simuPOP: a forward-time population genetics simulation environment

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For further information, please contact *bpeng@rice.edu*. More information about simuPOP can be found at *http://simupop.sourceforge.net*. A mailing list is devoted to the discussion of simuPOP, its email address is:*simupop-list@lists.sourceforge.net*. This work has been published in *bioinformatics* (Bo and Kimmel, 2005).

Introduction

simuPOP is a forward-time population genetics simulation environment. The core of simuPOP is a scripting language (Python) that provides a large number of objects and functions to manipulate populations, and a mechanism to evolve populations forward in time. Using this R/Splus-like environment, users can create, manipulate and evolve populations interactively, or write a script and run it as a batch file. Owing to its flexible and extensible design, simuPOP can simulate large and complex evolutionary processes with ease. At a more user-friendly level, simuPOP provides an increasing number of built-in scripts that perform simulations ranging from implementation of basic population genetics models to generating datasets under complex evolutionary scenarios.

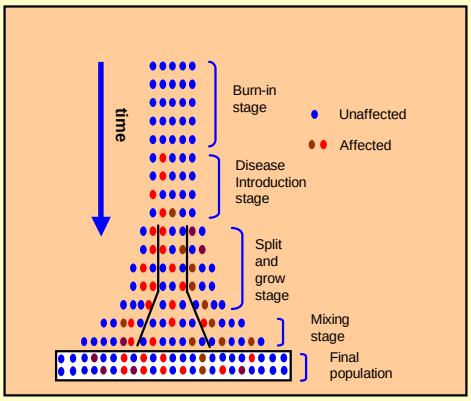
Why forward-time?

Backward-time (Coalescent, Kingman, 1982) based methods have dominated the area of genetic dataset generation because of their efficiency and flexibility. In contrast, forward-time simulations, although simpler as an idea, are computationally inefficient and have been used primarily for teaching purposes. Only recently, owing to the exponential growth of the power of personal computers, did the use of forward-time simulations begin in genetic studies.

	Backward-time (Coalescent)	Forward-time
Efficiency	**Yes**	No
Intuitiveness	No	**Yes**
Flexibility	Yes	**Excellent!**
Demographies	Yes	**Any**
Selection	No	**Any**
Recombination	Limited	**Any**
Migration	Yes	**Any**
Mutation	Yes	**Any**
Ancestral information	Discard	**Keep**
Trace the evolutionary process	No	**Yes**
Population information	No, sample based	**Yes**
Study population property	No	**Yes**
Resample from the resulting population	No	**Yes**
Test ascertainment methods	No	**Yes**
Application area	mostly around sample generation	**unlimited**
Avalability of programs	**Plenty**	EasyPOP (Balloux 2001) FPG (hey 2004) Now comes simuPOP!

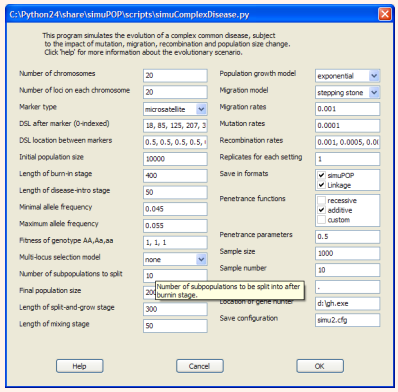
Features

- ✓ ***Scripting***. simuPOP is provided as a set of Python libraries, and is therefore backed by a full-blown object-oriented programming language.
- ✓ ***Flexibility***. simuPOP does not impose any limit on the size of genome, population, ploidy number, demographic model, mating type, etc. Using a large number of standard and hybrid (Python assisted) operators, plus the ability to extend simuPOP in Python, users can simulate almost arbitrarily complex evolutionary processes.
- ✓ ***Integration***. Owing to the 'glue language' nature of Python, it is easy to integrate simuPOP with other languages and programs.
- ✓ ***Comprehensiveness***. simuPOP consists of more than 70 operators and a lot more functions that cover all important aspects of genetic studies. These include mutation, migration, recombination, quantitative trait, selection, penetrance, ascertainment, statistics calculation, pedigree tracing, visualization and load/save in many popular formats.
- ✓ ***Efficiency***. simuPOP is written in *C++*. With extensive optimization, simuPOP can run very large simulations at reasonable speed.
- ✓ ***Openness.*** simuPOP is distributed free of charge under GPL license. The source code is available at simuPOP website.

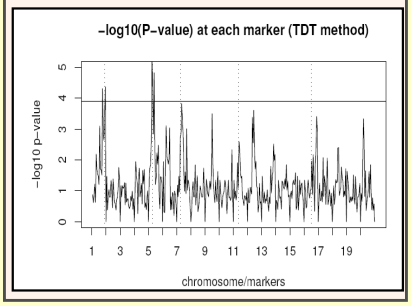


An example

The *simuComplexDisease.py* (above figure) script simulates the evolution of a complex disease under a four-stage evolutionary scenario. In a typical simulation, individuals have 20 chromosomes with 400 microsatellite markers and 5 binary disease susceptibility loci (DSL). The simulation starts with a small population and goes through four stages of evolution processes including burn-in, disease introduction, split and growth, and mixing. Recombination, mutation, selection, migration and demographic changes shape the genotype of the resulting large multi-generation populations in which samples can be drawn using different ascertainment methods.



Affected sibpair sample / TDT method



An interactive Python session

```
>>> from simuPOP import *
>>> from simuRPy import *
>>> simu = simulator(
...   population(size=1000, ploidy=2,
...     loci=[2]),
...   randomMating(),
...   rep=3 )
>>> simu.evolve(
...   preOps=[ initByValue([1,2,2,1])],
...   ops=[
...     recombinator( rate=0.1 ),
...     stat( LD=[0,1] ),
...     varPlotter("LD[0][1]",numRep=3,
...       ylim=[0,.25], xlab="generation",
...       ylab="D", title="LD Decay")
...     ],
...   end=100.
... )
```

The first two lines import simuPOP and simuRPy modules. The third command creates a **simulator** with three replicates of a diploid **population**. Random **mating** will be used to generate offspring. The last command uses the evolve function to evolve the populations for 100 generations, subject to four **operators**.

The first operator `initByValue` is applied to all populations before evolution. It initializes all individuals with the same genotype 12/21. The other three operators will be applied at every generation. `recombinator` will recombine parental chromosomes with the given recombination rate during the generation of offspring; `stat` will calculate standard linkage disequilibrium between the first and second loci. The result of this operator will be stored in a **local variable space** of each population and be retrieved and plotted by `varPlotter`, which uses R for plotting. When evolve is called, a graphics window will be fired and will display the dynamics of LD values for all three replicates.

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