

model

Complete
Genomes

Core
Genomes

Graph without
edit edges

Graph
model



possible
genomes

select core
genomes



core
genomes

sample a core
genome



reference
defining
distance
metric

edit the genome

chosen genome



choose start



SE: spell L letters

PE: spell 2L+gap letters



sequence (only for pair-end reads)

chop two reads

length L from both sides

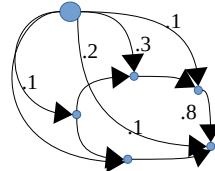


read w/ mutations

sequence



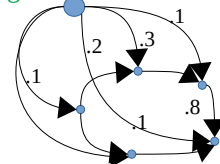
read w/ phred



reads from different
genomes

build a pVG

add edit edges



pVG

sample a
sequence

sample an
edited
sequence

edit the
sequence

**generative
process**

read → genome,
position,
seq.errors

read →
core genome,
position,
mutations,
seq.errors

read →
path (alignment),
mutations, seq.errors

read → path
(mutations, alignment),
seq.errors

**inference
task**

* color code:
deterministic
probabilistic
nondeterministic