in RP space. By simulating index of abundance data from the three parameter model, and fitting those data with the two parameter production model, we observe particular instances of how well RPs are inferred at the given misspecification of the two parameter model relative to the true three parameter production model. By gathering all of the simulated instances of how RPs are inferred (under the two parameter model), we form a set of example mappings to train a metamodel which represents the mapping of true RPs (under the three parameter model) to estimates of RPs under the misspecified two parameter production model. The metamodel is essentially a surrogate for inference under the misspecified two parameter production model that controls for the specific degree of model misspecification.

A flexible GP model is assumed for the structure of the metamodel to describe the mapping of RPs under misspecified two parameter models of productivity. A GP is a stochastic process generalizing the multivariate normal distribution to an infinite dimensional analog. GP models are often specified primarily through the choice of a covariance (or correlation) function which defines the relationship between locations in the input space. Typically corre-

179 lation functions are specified so that points closely related in space result in correlated effects
 180 in the model. In this setting the inputs to the GP metamodel are the space of reference points
 181 which define the simulated three parameter production models.

While index of abundance data are generated from three parameter models, at each design location of the simulation, fitting the restricted two parameter model results in a maximum likelihood estimate (MLE; and associated estimation uncertainty) of each of the productivity parameters (i.e. Schaefer: $[\log(r), \log(K)]$, BH: $[\log(\alpha), \log(\beta)]$). To simplify the specification of the metamodel, let \mathbf{y} be a vector collecting the fitted MLEs for one of the productivity parameters, and let $\boldsymbol{\omega}$ be a vector of estimates of the estimator variances (via the inverted Fisher information) at each \mathbf{y} . Each of the fitted productivity parameter estimates are then modeled using independent instances of the following GP metamodel.

$$\begin{aligned}\mathbf{y} &= \beta_0 + \mathbf{X}\boldsymbol{\beta} + \mathbf{v} + \boldsymbol{\epsilon} \\ \mathbf{v} &\sim N_n(\mathbf{0}, \tau^2 \mathbf{R}_\ell) \\ \boldsymbol{\epsilon} &\sim N_n(\mathbf{0}, \boldsymbol{\omega}' \mathbf{I})\end{aligned}\tag{1.13}$$

182 \mathbf{X} is the $n \times 2$ LHS design matrix of RPs for each simulated three parameter data
 183 generating model as described in Section (0.2). ϵ models independent normally distributed
 184 error, which provides an ideal mechanism for propagating uncertainty from inference in the
 185 simulation step into the metamodel. By matching each \mathbf{y}_i with an observed ω_i variance term,
 186 ϵ serves to down weight the influence of each \mathbf{y}_i in proportion to the inferred production model
 187 sampling distribution uncertainty. This has the effect of smoothing the GP model in a way
 188 similar to the nugget effect (Gramacy & Lee, 2012), although the application here models
 189 this effect heterogeneously.

The term, \mathbf{v} , contains spatially correlated GP effects. The correlation matrix, \mathbf{R}_ℓ describes how RPs close together in the simulation design are more correlated than those that are far away. This spatial effect is modeled with a squared exponential correlation function,

$$R(\mathbf{x}, \tilde{\mathbf{x}}) = \exp \left(\sum_{i=1}^2 \frac{-(x_i - \tilde{x}_i)^2}{2\ell_j^2} \right).\tag{1.14}$$

R has an anisotropic separable form which allows for differing length scales, ℓ_1 and ℓ_2 , in the different RP axes. The flexibility to model correlations separately in the different RP axes is key due to the differences in the extent of the RP domains marginally. The metamodel parameters β_0 , $\boldsymbol{\beta}$, τ^2 , ℓ_1 and ℓ_2 are fit via MLE against the observations \mathbf{y} , \mathbf{X} , and $\boldsymbol{\omega}$ from simulation fits.

Fitting the metamodel allows for a full predictive description of inference under the misspecified restricted models. Predictive estimates are obtained via kriging (Cressie, 2015)

$$\hat{y}(\mathbf{x}) = \beta_0 + \mathbf{x}\boldsymbol{\beta} + \mathbf{r}(\mathbf{x})'\mathbf{R}_\ell^{-1}\left(\mathbf{y} - (\beta_0 + \mathbf{X}\boldsymbol{\beta})\right) \quad (1.15)$$

$\hat{y}(\mathbf{x})$ is the predicted value of the modeled productivity parameter MLE under the two parameter production model, when the index of abundance is generated from the three parameter production model at RP location \mathbf{x} . $\mathbf{r}(\mathbf{x})$ is a vector-valued function of correlation function evaluations for the predictive location \mathbf{x} against all observations in \mathbf{X} (i.e. $\mathbf{r}(\mathbf{x}) = \mathbf{R}(\mathbf{x}, \mathbf{x}_i) \forall \mathbf{x}_i \in \mathbf{X}$).

While metamodeling occurs on the inferred productivity parameters of the restricted production model, the metamodel can also be used to build estimates of major biological RPs. For the BH model the relevant transformations for relating productivity parameters with RPs are given in Eqs. (2.5, 2.8) with γ fixed to -1; for the Schaefer model $\hat{B}^* = \frac{\hat{K}}{2}$ and $\hat{F}^* = \frac{\hat{r}}{2}$. Applying the metamodel predictive surfaces on the scale of RP estimates allows for the quantification of estimation bias that is induced by fitting a misspecified two parameter production model to indices of abundance generated under three parameter productivity.

2.3 Catch

It is known that contrast in the observed index and catch time series can effect inference on the productivity parameters (Hilborn & Walters, 1992). In this setting contrast refers to changes in the long term trends of index data. Figure (1.3, *right*) demonstrates an example of biomass that includes contrast induced by catch. It is not well understood how contrast may factor into inferential failure induced by model misspecification. Thus catch is parameterized so as to allow for a spectrum of possible contrast simulation settings.

216 Catch is parameterized so that $F(t)$ can be controlled with respect to F^* . Recall that
 217 catch is assumed to be proportional to biomass, so that $C(t) = F(t)B(t)$. To control $F(t)$
 218 with respect to F^* , $C(t)$ is specified by defining the quantity $\frac{F(t)}{F^*}$ as the relative fishing rate.
 219 $B(t)$ is defined by the solution of the ODE, and F^* is defined by the biological parameters
 220 of the model. By defining $\frac{F(t)}{F^*}$, catch can then be written as $C(t) = F^* \left(\frac{F(t)}{F^*} \right) B(t)$.

221 Intuitively $\frac{F(t)}{F^*}$ describes the fraction of F^* that $F(t)$ is specified to for the current $B(t)$.
 222 When $\frac{F(t)}{F^*} = 1$, $F(t)$ will be held at F^* , and the solution of the ODE brings $B(t)$ into
 223 equilibrium at B^* . When $\frac{F(t)}{F^*}$ is held constant in time biomass comes to equilibrium as an
 224 exponential decay from K approaching B^* . When $\frac{F(t)}{F^*} < 1$, $F(t)$ is lower than F^* and $B(t)$ is
 225 pushed toward $\bar{B} > B^*$. Contrarily, when $\frac{F(t)}{F^*} > 1$, $F(t)$ is higher than F^* and $B(t)$ is pushed
 226 toward $\bar{B} < B^*$; the precise values of \bar{B} can be calculated from the steady state biomass
 227 equations provided above and depend upon the specific form of the production function.

For the simulations presented here, a family of fishing behaviors are considered where the fishing rate accelerates as technology and fishing techniques improve rapidly until management practices are applied, which ultimately brings fishing into equilibrium at F^* . This is parameterized as three distinct phases, over a total of 45 units of time, with each phase lasting 15 time units. The specific form is given below.

$$\frac{F(t)}{F^*} = ae^{bt}\mathbf{1}_{0 \leq t < 15} + (d - ct)\mathbf{1}_{15 \leq t < 30} + \mathbf{1}_{30 \leq t \leq 45} \quad (1.16)$$

The first term of Eq(1.16) is an exponential increase in fishing, the second term is a linear decline in relative fishing as initial management practices are applied, and the third term, $\mathbf{1}_{30 \leq t \leq 45}$, simply holds the fishing rate at F^* there after. These three phases are controlled by the four parameters a , b , c , and d . By enforcing that the interface of the phases meet at χ_{max} and 1 respectively the relative fishing series is reduced to a two parameter family.

$$a = e^{\log(\chi_{max}) - 15b} \quad b = \frac{1}{t - 15} \log \left(\frac{\chi_{min}}{\chi_{max}} \right) \quad (1.17)$$

$$c = \frac{\chi_{max} - 1}{15 - 1} \quad d = 15c + \chi_{max} \quad (1.18)$$

228 By further specifying $\chi_{max} = 1.6^x$ and $\chi_{min} = 0.4^x$ the two parameters χ_{max} , and χ_{min}

can be reduced to the single parameter χ . The tuning parameter χ then singularly controls contrast that appears in time series data.

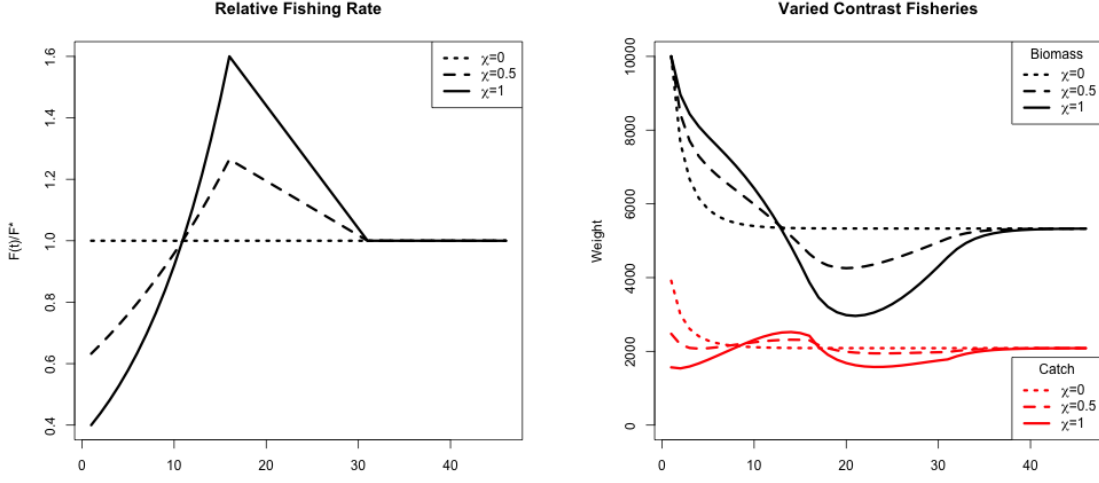


Figure 1.3: (left) Relative fishing with low, medium, and high contrast. (right) Population biomass and catch at each associated level of contrast.

When $\chi = 0$, the relative fishing rate is a constant at 1 to create a low contrast simulation environment. As χ increases Eq (1.16) induces more and more contrast in the observed index and catch time series until $\chi = 1$ which produces a high contrast simulation environment. Figure (1.3) demonstrates a spectrum of contrast simulation environments as well as the time series data they induce in the solution of the production model ODE.

2.4 Two Parameter Production Model Inference

The simulated mapping results from fitting an intentionally misspecified two parameter production model to index of abundance data that are generated from a more complex three parameter model of productivity. Thus, let I_t be an index of abundance simulated from the three parameter PT or Schnute production models at time $t \in \{1, 2, 3, \dots, T\}$. However the fitted model is specified to be intentionally misspecified so that the fitted model is driven by a two parameter Schaefer, or BH production model respectively.

The observation model for the fitted model is log-normal such that,

$$I_t|q, \sigma^2, \boldsymbol{\theta} \sim LN(qB_t(\boldsymbol{\theta}), \sigma^2). \quad (1.19)$$

$B_t(\boldsymbol{\theta})$ is defined by the solution of the ODEs defined by the Schaefer, or BH models. For

the Schaefer model $\boldsymbol{\theta} = [r, K]$, and for the BH model $\boldsymbol{\theta} = [\alpha, \beta]$. From the perspective of the fitted model, the observed I_t are assumed independent conditional on q , σ^2 , r , K and the two parameter ODE model for biomass. Thus the log likelihood can be written as

$$\log \mathcal{L}(q, \sigma^2, \boldsymbol{\theta}; I) = -\frac{T}{2} \log(\sigma^2) - \frac{1}{2\sigma^2} \sum_t \log \left(\frac{I_t}{qB_t(\boldsymbol{\theta})} \right)^2. \quad (1.20)$$

In this setting, q is fixed at the true value of 0.0005 to focus on the inferential effects of model misspecification on biological parameters. σ^2 and $\boldsymbol{\theta}$ are reparameterized to the log scale and fit via MLE. Reparameterizing the parameters to the log scale improves the reliability of optimization, in addition to facilitating the use of Hessian information for estimating MLE standard errors.

Given that the biological parameters enter the likelihood via a nonlinear ODE, and further the parameters themselves are related to each other nonlinearly, the likelihood function can often be difficult to optimize. A hybrid optimization scheme is used to maximize the log likelihood to ensure that a global MLE solution is found. The R package GA ([Scrucca, 2013, 2017](#)) is used to run a genetic algorithm to explore parameter space globally. Optimization periodically jumps into the L-BFGS-B local optimizer to refine optima within a local mode. The scheme functions by searching globally, with the genetic algorithm, across many initial values for starting the local gradient-based optimizer. The genetic algorithm serves to iteratively improve hot starts for the local gradient-based optimizer. Additionally, optimization is only considered to be converged when the optimum results in an invertible Hessian at the found MLE.

2.5 Continuous model formulation

An important (and often overlooked) implementation detail is the solution to the ODE which defines the progression of biomass through time. As a statistical model it is of paramount importance that this ODE not only have a solution, but also that the solution be unique. Of primary concern, uniqueness of the ODE solution is necessary for well conditioned inference.

If the form of $\frac{dB}{dt}$ is at least Lipschitz continuous, then the Cauchy-Lipschitz-Picard theorem provides local existence and uniqueness of $B(t)$. Recall from Eq(1.2) that $\frac{dB}{dt}$ is

separated into a term for biomass production, $P(B)$, and a term for removals, $Z(t)B(t)$. For determining Lipschitz continuity of $\frac{dB}{dt}$, the smallest Lipschitz constant of $\frac{dB}{dt}$ will be the sum of the constants for each of the terms $P(B)$ and $Z(t)B(t)$ separately. Typically any choice of $P(B)$ will be continuously differentiable, which implies Lipschitz continuity. At a minimum $Z(t)$ typically contains fishing mortality as a function of time $F(t)$ to model catch in time as $C(t) = F(t)B(t)$. $Z(t)$ may or may not contain M , but typically M is modeled as stationary in time and does not pose a continuity issue, unlike some potential assumptions for $C(t)$.

In practice $C(t)$ is determined by a series of observed, assumed known, catches. Catch observations are typically observed on a quarterly basis, but in practice may not be complete for every quarter (or year) of the modeled period. It is overwhelmingly common to discretize the ODE in time via Euler's method with integration step sizes to match the observation frequency of the modeled data. This is often computationally convenient when the underlying species dynamics are reasonably well behaved, however when the dynamics model is used as a statistical model, with the goal of inferring the behavior of the underlying species dynamics, the regularity of the dynamics are not guaranteed. An implicit assumption of continuity of catch in time provides the necessary regularity for the statistical model. Furthermore a continuous handling of the dynamics provides improved accuracy in evaluating the ODE, particularly when inferring productivity parameters which largely control the regularity of the dynamics.

While there are many ways to handle catch continuity, here I assume that catches accrue linearly between observed catches. This assumption defines the catch function as a piecewise linear function of time, with the smallest Lipschitz constant for the catch term defined by the steepest time segment of the catch function. This assumption represents one of the simplest ways of handling catch, while retaining Lipschitz continuity overall. Furthermore linearly interpolated catch is adequately parsimonious for the typical handling of catches.

Integration and Stiffness

As previously mentioned, the overwhelming majority of implementations of stock assessment models discretized the ODE using Euler's method with the integration step sized fixed so as to match the observation frequency. In this setting we explore model parameterizations that

explore the full extent of biologically relevant reference points. This exercise produces some combinations of parameters that result in numerically stiff ODEs.

The concept of stiffness in ODEs is hard to precisely characterize. [Wanner and Hairer \(1996, p.2\)](#) describe stiffness in the following pragmatic sense, “Stiff equations are problems for which explicit methods don’t work”. It is hard to make this definition more mathematically precise, but this is a consistent issue for models of fast growing species in the low contrast simulation. Euler’s method, as often implemented, is particularly poorly suited for these stiff regions of parameter space. In these stiff regions it is necessary to integrate the ODE with an implicit integration method.

Several of the most common implicit methods were tried including the Livermore Solver for ODEs (lsode), and the Variable Coefficient ODE Solver (vode) as implemented in the deSolve package of R ([Soetaert et al., 2010](#)). The difference between implicit solvers is negligible, while explicit methods result in wildly varying solutions to the ODE in stiff regions of parameter space. Results shown here are computed using the lsode integration since it runs relatively quickly and has a relatively smaller footprint in system memory.

3 Results

3.1 PT/Schaefer

An MSY -Optimal Catch History

When $F(t)$ is held constant at F^* , as it is in the "low contrast" simulation setting, $B(t)$ comes to equilibrium as an exponential decay from K to B^* . Understanding model misspecification bias is simplified in this setting due to the relative simplicity that this induces in $B(t)$. However this simplicity is known to poorly inform estimates of r , and thus F^* , due to the limited range of the production function that is observed (Hilborn & Walters, 1992).

