Computational Evolution & Digital Organisms

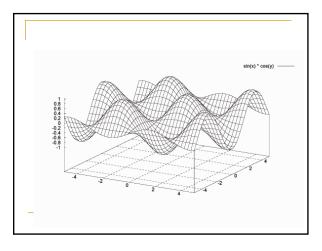
A look at a subset of Artificial Life By Daniel Weise

Computational Evolution

- Attempts to elucidate principles of evolution
 - Builds models of self-replicating organisms
 - Computational cost limits physical fidelity of the model.Digital or chemical models
 - Mutation creates variation in populations
 - Reproduction can be sexual or asexual
 - Ability to (out) reproduce its genome is the usual fitness measure
 - For some research, other fitness measures are used.

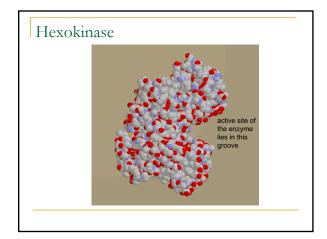
Not to be Confused With Evolutionary Computing

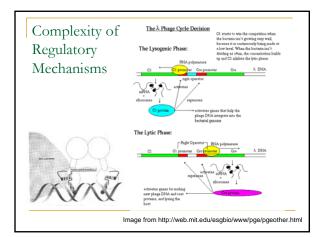
- A Search Technique inspired by biology
 - Points in search space represented as "genomes"
 - $\hfill\square$ Crossover produces new points in search space
 - Mutation ensures variety
 - Ensures more of search space is sampled
 - Fitness function determines which subset of population become progenitors
 - □ Larger populations increase coverage of space.
 - Search usually walks through "invalid" points

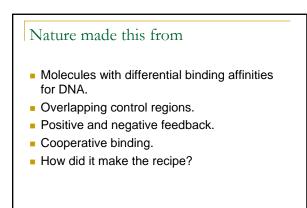


Overview of Talk

- Motivation: The complexity of cellular life
- Tierra and the evolution of digital organisms
- Avida and other Tierra inspired work
- Lessons/Future Research







Tierra, a Platform for Digital Evolution

- Design Requirements/Inventions:
 - Organisms must be self-reproductive
 - Ability to out-reproduce the competition only fitness criteria
 - Avoids "artificial" fitness functions.
 - Control (jumps/calls) is effected through *templates* and *targets*, which are complementary "bit strings"
 - Jump nop1 nop0 nop1 goes to nop0 nop1 nop0
 - Organisms sense the environment
 Dynamic "fitness" function

Tierra's Digital Organisms

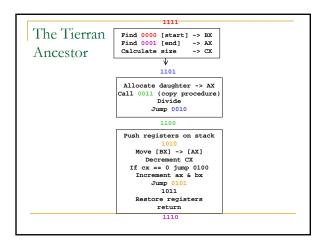
- Each organism (cpu) has
 - □ 4 registers (A, B, C, D)
 - Instruction pointer
 - 10 word stack
- Time slicing "implements" parallel organisms
- When space for new organisms is needed, the oldest organisms are reaped (as a rule).

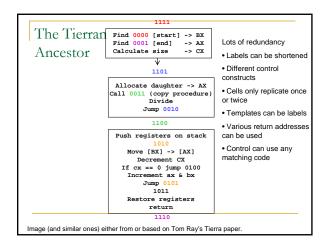
Tierra's Instruction Set

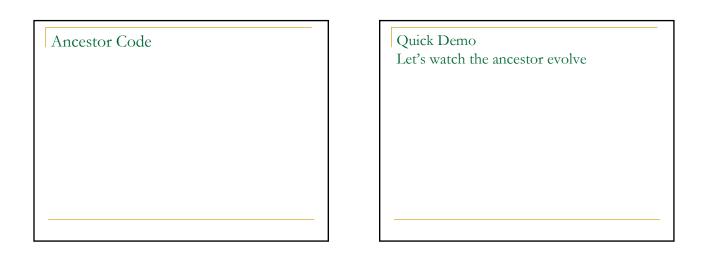
- Data Movement
 - PushA, PopA, PushB, PopB, etc for C and D
 - MOVDC (D <- C), MOVBA, COPY ([A] to [B])</p>
- Control
 - □ JumpO, JumpB, Call, Ret, IfZ, nop0, nop1
- Calculation
 - □ subcab, subaac, inca, incb, decc, incd, zero, not, shl
- Biological and Sensing
 adr, adrb, adrF, mal (allocate memory), divide

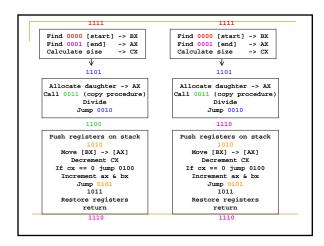
Mutational Sources

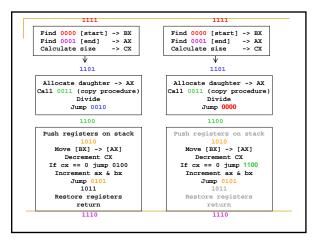
- A copy error every X copy instructions
- Cosmic rays
 - □ A bit in the soup gets flipped every Y instructions
 - Works because no cells are autosomes
 - Biased, not random
- Probabilistic results of instructions
- Every so often an instruction misfires
- E.g., incA adds 2
- No Insertion/deletions

