10. Genetic Algorithms

- General-purpose “black-box” optimisation method proposed by J. Holland (1975) and K. DeJong (1975).
- Method has attracted lots of interest, but theory is still incomplete and the empirical results inconclusive.
- Advantages: general-purpose, parallelisable, adapts incrementally to changing cost functions (“on-line optimisation”).
- Disadvantages: typically very slow – should be used with moderation for simple serial optimisation of a stable, easily evaluated cost function.
- Some claim that GA’s typically require fewer function evaluations to reach comparable results as e.g. simulated annealing. Thus the method may be good when function evaluations are expensive (e.g. require some actual physical measurement).

10.1 The Basic Algorithm

- We consider the so called “simple genetic algorithm”; also many other variations exist.
- Assume we wish to maximise a cost function $c$ defined on $n$-bit binary strings:

$$c : \{0, 1\}^n \rightarrow \mathbb{R}.$$ 

Other types of domains must be encoded into binary strings, which is a nontrivial problem. (Examples later.)
- View each of the candidate solutions $s \in \{0, 1\}^n$ as an individual or chromosome.
- At each stage (generation) $t$ the algorithm maintains a population of individuals $p_t = (s_1, \ldots, s_m)$.

Three operations defined on populations:

- selection $\sigma(p)$ (“survival of the fittest”)
- recombination $\rho(p)$ (“mating”, “crossover”)
- mutation $\mu(p)$

The Simple Genetic Algorithm:

```
function SGA($\sigma, \rho, \mu$):
  $p \leftarrow$ random initial population;
  while $p$ “not converged” do
    $p' \leftarrow \sigma(p)$;
    $p'' \leftarrow \rho(p')$;
    $p \leftarrow \mu(p'')$
  end while;
  return $p$ (or “fittest individual” in $p$).
end.
```

Selection (1/2)

- Denote $\Omega = \{0, 1\}^n$. The selection operator $\sigma : \Omega^m \rightarrow \Omega^m$ maps populations probabilistically: given an individual $s \in p$, the expected number of copies of $s$ in $\sigma(p)$ is proportional to the fitness of $s$ in $p$. This is a function of the cost of $s$ compared to the costs of other $s' \in p$.
- Some possible fitness functions:
  - Relative cost (⇒ “canonical GA”):
    $$f(s) = \frac{c(s)}{\frac{1}{m} \sum_{s' \in p} c(s')} \triangleq \frac{c(s)}{\bar{c}}.$$
Relative rank:

$$f(s) = \frac{r(s)}{m} = \frac{2}{m+1} \cdot r(s),$$

where $r(s)$ is the rank of individual $s$ in a worst-to-best ordering of all $s' \in p$.

## Selection (2/2)

Once the fitness of individuals has been evaluated, selection can be performed in different ways:

- **Roulette-wheel selection** ("stochastic sampling with replacement"):
  - Assign to each individual $s \in p$ a probability to be selected in proportion to its fitness value $f(s)$. Select $m$ individuals according to this distribution.
  - Pictorially: Divide a roulette wheel into $m$ sectors of width proportional to $f(s_1), \ldots, f(s_m)$. Spin the wheel $m$ times.

- **Remainder stochastic sampling**:
  - For each $s \in p$, select deterministically as many copies of $s$ as indicated by the integer part of $f(s)$. After this, perform stochastic sampling on the fractional parts of the $f(s)$.
  - Pictorially: Divide a fixed disk into $m$ sectors of width proportional to $f(s_1), \ldots, f(s_m)$. Place an outer wheel around the disk, with $m$ equally-spaced pointers. Spin the outer wheel once.

## Recombination (1/2)

- Given a population $p$, choose two random individuals $s, s' \in p$. With probability $p_p$, apply a crossover operator $\rho(s, s')$ to produce two new offspring individuals $t, t'$ that replace $s, s'$ in the population.
- Repeat the crossover throughout the population. Denote the total effect on the population as $\rho' = \rho(p)$.
- Practical implementation: choose $\frac{p_p}{2} \cdot m$ random pairs from $p$ and apply crossover deterministically.
- Typically $p_p \approx 0.7 \ldots 0.9$.

## Recombination (2/2)

Possible crossover operators:

- **1-point crossover**:

- **2-point crossover**:

- **uniform crossover**:
### Mutation

- Given population $p$, consider each bit of each individual and flip it with some small probability $p_\mu$. Denote the total effect on the population as $p' = \mu(p)$.
- Typically, $p_\mu \approx 0.001 \ldots 0.01$. Apparently good choice: $p_\mu = 1/n$ for $n$-bit strings.
- Theoretically mutation is disruptive. Recombination and selection should take care of optimisation; mutation is needed only to (re)introduce "lost alleles", alternative values for bits that have the same value in all current individuals.
- In practice mutation + selection = local search. Mutation, even with quite high values of $p_\mu$, can be efficient and is often more important than recombination.

### Hyperplane sampling (2/4)

Define:

- **order** of hyperplane $H$:
  \[ o(H) = \text{number of fixed bits in } H = n - \text{dim } H \]
- **average cost** of hyperplane $H$:
  \[ c(H) = \frac{1}{2^{n-o(H)}} \sum_{s \in H} c(s) \]
- $m(H, p) = \text{number of individuals in population } p \text{ that sample hyperplane } H.$

Heuristic claim: selection drives the search towards hyperplanes of higher average cost (quality).
Hyperplane sampling (3/4)

Consider e.g. the following cost function and partition of $\Omega$ into hyperplanes (in this case, intervals) of order 3:

Here the current population of 21 individuals samples the hyperplanes so that e.g. '000\*\*' and '010\*\*' are sampled by three individual each, and '100\*\*' and '101\*\*' by two individual each. Hyperplane '010\*\*' has a rather low average fitness in this population, whereas '111\*\*' has a rather high average fitness.