Figure (1.4) shows four of the most misspecified example production function fits as compared to the true data generating PT production functions. The rug plots below each set of curves show how the observed biomasses decay exponentially from K to B^* in each case. In particular, notice how observations only exist where the PT biomass is greater than B^* . Due to the leaning of the true PT curves, and the symmetry of the logistic parabola, the logistic curve only observes information about its slope at the origin from data observed on the right portion of the PT curves. The top two panels of Figure (1.4) shows PT data generated such that $\frac{B^*}{B(0)} > 0.5$; in these cases PT is steeper to the right of B^* than it is on the left, and so the the logistic curve over-estimates r , and consequently also over-estimates F^* . The

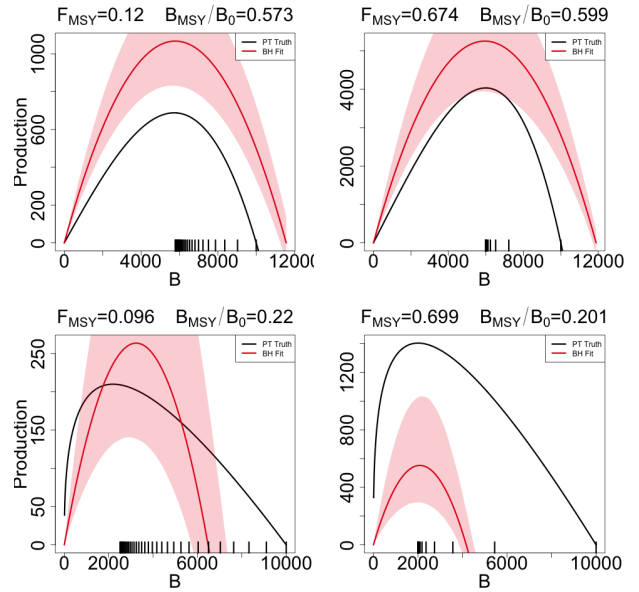


Figure 1.4: A comparison of the true PT production function (in black) and the estimated logistic curve (in red) with 95% CI shown. The examples shown represent the four corners of maximum model misspecification in the simulated RP-space. Observed biomasses are plotted in the rug plots below the curves.

bottom two panels of Figure (1.4) show PT data generated with $\frac{B^*}{B(0)} < 0.5$ and where the vice versa phenomena occurs. PT is shallower to the right of B^* than it is on the left

and so the logistic parabola estimate tends to under estimate F^* .

Metamodeled Trends

Each point in the space of the RPs F^* and $\frac{B^*}{B(0)}$ uniquely identifies a complete PT model with different combinations of parameters values. Recall that when $\gamma = 2$ for the PT model, the PT curve becomes a parabola and is equivalent to the logistic curve of the Schaefer model. Since the logistic curve is symmetric about B^* , the Schaefer model must fix the value of $\frac{B^*}{B(0)}$ at the constant 0.5 for any value of F^* . So the line through RP space defined by $\frac{B^*}{B(0)} = 0.5 \quad \forall \quad F^*$, defines the subset of RP space where $\gamma = 2$ and where the PT model is equivalent to the Schaefer model. For brevity this subset of RP where $\frac{B^*}{B(0)} = 0.5$ will be referred to as the ‘‘Schaefer set’’. Thus simulated data that are generated along the Schaefer set will be the only data that are not misspecified relative to the Schaefer model; as PT data are simulated farther and farther away from this line at $\frac{B^*}{B(0)} = 0.5$ model misspecification of the Schaefer model becomes worse and worse.

While Figure (1.4) demonstrates a real trend in simulation results, individual simulation runs will at best show jittery trends due to the stochastic nature of statistical inference. The GP process metamodel accounts for this stochasticity to focus analysis on the signal in the simulation results. Recall that metamodeling occurs on the scale of the inferred productivity parameters of the restricted production model, by transforming metamodel predictions via Eq. (1.11), metamodeled predictions are obtained for Schaefer RPs. By further subtracting the true data generating PT RPs from the predicted Schaefer RPs at each point in RP space a pattern of inferential RP bias, induced by model misspecification of the Schaefer model, can be seen.

Figure (1.5) shows the pattern of biases the Schaefer model creates when fit to PT data generated at each point of RP space. An equivalent way to think of Figure (1.5) is that since the Schaefer model must estimate RPs in the Schaefer set, the metamodel arrows indicate the mapping that is created by inferring RPs under a misspecified Schaefer model fit to PT data generated at each point over the pictured region.

Since $\frac{B^*}{B_0}$ must be 0.5 under the Schaefer model, biases in the $\frac{B^*}{B_0}$ direction must simply map vertically onto the Schaefer set. Due to this simplified RP geometry under the Schaefer

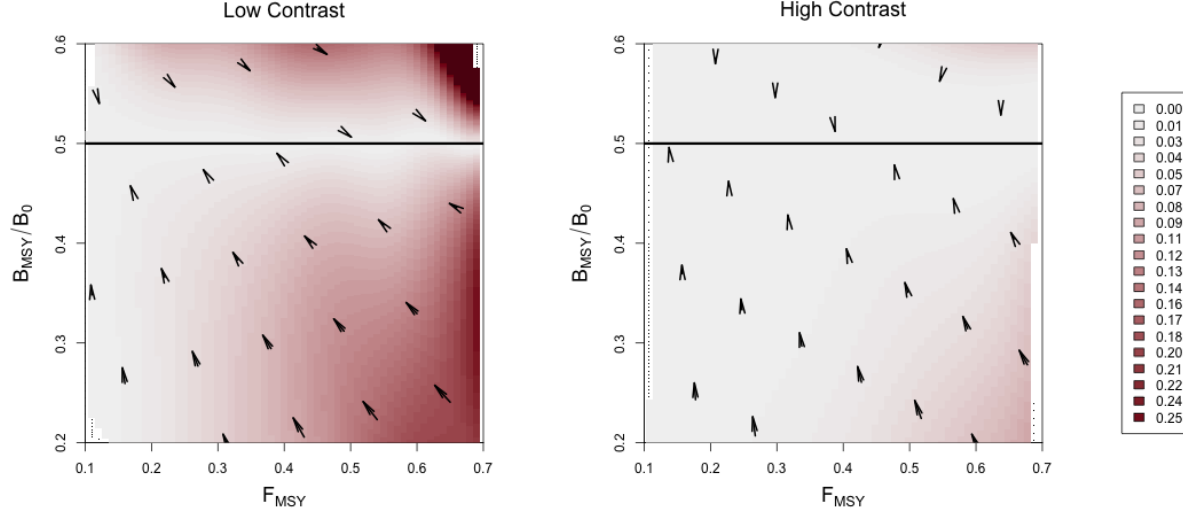


Figure 1.5: Joint bias direction for $(F^*, \frac{B^*}{B_0})$ estimates under the misspecified Schaefer Model. The intensity of color represents the excess bias relative to the shortest possible mapping. Results in the low contrast setting are shown *left*, and the high contrast setting is shown *right*.

model, the degree of bias in $\frac{B^*}{B_0}$ estimation is defined solely by the degree of model misspecification irrespective of F^* . Furthermore, the closest possible point along the Schaefer set that Schaefer model inference could map RPs would be the perfectly vertical mapping. This pattern only contains the strictly necessary bias present in $\frac{B^*}{B_0}$, and zero bias in F^* . Any deviation from this minimal bias pattern is necessarily due to added bias in F^* .

The two simulation settings shown in Figure (1.5) are identical except for the amount of contrast present in the simulated index. The left panel of Figure (1.5) shows RP biases in the low contrast setting, while the right panel shows the high contrast setting. Notice that in the low contrast setting the RP bias pattern is far from the minimum distance mapping, however when contrast is added the mapping becomes much closer to a minimal bias mapping. In the low contrast setting the observed bias is consistent with the pattern and mechanism described in Figure (1.4), where F^* is underestimated for data generated below the Schaefer line and overestimated above the Schaefer set. In the high contrast simulation the mapping is nearly minimal distance with the exception of PT data generated with simultaneously low $\frac{B^*}{B_0}$ and high F^* .

Figure (3.1) demonstrates how bias in F^* estimation decreases as contrast is added to

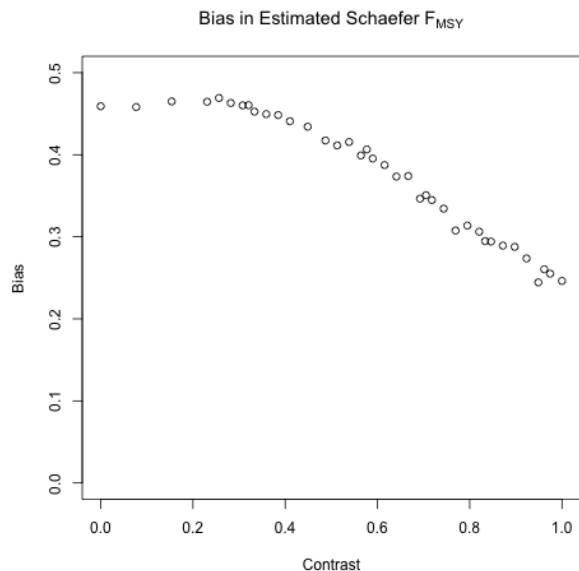


Figure 1.6: Bias in F^* under the Schaefer model when PT data are generated with increasing contrast so that F^* and $\frac{B^*}{B_0}$ are fixed at 0.699 and 0.201 respectively.

PT data as generated in the low $\frac{B^*}{B_0}$ and high F^* regime. By including additional contrast F^* bias is decreased, however parameterizing contrast so as to fully extinguish F^* bias may require a more complex model of fishing.

4 Discussion

Results presented here generally agree with what is known about estimating growth rate parameters (Lee et al., 2012; Conn et al., 2010; Magnusson & Hilborn, 2007). These study's appreciate the role of contrast for estimating growth rates, however they struggle to make generally extensible conclusions since they focus only on a handful of stocks that fall short of forming a random sample of the greater population of possible stock behaviors. The LHS design methods presented here are designed specifically to simulate a representative sample of stocks broadly across the space of possible RPs. Furthermore, the simulation design, taken together with the GP metamodel of productivity parameter estimates, allows this study to control the degree of model misspecification and generalize conclusions about the behavior of productivity estimation within the production model setting presented.

In the presence of contrast, F^* estimation can enjoy very low bias even for a wide range of poorly specified models; conversely in the absence of contrast F^* estimation can suffer very large bias even for slightly misspecified models. This pattern is particularly true for inference

under the Schaefer model where the geometry of the restricted RP set isolates estimation failure of F^* from $\frac{B^*}{B(0)}$. While contrast has a similar impact on F^* estimation under the BH model, the geometry of the BH RP set correlates estimation bias of F^* and $\frac{B^*}{B(0)}$. The GP metamodeling approach reveals a more general pattern that highly informative data sets (high contrast) produces a nearly minimal distance mapping of RPs onto the constrained RP set.

In all cases when model misspecification is removed, even with weakly informative data, RP estimation is unbiased and well estimated. Thus contrast alone is not the only factor leading to inferential failure. Model misspecification is a necessary but not sufficient condition for inducing RP estimation bias. The particular RP bias present depends on the RP geometry of the fitted model and how that geometry is misspecified relative to the data. The RP mapping is then oriented to the RP geometry of the fitted model.

While the relative fishing rate parameterized in Section (2.3) captures a usefully broad spectrum of relevant fishing behaviors, it is still limiting in the amount of information that it can induce. Improved methods for quantifying contrast in fisheries data, and/or methods of discovering more informative fishing behavior, could improve this analysis. In the absence of a maximally informative dataset simulation methods will not fully describe how inference fails, but the methods presented here tell the most complete picture yet, with explicit control of the degree model misspecification, contrast, and a simulation design that allows for uniform representative data generation across biologically meaningful stocks. The results presented here suggest the conjecture that under a maximally informative dataset, RP inference with a two parameter production function will be biased in the direction a shortest distance map from the true RPs onto restricted set of RPs under the two parameter model.

Given the potential for model misspecification of RPs, a minimal distance mapping of RPs represents a best-case scenario where the total bias of RPs, when measured jointly, is minimized. That said, without recognizing the geometry of how two parameter models of productivity limit RP space this may lead to unintuitive implications in RP estimation. For example, due to the shape of the BH RP set a minimal distance mapping ensures that if there is bias in one of $\frac{B^*}{B_0}$ or F^* , there will necessarily be bias in the other RP. However under the Schaefer model, since the RP set is a constant in $\frac{B^*}{B_0}$, bias in F^* is not adulterated in the

same way by bias in $\frac{B^*}{B_0}$ estimation. While models with constant RPs, such as the logistic model $\frac{B^*}{B_0} = \frac{1}{2}$ or the Fox model $\frac{B^*}{B_0} = \frac{1}{e}$, are extremely limited, they can be valuable tools for developing intuition precisely because they isolate RP estimation in their free RPs from the correlated RP biases present in models like the BH or Ricker model.

When one considers the implications of RP bias, overestimation of RPs carries the severe implication of management recommendations potentially leading to overfishing, while underestimation of RP leads to overly conservative management. In this sense, when the true model is not known, the geometry of the BH set together with the metamodeled bias trends makes the BH model a naturally conservative estimator of RPs for most stocks. For most non-BH populations the BH model is likely to make conservative errors in its estimates of F^* and $\frac{B^*}{B_0}$. The one notable exception to the conservatism of the BH model stands for data generated in the Cushing-like regime of Schnute RPs. In this regime the BH model tends to be fairly unbiased overall, however the bias that is present for these populations tends to be overestimation in both RPs, leading to much more severe management consequences for those populations.

The RP bias trends of the Schaefer model demonstrate much less conservatism than the BH overall. For any population with $\frac{B^*}{B_0} < 0.5$, $\frac{B^*}{B_0}$ will be overestimated. When the population comes from the regime where $\frac{B^*}{B_0} > 0.5$, $\frac{B^*}{B_0}$ will be under estimated, but F^* is likely to be overestimated depending on the degree of contrast present in the data. So while the Schaefer model is an intuitive model, it tends to lead to much less conservative RP estimation.

While it is important to recognize these limitations of two parameter models of productivity, we should not solely accept conservatism as a rationale of choosing a BH model of productivity. Increasing the flexibility of the production function by moving toward three parameter models would release the underlying structural limitations (Mangel et al., 2013) that cause these RP biases in the first place. Punt and Cope (2019) considers a suite of possible three parameter curves which could be used instead of current two parameter curves. For all of their benefits, three parameter production functions have their own complicating factors, and the structure present in the Schnute model explored here makes it an intuitive bridge model for developing three parameter models going forward.

- show a schnute fit to data? (Yeakel & Mangel, 2015) Prior

- summary of σ over RP space comparing between models (PT, Schnute, Schnute DD) to show areas of model breakdown.

 - miss-identifying signal for noise.
 - It happens more as the dynamics get more complex.
 - point to the full age structured models.
- show the constrained BH space over a grid of $M, \kappa, \omega, W_\infty$
- Show that the constrained spaces vary only slightly as compared with the consequences of misspecifying the functional form.
- estimating these other quantities (while they can create quite different Biomass series) can only do so much to improve (expand) RP inference as compared with correctly modeling P .
- mapping distance as a function of contrast at $(3.5, 0.5)$
- for LHS grid locations show $\frac{B^*}{B_0}$ and F^* biases for grids in $M \in (0, 0.5)$ For sure in High Contrast, maybe also in Low??.

5 Appendix: Inverting $\frac{B^*}{B(0)}$ and γ for the PT Model

For brevity let $\zeta = \frac{B^*}{B(0)}$.

$$\begin{aligned}\zeta &= \left(\frac{1}{\gamma}\right)^{\frac{1}{\gamma-1}} \\ \zeta &= \gamma \zeta^\gamma \\ \zeta &= \gamma e^{\gamma \log(\zeta)} \\ \zeta \log(\zeta) &= \gamma \log(\zeta) e^{\gamma \log(\zeta)}\end{aligned}$$

The Lambert product logarithm, W , is defined as the inverse function of $z = xe^x$ such that $x = W(z)$. Applying this definition allows for the isolation of γ .

$$\begin{aligned}\gamma \log(\zeta) &= W(\zeta \log(\zeta)) \\ \gamma &= \frac{W(\zeta \log(\zeta))}{\log(\zeta)}\end{aligned}\tag{1.21}$$

The Lambert product logarithm is a multivalued function with a branch point at $-\frac{1}{e}$. The principal branch, $W_0(z)$, is defined on $z \in (-\frac{1}{e}, \infty)$, and the lower branch, $W_{-1}(z)$, is defined on $z \in (-\frac{1}{e}, 0)$. Taken individually, each respective branch is analytic, but cannot be expressed in terms of elementary functions.

When $\zeta \in (0, \frac{1}{e})$ the solution of interest in Eq. (1.12) comes from W_0 . When $\zeta \rightarrow \frac{1}{e}$, the Fox Model emerges as $\gamma \rightarrow 1$. When $\zeta \in (\frac{1}{e}, 1)$ the solution of interest comes from W_{-1} . For the use case presented here, Eq. (1.12) is to be interpreted as,

$$\gamma = \begin{cases} \frac{W_0(\zeta \log(\zeta))}{\log(\zeta)} & \zeta \in (0, \frac{1}{e}) \\ \frac{W_{-1}(\zeta \log(\zeta))}{\log(\zeta)} & \zeta \in (\frac{1}{e}, 1) \end{cases}.\tag{1.22}$$

Prager 2002, Figure(2).

<https://math.stackexchange.com/questions/3004835/is-the-lambert-w-function-analytic-if-not-everywhere-then-on-what-set-is-it-an> <https://researchportal.bath.ac.uk/en/publications/algebraic-properties-of-the-lambert-w-function-from-a-result-of-r>

489 Chapter 2

490 The Schnute Model

0.1 Schnute Model

The Schnute production function is a three parameter generalization of many of the most common two parameter production functions (Deriso, 1980; J. Schnute, 1985). It can be written in the following form, with parameters α , β , and γ ,

$$P_s(B; [\alpha, \beta, \gamma]) = \alpha B(1 - \beta\gamma B)^{\frac{1}{\gamma}}. \quad (2.1)$$

The BH and Logistic production functions arise when γ is fixed to -1 or 1 respectively. The Ricker model is a limiting case as $\gamma \rightarrow 0$. For $\gamma < -1$ a family of strictly increasing Cushing-like curves arise, culminating in linear production as $\gamma \rightarrow -\infty$. These special cases form natural regimes of similarly behaving production functions as seen in Figure (2.1).

The behavior of RP inference under the BH model is of particular interest due to the overwhelming popularity of the BH assumption in fisheries models. Since Schnute production models can represent a quantifiably wide variety of possible productivity behaviors, they present an ideal simulation environment for inquiry of the reliability of inference under the BH assumption.