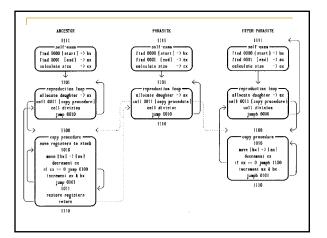


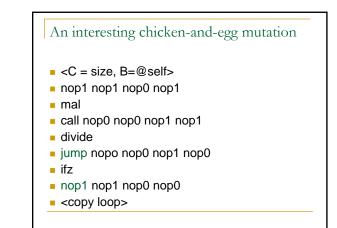










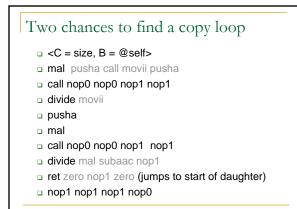


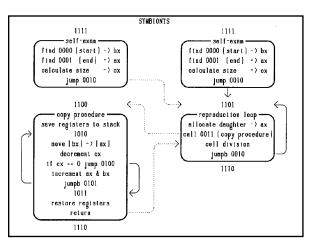
An interesting chicken-and-egg mutation

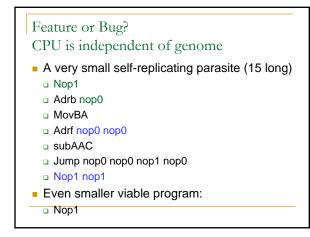
- <C = size, B=@self>
- nop1 nop1 nop0 nop1
- mal
- call nop0 nop0 nop1 nop1
- divide
- pushb (was jump) nopo nop0 nop1 nop0
- ifz
- ret (was nop1) nop1 nop0 nop0
- <copy loop>

A Copy-Once Parasite

- Stays just ahead of the reaper
- nop1 nop1 zero not0 shl shl movdc
- adrb nop0 nop0 pushc nop0
- □ subaac
- movba pushd nop0
- adr nop0 nop1
- inca
- subcab pusha nop1 pushd nop1
- mal
- a call nop0 nop0 nop1 nop0
- divide







Feature or Bug? Non-local effects

- A template can match any nearby target
- A request for memory can kill *any* organism, even one "fitter"
- A daughter cell can be placed anywhere
- Allocating a large amount of memory for a daughter can kill tens of organisms, creating a dieoff

Feature or Bug?

Spaghetti Code is a Frequent Occurrence

- Symbionts arise quite frequently
- When a target is mutated, the target in another cell is used.

Bug or Feature? Parasites require necrophilia

- Instructions are left in memory when an organism is reaped.
- "Parasites" keep using these instructions.

Bug or Feature? Sloppy replicators instead of Indels

- Tierra lacks insertion/deletion mutations
 - Biology uses indels
 - $\hfill\square$ Harder to remove instructions without deletions
 - □ Harder to make room for new instructions
- Tierra makes up for it with sloppy replicators that move instructions around willy nilly
 - Buy maybe this is needed anyway?

Is Sloppiness needed to Bootstrap Complexity?

- Sloppiness (ad-hoc) mixing gave us
 - Mitochondia (ingestion without digestion)
 - Cloroplasts in bacteria (same story)
 - Gene mixing (via viruses)
 - Diploidy from Haploidy

Avida

- Inspired by Tierra, but
 - Controlled instruction pointers (less slopiness)
 - Insertion/Deletion mutations
 - 2 dimensional grid of organisms, not instructions
 - Only local next-neighbor effects
 - Fitness functions to augment reproduction
- Experiments to test biological theories
 - Evolution of Complexity
 - Evolution of Complex Functions
 - Relationship among evolution rate and landscape

Lots of questions raised by Avida paper we read.

- What happens when treated as search problem without using populations?
 - a How does the system walk through deleterious steps in the search space?
- What insights are gained by treating the reduced trace of a program as its phenotype?
- Does this remove epistatic measurement effects?
- What about sexual reproduction?
- What is the density of paths thru mutation space?
- Would a more Tierra-like system be better?
- What sized rewards would work?

Digital Biosphere

- Inspired by Tierra/Avida but
 - Want to design open-ended evolutionary frameworks
 - □ Focus is on evolutionary trajectories.
 - Are there principles regarding these trajectories?
 - Will exploit the constraints of physics
 - Conservation Laws!
 - Energy requirements and metabolism
 - Will eventually move to chemical modeling to get closer to biology.

Lessons

- Evolution finds corners of the search space
 If you build it, they will exploit it
 - Complexity comes from exploiting environment
- Co-evolution makes the problem interesting and different
 - Changing fitness functions
- Designing a system for open-ended evolution is still very much an open-ended problem.

What's it all mean? We have a source of new insights

- Watching evolving dynamical systems give insight and ideas.
- Biologists aren't trained to do this.
- Many insights will be gained that will eventually transfer over to biological thinking

More information

- Me: <u>http://cs.washington.edu/homes/weise</u>
- Reading course: <u>http://cs.washington.edu/homes/weise/590ce.</u> <u>html</u>
- Course will have a project based 3 credit option.

Open Questions/Future Research

- Investigate the "Worm-hole" hypothesis: no interesting genomes arise solely from singlestep changes to existing genomes.
- Define phenotype as an organisms birthing trace. Now re-explain all subsequent papers in this light.
- How do we get true diversity s.t. environment changes kill half of everything?
- How do we automatically detect novelty?

On "designing" open-ended evolutionary systems.

- In the days when Sussman was a novice, Minsky once came to him as he sat hacking at the PDP-6.
- "What are you doing?", asked Minsky.
- "I am training a randomly wired neural net to play Tic-tac-toe" Sussman replied.
- "Why is the net wired randomly?", asked Minsky.
- "I do not want it to have any preconceptions of how to play", Sussman said.
- Minsky then shut his eyes.
- "Why do you close your eyes?", Sussman asked his teacher. "So that the room will be empty."
- At that moment, Sussman was enlightened.

The Value of Diploidy?

- Most of the genes perform a walk from viable organism to viable organism.
- Some of the genes walk through non-viable points in the search space.