Utah Chub CR

POPULATION SIZE, MIGRATION, DIVERGENCE, ASSIGNMENT, HISTORY

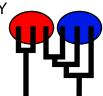
Bayesian inference using the structured coalescent

Migrate-n version 4.4.4(git:v4-series-26-ge85c6ff) [June-1-2019]

Compiled for a SYMMETRIC multiprocessors (Grandcentral)

Program started at Wed May 19 21:08:40 2021

Program finished at Wed May 19 23:19:17 2021 [Runtime:0000:02:10:37]



Options

Inheritance scalers in use for Thetas:

All loci use an inheritance scaler of 1.0

[The locus with a scaler of 1.0 used as reference]

Random number seed: (with internal timer) 925931068

Start parameters:

Theta values were generated RANDOM start value from the prior

M values were generated RANDOM start value from the prior

Connection matrix:

m = average (average over a group of Thetas or M,

s = symmetric migration M, S = symmetric 4Nm,

0 = zero, and not estimated,

* = migration free to vary, Thetas are on diagonal

d = row population split off column population, D = split and then migration

Population 1 2 3 1 GSL * d 0 2 GSLD 0 * d 3 SEV 0 0 *

Order of parameters:

1	Θ_1	<displayed></displayed>
2	Θ_2	<displayed></displayed>
3	Θ_3^-	<displayed></displayed>

							U	tah Chub CR:
4	Δ 2-	->1		<displa< td=""><td>yed></td><td></td><td></td><td></td></displa<>	yed>			
5	σ 2.	->1		<displa< td=""><td>yed></td><td></td><td></td><td></td></displa<>	yed>			
6	٨			<displa< td=""><td>ved></td><td></td><td></td><td></td></displa<>	ved>			
•	△ 3.	->2		\\ \alpha\\ \ext{opia}	your			
7	σ 3.	->2		<displa< td=""><td>yed></td><td></td><td></td><td></td></displa<>	yed>			
Mutation rate an	nong la	oci:					Mutation ra	ate is constant
Analysis strategy	y:						Baye	esian inference
-Population size	estim	ation:					Exponen	tial Distribution
-Geneflow estim	nation:						Exponen	tial Distribution
-Divergence tim	e estin	nation:		No	rmal Distrib	oution Shortcu	t (mean and	standard dev.)
Proposal distribu	ıtions f	or paramete	er					
Parameter			Prop	oosal				
Theta		Metropolis sampli		pling				
M		M	1etropolis sam	pling				
Divergence		M	1etropolis sam	pling				
Divergence Sprea	ad	M	1etropolis sam	pling				
Genealogy		N	/letropolis-Has	tings				
Prior distribution	for pa	rameter						
Parameter		Prior	Minimum	MeanMa	aximum	Delta	Bins U	JpdateFreq
1 The	ta **	Uniform	0.000000	0.010	0.100	0.010	1500	0.07143
2 The	ta **	Uniform	0.000000	0.010	0.100	0.010	1500	0.07143
3 The	ta **	Uniform	0.000000	0.010	0.100	0.010	1500	0.07143
4 Splittime mea	an **	Uniform	0.000000	0.010	0.500	0.050	1500	0.07143
5 Splittime s	td **	Uniform	0.000000	0.010	0.500	0.050	1500	0.07143
6 Splittime mea	an **	Uniform	0.000000	0.010	0.500	0.050	1500	0.07143
7 Splittime s	td **	Uniform	0.000000	0.010	0.500	0.050	1500	0.07143
[* * means priors	were	set globally]					
Markov chain se	ttings:							Long chain
	-							-

Markov chain settings:Long chairNumber of chains1Recorded steps [a]10000Increment (record every x step [b]1000Number of concurrent chains (replicates) [c]2Visited (sampled) parameter values [a*b*c]20000000Number of discard trees per chain (burn-in)1000

Multiple Markov chains:

Static heating scheme		4 chains	with temperatures
	1000000.00	3.00	1.50 1.00
		Swa	apping interval is 1
Print options:			
Data file:			infile
Haplotyping is turned on:			NO
Output file:			outfile
Log file:			logfile
Posterior distribution raw histogram file:			bayesfile
Raw data from the MCMC run:			bayesallfile.gz
Print data:			No
Print genealogies [only some for some data type]:			None

Data summary

Data file: infile

Datatype: Haplotype data

Number of loci:

Mutationmodel:

Locus Sublocus Mutationmodel Mutationmodel parameters

1 1 HKY [Bf:0.32 0.21 0.14 0.32, kappa=4.610]