Under Schnute production, biomass dynamics evolve according to the following ODE,

$$\frac{dB}{dt} = P_s(B; \theta) - (M + F)B. \quad (2.2)$$

This equation largely takes the same form as previously described, except that P_s is the Schnute production function and natural mortality, M , is modeled explicitly here. Natural

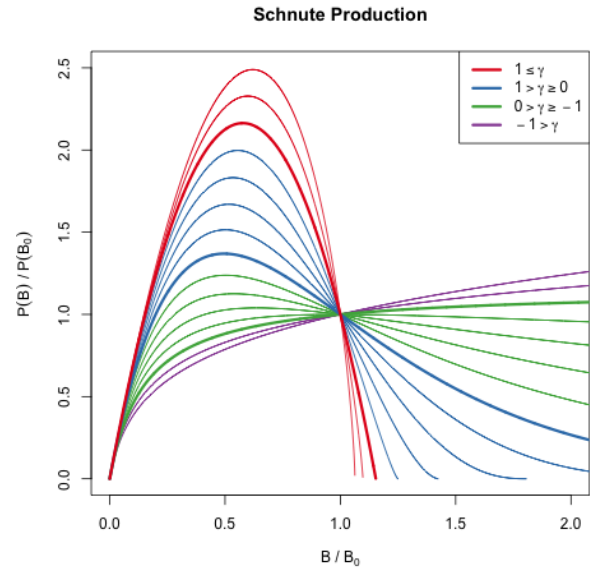


Figure 2.1: The Schnute production function plotted across a variety of parameter values. Regimes of similarly behaving curves are grouped by color.

511 mortality models the instantaneous rate of mortality from all causes outside of fishing. Ex-
 512 plicitly modeling natural mortality is not only a typical assumption of fisheries models, but
 513 is also key to the making RPs well defined over the relevant domain of γ .

The derivation of RPs under Eq. (2.2) follows a similar logic as under the PT model. An expression for equilibrium biomass is attained by setting $\frac{dB}{dt} = 0$ and rearranging the resulting expression to solve for B

$$\bar{B}(F) = \frac{1}{\gamma\beta} \left(1 - \left(\frac{M+F}{\alpha} \right)^\gamma \right). \quad (2.3)$$

The above expression quickly yields B_0 , B^* by evaluation at $F = 0$ and F^* respectively,

$$B_0 = \frac{1}{\gamma\beta} \left(1 - \left(\frac{M}{\alpha} \right)^\gamma \right) \quad (2.4)$$

$$\frac{B^*}{B_0} = \frac{1 - \left(\frac{M+F^*}{\alpha} \right)^\gamma}{1 - \left(\frac{M}{\alpha} \right)^\gamma}. \quad (2.5)$$

Attaining an expression for F^* requires maximization of equilibrium yield, $\bar{Y} = F\bar{B}(F)$, with respect to F . Analytically maximizing proceeds by differentiating \bar{Y} to produce

$$\frac{d\bar{Y}}{dF} = \bar{B}(F) + F \frac{d\bar{B}}{dF} \quad (2.6)$$

$$\frac{d\bar{B}}{dF} = -\frac{1}{\beta} \left(\frac{\left(\frac{M+F}{\alpha} \right)^\gamma}{F+M} \right). \quad (2.7)$$

Setting $\frac{d\bar{Y}}{dF} = 0$, filling in the expressions for $\bar{B}(F)$ and $\frac{d\bar{B}}{dF}$, then rearranging to solve for F^* is less yielding here than it was in the case of the PT model. This procedure falls short of providing an analytical solution for F^* directly in terms of θ , but rather shows that F^* must respect the following expression,

$$0 = \frac{1}{\gamma} - \left(\frac{1}{\gamma} + \frac{F^*}{F^* + M} \right) \left(\frac{F^* + M}{\alpha} \right)^\gamma. \quad (2.8)$$

514 The lack of an analytical solution here is understood. [J. T. Schnute and Richards \(1998,](#)
 515 [pg. 519\)](#) specifically points out that F^* cannot be expressed analytically in terms of produc-
 516 tivity parameters, but rather gives a partial analytical expression for the inverse relationship.

Although parameterized slightly differently, [J. T. Schnute and Richards \(1998\)](#) derives expressions for α and β as a function of RPs and γ .

Since RPs are left without a closed form expression, computing RPs from productivity parameters amounts to numerically solving the system formed by collecting the expressions (2.8), (2.4), and (2.5).

Simulation

For the purposed of simulation, it is not necessary to completely know the precise relationships mapping RPs $\mapsto \theta$ or $\theta \mapsto$ RPs. Simulation only requires enough knowledge of these mappings to gather a list of (α, β, γ) tuples, for data generation under the Schnute model, and the corresponding RPs in some reasonable space-filling design over RP space.

Similarly to [J. T. Schnute and Richards \(1998\)](#), expressions (2.8) and (2.4) are solved for α and β respectively. This leads to the partial mapping $(F^*, B_0) \mapsto (\alpha(\cdot, \gamma), \beta(\cdot, \gamma))$ in terms of RPs and γ . By further working with Eq. (2.5), to identify γ , the following system is obtained,

$$\begin{aligned}\alpha &= (M + F^*) \left(1 + \frac{\gamma F^*}{M + F^*} \right)^{1/\gamma} \\ \beta &= \frac{1}{\gamma B_0} \left(1 - \left(\frac{M}{\alpha} \right)^\gamma \right) \\ \frac{B^*}{B_0} &= \frac{1 - \left(\frac{M + F^*}{\alpha} \right)^\gamma}{1 - \left(\frac{M}{\alpha} \right)^\gamma}.\end{aligned}\tag{2.9}$$

For a population experiencing natural mortality M , by fixing F^* , B_0 , and $\frac{B^*}{B_0}$ the above system can fully specify α and β for a given γ . Notice for a given γ a cascade of closed form solutions for α and β can be obtained. First $\alpha(\gamma)$ can be computed, and then $\beta(\alpha(\gamma), \gamma)$ can be computed. If $\alpha(\gamma)$ is filled back into the expression for $\frac{B^*}{B_0}$, the system collapses into a single onerous expression for $\frac{B^*}{B_0}(\alpha(\gamma), \gamma)$. For brevity, define the function $\zeta(\gamma) = \frac{B^*}{B_0}(\alpha(\gamma), \gamma, F^*, M)$ based on Eq. (2.5).

Inverting $\zeta(\gamma)$ for γ , and computing the cascade of $\alpha(\gamma)$, and then $\beta(\alpha(\gamma), \gamma)$, fully defines the Schnute model for a given $(\frac{F^*}{M}, \frac{B^*}{B_0})$. However inverting ζ accurately is extremely difficult. Inverting ζ analytically is not feasible, and numerical methods for inverting ζ are unstable

and can be computationally expensive. Rather than numerically invert precise values of $\zeta(\gamma)$, γ is sampled so that the overall simulation design is space filling as described in Section (0.2).

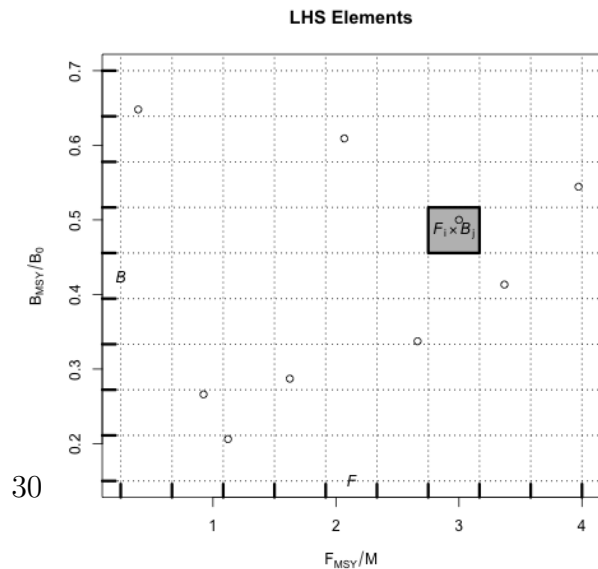
Each design location defines a complete Schnute production model with the given RP values. Indices of abundance are simulated from the Schnute model at each design location, a small amount of residual variation, $\sigma = 0.01$, is added to the simulated index, and the data are then fit with a misspecified BH production model. The design at large captures various degrees of model misspecification relative to the BH model, so as to observe the effect of productivity model misspecification upon RP inference.

0.2 Latin Hypercube Sampling

The goal of space filling design in this setting is to extend the notion of the random sample (and its desirable parameter estimation properties) across the simulated RP domain so as to represent the simulated space as well as possible (Gramacy, 2020). The simple random sample is the classical approach to unbiased parameter estimation, however simple randomness is patchy, often sampling some regions of design space quite densely, while leaving other regions of design space empty. Space filling designs aim to preserve (or enhance) parameter estimation properties across the simulated domain (Devon Lin & Tang, 2015; Stein, 1987), while constraining samples to be spaced in some notion of spread over the entire space. Latin hypercube sampling (McKay et al., 2000, LHS) is among the most foundational of space filling designs used in computer experiments.

A LHS of size n , in the 2 dimensional space defined by RPs, distributes samples so as to spread points across a design region in a broadly representative way. A LHS design extends the notion of a univariate random uniform sample across multiple dimensions so that each margin of the design space enjoys a uniform distribution.

LHS designs achieve this notion of uni-



formity by first partitioning each dimension of the design space into regular grids of size n . By intersecting the grids of each dimension, cells are produced that evenly partition the design space. In two dimensions n^2 cells are produced, from which a total of n samples are taken. Crucially only one sample is taken from a given element of each grid in each dimension so as to reduce clumping of the n samples across the design space.

Schnute Design

Due to the lack of an analytical relationship mapping RPs $\mapsto \theta$, analogous to the PT model's Eq. (1.12), producing a LHS design over Schnute RPs requires a more tactful approach. The structured relationship between the RPs and productivity parameters, described in Section (0.1), allows an approximate LHS to be obtained by a careful navigation of the system of equations seen in Eq. (2.9).

Under the Schnute model, let \mathcal{F} and \mathcal{B} represent regular grids on $\frac{F^*}{M} \in (0.25, 4)$ and $\frac{B^*}{B_0} \in (0.15, 0.7)$ respectively which can serve as the scaffolding for computing an approximate LHS.

Since it is not practical to invert $\zeta(\gamma)$, a uniform sample in $\frac{B^*}{B_0}$ can be obtained by modeling γ as a random variable, with realization γ^* , and thinking of $\zeta(\gamma)$ as its cumulative distribution function (CDF). The aim is to model γ as an easily sampled random variable with a CDF that closely approximates ζ , so that $\zeta(\gamma^*) \sim U(\zeta_{min}, 1)$ as closely as possible. There may be many good models for the distribution of γ , but in this setting the

Given B_0 , M , and F^* :

- 1) Draw $\gamma^* \sim \gamma|F^*, M$.
- 2) Compute $\frac{B^*}{B_0} = \zeta(\gamma^*)$
- 3) Compute $\alpha^* = \alpha(\gamma^*, F^*, M)$
- 4) Compute $\beta^* = \beta(\alpha^*, \gamma^*, M, B_0)$

Figure 2.3: An outline of the sampling procedure for γ given B_0 , M , and F^* .

following distribution is very effective,

$$\gamma \sim \zeta_{min} \delta(\gamma_{min}) + t(\mu, \sigma, \nu) \mathbf{1}_{\gamma > \gamma_{min}}. \quad (2.10)$$

585 Above, t is the density of the three pa-
 586 rameter location-scale family Student's t dis-
 587 tribution with location μ , scale σ , and de-
 588 grees of freedom ν . $\mathbf{1}_{\gamma > \gamma_{min}}$ is an indica-
 589 tor function that serves to truncate the Stu-
 590 dent's t distribution at the lower bound γ_{min} .
 591 $\delta(\gamma_{min})$ is the Dirac delta function evaluated
 592 at γ_{min} , which is scaled by the known value
 593 ζ_{min} ; this places probability mass ζ_{min} at
 594 the point γ_{min} . Since sampling from a Stu-
 595 dent's t distribution is readily doable, sam-
 596 pling from a truncated Student's t mixture
 597 only requires slight modification.

Let T be the CDF of the modeled distri-
 bution of γ . Since the point $(\gamma_{min}, \zeta_{min})$ is
 known from the dynamics of the Schnute model at a given RP, full specification of Eq. (2.10)
 only requires determining the values for μ , σ , and ν which make T best approximate $\zeta(\gamma)$.
 Thus, the values of μ , σ , and ν are chosen by minimizing the L^2 distance between $T(\gamma)$ and
 $\zeta(\gamma)$.

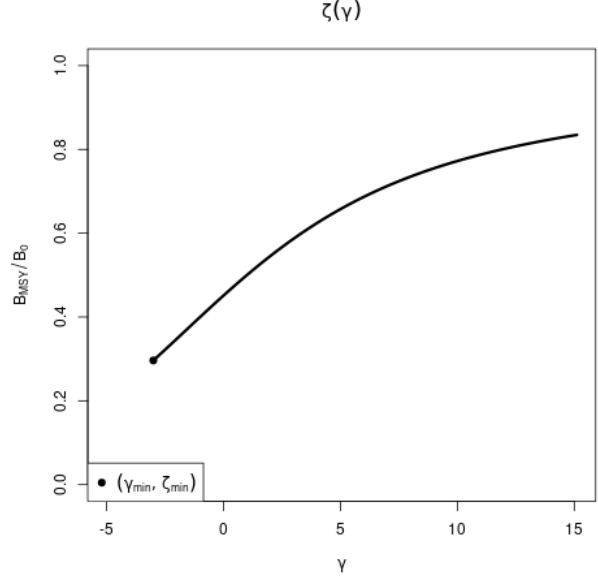


Figure 2.4: $\zeta(\gamma)$ Plotted for $F^* = 0.1$ and $M = 0.2$. The point $(\gamma_{min}, \zeta_{min})$ shows the lowest biologically meaningful value of γ ; below which productivity is negative.

$$[\hat{\mu}, \hat{\sigma}, \hat{\nu}] = \arg \min_{[\mu, \sigma, \nu]} \int_{\Gamma} (T(\gamma; \mu, \sigma, \nu) - \zeta(\gamma))^2 d\gamma \quad (2.11)$$

The distribution $T(\gamma|\hat{\mu}, \hat{\sigma}, \hat{\nu})$ is fit for use in generating γ^* random variates at a specific F^* and M . This approximation releases the need to invert ζ w.r.t γ by using samples of γ^* values to generate approximately uniform samples of $\zeta(\gamma^*)$. By sampling approximately uniform $\zeta(\gamma^*)$ random variates in this way, and making use of the structure in Eq. (2.9), an approximate LHS sample can be collected via Algorithm (1).

$\frac{F^*}{M}$ is drawn uniformly from \mathcal{F}_i . Conditioning on the sample of F^* , and M , $T(\gamma|\hat{\mu}, \hat{\sigma}, \hat{\nu})$ is fit and γ^* is sampled. ζ^* is then computed and placed into the appropriate grid element \mathcal{B}_j . Given γ^* , the cascade $\alpha(\gamma^*)$, and $\beta(\alpha(\gamma^*), \gamma^*)$, can be computed.

The algorithm continues until all of the de-

sign elements, $(\frac{F^*}{M}, \zeta^*) \Leftrightarrow (\alpha^*, \beta^*, \gamma^*)$, have been computed for all $i \in [1, \dots, n]$.

Design Refinement

Since the behavior of RP inference, under misspecified models, will vary in yet-unknown ways, the exact sampling design density may be hard to know a priori. Several factors, including the particular level of observation uncertainty, high variance (i.e. hard to resolve) features of the response surface, or simply "gappy" instantiations of the initial LHS design may necessitate adaptive design refinement, to accurately describe RP biases. Given the temperamental relationship between RPs and productivity parameters in the Schnute model, a recursive refinement algorithm that makes use of the previously described LHS routine, is developed.

While LHS ensures uniformity in the design margins, and a certain degree of spread, it

Algorithm 1 LHS of size n on rectangle R .

```

1: procedure  $LHS_n(R)$ 
2:   Define  $n$ -grids  $\mathcal{F}, \mathcal{B} \in R$ 
3:   for each grid element  $i$  do
4:     Draw  $\frac{F^*}{M} \sim Unif(\mathcal{F}_i)$ 
5:     Compute  $[\hat{\mu}, \hat{\sigma}, \hat{\nu}]$  given  $F^*$  &  $M$ 
6:     while  $\mathcal{B}_j$  not sampled do
7:       Draw  $\gamma^* \sim T(\gamma|\hat{\mu}, \hat{\sigma}, \hat{\nu})$ 
8:       Compute  $\zeta^* = \zeta(\gamma^*)$ 
9:       Compute  $j$  such that  $\zeta^* \in \mathcal{B}_j$ 
10:    end while
11:    Compute  $\alpha^* = \alpha(\gamma^*, F^*, M)$ 
12:    Compute  $\beta^* = \beta(\alpha^*, \gamma^*, M, B_0)$ 
13:    Save  $(\frac{F^*}{M}, \zeta^*) \Leftrightarrow (\alpha^*, \beta^*, \gamma^*)$  in  $\mathcal{F}_i \times \mathcal{B}_j$ 
14:  end for
15: end procedure

```

is widely recognized that particular LHS instantiations may leave substantive gaps in the simulation design. To correct this, LHS is often paired with design elements of maximin design (Morris & Mitchell, 1995; Devon Lin & Tang, 2015). Maximin designs sample the design space by maximizing the minimum distance between sampled points. This has the advantage of definitionally filling holes in the design, however because no points are ever drawn outside of the design domain, samples tend to clump around edges (particularly corners) of the design domain. Since LHS ensures uniformity in the margins and maximin designs enjoys a certain sense of optimality in how they define and fill gaps (Johnson et al., 1990), the methods are quite complimentary when combined.

Making use of this complimentary relationship, holes in the existing LHS design of RPs are identified based on maximin design principles. New design points are collected based on areas of the RP design space which maximizes the minimum distance between all pairs of points in the current design, based on the following distance function

$$d(\mathbf{x}, \mathbf{x}') = \sqrt{(\mathbf{x} - \mathbf{x}')^T \mathbf{D}^{-1} (\mathbf{x} - \mathbf{x}')} \quad (2.12)$$

$$\mathbf{D} = \mathbf{diag} \left[\left(\max(\mathcal{F}) - \min(\mathcal{F}) \right)^2, \left(\max(\mathcal{B}) - \min(\mathcal{B}) \right)^2 \right].$$

Above, d is a scaled distance function that defines the distance between points in the differing scales of $\frac{B^*}{B_0}$ and $\frac{F^*}{M}$. \mathbf{D} is a diagonal matrix that measures the squared size of the domain in each axis of so as to normalize distances to a common scale.

