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## 1 Halibut Drafty Code

### 1.1 Types of Sample

- Sex
- Year caught
- Length
- May or may not have otolith age
- Otolith age because kin pair member
- DNAge (Not on table now)
- Should be non-lethal variable somewhere

Secret R tip: ?Quotes

### 1.2 The code we wrote

```
enum sex{
  male == 0,
  female == 1
};

enum mt{
  shared == 0,
  unshared == 1
};

template<class Type>
get_fec(Type length, int sex){
  Type fec;
  //alpha, wexp are inputs
  Type weight = alpha[sex]*pow(weight,wexp[sex]);
  //ppnmature is an input, proportion mature at length given sex
  Type fec = pow(weight,omega[sex])*ppnmature(length,sex);
  return fec;
}
```

```

array<Type> meanfec_sa(sexes,A);
for(int s = 0; s < sexes; ++s){
    for(int a = 0; a < A; ++A){
        Type runningfec = 0.0;
        for(int l = 0; l < L_max; ++l){
runningfec += Pr_l_given_as(a,s)*get_fec(l,s);
        }
        meanfec_sa(s,a) = runningfec;
    }
}

```

```

array<Type> TR0(S,Y);
for(int s = 0; s < sexes; ++s){
    for(int y = 0; y < Y; ++y){
        Type partialR0 = 0.0;
        for(int a = 0; a < A; ++a){
            partialR0 += N(a,y)*meanfec_sa(s,a);
        }
        TR0(s,y) = partialR0;
    }
}

```

```

for(int s1 = 0; s1 < blah.dim(s1); s1++){
    for(int y1 = 0; y1 < blah.dim(y1); y1++){
        for(int l1 = 0; l1 < blah.dim(l1); l1++){
            for(int a1 = 0; a1 < blah.dim(a1); a1++){
for(int y2 = 0; y2 < blah.dim(y2); y2++){
            for(int a2 = 0; a2 < blah.dim(a2); a2++){
                int B2 = y2-a2;
                int B1 = y1-a1;
                Type Prob;
                if(y1 < B2){
                    Prob = 0;
                }
                else if(B1 >= B2){
                    Prob = 0;
                }
                else{
                    Type L_inf_A = l1/(1-exp(-k_[s1]*(a1-t0[s1])));
                    //age at juv birth year
                    int a_b2 = a1-(y1-B2);
                    Type l_b2 = L_inf_A*(1-exp(-k_[s1]*(a1-t0[s1])));
                    //what is get_fec and TR0?
                    Prob = get_fec(s1,l_b2)/TR0(s1,B2);
                }
            }
        }
    }
}

```

```

        Pr_MOP_SYLAYA[s1,y1,l1,a1,y2,a2] = Prob;
    }
    }
}
}

array<Type> Pr_A_SYL(a,l,s,y);
for(int l = 0; l < blah.dim(l); l++){
    for(int s = 0; s < blah.dim(s); s++){
        for(int y = 0; y < blah.dim(y); y++){
Type Prob_denom = 0.0;
for(int a = 0; a < blah.dim(a); a++){
    Type Prob_num = (Pr_l_given_as(a,s)*(N(a,s,y)));
    Prob_denom += Prob_num
    Pr_A_SYL(a,l,s,y) = Prob_num;
}
for(int a = 0; a < blah.dim(a); a++){
    Pr_A_SYL(a,l,s,y) /= Prob_denom;
}
    }
}
}

```

```

//uncertain adult ages
for(int s1 = 0; s1 < blah.dim(s1); s1++){
    for(int y1 = 0; y1 < blah.dim(y1); y1++){
        for(int l1 = 0; l1 < blah.dim(l1); l1++){
for(int y2 = 0; y2 < blah.dim(y2); y2++){
    for(int a2 = 0; a2 < blah.dim(a2); a2++){
        Pr_POP_running = 0.0;
        for(int a1 = 0; a1 < someblah.dim(a1); a1++){

            Pr_POP_running += Pr_MOP_SYLAYA[s1,y1,l1,a1,y2,a2]*Pr_A_SYL[a1,l1,s1,y1];
        }
        Pr_POP_sylya[s1,y1,l1,y2,a2] = Pr_POP_running;
    }
}
    }
}
}