Sites per locus

Locus Sites

1 935

Site rate variation and probabilities:

Locus Sublocus Region type Rate of change Probability Patch size

1	1	1	1.000	1.000	1.000		
Populat	tion				Locus	Gene co	ppies
						data	(missing)
1 GSL					1	22	
2 GSLE)				1	15	
3 SEV					1	10	
Total of	all popu	lations			1	47	(0)

Bayesian Analysis: Posterior distribution table

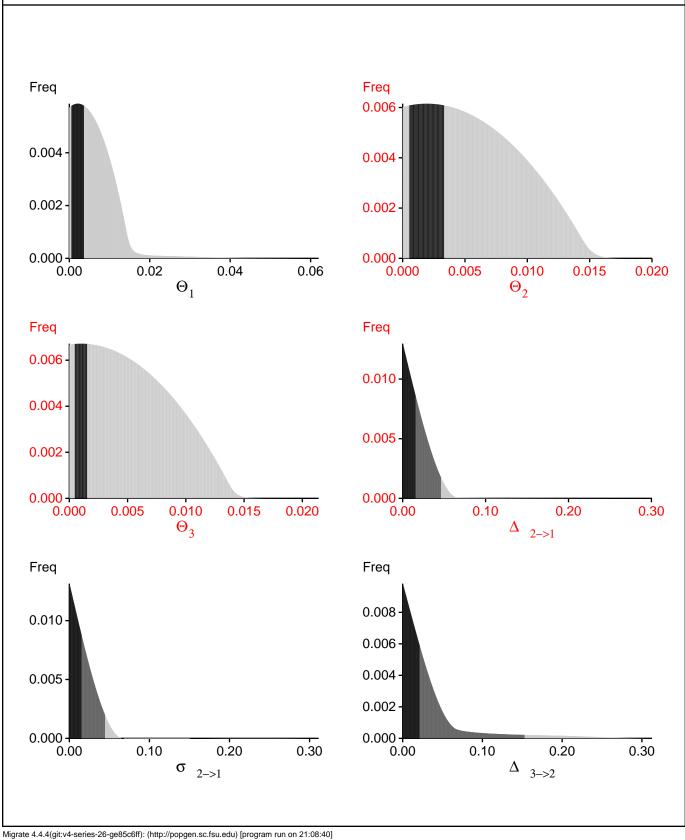
Locus	Parameter	2.5%	25.0%	Mode	75.0%	97.5%	Median	Mean
1	Θ1	0.00047	0.00047	0.00210	0.00367	0.00367	0.00583	0.00394
1	Θ_2	0.00047	0.00047	0.00197	0.00333	0.00333	0.00557	0.00194
1	Θ_3	0.00040	0.00040	0.00103	0.00153	0.00153	0.00517	0.00102
1	D _{2->1}	0.00000	0.00000	0.00017	0.01567	0.04633	0.01583	0.03139
1	S _{2->1}	0.00000	0.00000	0.00017	0.01533	0.04467	0.01550	0.14419
1	D _{3->2}	0.00000	0.00000	0.00017	0.02133	0.15300	0.02150	0.08718
1	S _{3->2}	0.00000	0.00000	0.00017	0.04567	0.22300	0.04583	0.16529

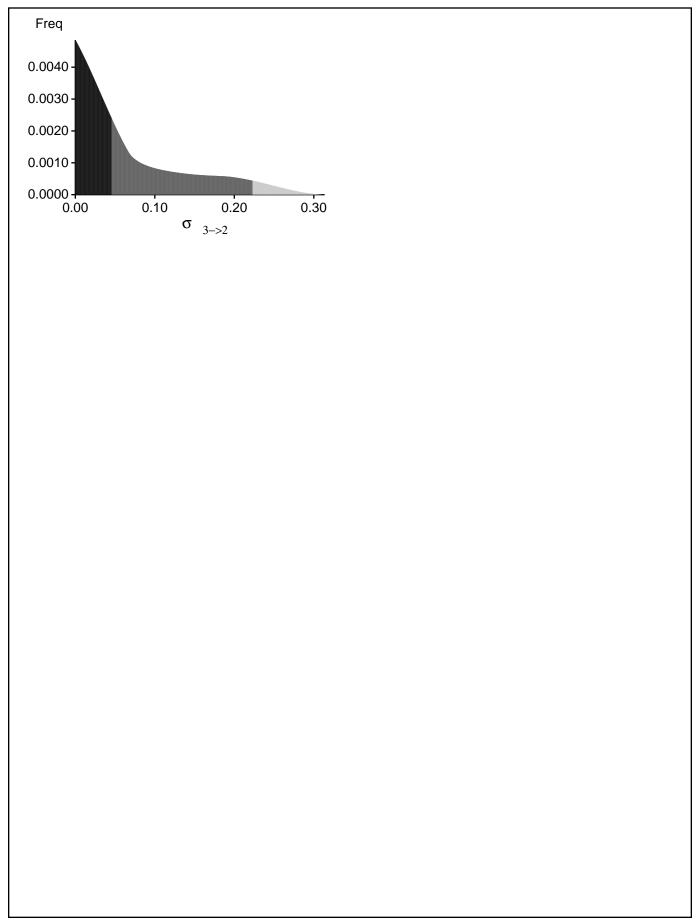
Citation suggestions:

Beerli P., 2006. Comparison of Bayesian and maximum-likelihood inference of population genetic parameters. Bioinformatics 22:341-345

Beerli P., 2009. How to use MIGRATE or why are Markov chain Monte Carlo programs difficult to use? In Population Genetics for Animal Conservation, G. Bertorelle, M. W. Bruford, H. C. Hauffe, A. Rizzoli, and C. Vernesi, eds., vol. 17 of Conservation Biology, Cambridge University Press, Cambridge UK, pp. 42-79.

Bayesian Analysis: Posterior distribution for locus 1





Log-Probability of the data given the model (marginal likelihood)

Use this value for Bayes factor calculations:

BF = Exp[ln(Prob(D | thisModel) - ln(Prob(D | otherModel) or as LBF = 2 (ln(Prob(D | thisModel) - ln(Prob(D | otherModel)) shows the support for thisModel]

Method	In(Prob(D Model))	Notes
Thermodynamic integration	-2177.972899	(1a)
	-1619.180224	(1b)
Harmonic mean	-1482.053864	(2)

(1a, 1b and 2) are approximations to the marginal likelihood, make sure that the program run long enough! (1a, 1b) and (2) should give similar results, in principle.

But (2) is overestimating the likelihood, it is presented for historical reasons and should not be used (1a, 1b) needs heating with chains that span a temperature range of 1.0 to at least 100,000.

(1b) is using a Bezier-curve to get better approximations for runs with low number of heated chains

Citation suggestions:

Beerli P. and M. Palczewski, 2010. Unified framework to evaluate panmixia and migration direction among multiple sampling locations, Genetics, 185: 313-326.

Acceptance ratios for all parameters and the genealogies

Parameter	Accepted changes	Ratio
Θ_1	541852/1426596	0.37982
Θ_2	857176/1429977	0.59943
Θ_3	506287/1430550	0.35391
$\Delta_{2\rightarrow 1}$	625280/1426991	0.43818
$\sigma_{2\rightarrow 1}$	980039/1427854	0.68637
Δ $3->2$	670650/1429388	0.46919
$\sigma_{3\rightarrow 2}$	750838/1428062	0.52577
Genealogies	1724425/10000582	0.17243

MCMC-Autocorrelation and Effective MCMC Sample Size

Parameter	Autocorrelation	Effective Sampe Size
Θ_1	0.57334	7371.17
Θ_2	0.66891	4610.59
Θ_3^2	0.33509	9977.30
$\Delta^{\circ}_{2\rightarrow 1}$	0.16873	14627.63
$\sigma_{2\rightarrow 1}$	0.69668	4348.62
$\Delta_{3\rightarrow 2}$	0.19353	13512.82
$\sigma_{3\rightarrow 2}$	0.28301	11175.63
Genealogies	0.28301	11175.63

Potential Problems

This section reports potential problems with your run, but such reporting is often not very accurate. Whith many parameters in a multilocus analysi s, it is very common that some parameters for some loci will not be very informative, triggering suggestions (for example to increase the prior ran ge) that are not sensible. This suggestion tool will improve with time, therefore do not blindly follow its suggestions. If some parameters are fla

gged, inspect the tables carefully and judge wether an action is required. For example, if you run a Bayesian inference with sequence data, for mac roscopic species there is rarely the need to increase the prior for Theta
beyond 0.1; but if you use microsatellites it is rather common that your prior distribution for Theta should have
a range from 0.0 to 100 or more. With many populations (>3) it is also very common that some migration rou
tes are estimated poorly because the data contains little or no information for that route. Increasing the range will
not help in such situations, reducing number of parameters may help in such situations.
No warning was recorded during the run