If \mathbf{X}_n is the initial design, computed on R_{full} , let \mathbf{x}_a be the augmenting point which maximizes the minimum distance between all of the existing design points,

$$\mathbf{x}_a = \underset{\mathbf{x}'}{\operatorname{argmax}} \min \{ d(\mathbf{x}_i, \mathbf{x}') : i = 1, \dots, n \}. \quad (2.13)$$

The point \mathbf{x}_a is used as an anchor for augmenting \mathbf{X}_n . An additional $LHS_{n'}$ (via Algorithm (1)) is collected, adding n' design points, centered around \mathbf{x}_a , to the overall design. The augmenting region, $R_{(\mathbf{x}_a, d_a)}$, for collecting $LHS_{n'}$ is defined based on the square centered at \mathbf{x}_a with side length $2d_a$, where $d_a = \min \{ d(\mathbf{x}_i, \mathbf{x}_a) : i = 1, \dots, n \}$, in the space defined by the metric d .

Due to the tendency of maximin sampling to cluster augmenting points on the edges of the design space, $R_{(x_a, d_a)}$ is truncated by the outer most limits of R_{full} so as to focus design augmentation within the specified domain of the simulation. Furthermore, since the design space has a nonlinear constraint at low values of $\frac{B}{B_0}$, the calculation of x_a is further truncated based on a convex hull defined by the existing samples in the overall design.

Design refinement then proceeds as follows. An initial design is computed, $X_n = LHS_n(R_{full})$, based on an overall simulated region of RPs R_{full} . The maximin augmenting point, x_a , is computed at a maximin distance of d_a from the existing samples. An augmenting design $X_{n'} = LHS_{n'}(R_{(x_a, d_a)})$ is collected and added to X_n . Design refinement carries on recursively collecting augmenting designs in this way until the maximin distance falls below the desired level.

0.3 Gaussian Process Metamodel

At its core, a metamodel is simply a model of some mapping of inputs to outputs (the mapping itself is typically defined by a computer model). By modeling the mapping with a statistical model (that explicitly defines the relevant features of the mapping) a metamodel defines a specific ontology for the mapping. By simulating examples of the mapping, the inferential infrastructure of the statistical model is used to empirically learn an effective emulation of the mapping within the ontology defined by the statistical model. The predictive infrastructure of the statistical model is then useful as an approximate abstraction of the system itself to better understand the system through further data collection, cheap approximation of the mapping, and/or study of the mapping itself.

In this setting, the aim of metamodeling is to study how well RPs are inferred when typical two parameter models of productivity (Logistic and BH) are misspecified for populations that are actually driven by more complicated dynamics. The simulation design, \mathbf{X} , provides a sample of different population dynamics that are driven by three parameter production functions broadly in RP space. By simulating index of abundance data from the three parameter model, and fitting those data with the two parameter production model, we observe particular instances of how well RPs are inferred at the given misspecification of the two parameter model relative to the true three parameter production model. By gathering

all of the simulated instances of how RPs are inferred (under the two parameter model), we form a set of example mappings to train a metamodel which represents the mapping of true RPs (under the three parameter model) to estimates of RPs under the misspecified two parameter production model. The metamodel is essentially a surrogate for inference under the misspecified two parameter production model that controls for the specific degree of model misspecification.

A flexible GP model is assumed for the structure of the metamodel to describe the mapping of RPs under misspecified two parameter models of productivity. A GP is a stochastic process generalizing the multivariate normal distribution to an infinite dimensional analog. GP models are often specified primarily through the choice of a covariance (or correlation) function which defines the relationship between locations in the input space. Typically correlation functions are specified so that points closely related in space result in correlated effects in the model. In this setting the inputs to the GP metamodel are the space of reference points which define the simulated three parameter production models.

While index of abundance data are generated from three parameter models, at each design location of the simulation, fitting the restricted two parameter model results in a maximum likelihood estimate (MLE; and associated estimation uncertainty) of each of the productivity parameters (i.e. Schaefer: $[\log(r), \log(K)]$, BH: $[\log(\alpha), \log(\beta)]$). To simplify the specification of the metamodel, let \mathbf{y} be a vector collecting the fitted MLEs for one of the productivity parameters, and let $\boldsymbol{\omega}$ be a vector of estimates of the estimator variances (via the inverted Fisher information) at each \mathbf{y} . Each of the fitted productivity parameter estimates are then modeled using independent instances of the following GP metamodel.

$$\begin{aligned}\mathbf{y} &= \beta_0 + \mathbf{X}\boldsymbol{\beta} + \mathbf{v} + \boldsymbol{\epsilon} \\ \mathbf{v} &\sim N_n(\mathbf{0}, \tau^2 \mathbf{R}_\ell) \\ \boldsymbol{\epsilon} &\sim N_n(\mathbf{0}, \boldsymbol{\omega}' \mathbf{I})\end{aligned}\tag{2.14}$$

\mathbf{X} is the $n \times 2$ LHS design matrix of RPs for each simulated three parameter data generating model as described in Section (0.2). ϵ models independent normally distributed error, which provides an ideal mechanism for propagating uncertainty from inference in the

simulation step into the metamodel. By matching each y_i with an observed ω_i variance term, ϵ serves to down weight the influence of each y_i in proportion to the inferred production model sampling distribution uncertainty. This has the effect of smoothing the GP model in a way similar to the nugget effect (Gramacy & Lee, 2012), although the application here models this effect heterogeneously.

The term, \mathbf{v} , contains spatially correlated GP effects. The correlation matrix, \mathbf{R}_ℓ describes how RPs close together in the simulation design are more correlated than those that are far away. This spatial effect is modeled with a squared exponential correlation function,

$$R(\mathbf{x}, \tilde{\mathbf{x}}) = \exp \left(\sum_{i=1}^2 \frac{-(x_i - \tilde{x}_i)^2}{2\ell_j^2} \right). \quad (2.15)$$

R has an anisotropic separable form which allows for differing length scales, ℓ_1 and ℓ_2 , in the different RP axes. The flexibility to model correlations separately in the different RP axes is key due to the differences in the extent of the RP domains marginally. The metamodel parameters β_0 , $\boldsymbol{\beta}$, τ^2 , ℓ_1 and ℓ_2 are fit via MLE against the observations \mathbf{y} , \mathbf{X} , and $\boldsymbol{\omega}$ from simulation fits.

Fitting the metamodel allows for a full predictive description of inference under the misspecified restricted models. Predictive estimates are obtained via kriging (Cressie, 2015)

$$\hat{y}(\mathbf{x}) = \beta_0 + \mathbf{x}\boldsymbol{\beta} + \mathbf{r}(\mathbf{x})'\mathbf{R}_\ell^{-1}(\mathbf{y} - (\beta_0 + \mathbf{X}\boldsymbol{\beta})) \quad (2.16)$$

$\hat{y}(\mathbf{x})$ is the predicted value of the modeled productivity parameter MLE under the two parameter production model, when the index of abundance is generated from the three parameter production model at RP location \mathbf{x} . $\mathbf{r}(\mathbf{x})$ is a vector-valued function of correlation function evaluations for the predictive location \mathbf{x} against all observations in \mathbf{X} (i.e. $\mathbf{r}(\mathbf{x}) = \mathbf{R}(\mathbf{x}, \mathbf{x}_i) \forall \mathbf{x}_i \in \mathbf{X}$).

While metamodeling occurs on the inferred productivity parameters of the restricted production model, the metamodel can also be used to build estimates of major biological RPs. For the BH model the relevant transformations for relating productivity parameters with RPs are given in Eqs. (2.5, 2.8) with γ fixed to -1; for the Schaefer model $\hat{B}^* = \frac{\hat{K}}{2}$ and

710 $\hat{F}^* = \frac{\hat{r}}{2}$. Applying the metamodel predictive surfaces on the scale of RP estimates allows for
711 the quantification of estimation bias that is induced by fitting a misspecified two parameter
712 production model to indices of abundance generated under three parameter productivity.

Results

1.1 Schnute/BH

Design

Algorithm (1) enforces uniform marginals in $\frac{F^*}{M}$ directly, as well as the adherence of the overall design to latin squares. Figure (2.5) shows a uniform Q-Q plot for sampled ζ , using Algorithm (1), against theoretical uniform quantiles. As evidence by the excellent coherence to the theoretical uniform quantiles, the approximation in Section (0.2) for sampling γ (and therefore $\zeta(\gamma)$), is very effective. Furthermore since numerical inversion of $\zeta(\gamma)$ is costly and unreliable, the relative speed and accuracy that this approximate LHS sampling method provides is pivotal for the rest of the work presented here.

Similarly to the PT model, the three parameter Schnute model is uniquely identified by each point in the space of $\frac{F^*}{M}$ and $\frac{B^*}{B_0}$ RPs. As seen in Figure (2.6), Schnute production has different behaviors in different ranges of RPs space, which are entirely defined by the value of γ (shown in Figure (2.1)). When $\gamma \geq 1$ the Schnute model produces a family of Logistic-like curves that are increasingly right leaning as γ increases. For $1 > \gamma \geq 0$, Schnute production takes a family of left leaning Ricker-like curves that all, at least, approach the x-axis. For $0 > \gamma > -1$ there are a family of BH-like curves that do not

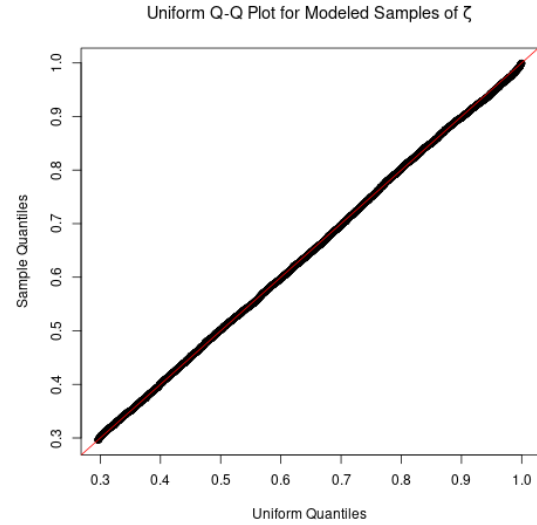


Figure 2.5: Uniform Q-Q plot for ζ plotted for $F^* = 0.1$ and $M = 0.2$.

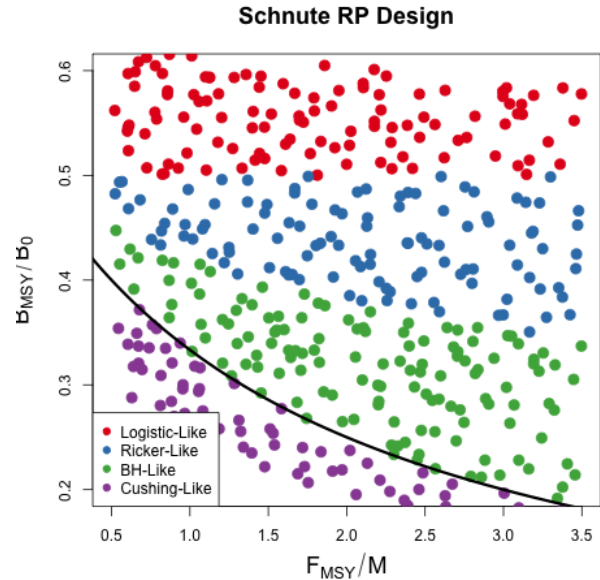


Figure 2.6: A Schnute RP design. Colors indicate different regimes of Schnute production. The black curve shows the BH set.

approach the x-axis but still have decreas-

ing productivity for large biomass stocks. When γ is exactly -1 Schnute reduces to BH production which has asymptoting production for large biomass. Finally when $-1 > \gamma$ Schnute produces a family of increasing Cushing-like curves that do not asymptote, and produces linear production as $\gamma \rightarrow -\infty$.

Modeling index data that are simulated broadly over the theoretical space of RPs with misspecified BH production greatly limits the range of possible RPs that can be inferred. Under BH production the full theoretical space of RPs are limited to the curve $\frac{B^*}{B_0} = \frac{1}{F^*/M+2}$. Define the “BH set” as the set of RPs defined by this limited space, i.e. the curve $\left\{ \left(\frac{F^*}{M}, \frac{B^*}{B_0} \right) \mid \frac{B^*}{B_0} = \frac{1}{F^*/M+2} \right\}$. as seen in the black curve in Figure (2.6). The farther away from this set that Schnute data are simulated, the worse the BH model is misspecified for those data.

Metamodeled Trends

Unlike the Schaefer model, the BH set is not a constant in $\frac{B^*}{B_0}$. Under the BH model, bias in $\frac{B^*}{B_0}$ is no longer entirely defined by the degree of model misspecification, but rather the structure of BH RPs allows bias in both $\frac{B^*}{B_0}$ and $\frac{F^*}{M}$ to interact as a function of contrast in the data.

High Contrast Figure (2.7) shows metamodeled RP bias surfaces for inference under the BH model in the high contrast setting. The (*left*) and (*bottom*) panels focus only on the $\frac{B^*}{B(0)}$ and $\frac{F^*}{M}$ components of bias respectively. In these panels bias is shown as relative bias, $\frac{\widehat{RP}-RP}{RP}$, similar to a percent error calculation. Where RP represents the true value of the three parameter RP, and \widehat{RP} refers to the metamodel estimate.

Figure (2.7, *top-right*) combines the components of bias to show the overall mapping of RPs under BH inference in the high contrast simulation setting. Unlike high contrast RP inference under the Schaefer model, the BH model does shows bias in both RPs here. Despite the bias in $\frac{B^*}{B(0)}$ and $\frac{F^*}{M}$ these results are similar to that of the Schaefer model in that the overall mapping of RPs is very nearly a minimal distance mapping onto the constrained set of RPs. The primary difference between Schaefer model and BH RP inference is the geometry of their limited RP spaces. Unlike the Schaefer model the BH set encourages bias

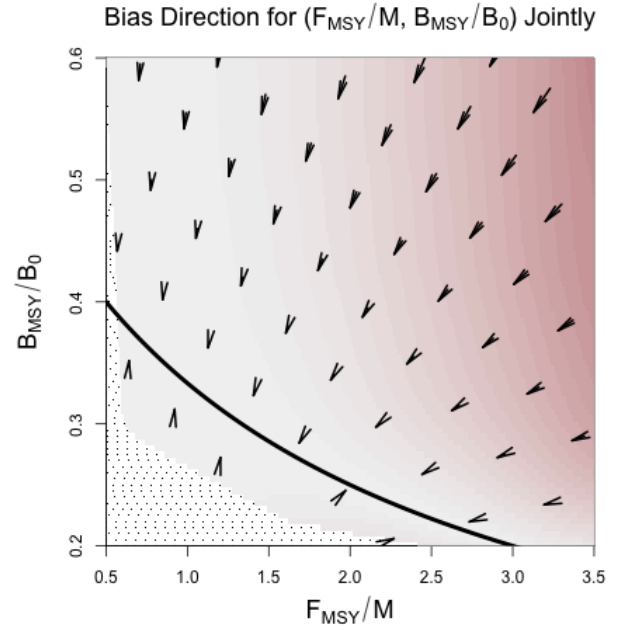
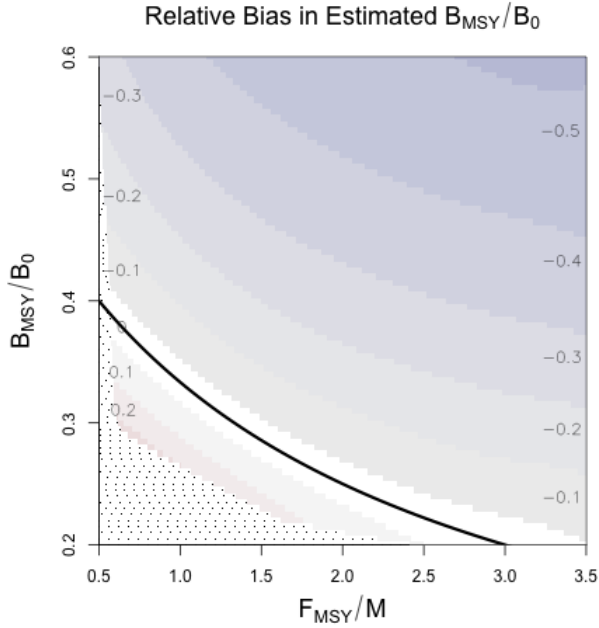
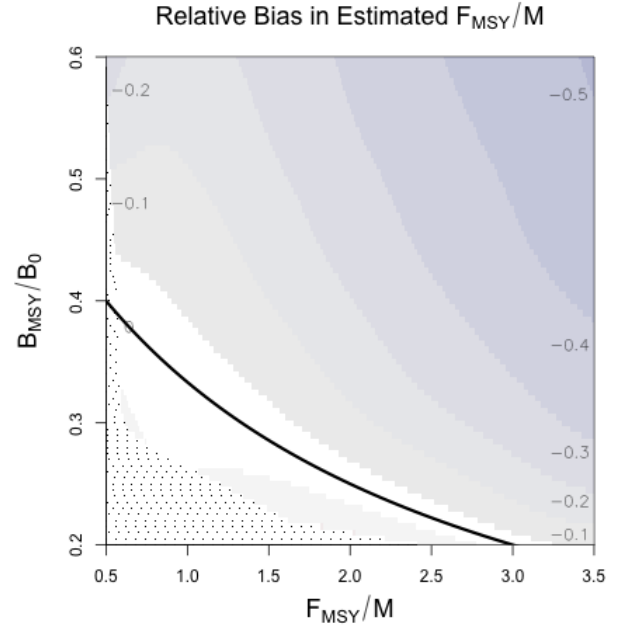
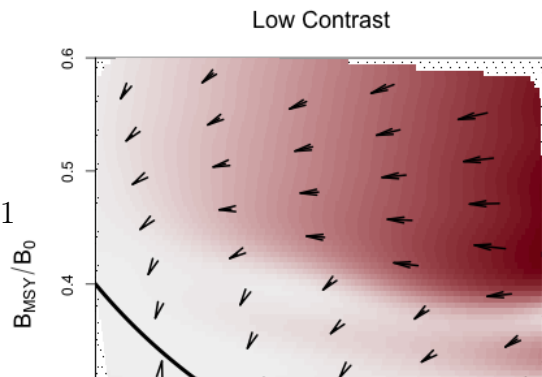


Figure 2.7: Heatplots showing the bias in RP estimation induced by model misspecification of the BH model in the high contrast simulation setting. In all cases the restricted RP-space of the BH set is shown as the black curve. (*left*) Relative bias in $\frac{B^*}{B(0)}$. (*top-right*) Bias in RP-space shown directionally. Arrows point from the location where data is generated, toward the location in the BH set where MLE projects estimated RPs. The intensity of color represents the excess bias relative to the shortest possible mapping. (*bottom*) Relative bias in F^* .