```

```

//uncertain juv ages
for(int s1 = 0; s1 < blah.dim(s1); s1++){
    for(int y1 = 0; y1 < blah.dim(y1); y1++){
        for(int l1 = 0; l1 < blah.dim(l1); l1++){

```

```

for(int a1 = 0; a1 < blah.dim(a1); a1++){
    for(int l2 = 0; l2 < blah.dim(l2); l2++){
        for(int s2 = 0; s2 < blah.dim(s2); s2++){
            for(int y2 = 0; y2 < blah.dim(y2); y2++){
//summing over possible juv ages
Type Pr_POP_running = 0.0;
for(int a2p = 0; a2p < A; a2p++){
    Pr_POP_running += Pr_A_SYL[a2p,l2,s2,y2]*Pr_MOP_SYLAYA[s1,y1,l1,a1,y2,a2p]
}
Pr_POP_sylalsy[s1,y1,l1,a1,l2,s2,y2] = Pr_POP_running;
}
}
}
}

//HSPs

// \ (Pr[OHSP
array<Type> Pr_OHSP_sbb(sexes,Y,Y);
for(int sp = 0; sp < sexes; sp++){
for(int B1 = 0; B1 < Y; B1++){
    for(int B2 = B1; B2 < Y; B2++){
        //Handle same cohort case at the end
        Type Pr_HSP_running = 0.0;
        //potential ages of parent
        for(int app = 0; app < A; app++){
            Type Pr_surv = N(app+(B2-B1),sp,B2)/N(app,sp,B1);
            Type pp_ERO_B2 = 0.0;
            Type denom = 0.0;
            for(int q = 0; q < num_qs; q++){
//length_at_age_q gets length at age given quantile
Pr_pp_is_qth_q_1 = 1/num_qs*get_fec(length_at_age_q(q,app,sp),sp);
denom += Pr_pp_is_qth_q_1;
pp_ERO_B2_if_q = get_fec(length_at_age_q(a,app+(B2-B1),sp),sp);
pp_ERO_B2 += Pr_pp_is_qth_q_1*pp_ERO_B2_if_q;
}
pp_ERO_B2 /= denom;
Pr_pp_is_1s_mum = (meanfec_sa(sp,app)/TRO(sp,B1));
Pr_HSP_running += N(app,sp,B1)*Pr_pp_is_1s_mum*
Pr_surv*
pp_ERO_B2/TRO(sp,B2);

        //compensate for lucky litter (same cohort)
        if(B2 == B1){
//lucky_litter_factor is a PARAMETER_VECTOR
Pr_HSP_running *= lucky_litter_factor(sp);

```

```

    }
    Pr_OHSP_sbb[sp,B1,B2] = Pr_HSP_running;
  }
}

  }
}

  }

// unordered HSPs (should really be done on SO_Kins)
// Should deal with ordering of POPs
for(int sp = 0; sp < sexes; sp++){
  for(int B1 = 0; B1 < Y; B1++){
    for(int B2 = B1; B2 < Y; B2++){
//p_false_neg is a data input from kinference or CKMRsim
Pr_HSP_sbb[sp,B2,B1] = Pr_OHSP_sbb[sp,B1,B2];
Pr_HSP_sbb[sp,B1,B2] = Pr_OHSP_sbb[sp,B1,B2];
    }
  }
}

array<Type> Pr_HSP_slysb(sexes,L_max,Y,sexes,Y);

for(int sp = 0; sp < sexes; sp++){
  for(int l1 = 0; l1 < L_max; l1++){
    for(int y1 = 0; y1 < Y; y1++){
for(int s1 = 0; s1 < sexes; s1++){
  for(int B2 = 0; B2 < Y; B2++){

    //summing over possible l1 ages
    Type Pr_HSP_running = 0.0;
    for(int a1 = 0; a1 < A; a1++){
      Pr_HSP_running += Pr_A_SYL[a1,l1,s1,y1]*Pr_HSP_sbb[sp,y1-a1,B2];
    }
    Pr_HSP_slysb[sp,l1,y1,s1,B2] = Pr_HSP_running;
  }
}
  }
}

//when we don't know both hsp ages

//grandparent grandoffspring
array<Type> Pr_GGP_syabs(sexes,Y,A,Y,sexes);
for(int s1 = 0; s1 < sexes; s1++){
  for(int y1 = 0; y1 < Y; y1++){