Hyperplane sampling (3/4)

Then the result of e.g. roulette wheel selection on this population might lead to elimination of some individuals and duplication of others:

Then, in terms of expected values, one can show that

$$m(H, \sigma(p)) = m(H, p) \cdot f(H, p).$$
The effect of crossover on schemata (2/2)

Generally, the probability that in 1-point crossover a schema \( H = \{0, 1, \ast\}^n \) is retained, is (ignoring the possibility of “lucky combinations”)

\[
\Pr(\text{retain } H) \approx 1 - \frac{\Delta(H)}{n-1},
\]

where \( \Delta(H) \) is the defining length of \( H \), i.e. the distance between the first and last fixed bit in \( H \).

More precisely, if \( H \) has \( m(H,p) \) representatives in population \( p \) of total size \( m \):

\[
\Pr(\text{retain } H) \geq 1 - \frac{\Delta(H)}{n-1} \left( 1 - \frac{m(H,p)}{m} \right).
\]

The Schema “Theorem” (1/2)

Heuristic estimate of the changes in representation of a given schema \( H \) from one generation to the next. Proposed by J. Holland (1975).

Denote:

\[
m(H,t) = \text{number of individuals in population at generation } t \text{ that sample } H.
\]

Then:

(i) Effect of selection:

\[
m(H,t') \approx m(H,t) \cdot f(H)
\]

(ii) Effect of recombination:

\[
m(H,t'') \approx (1 - p_p)m(H,t') + p_p \left( m(H,t') \Pr(\text{retain } H) + m \cdot \Pr(\text{luck}) \right)
\]

\[
\geq (1 - p_p)m(H,t') + p_p m(H,t') \left( 1 - \frac{\Delta(H)}{n-1} \left( 1 - \frac{m(H,t')}{m} \right) \right)
\]

\[
= m(H,t') \left( 1 - p_p \frac{\Delta(H)}{n-1} \left( 1 - \frac{m(H,t')}{m} \right) \right)
\]

(iii) Effect of mutation:

\[
m(H,t+1) \approx m(H,t'') \cdot (1 - p_p)^{\alpha(H)}
\]
The Schema “Theorem” (2/2)

In summary, then:

\[ m(H, t + 1) \geq m(H, t) \cdot f(H) \cdot \left( 1 - p_0 \frac{\Delta(H)}{n - 1} \left( 1 - \frac{m(H, t)}{m} \right) \right) \cdot \left( 1 - p_0 \right)^{m(H)} \]

The formula leads to so called “Building Block Hypothesis”:

- In a genetic search, short, above-average fitness schemata of low order (“building blocks”) receive an exponentially increasing representation in the population.

- Proper treatment: analyse the genetic search as a stochastic process (Markov chain). This is unfortunately very difficult.

Criticisms

- Many of the approximations used in deriving the “Schema Theorem” implicitly assume that the population is very large. In particular, it is assumed that all the relevant schemata are well sampled. This is clearly not possible in practice, because there are \( 3^n \) possible schemata of length \( n \).

- The “Schema Theorem” cannot be used to predict the development of the population for much more than one generation, because:
  1. the long-term development depends on the coevolution of the schemata, and the “theorem” considers only one schema in isolation;
  2. an “exponential growth” cannot continue for long in a finite population.

10.3 Applications of Genetic Algorithms

General comments on coding:

- If the function to be optimised is not naturally defined on binary strings, then the domain must be coded. This is a nontrivial task for GA’s, because the representation influences the computation.

- Real numbers can be block-coded into sequences of integers.

- For integers, the Gray code should be considered as an alternative to the standard binary representation. In the Gray code the binary representation of integer \( k + 1 \) differs from that of integer \( k \) in only one bit. Thus, mutating a Gray coded integer by one bit can only change its value by \( \pm 1 \).
Gray code conversion

<table>
<thead>
<tr>
<th>integer</th>
<th>standard</th>
<th>Gray</th>
</tr>
</thead>
<tbody>
<tr>
<td>(k)</td>
<td>(a₁a₂a₃)</td>
<td>(b₁b₂b₃)</td>
</tr>
<tr>
<td>0</td>
<td>000</td>
<td>000</td>
</tr>
<tr>
<td>1</td>
<td>001</td>
<td>001</td>
</tr>
<tr>
<td>2</td>
<td>010</td>
<td>011</td>
</tr>
<tr>
<td>3</td>
<td>011</td>
<td>010</td>
</tr>
<tr>
<td>4</td>
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<td>110</td>
</tr>
<tr>
<td>5</td>
<td>101</td>
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<td>110</td>
<td>101</td>
</tr>
<tr>
<td>7</td>
<td>111</td>
<td>100</td>
</tr>
</tbody>
</table>

- standard → Gray conversion: \( b_i = \begin{cases} a_i & i = 1, \\ a_{i-1} \oplus a_i & i > 1 \end{cases} \)
- Gray → standard conversion: \( a_i = \bigoplus_{j=1}^{i} b_j \)

Other coding issues
- Cycles/permutations
- Trees
- Graphs
- ...