771 in both RPs for misspecified models even in very well informed setting.

772 **Low Contrast** Figure (2.8) shows the
 773 mapping of RPs in the low contrast simu-
 774 lation setting. Figures (2.8) and (2.7, *top-*
 775 *right*) share a common scale for the inten-



sity of color to facilitate comparison. In Figure (2.8) notice that the mildly misspecified area around the BH set produces mappings onto the BH set which resemble the minimal distance mapping seen in the high contrast setting. The primary difference in this low contrast setting, is the break point around $\frac{B^*}{B(0)} = 0.4$ above which $\frac{F^*}{M}$ is sharply underestimated.

The region of RPs where the BH model manages to recover the minimal distance mapping may be considered a “safe regime” of data types that are reasonably well modeled by a BH model. By comparison of Figure (2.8), with Figure (2.6), this safe regime of the BH model occurs for data generated for Cushing-like or BH-like production. While bias of the RPs can still become concerningly large, this region can be considered safe in the sense that even for low contrast data RP estimation under the the BH model recovers the minimal distance mapping.

Outside of this safe regime, RP estimation breaks from the minimal distance mapping at the interface between BH-Like and Ricker-Like regimes of the Schnute model (again see Figure (2.6)). The Ricker model lies along this regime interface, and represents the first model to approach the x-axis for large biomasses as γ increases. This markedly unBH-like productivity in the low information simulation setting breaks MLE inference from the minimal distance mapping and instead maps RPs to extremely low values of F^* ; consequently $\frac{B^*}{B(0)}$ is estimated

Estimated Yield Curves For Poorly Specified BH

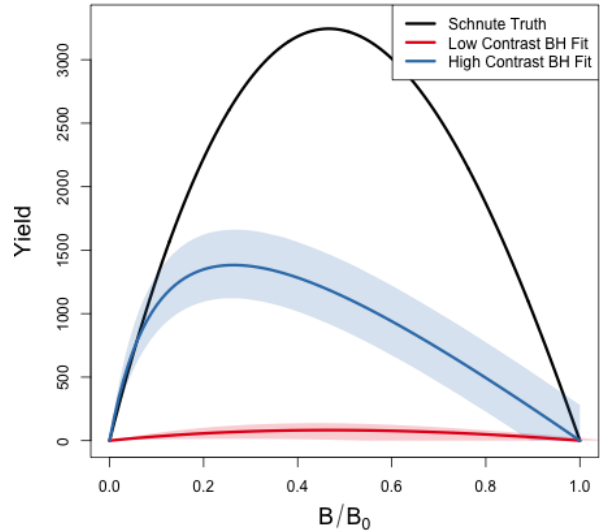


Figure 2.9: Yield curves for data generated with $\frac{F^*}{M} = 3.48$ and $\frac{B^*}{B(0)} = 0.48$.

near the limiting value under the BH (i.e.

$\lim_{F^* \rightarrow 0} \frac{1}{F^*/M+2} = 0.5$). Similarly the set of Ricker RPs (as well as the Schaeffer set) include this trivial limiting point in common ($\frac{F^*}{M} = 0$, $\frac{B^*}{B(0)} = 0.5$).

Interestingly, in the high contrast setting this trivial mapping for highly misspecified BH models is not present. This suggests that, under a misspecified BH model, the presence of adequate information in the data to produce reasonable estimates of $\frac{F^*}{M}$, drives $\frac{B^*}{B(0)}$ below 0.5 in accordance with $\frac{B^*}{B(0)} = \frac{1}{F^*/M+2}$, even when the true $\frac{B^*}{B(0)} > 0.5$. This phenomena balances RP estimation within the constrained BH set as mediated by the information content of the data and the degree of model misspecification. When the information content in the data is too small to drive a compromised RP estimate, inference completely disregards accurate estimation of F^* in order to better estimate $\frac{B^*}{B(0)}$ by exploiting the common limiting behavior of the BH set and that of Ricker-like and Logistic-like models.

2 Discussion

Results presented here generally agree with what is known about estimating growth rate parameters (Lee et al., 2012; Conn et al., 2010; Magnusson & Hilborn, 2007). These study's appreciate the role of contrast for estimating growth rates, however they struggle to make generally extensible conclusions since they focus only on a handful of stocks that fall short of forming a random sample of the greater population of possible stock behaviors. The LHS design methods presented here are designed specifically to simulate a representative sample of stocks broadly across the space of possible RPs. Furthermore, the simulation design, taken together with the GP metamodel of productivity parameter estimates, allows this study to control the degree of model misspecification and generalize conclusions about the behavior of productivity estimation within the production model setting presented.

In the presence of contrast, F^* estimation can enjoy very low bias even for a wide range of poorly specified models; conversely in the absence of contrast F^* estimation can suffer very large bias even for slightly misspecified models. This pattern is particularly true for inference under the Schaeffer model where the geometry of the restricted RP set isolates estimation failure of F^* from $\frac{B^*}{B(0)}$. While contrast has a similar impact on F^* estimation under the BH model, the geometry of the BH RP set correlates estimation bias of F^* and $\frac{B^*}{B(0)}$. The

GP metamodeling approach reveals a more general pattern that highly informative data sets (high contrast) produces a nearly minimal distance mapping of RPs onto the constrained RP set.

In all cases when model misspecification is removed, even with weakly informative data, RP estimation is unbiased and well estimated. Thus contrast alone is not the only factor leading to inferential failure. Model misspecification is a necessary but not sufficient condition for inducing RP estimation bias. The particular RP bias present depends on the RP geometry of the fitted model and how that geometry is misspecified relative to the data. The RP mapping is then oriented to the RP geometry of the fitted model.

While the relative fishing rate parameterized in Section (2.3) captures a usefully broad spectrum of relevant fishing behaviors, it is still limiting in the amount of information that it can induce. Improved methods for quantifying contrast in fisheries data, and/or methods of discovering more informative fishing behavior, could improve this analysis. In the absence of a maximally informative dataset simulation methods will not fully describe how inference fails, but the methods presented here tell the most complete picture yet, with explicit control of the degree model misspecification, contrast, and a simulation design that allows for uniform representative data generation across biologically meaningful stocks. The results presented here suggest the conjecture that under a maximally informative dataset, RP inference with a two parameter production function will be biased in the direction a shortest distance map from the true RPs onto restricted set of RPs under the two parameter model.

Given the potential for model misspecification of RPs, a minimal distance mapping of RPs represents a best-case scenario where the total bias of RPs, when measured jointly, is minimized. That said, without recognizing the geometry of how two parameter models of productivity limit RP space this may lead to unintuitive implications in RP estimation. For example, due to the shape of the BH RP set a minimal distance mapping ensures that if there is bias in one of $\frac{B^*}{B_0}$ or F^* , there will necessarily be bias in the other RP. However under the Schaefer model, since the RP set is a constant in $\frac{B^*}{B_0}$, bias in F^* is not adulterated in the same way by bias in $\frac{B^*}{B_0}$ estimation. While models with constant RPs, such as the logistic model $\frac{B^*}{B_0} = \frac{1}{2}$ or the Fox model $\frac{B^*}{B_0} = \frac{1}{e}$, are extremely limited, they can be valuable tools for developing intuition precisely because they isolate RP estimation in their free RPs from

the correlated RP biases present in models like the BH or Ricker model.

When one considers the implications of RP bias, overestimation of RPs carries the severe implication of management recommendations potentially leading to overfishing, while underestimation of RP leads to overly conservative management. In this sense, when the true model is not known, the geometry of the BH set together with the metamodeled bias trends makes the BH model a naturally conservative estimator of RPs for most stocks. For most non-BH populations the BH model is likely to make conservative errors in its estimates of F^* and $\frac{B^*}{B_0}$. The one notable exception to the conservatism of the BH model stands for data generated in the Cushing-like regime of Schnute RPs. In this regime the BH model tends to be fairly unbiased overall, however the bias that is present for these populations tends to be overestimation in both RPs, leading to much more severe management consequences for those populations.

The RP bias trends of the Schaefer model demonstrate much less conservatism than the BH overall. For any population with $\frac{B^*}{B_0} < 0.5$, $\frac{B^*}{B_0}$ will be overestimated. When the population comes from the regime where $\frac{B^*}{B_0} > 0.5$, $\frac{B^*}{B_0}$ will be under estimated, but F^* is likely to be overestimated depending on the degree of contrast present in the data. So while the Schaefer model is an intuitive model, it tends to lead to much less conservative RP estimation.

While it is important to recognize these limitations of two parameter models of productivity, we should not solely accept conservatism as a rationale of choosing a BH model of productivity. Increasing the flexibility of the production function by moving toward three parameter models would release the underlying structural limitations (Mangel et al., 2013) that cause these RP biases in the first place. Punt and Cope (2019) considers a suite of possible three parameter curves which could be used instead of current two parameter curves. For all of their benefits, three parameter production functions have their own complicating factors, and the structure present in the Schnute model explored here makes it an intuitive bridge model for developing three parameter models going forward.

- [show a schnute fit to data?](#) (Yeakel & Mangel, 2015) Prior

- 893 • summary of σ over RP space comparing between models (PT, Schnute, Schnute DD)

894 to show areas of model breakdown.

 - 895 – miss-identifying signal for noise.
 - 896 – It happens more as the dynamics get more complex.
 - 897 – point to the full age structured models.
- 898 • show the constrained BH space over a grid of $M, \kappa, \omega, W_\infty$
- 899 • Show that the constrained spaces vary only slightly as compared with the consequences

900 of misspecifying the functional form.
- 901 • estimating these other quantities (while they can create quite different Biomass series)

902 can only do so much to improve (expand) RP inference as compared with correctly

903 modeling P .
- 904 • mapping distance as a function of contrast at (3.5, 0.5)
- 905 • for LHS grid locations show $\frac{B^*}{B_0}$ and F^* biases for grids in $M \in (0, 0.5)$ For sure in High

906 Contrast, maybe also in Low??.

907 Chapter 3

908 A Delay Differential Model

- 909 • Introduction
- 910 – piggy back intro off of simpleModel
- 911 – problem statement and motivation
- 912 – introduce reference point and management decision making
- 913 – new dynamics of cohorting.
- 914 • Methods
- 915 – state and describe model
- 916 – Reference Point Derivation
- 917 – layout data generation/space filling problem
- 918 – how far to get the math for inputting into CAS
- 919 – method of CAS.
- 920 – describe and plot ζ .
- 921 – constrained BH space (method for visualizing)
- 922 – appendix for RP CAS calculation
- 923 • Results
- 924 • summary of σ over RP space comparing between models (PT, Schnute, Schnute DD)
- 925 to show areas of model breakdown.
- 926 – miss-identifying signal for noise.
- 927 – It happens more as the dynamics get more complex.
- 928 – point to the full age structured models.
- 929 • Show that the constrained spaces vary only slightly as compared with the consequences
- 930 of misspecifying the functional form.
- 931 • ?Discussion?

- 932 • summary of σ over RP space comparing between models (PT, Schnute, Schnute DD)

933 to show areas of model breakdown.

 - 934 – miss-identifying signal for noise.
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 - 936 – point to the full age structured models.
- 937 • show the constrained BH space over a grid of $M, \kappa, \omega, W_\infty$
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- 939 of misspecifying the functional form.
- 940 • estimating these other quantities (while they can create quite different Biomass series)
- 941 can only do so much to improve (expand) RP inference as compared with correctly
- 942 modeling P .

1 Introduction

- the delay model: [J. Schnute \(1985\)](#) [J. Schnute \(1987\)](#) [Fournier and Doonan \(1987\)](#).
- discrete: [Hilborn and Walters \(1992, pg. 334\)](#)
- [Walters \(2020\)](#)
- automatic accounting for cohort cycles

2 Methods

2.1 Delay Differential Model

Age structured fisheries models typically assume [Von Bertalanffy \(1938, VB\)](#) growth in length with age. To model weight the assumption of VB growth in length is composed with a power law relating length to weight, $w = al^b$. Since b is usually ~ 3 this composition of assumed functional forms typically results in a monotonically increasing sigmoidal curve of weight with age. When $b \leq 1$ weight at age takes a VB-like form with $b = 1$ resulting in an exact correspondence of simultaneous VB-growth in length and weight.

The delay model slightly abridges these relationships by directly assuming VB growth in weight as follows,

$$w(a) = w_{\infty}(1 - e^{-\kappa(a-a_0)}). \quad (3.1)$$

κ is a parameter that controls the instantaneous rate of individual growth (in weight) with age. w_{∞} is the maximum weight of individuals in the population, and $w(a)$ is the average

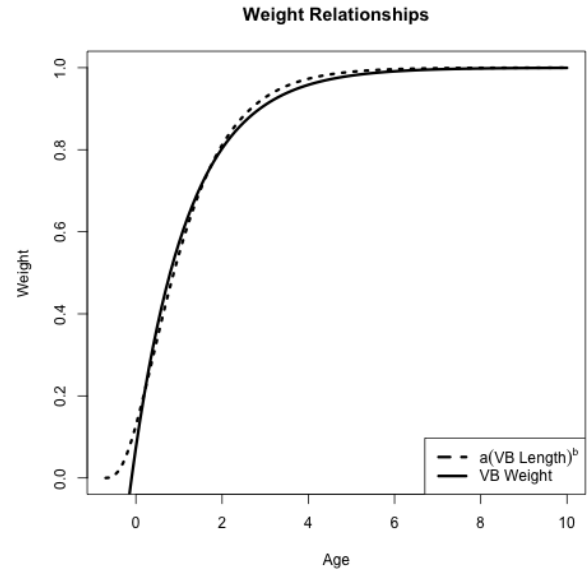


Figure 3.1: The typical composition of allometric weight ($b = 3$) with VB growth in length, as approximated by VB growth in weight directly.

weight of an individual at age a . The parameter a_0 controls the age at which individuals are assumed to have zero weight; by letting $a_0 < 0$ this allows fish of age zero to have positive weight. Rather than taking a sigmoidally increasing function, VB growth directly in weight results in an monotonically inceasing curve that asymptotes with a strictly decreasing growth rate with age. (only a good approximation for older ages where growth begins to decline)

Together with VB growth, the delay model is derived from the assumption that both natural mortality and fishing selectivity are separately propotional to a common heavyside step function with age. That is to say, before a threshold age of selectivity, a_s , the population is assumed not to experience any mortality whatsoever, but all fish older then a_s experience the same rate of natural mortaility. Simulaneously all fish older than a_s are equally vulnerable to fishing (i.e. knife edge selectivity at age a_s), although fishing effort may vary from through time.

Walters (2020) shows that within these assumptions the following delay differential system of equations exactly models the population dynamics of the total exploitable biomass $B(t)$ and number of individuals $N(t)$ through time.

$$\frac{dB}{dt} = w(a_s)R(B; \theta) + \kappa [w_\infty N - B] - (M + F)B \quad (3.2)$$

$$\frac{dN}{dt} = R(B; \theta) - (M + F)N \quad (3.3)$$

This formulation separates the number of individuals in the population from the biomass of the population. The dynamics of N , as seen in Eq (3.3), are very similar to that of the production models previously presented, however the role of the production function is now filled by a "recruitment" function, $R(B)$, which describes the number of new individuals recruiting into the expoitable population as a function of exploitable biomass. In turn, the biomass dynamics are coupled to the numbers dynamics by the assumption of VB growth with growth parameters appearing in Eq (3.2), converting population numbers into biomass and accounting for the growth of biomass with age.

Eq (3.2) of the above model expands the notion of biomass production into the processes of recruitment, individual growth, and maturity. The term $w(a_s)R(B; \theta)$ represents the

986 biomass of new recruits; with $w(a_s)$ representing the weight of individuals at the age of
 987 maturity, a_s , and $R(B; \theta)$ representing the number of new recruits entering the exploitable
 988 population at time t . The negative term, $(M + F)B$, represents all causes of mortality as
 989 it is applied to biomass. Finally, the term $\kappa[w_\infty N - B]$ accounts for the net growth of the
 990 existing biomass by discounting the limiting maximal individual growth rate by metabolic
 991 weight loss proportional to $B(t)$. This term, together with the delay structure in R , provides
 992 the major computational savings of the delay differential setting, as compared with full age
 993 structured models, by automatically keeping track of changes in the mean size and growth
 994 associated with changes in recruitment as cohorts mature into the population.

Often a BH functional form is assumed for the stock recruitment relationship, but any adequately flexible family of functions may model this relationship. For the sake of evaluating the adequacy of assumed BH recruitment the simulation setting below is derived for the delay model under the assumption of the generalized three parameter Schnute recruitment as follows.

$$R(B; [\alpha, \beta, \gamma]') = \alpha B(t - a_s)(1 - \beta \gamma B(t - a_s))^{\frac{1}{\gamma}} \quad (3.4)$$

995 The parameters $\theta' = [\alpha, \beta, \gamma]$ function similarly in this setting as previously described in
 996 Section (??). That said, since the delay model explicitly parses out growth in it's dynamics,
 997 these parameters only describe the net processes of larval production, and maturation into
 998 the population, where as the production model used these parameters to also model the net
 999 effects of growth on biomass production. The γ parameter generalizes the family to model
 1000 varying degrees of decreasing recruitment for large biomasses as γ increases. The Schnute
 1001 function is exactly equivalent to BH recruitment at the special case when $\gamma = -1$, it passes
 1002 through the Ricker model as $\gamma \rightarrow 0$, and Logistic recruitment occurs when $\gamma = 1$.

1003 Since the delay model assumes knife edge selectivity, at age a_s , the term $B(t - a_s)$ appears
 1004 in R . That is to say fish recruiting into the exploitable population are the result of larval
 1005 production of biomass a_s time units in the past. This is because fishing selectivity is only
 1006 assumed to occur for fish that are at least a_s time units old and thus fish younger than a_s
 1007 are not exploitable. This waiting period requires that new recruits be the result of spawning

1008 biomass a_s time units in the past. Modeling maturity in this way results in dynamics
 1009 equations which are a system of delay differential equations as opposed to the simple ODEs
 1010 that arise in the production model setting.