```

```

        for(int a1 = 0; a1 < A; a1++){
for(int B2 = 0; B2 < Y; B2++){
    for(int spp = 0; spp < sexes; spp++){
        int B1 = y1-a1;
        //year 1 mature
        int mat_1_y = min_age_mature(s1)+B1;
        Type Pr_GGP_running = 0.0;
        for(int app = B2-y1; app < B2-mat_1_y; app++){
            int Bpp = B2-app;
            Type Pr_pp_is_2s_mum = (meanfec_sa(spp,app)/TRO(spp,B2));
            int a1_bpp = Bpp-B1;
            Type Pr_1_is_pps_mum = (meanfec_sa(s1,a1_bpp)/TRO(s1,Bpp));
            Pr_GGP_running += Pr_pp_is_2s_mum*Pr_1_is_pps_mum;
        }
        Pr_GGP_syabs(s1,y1,a1,B2,spp) = Pr_GGP_running;
    }
}
    }
}

array<Type> Pr_SO_kins(smtdna,sexes,Y,A,sexes,Y,A);
    for(int s1 = 0; s1 < sexes; s1++){
        for(int y1 = 0; y1 < sexes; y1++){
for(int a1 = 0; a1 < A; a1++){
    for(int s2 = 0; s2 < sexes; s2++){
        for(int y2 = 0; y2 < Y; y2++){

            for(int a2 = 0; a2 < A; a2++){
int B1 = y1 - a1;
int B2 = y2 - a2;
//shared mt case first
Type Pr_SO_kins_temp = (Pr_HSP_sbb[female,B1,B2]);
if((B2 < B1) & s2 == female){
    Pr_SO_kins_temp += Pr_GGP_syabs[female,y2,a2,B1,female];
}else if((B1 < B2) & s1 == female){
    Pr_SO_kins_temp += Pr_GGP_syabs[female,y1,a1,B2,female];
}
//shared is enum mt var
Pr_SO_kins(shared,s1,y1,a1,s2,y2,a2) = (1-p_false_neg)*Pr_SO_kins_temp;

//unshared mt case
else if((B2 < B1) & s2 == male){
    Pr_SO_kins_temp += Pr_GGP_syabs[male,y2,a2,B1,female]+Pr_GGP_syabs[male,y2,a2,B1,male];
}else if((B1 < B2) & s1 == male){
    Pr_SO_kins_temp += Pr_GGP_syabs[male,y1,a1,B2,female]+Pr_GGP_syabs[male,y1,a1,B2,male];
}
Pr_SO_kins(unshared,s1,y1,a1,s2,y2,a2) = (1-p_false_neg)*Pr_SO_kins_temp;

```

```

    }
  }
}
  }
}
}

```

```

// Age composition data
for(int s = 0; s < sexes; s++){
  for(int y = 0; y < Y; y++){
    for(int l = 0; l < L_max; l++){
for(int a = 0; a < A; a++){
  //change distro
  nll -= dpois(n_otos_at_age(a,l,s,y),
    //tot number of otoliths
n_otos(l,s,y)*Pr_A_SYL[s,y,l],true);
}
  }
}
}

