## **AUTO**

POPULATION SIZE, MIGRATION, DIVERGENCE, ASSIGNMENT, HISTORY

Bayesian inference using the structured coalescent

Migrate-n version 5.0.0a [May-20-2017]

Using Intel AVX (Advanced Vector Extensions)

Compiled for PARALLEL computer architectures

One master and 20 compute nodes are available.

Program started at Mon Nov 6 22:06:06 2017

Program finished at Mon Nov 6 23:43:34 2017 [Runtime:0000:01:37:28]



#### **Options**

Datatype: DNA sequence data

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) 974312299

Start parameters:

Theta values were generated

Using a percent value of the prior

M values were generated Using a percent value of 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 1 1 Romanshorn\_0 \* \* 1 Arbon 1 \* \*

Order of parameters:

1  $\Theta_1$  <displayed>

Mutation rate among loci: Mutation rate is constant for all loci

Analysis strategy: Bayesian inference -Population size estimation: **Exponential Distribution** 

Proposal distributions for parameter

Parameter Proposal Theta Metropolis sampling M Metropolis sampling Divergence Metropolis sampling Divergence Spread Metropolis sampling Genealogy Metropolis-Hastings

Prior distribution for parameter

Parameter Prior Minimum Mean Maximum Delta Bins UpdateFreq Theta -11 Uniform 1500 0.20000 0.000000 0.050 0.100 0.010

[-1 -1 means priors were set globally]

Markov chain settings: Long chain

Number of chains

50000 Recorded steps [a] Increment (record every x step [b] 200 2 Number of concurrent chains (replicates) [c]

20000000 Visited (sampled) parameter values [a\*b\*c]

10000 Number of discard trees per chain (burn-in)

Multiple Markov chains:

Static heating scheme 4 chains with temperatures

> 1000000.00 3.00 1.50 1.00

Swapping interval is 1

Print options:

Data file: infile.1.0

Haplotyping is turned on: NO

Output file: outfile\_1.0\_0.5

bayesfile Raw data from the MCMC run: bayesallfile\_1.0\_0.5

Print data: No

Print genealogies [only some for some data type]: None

Posterior distribution raw histogram file:

## Data summary

Data file: infile.1.0
Datatype: Sequence data

Number of loci:

Mutationmodel parameters

#### Mutationmodel: Locus Sublocus

1	1	Jukes-Cantor	[Basefreq: =0.25]
2	1	Jukes-Cantor	[Basefreq: =0.25]
3	1	Jukes-Cantor	[Basefreq: =0.25]
4	1	Jukes-Cantor	[Basefreq: =0.25]
5	1	Jukes-Cantor	[Basefreq: =0.25]
6	1	Jukes-Cantor	[Basefreq: =0.25]
7	1	Jukes-Cantor	[Basefreq: =0.25]
8	1	Jukes-Cantor	[Basefreq: =0.25]
9	1	Jukes-Cantor	[Basefreq: =0.25]
10	1	Jukes-Cantor	[Basefreg: =0.25]

Mutationmodel

#### Sites per locus

Locus	Sites
1	10000
2	10000
3	10000
4	10000
5	10000
6	10000
7	10000
8	10000
9	10000
10	10000

Site rate variation and probabilities:

Locus Sublocus Region type Rate of change Probability Patch size

1	1	1	1.000	1.000	1.000
2	1	1	1.000	1.000	1.000
3	1	1	1.000	1.000	1.000
4	1	1	1.000	1.000	1.000
5	1	1	1.000	1.000	1.000
6	1	1	1.000	1.000	1.000

		4	1.000	4.000	4.000	
7	1	1	1.000	1.000	1.000	
8	1	1	1.000	1.000	1.000	
9	1	1	1.000	1.000	1.000	
10	1	1	1.000	1.000	1.000	0
Popula					Locus	Gene copies
1 Rom	anshorn_	0			1	10
					2	10
					3	10
					4	10
					5	10
					6	10
					7	10
					8	10
					9	10
					10	10
1 Arbo	n_1				1	10
					2	10
					3	10
					4	10
					5	10
					6	10
					7	10
					8	10
					9	10
					10	10
Total o	f all popul	lations			1	20
					2	20
					3	20
					4	20
					5	20
					6	20
					7	20
					8	20
					9	20
					10	20