1011 \sim interpretation of recruitment (larval production, recruitment) [growth external] vs.
 1012 production (larval production, recruitment, growth)

1013 • general structure: [Walters \(2020\)](#) [Hilborn and Walters \(1992, pg. 334\)](#)

1014 • growth: [Von Bertalanffy \(1938\)](#)

1015 • recruitment: [J. Schnute \(1985\)](#); [J. T. Schnute and Richards \(1998\)](#)

1016 2.2 Reference Points

1017 Deriving reference points for the delay model under Schnute recruitment is conceptually
 1018 similar to the production model setting. The additional nonlinear VB growth assumptions
 1019 along side Schnute recruitment quickly make the expressions look somewhat unweildy, al-
 1020 though analytical solutions can still be derived for most of the same quantities (although
 1021 complicated by growth parameters).

Starting from Eqs. (3.2) and (3.3), setting both $\frac{dB}{dt}$ and $\frac{dN}{dt}$ simultaneously equal to zero, and solving for B and N as a function of fishing, gives the equilibrium biomass and numbers equations.

$$\bar{B}(F) = \frac{1}{\beta\gamma} \left(1 - \left(\frac{(F+M)(F+M+\kappa)}{\alpha w(a_s)(F+M+\frac{\kappa w_\infty}{w(a_s)})} \right)^\gamma \right) \quad (3.5)$$

$$\bar{N}(F) = \frac{\alpha \bar{B}(F)(1 - \beta\gamma \bar{B}(F))^{1/\gamma}}{F+M} \quad (3.6)$$

1022 Eq. (3.6) is just $\frac{R(\bar{B})}{F+M}$, and is coupled to $\bar{B}(F)$ where most of the dynamics appear. Eq. (3.5)
 1023 resembles Eq (2.3) from the simple production model setting although the growth parameters
 1024 κ , w_∞ and $w(a_s)$, make slight adjustments to the balance of the maximum rate of recruitment
 1025 and mortality rate to give an expression for equilibrium biomass that accounts for the factors
 1026 of individual growth.

Expressions for B_0 and B^* are attained by evaluating $\bar{B}(F)$ at $F = 0$ and $F = F^*$ respectively. Calculation of F^* typically involves maximization of equilibrium yield, $\bar{Y} = F\bar{B}(F)$. While it was not possible to analytically maximize \bar{Y} , stable numerical solutions for calculating F^* were obtained by numerically solving for the roots of the analytical derivative of equilibrium yield with respect to F . Below a greatly simplified expression for $\frac{d\bar{Y}}{dF}$ is shown; the substitution $Z = F + M$ (total mortality rate) has been made to produce a more compact expression.

$$\frac{d\bar{Y}}{dF} = \frac{1}{\beta\gamma} \left[1 - \left(\frac{Z(Z + \kappa)}{\alpha w(a_s)(Z + \frac{\kappa w_\infty}{w(a_s)})} \right)^\gamma - \left(\frac{\gamma F}{\alpha w(a_s)} \right) \left(\frac{Z(Z + \kappa)}{\alpha w(a_s)(Z + \frac{\kappa w_\infty}{w(a_s)})} \right)^{\gamma-1} \left(1 + \frac{\left(\frac{\kappa w_\infty}{w(a_s)} \right) \left(\kappa - \frac{\kappa w_\infty}{w(a_s)} \right)}{\left(Z + \frac{\kappa w_\infty}{w(a_s)} \right)^2} \right) \right] \quad (3.7)$$

F^* is calculated as the numerical root, w.r.t. F , of the above expression. The numerical root is calculated using the base R uniroot function which employs a derivative free search given by [Brent \(1973\)](#).

BH Constraint

In the simple production model the BH constrained RPs are fixed to $\frac{1}{x+2}$. In the delay differential modeling setting the constrained BH RP set is complicated by the growth parameters a_s and κ . Under BH recruitment these parameters of the delay model slightly influence this relationship as seen in Figure (3.2). That said, the influence of a_s and κ on RPs is still largely limited to a confined region of reference point space which resembles the $\frac{1}{x+2}$ form. In fact the confined region of RPs is bounded above by $\frac{1}{x+2}$. In Figure (3.2) notice that for values of a_s and κ that result in high $w(a_s)$ (high values of κ and small values of a_s seen in red) the BH RP space converges to $\frac{1}{x+2}$ as derived in the simple production model setting. In opposition to

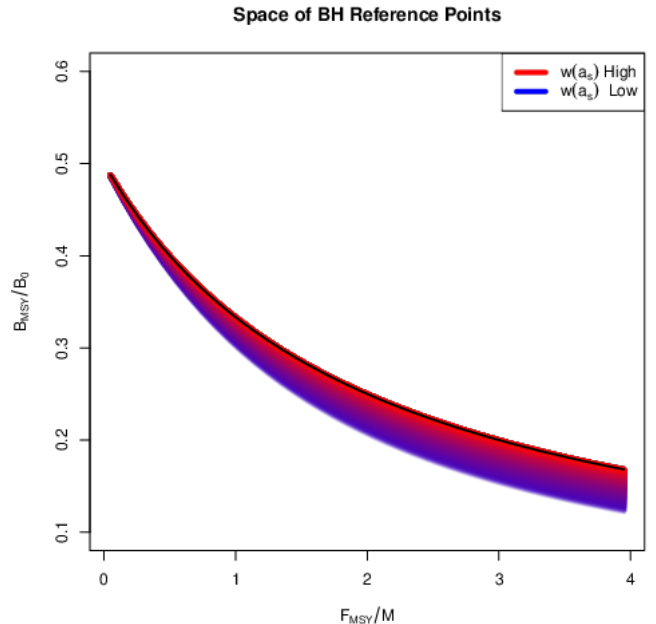


Figure 3.2: The space of BH RPs for the delay model as a function of κ and a_s . The RP space is plotted for 80×80 combinations of $\kappa \in [0.1, 2]$ and $a_s \in [0.1, 10]$. The color drawn is the resulting value of $w(a_s)$ mapped between blue and red. $\frac{1}{x+2}$ is plotted in black for reference.

1054 the simple production model limit, when $w(a_s)$ is low (as seen in the more blue region of
1055 Figure(3.2)), RPs decrease as the influence of growth in the dynamics increases.

1056 2 .3 Delay Differential Integration

1057 The delay model belongs to a class of differential equations known as delay differential
1058 equations (DDE). The delay arises from the $B(t - a_s)$ terms found in the recruitment
1059 function. Solving DDEs require special care which depends on the nature of the time delay.
1060 The addition of time-varying delays, many different delays, or very small delays (delays
1061 below the step size of the numerical integrator) results in some of the more challenging
1062 settings for solving DDEs. However with a single stationary model of the age of selectivity,
1063 the delay model in this setting represents one of the most straight forward DDE structures.
1064 The most numerically challenging case presented here arises in the case of the limiting
1065 production model when $a_s \rightarrow 0$ while $\kappa \rightarrow \infty$. That said the limiting production model can
1066 be approximated for values of $a_s \approx 0.1$, and it was straightforward to ensure that the step
1067 size of the integrator remained reasonably below 0.1.

1068 The DDE presented here is integrated with the initial values fixed at B_0 and N_0 as given
1069 by Eqs. (3.5) and (3.6) with $F = 0$ at any given configuration of θ and growth parameters.
1070 The system given in Eqs. (3.2) and (3.3) are then solved numerically using the implicit
1071 Livermore Solver (lsode) as implemented in the `dede` function of the R package `deSolve`
1072 (Soetaert et al., 2010). The `dede` solver provides many methods for integrating DDEs, but
1073 lsode was chosen because it is an implicit method that runs relatively quickly with a relatively
1074 smaller footprint in system memory as compared with other methods. The `radau` method
1075 was also tried in more computationally challenging settings with good results (albeit running
1076 more slowly than lsode). Ultimately the simulated parameter space did not produce DDEs
1077 that require the more expensive `radau` integrator to solve accurately.

1078 2 .4 Simulation Design

1079 Similarly as previously described in Section (0 .1) the relationship between RPs $\mapsto \theta$ cannot
1080 be fully expressed analytically for the Schnute delay model. However, just as in the produc-
1081 tion model setting, simulation only requires enough knowledge of these mappings to gather

1082 a list of (α, β, γ) tuples and the corresponding RPs in some reasonable space-filling design
 1083 over RP space.

1084 In the delay model a partial mapping for $(F^*, B_0) \mapsto (\alpha(\cdot, \gamma), \beta(\cdot, \gamma))$ can be derived
 1085 analytically in terms of RPs and γ . The substitution $Z^* = F^* + M$ is made where F^* and
 1086 M appear together to produce a more compact expression.

$$\alpha = \left[\left(\frac{Z^*(Z^* + \kappa)}{w(a_s)(Z^* + \frac{\kappa w_\infty}{w(a_s)})} \right)^\gamma + \left(\frac{\gamma F^*}{w(a_s)} \right) \left(\frac{Z^*(Z^* + \kappa)}{w(a_s)(Z^* + \frac{\kappa w_\infty}{w(a_s)})} \right)^{\gamma-1} \left(1 + \frac{\left(\frac{\kappa w_\infty}{w(a_s)} \right) \left(\kappa - \frac{\kappa w_\infty}{w(a_s)} \right)}{(Z^* + \frac{\kappa w_\infty}{w(a_s)})^2} \right) \right]^{\frac{1}{\gamma}} \quad (3.8)$$

$$\beta = \frac{1}{\gamma B_0} \left(1 - \left(\frac{M(M + \kappa)}{\alpha w(a_s)(M + \frac{\kappa w_\infty}{w(a_s)})} \right)^\gamma \right) \quad (3.9)$$

Above Eq. (3.8) results from setting Eq. (3.7) equal to zero and solving for α , and
 Eq. (3.9) results from solving the $\bar{B}(0)$ expression, as derived from Eq. (3.5), for β . The
 system is completed by further working with the $\frac{\bar{B}(F^*)}{\bar{B}(0)}$ expression, as seen below, to identify
 γ .

$$\frac{B^*}{B_0} = \frac{1 - \left(\frac{(F^* + M)(F^* + M + \kappa)}{\alpha w(a_s)(F^* + M + \frac{\kappa w_\infty}{w(a_s)})} \right)^\gamma}{1 - \left(\frac{M(M + \kappa)}{\alpha w(a_s)(M + \frac{\kappa w_\infty}{w(a_s)})} \right)^\gamma} \quad (3.10)$$

1087 The system formed by collecting Eqs. (3.8), (3.9), and (3.10) can be navigated similarly
 1088 to Eq. (2.9) in the Schnute production model setting. For a population experiencing natural
 1089 mortality M , VB growth with paramters κ and w_∞ , and age of selectivity a_s the above
 1090 system can fully specify α and β for a given γ , by fixing F^* , B_0 , and $\frac{B^*}{B_0}$. For a given γ a
 1091 cascade of closed form solutions for α and β can be obtained, just as in Section (0.1). First
 1092 $\alpha(\gamma)$ can be computed, and then $\beta(\alpha(\gamma), \gamma)$ can be computed. If $\alpha(\gamma)$ is filled back into the
 1093 expression for $\frac{B^*}{B_0}$, the system collapses into a single onerous expression for $\frac{B^*}{B_0}(\alpha(\gamma), \gamma)$. For
 1094 brevity, define the function $\zeta(\gamma) = \frac{B^*}{B_0}(\alpha(\gamma), \gamma, F^*, M)$ based on Eq. (3.10).

1095 Again rather than inverting $\zeta(\gamma)$ for γ , γ is the sampled so that the overall simulation
 1096 design is space filling as described in Section (0.2). Given the sampled γ , the cascade of
 1097 $\alpha(\gamma)$, and then $\beta(\alpha(\gamma), \gamma)$, can be computed, and the Schnute delay model is fully defined
 1098 by a given $(\frac{F^*}{M}, \frac{B^*}{B_0})$. While conceputally this framing is similar to the Schnute production
 1099 model, the analytical expressions are more complex, and numerically trecherous, since growth

parameters appear explicitly here. Other ways of navigating the RPs $\mapsto \theta$ system are possible, but for the sake of numerical stability this strategy has proven the most reliably accurate by limiting exposure to numerical error propagation.

Each design location defines a complete Schnute delay differential model with the given RP values. Indices of abundance are simulated from the Schnute model at each design location, a small amount of residual variation, $\sigma = 0.01$, is added to the simulated index, and the data are then fit with a misspecified BH model. The design captures various degrees of model misspecification relative to the BH model, so as to observe the effect of recruitment misspecification upon RP inference.

point to catch, and LHS design, and Metamodel.

2.5 Parameter Estimation

- I use B only here
- quick statement of inference, and reference to previous section

Let I_t , $t \in \{1, 2, 3, \dots, T\}$, be a series of indices of abundance, proportional to biomass, as simulated from the Schnute Delay model. These data are modelled with the following log-normal observation model that has been intentionally constrained to BH recruitment,

$$I_t \sim LN(qB_t(\boldsymbol{\theta}, \boldsymbol{\phi}), \sigma^2). \quad (3.11)$$

$B_t(\boldsymbol{\theta}, \boldsymbol{\phi})$ is the biomass solution of the BH constrained DDE system. The BH constraint is implemented by fixing $\gamma = -1$ so that $\boldsymbol{\theta}' = [\alpha, \beta, \gamma = -1]$. $\boldsymbol{\phi}$ is a vector of growth and maturity parameters, $\boldsymbol{\phi}' = [\kappa, w_\infty, a_0, a_s]$. The nuisance parameter q models the proportionality constant of the index with process biomass, and σ^2 models residual variation of the index.

In this setting, $\boldsymbol{\phi}$ and q are fixed to focus on the inferential affects of model misspecification on recruitment parameters and RPs. Without an explicit mechanism for the delay model to incorporate age data, under the BH model $\boldsymbol{\phi}$ is not well informed and would typically be estimated externally for data limited stocks. Under BH recruitment $\boldsymbol{\phi}$ can only slightly impact RPs as seen in Figure (3.2).

1123 σ^2 and θ are reparameterized to the log scale and fit via MLE. Reparameterizing the
 1124 parameters to the log scale improves the reliability of optimization, in addition to facili-
 1125 tating the use of Hessian information for estimating MLE standard errors. Given that the
 1126 biological parameters enter the likelihood via a nonlinear differential equation, and further
 1127 the parameters themselves are related to each other nonlinearly, the likelihood function can
 1128 often be difficult to optimize. A hybrid optimization scheme is used to maximize the log
 1129 likelihood to ensure that a global MLE solution is found. The R package GA ([Scrucca, 2013](#),
 1130 [2017](#)) is used to run a genetic algorithm to explore parameter space globally. Optimization
 1131 periodically jumps into the L-BFGS-B local optimizer to refine optima within a local mode.
 1132 The scheme functions by searching globally, with the genetic algorithm, across many initial
 1133 values for starting the local gradient-based optimizer. The genetic algorithm serves to iter-
 1134 atively improve hot starts for the local gradient-based optimizer. Additionally, optimization
 1135 is only considered to be converged when the optimum results in an invertible Hessian at the
 1136 found MLE.

- 1137 • fixed $M = 0.2$, $a_0 = -1$, $w_\infty = 1$
- 1138 • play with κ and age of selectivity a_s

1139 Numbers Indices

While not utilized here, age structured models may commonly model indices as proportional
 to numbers rather than (or simultaneously to) biomass. When solving the DDE, Eq. (3.3)
 points out that the full DDE solution will expose a numbers solution simultaneously with
 a biomass solution that may be used for these purposes. These solutions are often quite
 similar since the main driver of process behavior comes from the form of R which is shared
 among N and B . However, it is common on the west coast of the US that indices derived
 from commercial fisheries are measured as weights while indices derived from recreational
 fisheries are often measured as counts. If a numbers index, J_t , is observed alongside the
 previously mentioned biomass index, the following likelihood component is often added as a
 conditionally independent component of the likelihood,

$$J_t \sim LN(pN_t(\boldsymbol{\theta}, \boldsymbol{\phi}), \tau^2). \quad (3.12)$$

1140 $N_t(\boldsymbol{\theta}, \boldsymbol{\phi})$ is the numbers solution of the DDE system. $\boldsymbol{\theta}$ and $\boldsymbol{\phi}$ are the productivity and
 1141 growth parameters shared in common with the biomass component. p and τ^2 are then the
 1142 analogous proportionality constant and residual variation of the numbers index respectively.

1143 2.6 GP Metamodel

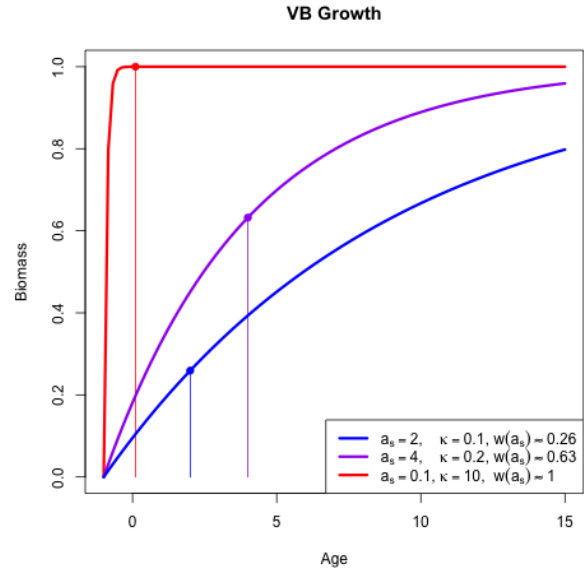
1144 point to catch, and LHS design, and Metamodel.

1145 2.7 Clustering Catastrophic Model Failure

1146 When

1147 3 Results

1148 Figure (3.3) shows three hypothetical
 1149 individual-growth/maturity curves that span
 1150 a wide range of RPs. As seen in Figure (3.2),
 1151 the larger values of $w(a_s)$ correspond to less
 1152 dramatic growth with the red curve demon-
 1153 strating the simle (no growth) production
 1154 model limit ($a_s \rightarrow 0$ and $\kappa \rightarrow \infty$). The cases
 1155 with smaller $w(a_s)$ values (blue and purple
 1156 curves) correspond to more dramatic growth
 1157 behaviors, with the blue curve where $a_s = 2$
 1158 and $\kappa = 0.1$ representing the most dramatic
 1159 growth shown here.