```

```

for(int y1 = 0; y1 < blah.dim(y1); y1++){
  for(int a1 = 0; a1 < blah.dim(a1); a1++){
    for(int y2 = 0; y2 < blah.dim(y2); y2++){
      for(int a2 = 0; a2 < blah.dim(a2); a2++){
for(int sp = 0; sp < sexes; sp++){
  int n_comps = n_comps_HS_yaya[y1,a1,y2,a2];
  int n_S0_kins = n_S0_kins_syaya[sp,y1,a1,y2,a2];
  nll -= dpois(n_S0_kins,n_comps*Pr_S0_kins,true);
}
  }
}
}
}

```

```

for(int s1 = 0; s1 < blah.dim(s1); s1++){
  for(int y1 = 0; y1 < blah.dim(y1); y1++){
    for(int l1 = 0; l1 < blah.dim(l1); l1++){
      for(int a1 = 0; a1 < blah.dim(a1); a1++){
for(int y2 = 0; y2 < blah.dim(y2); y2++){
  for(int a2 = 0; a2 < blah.dim(a2); a2++){

```

```

    int n_comps = n_comps_POP_sylaya[s1,y1,l1,a1,y2,a2];
    int n_POPs = n_POP_sylaya[s1,y1,l1,a1,y2,a2];
    //negative log likelihood
    nll -= dpois(n_POPs,n_comps*Pr_MOP_SYLAYA[s1,y1,l1,a1,y2,a2],true);
  }
}
  }
}
}
}

//Post-hoc

return nll;
}

```

## 2 A Basic CKMR model in TMB

Here is a simple CKMR model using TMB based off what I've done to maybe help show how the peices fit together. This uses some data from one of my simulations so there's 3 age classes and sampling is done non-lethally.

This model only uses POPs so it can't estimate survival and that's assumed known and passed into the model (the Z)). It's also assumed to be the same every year. If HSPs were added it is possible to estimate Z.

```

// This is needed to actually load in the TMB library
#include <TMB.hpp>

//Your TMB objective function (the model) will always start with these two lines
template<class Type>
Type objective_function<Type>::operator() (){

  //Parameter and data setup

  //The optimizer works best on -Inf to Inf.
  // By passing in parameters on log space and transforming inside the model
  // we keep the optimizer happy while ensuring things that should be positive are
  // always postive.

  PARAMETER(log_init_tot_N);
  PARAMETER_VECTOR(log_fecundity);

  vector<Type> fecundity = exp(log_fecundity);

  DATA_INTEGER(Y);
  DATA_INTEGER(A);
  DATA_VECTOR(Z);

```



```

DATA_VECTOR(init_frac);
DATA_IMATRIX(POPs);

//Building the Numbers at age matrix
matrix<Type> log_N(A,Y);

//setting initial numbers
for(int a = 0; a < A; ++a){
    log_N(a,0) = log_init_tot_N+log(init_frac(a));
}

//Our negative log likelihood, what we are maximizing
Type nll = 0.0;

//Population dynamics
//We need the pop dynamics to figure out the numbers at age for our expected probabilities

//Propagate the numbers at age for each year:
for(int y = 1; y < Y; ++y){
    //Recruitment, probably should be a random effect...
    //Here I just set it to be the number of offspring from last year
    Type TR0 = 0.0;
    for(int a = 0; a < A; ++a){
        TR0 += fecundity(a)*exp(log_N(a,y-1));
    }
    log_N(0,y) = log(TR0);

    for(int a = 1; a < A; ++a){
        //fraction of fish that survive from last year to the current year
        log_N(a,y) = log_N(a-1,y-1)-Z(a-1);
    }
}

//The actual CKMR bit
// I do one matrix rather than a million nested for loops
// The POPs matrix contains all the data we need, like a flattened array
for(int i = 0; i < POPs.rows(); ++i){
    //pull out the correct covariates/data from each row:
    int juv_by = POPs(i,0);
    int par_by = POPs(i,1);
    int par_sy = POPs(i,3);
    //How many real POPs we found
    Type n_POPs = POPs(i,5);
    Type n_comps = POPs(i,4);

    //Finding the expected probability
    //Get the TR0:
    Type TR0 = 0.0;