## Bayesian Analysis: Posterior distribution table

Locus	Parameter	2.5%	25.0%	Mode	75.0%	97.5%	Median	Mean
1	$\Theta_1$	0.04653	0.04760	0.04863	0.04960	0.05067	0.04870	0.09426
2	$\Theta_1$	0.00000	0.00000	0.00003	0.09993	0.09993	0.00003	0.09415
3	$\Theta_1$	0.04407	0.04680	0.04770	0.04860	0.05133	0.04617	0.09409
4	$\Theta_1$	0.00000	0.00000	0.00003	0.09993	0.09993	0.00003	0.09400
5	$\Theta_1$	0.00000	0.00000	0.00003	0.09993	0.09993	0.00003	0.09408
6	$\Theta_1$	0.04740	0.04847	0.04950	0.05047	0.05153	0.04957	0.09383
7	$\Theta_1$	0.00000	0.00000	0.00003	0.09993	0.09993	0.00003	0.09420
8	$\Theta_1$	0.04327	0.04500	0.04603	0.04707	0.04880	0.04610	0.09353
9	$\Theta_1$	0.04167	0.04300	0.04403	0.04507	0.04640	0.04410	0.09383
10	$\Theta_1$	0.04347	0.04507	0.04623	0.04733	0.04893	0.04630	0.09381
All	$\Theta_1$	0.04100	0.04440	0.04557	0.04827	0.05127	0.04630	0.09931

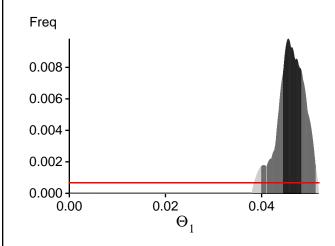
#### Citation suggestions:

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

Beerli P., 2007. Estimation of the population scaled mutation rate from microsatellite data, Genetics, 177:1967-1968.

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 over all loci



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

Use this value for Bayes factor calculations:

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

Locus	TI(1a)	BTI(1b)	SS(2)	HS(3)
1	-22014.49	-18794.42	-18270.41	-18404.24
2	-26344.73	-20484.57	-nan	-19578.64
3	-20480.97	-18445.16	-nan	-18256.42
4	-24213.67	-20433.95	-nan	-19954.58
5	-23598.84	-19384.81	-nan	-18769.42
6	-26896.63	-21937.30	-nan	-21227.48
7	-24662.58	-19586.34	-nan	-18823.77
8	-20400.24	-17727.10	-nan	-17397.84
9	-22590.83	-19108.63	-nan	-18639.50
10	-22546.66	-19045.16	-nan	-18568.44
All	-233736.85	-194934.64	-nan	-189607.54

- (1a) TI: Thermodynamic integration: log(Prob(D|Model)): Good approximation with many temperatures
- (1b) BTI: Bezier-approximated Thermodynamic integration: when using few temperatures USE THIS!
- (2) SS: Steppingstone Sampling (Xie et al 2011)
- (3) HS: Harmonic mean approximation: Overestimates the marginal likelihood, poor variance [Scaling factor = 12.792594]

Citation suggestions:

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

Palczewski M. and P. Beerli, 2014. Population model comparison using multi-locus datasets.

In M.-H. Chen, L. Kuo, and P. O. Lewis, editors, Bayesian Phylogenetics: Methods, Algorithms, and Applications, pages 187-200. CRC Press, 2014.

Xie W., P. O. Lewis, Y. Fan, L. Kuo, and M.-H. Chen. 2011. Improving marginal likelihood

estimation for Bayesian phylogenetic model selection. Systematic Biology, 60(2):150â 160, 2011.

# Acceptance ratios for all parameters and the genealogies

Parameter	Accepted changes	Ratio
$\Theta_1$ Genealogies	33118443/39994547 3553106/160005453	0.82807 0.02221

# MCMC-Autocorrelation and Effective MCMC Sample Size

Parameter	Autocorrelation	Effective Sampe Size
$\Theta_1$	0.11419	792397.57
Genealogies	0.92477	36568.13

# Average temperatures during the run

# Chain Temperatures 1 0.00000 2 0.00000 3 0.00000 4 0.00000

Adaptive heating often fails, if the average temperatures are very close together try to rerun using static heating! If you want to compare models using marginal likelihoods then you MUST use static heating

#### 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