1157 Figure 3.3: Three hypothetical individual-
 1158 growth curves, showing $w(a_s)$ on each curve.
 1159 growth shown here.

1160 Figure (3.4) demonstrates a range of biomass dynamics that the Schnute delay model
 1161 can display under a spectrum of growth behaviors with fishing held consistent at F_{MSY} . The
 1162 three special cases of $\gamma = -1$ (BH), $\gamma \rightarrow 0$ (Ricker), and $\gamma = 1$ (Logistic) recruitment are
 1163 shown in each of the above shown growth configurations. Notice under the most dramatic
 1164 growth ($a_s = 2$ and $\kappa = 0.1$) setting, biomass of the Logistic model comes into equilibrium at
 1165 B_{MSY} as an oscillating curve. This effect occurs here due to the Logistic model's relatively
 1166 high $\frac{B^*}{B_0}$ interacting with the lag in selectivity upon the sudden onset of fishing; this produces

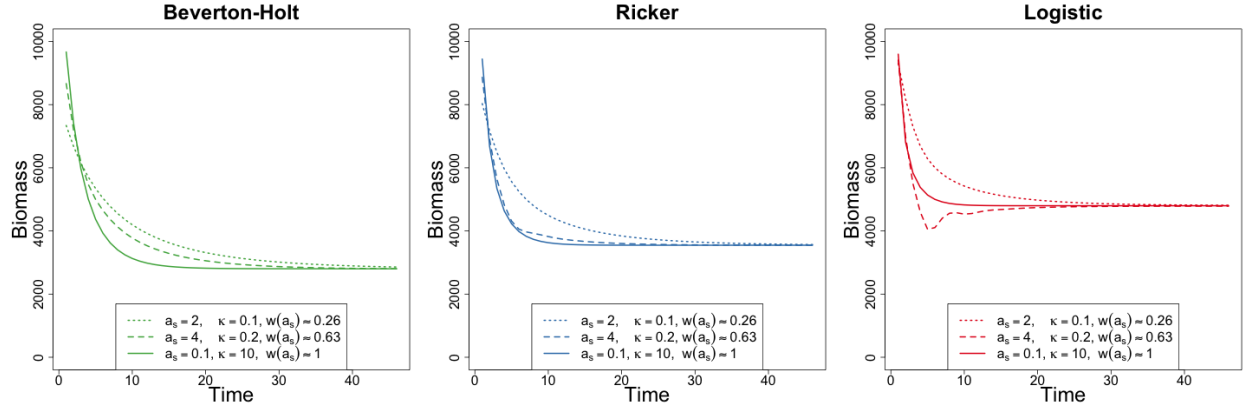


Figure 3.4: Biomass dynamics of BH (*left*), Ricker (*center*), and Logistic (*right*) delay differential models in the low contrast simulation setting. In all cases $\alpha = 1.2$ and β is chosen so that each model shares the same B_{MSY} within each given γ .

1167 a shock that pushes biomass past B_{MSY} setting up an oscillatory pattern of recruitment. One
 1168 may also observe these oscillations under the Ricker model by exaggerating the a_s lag as well
 1169 as the steepness of the Ricker curve. The BH model may also demonstrate these oscillations,
 1170 in a heavily lagged setting, by shocking the population past its relatively low B_{MSY} as a
 1171 sudden release in fishing applied to a heavily :

1172 Figure (3.5) shows the range of RPs that
 1173 can be modeled with each of the BH, Ricker,
 1174 and Logistic recruitments over the spectrum
 1175 of individual-growth/maturity models simulated here. Notice that the more dramatic
 1176 the growth, the further the RP curve lies
 1177 from the simple production model, but each
 1178 recruitment model reacts differently under
 1179 each of the given growth parameters. The
 1180 Ricker and BH RP-spaces are qualitatively
 1181 similar in shape with more dramatic growth
 1182 settings decreasing $\frac{B_{MSY}}{B_0}$ relative to the simple
 1183 production model setting. The Logistic model on the other hand increases $\frac{B_{MSY}}{B_0}$ relative
 1184 to the simple production model setting as growth parameters become more dramatic. It is
 1185

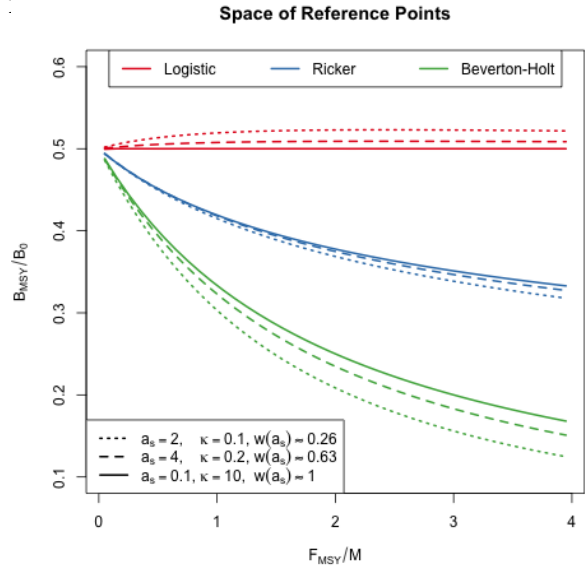


Figure 3.5: Restricted RP-space under each recruitment models, with each growth curve.

1186 also worth noting that the Ricker model's RPs are much less influenced by growth parameters
1187 as compared with that of the BH or Logistic model.

3.1 Simple Production Model Limit

Under the delay differential's limiting simple production model ($a_s = 0.1$ and $\kappa = 10$), the expectation is that RP inference should be identical to that of the model seen in Chapter (2). By way of verifying this equivalence, Figure (3.6) demonstrates a virtually identical pattern of RP biases as previously seen in Figures (2.7) and (2.8) (under both of the high and low contrast settings).

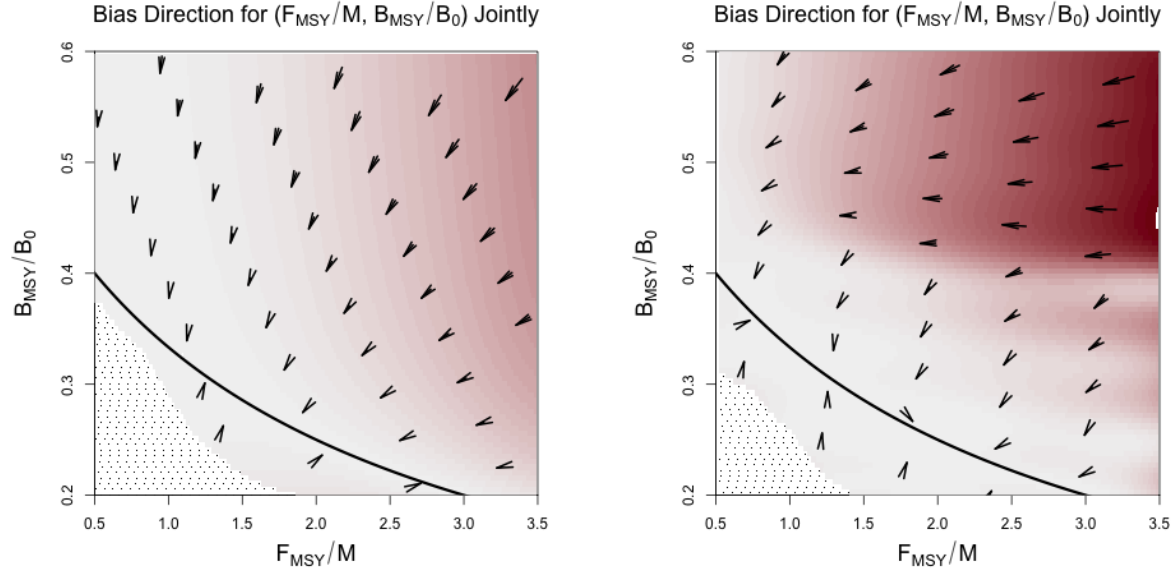


Figure 3.6: RP mapping of BH delay model fit to Schnute delay data under the simple (no growth) production model limit. *left* : High contrast simulation. *right* : Low contrast simulation.

Indeed in the high contrast setting, Figure (3.6, *left*) shows how the BH model induces the same pattern of bias as seen in Chapter (2). There is bias in both RPs (in accordance with the $\frac{B^*}{B(0)} = \frac{1}{F^*/M+2}$ RP-set) so as to produce a nearly minimal distance mapping of RPs onto the constrained BH set of RPs. Similarly, in the low contrast setting, Figure (3.6, *right*) again shows the same two regimens pattern of RP inference. Firstly, there is a region of relatively small model misspecification where the minimal distance mapping is preserved. Secondly, as model misspecification becomes greater (around the Ricker set) $\frac{F^*}{M}$ begins to be sharply underestimated. Above this break point in RP estimation inference appears to be driven to the trivial RP $\frac{F^*}{M} = 0, \frac{B^*}{B(0)} = 0.5$) that is shared in common among all of the

1203 two-parameter models described here.

1204 These results merely confirm that the theoretical limiting dynamics do indeed replicate
 1205 expected RP inference patterns as previously observed in Chapter (2).

1206 3.2 Moderate Growth

1207 Moving past verification of the simple production model, other values of a_s and κ provide
 1208 a probe into the effects individual growth dynamics may have on RP inference. Individual
 1209 growth is a multifaceted phenomena that is not easily reduced to a single number, but for the
 1210 purposes of this model $w(a_s)$ serves as a decent proxy for the extent of the model dynamics
 1211 that are due to individual growth. This follows from the intuition that individuals maturing
 1212 at a smaller fraction of w_∞ demonstrate the dynamics of growth during an observable (to
 1213 the model) phase rather than growth occurring prior to selection.

1214 That said, $w(a_s)$ is not a one-to-one map of κ and a_s . A level curve of $w(a_s; \kappa) = c$ is
 1215 attained by increasing the value of a_s and decreasing κ correspondingly, or vice versa. The
 1216 case where $a_s = 4$ and $\kappa = 0.2$ (resulting in $w(a_s) \approx 0.6$) represents a plausibly biological
 1217 example of moderate growth. Similar examples of the $w(a_s) = 0.6$ level curve result in much
 1218 larger lags (discussed in Section (3.5)) or larger κ 's which quickly tend toward behaviors
 1219 previously described in the simple production model setting.

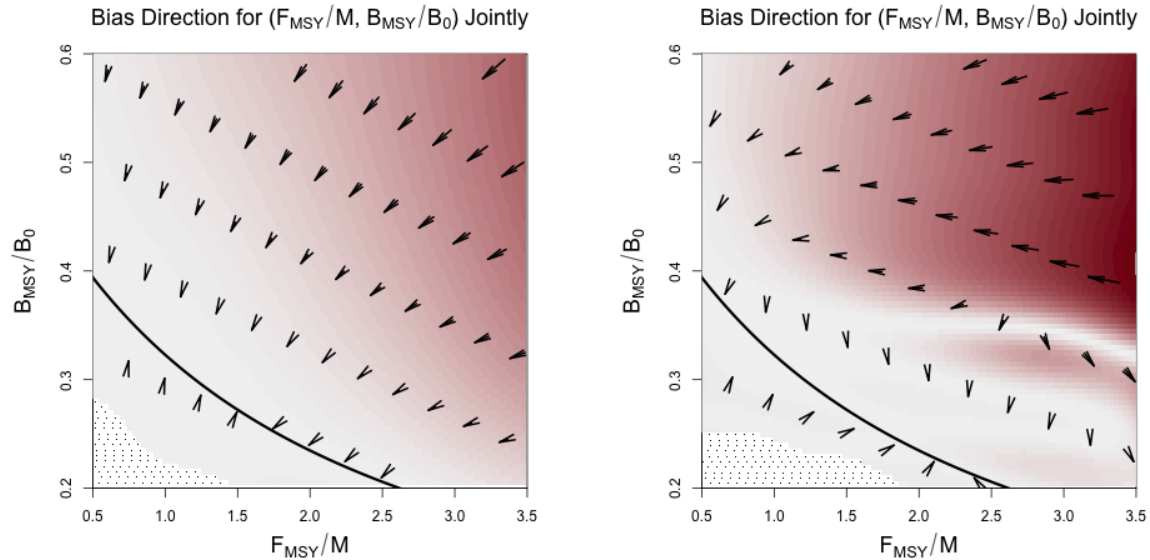


Figure 3.7: RP mapping of BH delay model fit to Schnute delay data under moderate growth ($a_s = 4$ and $\kappa = 0.2$). *Left* : High contrast simulation. *Right* : Low contrast simulation.

1220 The RP mappings seen in Figure (3.7) show very similar RP mappings to that of the
1221 simple production model, with the biggest differences occuring around the location of the
1222 break point where the low contrast model begins to dramatically underestimate $\frac{F^*}{M}$. In the
1223 high contrast simulation setting Figure (3.7; *left*), the RP mappings again demonstrate a
1224 nearly identical minimal distance mapping of RPs onto the constrained BH RP set. In the
1225 low contrast setting Figure (3.7; *right*) a very similar two regiem pattern of RP inference is
1226 observed, however the location of the break between these regiem appears at lower values
1227 of $\frac{B^*}{B(0)}$. In this moderate growth setting the break point occures around values of $\frac{B^*}{B(0)}$ just
1228 below 0.4. as opposed to the simple production model where the break point occurs at $\frac{B^*}{B(0)}$
1229 just above 0.4.

1230 3 .3 Emphatic Growth Dynamics

1231 The emphatic growth setting simulated here fixes $a_s = 2$ and $\kappa = 0.1$, to simulate a species
1232 that grows quite slowly and yet matures into the reproducing stock at a relatively early
1233 age. This combination has the effect of exaggerating the components of the model dynamics
1234 which are related to individual growth since individuals recruit at a small size and slowly
1235 grow over the extent of the modeled period.

1236 The slow growth of these dynamics oppose the simple production model setting in the
1237 sense that they move the constrained RP set a large distance (largest among the spectrum
1238 of decreasing $w(a_s)$ populations simulated here) away from the $\frac{1}{x+2}$ limiting case. It is
1239 interesting to note that this is true for all of the two parameter constrained constrained RP
1240 sets as seen in Figure (3.5).

1241 Despite the emphaticly growth driven dynamics in this setting, the RP mappings seen
1242 in Figure (3.8) obviously bare a huge resemblance to the previously seen RP mappings.
1243 Again the biggest differences in the RP mappings occur around the location of the break
1244 point where the low contrast model begins to dramatically underestimate $\frac{F^*}{M}$. In this low
1245 contrast setting the break point in RP estimation occures around values of $\frac{B^*}{B(0)}$ well below
1246 0.4 with the behaviour extending as far down as $\frac{B^*}{B(0)} = 0.3$. This regiem shift occurs well
1247 below that of the Ricker set, as initially observed in the production model setting. This
1248 reduced range of acceptable RP inference indicates that under increasingly dramatic growth

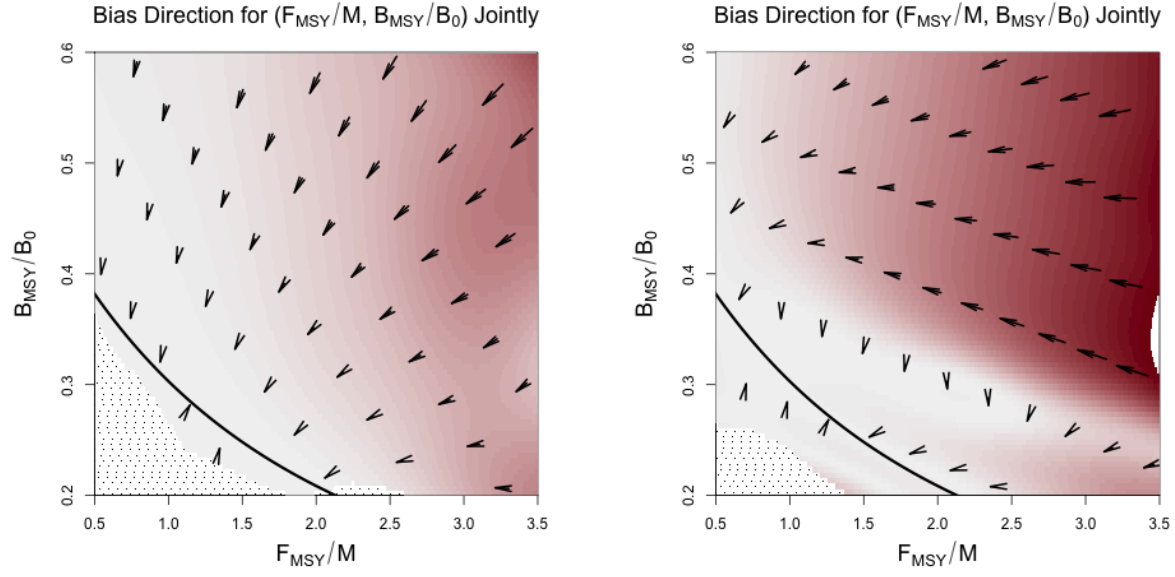


Figure 3.8: RP mapping of BH delay model fit to Schnute delay data under dramatic growth ($a_s = 2$ and $\kappa = 0.1$). *Left* : High contrast simulation. *Right* : Low contrast simulation.

the model misspecification issue of the BH model becomes an increasingly brittle assumption with respect to RPs.

Interestingly this pattern only follows for the low contrast setting. In the high contrast setting inference returns to a pattern resembling the minimal distance mapping onto BH RP set. Further pointing to the importance of contrast for informing these models.

3.4 Clustering?

- validity of clustering method?
- compare clustering thresholds on one figure

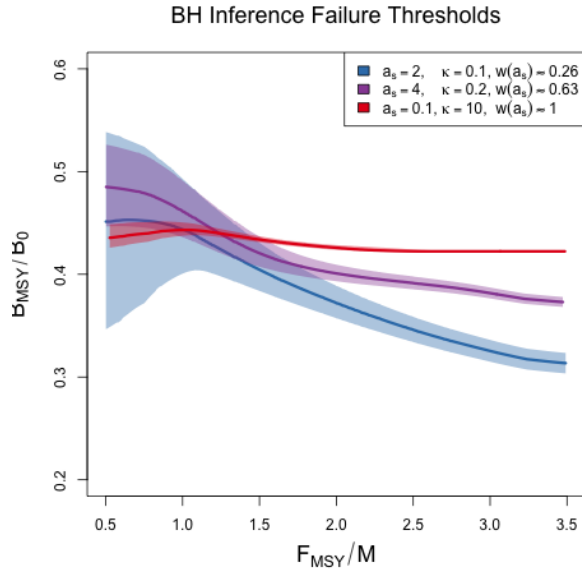


Figure 3.9: BH RP estimation failure thresholds with increasingly emphatic individual growth dynamics.