```

```

    for(int a = 0; a < A; ++a){
        TR0 += fecundity(a)*exp(log_N(a,juv_by));
    }
    int p_age = juv_by-par_by-1; //the 1 is because C++ starts everything at 0
    Type eprob = (2.0*fecundity(p_age))/TR0;

    // Since non-lethal sampling adjust for survival of parent
    if(par_sy < juv_by){
        //parents age at sampling
        int p_age_sy = par_sy-par_by-1;
        Type Pr_surv = exp(log_N(p_age,juv_by)-log_N(p_age_sy,par_sy));
        eprob *= Pr_surv;
    }

    //Add the contribution to the likelihood
    nll -= dpois(n_POPs,n_comps*eprob,true);
}

matrix<Type> N(A,Y);
N = log_N.array().exp();

//Get anything we want out of the model and back into R
REPORT(N);
REPORT(fecundity);
//Get the standard error for that quantity:
ADREPORT(log_N);
ADREPORT(log_fecundity);

return nll;
}

```

## 2.1 Running this model from R

```

library(TMB)

##load in the required data for the model, an object called stuff

load("basicCKMRdat.Rdata")

##Compile the model
compile("basicCKMR.cpp")

##Load the compiled model
dyn.load(dynlib("basicCKMR"))

```

```

##annoying bit because there isn't enough data to freely estimate the initial numbers at age
##It's basically the fraction of the numbers at age in the population
##Can be done inside with Z if estimated but easier here since Z isn't
init_frac = c(1,exp(-stuff$Z[1]),exp(-stuff$Z[1]-stuff$Z[2]))
init_frac = init_frac/sum(init_frac)

##set up data and parameters to give to TMB model, must be as named lists
## matching what's in the model
dat = list()
dat$Y = 10
dat$A = 3
dat$Z = c(0.9395,1.670,0)
dat$init_frac = init_frac
##The actual important data, number of POPs found and number of comparisons