3.5 Oscillatory Growth Influence

While the above patterns of RP estimation follow for biological regimens of the $w(a_s; \kappa) = c$ level curve, as a_s increases an oscillatory regimen also exists within these dynamics. While RP estimation behaves similarly in this oscillatory regimen there are unique features in this setting that are not present in the more biological regimens. Below consider the oscillatory example of a logistic delay model with $a_s = 10$ fixing fishing at F_{MSY} .

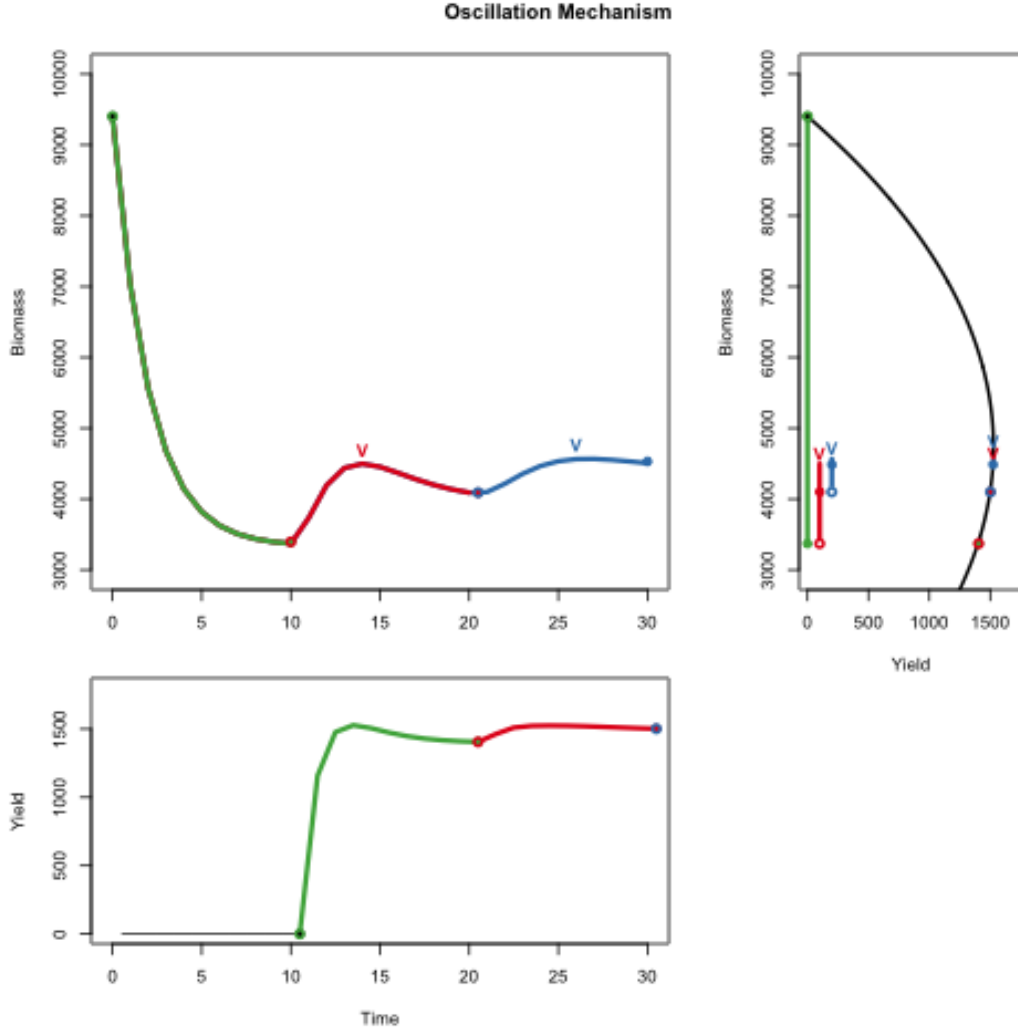


Figure 3.10: *top left* : Logistic biomass over 30 epochs of time with $a_s = 10$. Green, red, and blue colors indicate three 10 epoch long windows of biomass. v indicates local biomass oscillation maxima. *top right* : Yield plotted over the range of biomasses shown. The biomass range of each 10 epoch window is shown in the vertical colored lines. *bottom left* : Yield plotted through time. Colors correspond to the lagged biomass region that results in the evaluated yield. The black horizontal line demonstrates the pre-model assumption of biomass fixed at B_0 .

Figure (3.10) demonstrates the mechanism of how these oscillatory dynamics form. Oscillatory dynamics appear when fishing pushes biomass past B_{MSY} within the lagged a_s window of recruitment. The delay model assumes that biomass is fixed in equilibrium at B_0 , for $t \leq 0$. Therefore in the green region of the biomass series, $0 < t < 10$, the population recruits at $R(B_0)$. Figure (3.10) shows that in this initial period $R(B_0)$ results in zero yield for that period, and biomass falls as a result.

Once t exceeds a_s , the lagged recruitment refers to the integrated biomass series to evaluate recruitment based on $R(B_{t-a_s})$. The red region of the biomass series is the result of yield over the initial green biomasses. Figure (3.10) shows that the yield over the green biomass series first increases, as biomass approaches B_{MSY} and then decreases as biomass passes B_{MSY} . This creates the local maximum in the red biomass series.

Furthermore, the blue region of the biomass series is then based on yield over the red biomasses. Notice that since the red biomasses first increase and then decrease, yield increases as the red biomass increases toward B_{MSY} , and yield subsequently decreases following the descending leg of the red biomass series. This yield pattern carries the oscillation of the red biomass region forward into the blue region.

This process of biomass oscillation carries on in this manner nonetheless approaching equilibrium at B_{MSY} . Equilibrium is reached in an oscillatory manner set off by the green biomass series crossing over from above B_{MSY} to below it. The example shown in Figure (3.10) exemplifies the oscillatory phenomena simulated here, but the mechanism that produces these oscillations may occur with other forms of recruitment outside of logistic recruitment whenever fishing cases biomass to cross over B_{MSY} within the lagged recruitment window.

RP Estimation

Statistical inference in the oscillatory regime can be challenging. Depending on the parameters inferred, the likelihood can have multiple local modes which require global optimization techniques to distinguish. Furthermore, parameter estimation is more uncertain in this setting as the likelihood may confuse oscillations with residual noise.

Figure (3.11) shows the BH RP mapping fixing $w(10;0.1) \approx 0.6$ in the high contrast

simulation setting. This places the dynamics firmly in the oscillatory regiem, but the high contrast setting provides significant information for inferring recruitment parameters.

Interestingly in this high contrast setting, a very similar two regiem pattern of RP inference is observed as previously seen in low contrast settings. That said the boundary between the regiems in this setting is much smoother and the location of the break between these regiems appears around higher values of $\frac{B^*}{B(0)}$.

This higher $\frac{B^*}{B(0)}$ break point, hovering around 0.5, is consistent with the mechanism which induces ocillation. Starting the biomass at $\bar{B}(0)$ in the ocillatory regiem, increased $\frac{B^*}{B(0)}$ will tend to exasterbate oscillatory behavior by increasing B_{MSY} so that biomass is more easily pushed past B_{MSY} within the initial lagged as window of recruitment. This produces more dramatic oscillations in the higher $\frac{B^*}{B(0)}$ region of RP space.

The fitted BH model does not produce significant ocillations because under the BH model $\frac{B^*}{B(0)}$ is constrained below 0.5 with the majority of the simulation BH $\frac{B^*}{B(0)}$ RPs falling between 0.4 and 0.2. Therefore, the fitted BH model will not tend to push biomass past B_{MSY} and thus is incapable of modeling oscillatory biomass series. Figure (3.12) shows a subset of example BH fits, which demonstrats the limited oscillatory capacity of the BH fits. Furthermore, since the BH model has a limited oscillatory

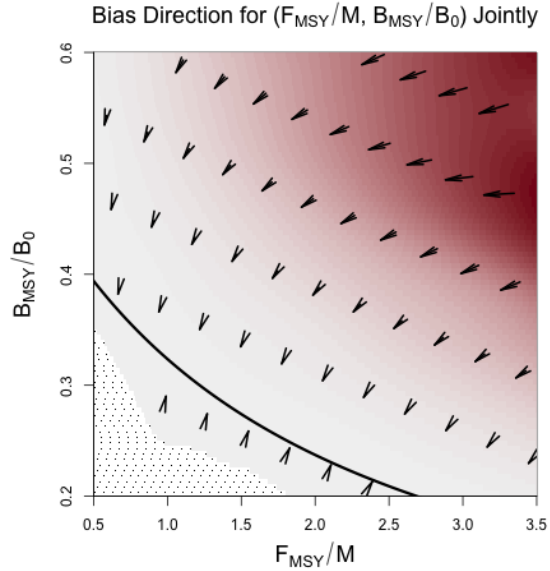


Figure 3.11: RP mapping of BH delay model fit to high contrast Schnute delay data under oscillatory growth ($a_s = 10$ and $\kappa = 0.1$).

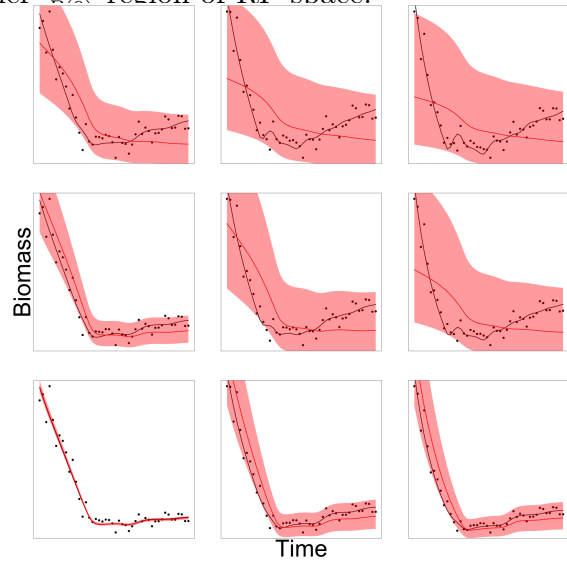


Figure 3.12: Example BH fits (red) to Schnute data (black). Each example plot is arranged to mirror its location in RP space.

capacity in this setting, the BH model tends to explain the oscillations with artificially high residual variation and artificially low steepness focusing on overly simplistic trends in the data.

Estimating More

Figure (3.13) shows a subset of example model fits broadly over RP space. Model fits are shown both under the two-parameter BH model as well as under the three parameter Schnute model, each model estimating all of its recruitment parameters as well as the growth and maturity parameters κ and a_s . Notice that the BH model, even when additionally estimating κ and a_s , does not gain the flexibility to properly model Schnute data.

The lack of oscillatory dynamics produced by the BH model causes the misspecified BH fits in Figure (3.13) to largely estimate κ and a_s so as to approximate the production model limiting case. The fitted Schnute model can produce the oscillatory dynamics and thus the information in the oscillatory data well inform estimates of κ and a_s under the Schnute model. Furthermore, the Schnute model has no issue learning its γ parameter.

While Statistical inference in the oscillatory regiem can be challenging in the highly constrained BH model, the Schnute model can easily estimate its extra γ parameter. The flexibility of estimating γ simplifies inference by correctly specifying RPs, and also by opening up the model dynamics to reveal additional information about κ and a_s in the data.

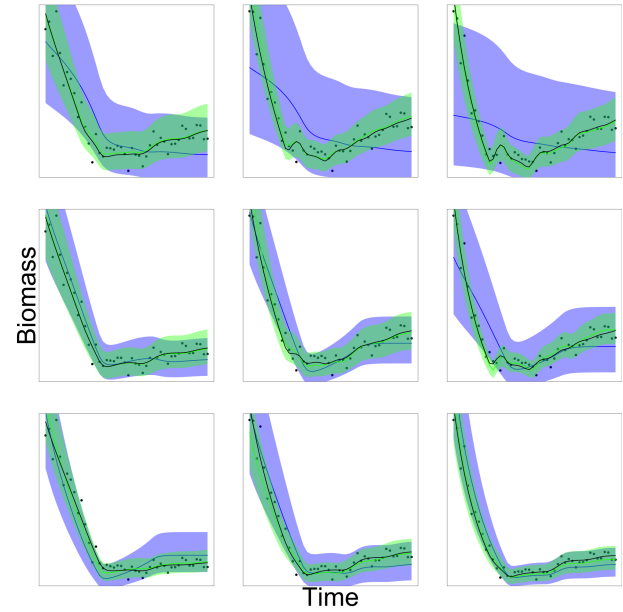


Figure 3.13: κ and a_s estimation under BH (*blue*) and Schnute (*green*) fits to Schnute data (*black*) arranged to mirror RP space.

4 Discussion

- break point decreases with growth
- inference becomes more brittle with more dramatic growth.
 - interaction between assumed form of growth and stock recruitment.
 - low-side steepness bias masks oscillatory/shock patterns induced by growth and maturity parameters
- misspecified BH prevents learning growth
- increasing growth accelerates model misspecification
- statistical evidence of minimum distance mapping within acceptable region, although float idea of PT-like pattern as BH set flattens. (explaining perturbations)

5 old ideas

- show production model limit (contrast
 - $a_s \rightarrow 0$: instant maturity
 - $\kappa \rightarrow \infty$: recruit as an adult ()
- describe second order shapes of growth/maturity (and cause)
 - weight of recruits \Rightarrow scaling biomass (q , β , and w_∞)
 -
- describe RP bias
- flat

- 1368 • summary of σ over RP space comparing between models (PT, Schnute, Schnute DD)
- 1369 to show areas of model breakdown.
- 1370 – miss-identifying signal for noise.
- 1371 – It happens more as the dynamics get more complex.
- 1372 – point to the full age structured models.
- 1373 • show the constrained BH space over a grid of $M, \kappa, \omega, W_\infty$
- 1374 • Show that the constrained spaces vary only slightly as compared with the consequences
- 1375 of misspecifying the functional form.
- 1376 • estimating these other quantities (while they can create quite different Biomass series)
- 1377 can only do so much to improve (expand) RP inference as compared with correctly
- 1378 modeling P .
- 1379 • mapping distance as a function of contrast at (3.5, 0.5)
- 1380 • for LHS grid locations show $\frac{B^*}{B_0}$ and F^* biases for grids in $M \in (0, 0.5)$ For sure in High
- 1381 Contrast, maybe also in Low??.

1382 6 Appendix: Inverting $\frac{B^*}{B(0)}$ and γ for the PT Model

For brevity let $\zeta = \frac{B^*}{B(0)}$.

$$\begin{aligned}\zeta &= \left(\frac{1}{\gamma}\right)^{\frac{1}{\gamma-1}} \\ \zeta &= \gamma \zeta^\gamma \\ \zeta &= \gamma e^{\gamma \log(\zeta)} \\ \zeta \log(\zeta) &= \gamma \log(\zeta) e^{\gamma \log(\zeta)}\end{aligned}$$

The Lambert product logarithm, W , is defined as the inverse function of $z = xe^x$ such that $x = W(z)$. Applying this definition allows for the isolation of γ .

$$\begin{aligned}\gamma \log(\zeta) &= W(\zeta \log(\zeta)) \\ \gamma &= \frac{W(\zeta \log(\zeta))}{\log(\zeta)}\end{aligned}\tag{3.13}$$

1383 The Lambert product logarithm is a multivalued function with a branch point at $-\frac{1}{e}$. The
1384 principal branch, $W_0(z)$, is defined on $z \in (-\frac{1}{e}, \infty)$, and the lower branch, $W_{-1}(z)$, is
1385 defined on $z \in (-\frac{1}{e}, 0)$. Taken individually, each respective branch is analytic, but cannot
1386 be expressed in terms of elementary functions.

1387 When $\zeta \in (0, \frac{1}{e})$ the solution of interest in Eq. (1.12) comes from W_0 . When $\zeta \rightarrow \frac{1}{e}$, the
1388 Fox Model emerges as $\gamma \rightarrow 1$. When $\zeta \in (\frac{1}{e}, 1)$ the solution of interest comes from W_{-1} . For
1389 the use case presented here, Eq. (1.12) is to be interpreted as,

$$\gamma = \begin{cases} \frac{W_0(\zeta \log(\zeta))}{\log(\zeta)} & \zeta \in (0, \frac{1}{e}) \\ \frac{W_{-1}(\zeta \log(\zeta))}{\log(\zeta)} & \zeta \in (\frac{1}{e}, 1) \end{cases}.\tag{3.14}$$

7 Appendix: The Replacement Line for the Delay Model

The replacement line is the rate of productivity which exactly balances biomass loss in the absence of fishing. In the simple production model, productivity must simply balance biomass loss due to M . Thus when $R(B; \theta) > MB$ there will be some surplus productivity to enable fishing.

In the delay model, productivity is complicated by biomass changing, both with the recruitment of young into the reproducing population, as well as biomass accumulation due to the growth of existing individuals in the population. To derive the replacement line in the case of the delay model the equilibrium equations in the absence of fishing are considered and the $R(B)$ that this implies is then isolated.

$$0 = \frac{dB}{dt} = w(a_s)R(B) + \kappa [w_\infty N - B] - MB \quad (3.15)$$

$$0 = \frac{dN}{dt} = R(B) - MN. \quad (3.16)$$

Eq(3.16) quickly gives $\bar{N} = R(B)/M$. Substituting this equilibrium value into Eq(3.15) to rewrite N in terms of B ,

$$0 = w(a_s)R(B) + \kappa \left[w_\infty \frac{R(B)}{M} - B \right] - MB. \quad (3.17)$$

Collecting like terms,

$$R(B) \left[w(a_s) + \frac{\kappa w_\infty}{M} \right] = [M + \kappa] B. \quad (3.18)$$

Finally solving for $R(B)$, and simplifying, gives the equation of the replacement line as,

$$R(B) = \left[\frac{M(M + \kappa)}{w(a_s)M + \kappa w_\infty} \right] B. \quad (3.19)$$

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