dat$POPs = as.matrix(stuff$POPs)

##Our initial guess for parameters
parm = list()
parm$log_init_tot_N = log(500)
parm$log_fecundity = log(c(1.1,1.1,1.1))

##Make the objective function with TMB
obj = MakeADFun(dat,parm)

##optimize the model to get the most likely parameters given the data
opt = nlminb(obj$par,obj$fn,obj$gr)

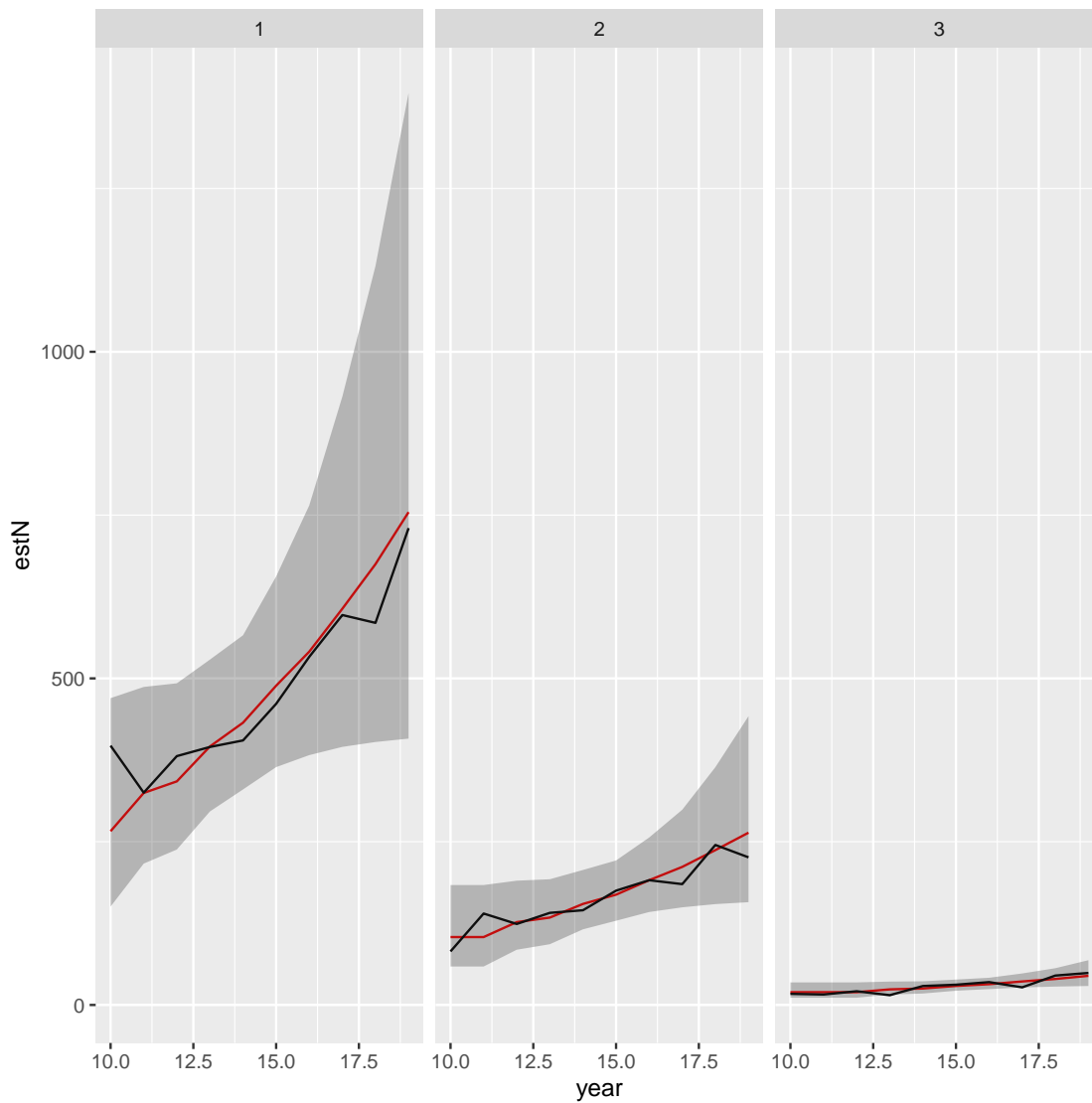
##get the data out from the model, make confidence intervals
report = obj$report()
sdr = sdreport(obj)
logN = as.data.frame(summary(sdr)[which(rownames(summary(sdr)) == "log_N"),])
logN$year = sort(rep(10:19,3))
logN$age = rep(1:3,10)
logN$lower = exp(logN[,1] - 1.96*logN[,2])
logN$upper = exp(logN[,1] + 1.96*logN[,2])
logN$estN = exp(logN[,1])

logFec = as.data.frame(summary(sdr)[which(rownames(summary(sdr)) == "log_fecundity"),])
logFec = logFec[1:3,]
logFec$fec = exp(logFec[,1])
logFec$lower = exp(logFec[,1] - 1.96*logFec[,2])
logFec$upper = exp(logFec[,1] + 1.96*logFec[,2])

##compare with true N and fecundity
trueN = stuff$trueN
logN$trueN = as.vector(trueN)
trueFec = stuff$trueFec

```

```
library(ggplot2)
Nplot = ggplot(logN) + geom_line(aes(x=year,y=estN),color="red") + geom_line(aes(x=year,y=trueN))
```



The red line is the estimate from the model, black is simulation and the ribbon is the 95% CI. The model gets the abundance back well over the years.

Note that the fecundity for age 3 estimate is too small and it still lies outside the 95% CI. This isn't a big sample from a small population and age 3's are particularly rare (there's about 1 or 2 a year in the sample). I also didn't place any constraints on fecundity other than it being positive.