



# Evolution and Genetic Algorithms

# Exponential growth and *per capita* growth rate

- ▶  $R$  : population size (number of individuals, or, rather, *density*, i.e. individuals per area or volume unit)
- ▶  $b$  : *per capita* birth rate ("how many offspring each individual gets, per time unit")
- ▶  $d$  : per capita death rate ("how many times each individual dies, per time unit")
- ▶ Total births per time unit:  $bR$   
Total deaths per time unit:  $dR$
- ▶ Population growth rate:  $\frac{dR}{dt} = bR - dR = (b - d)R = rR$
- ▶  $r$  : intrinsic growth rate
- ▶ The *per capita growth rate*, defined as  $\frac{dR}{dt} / R$  is in this model equal to  $\frac{dR}{dt} / R = rR / R = r$ .
- ▶ The *per capita* growth rate is the growth rate per individual. In this model it is given by the model parameter  $r$ .
- ▶ The equation  $\frac{dR}{dt} = rR$  describes *exponential growth*.

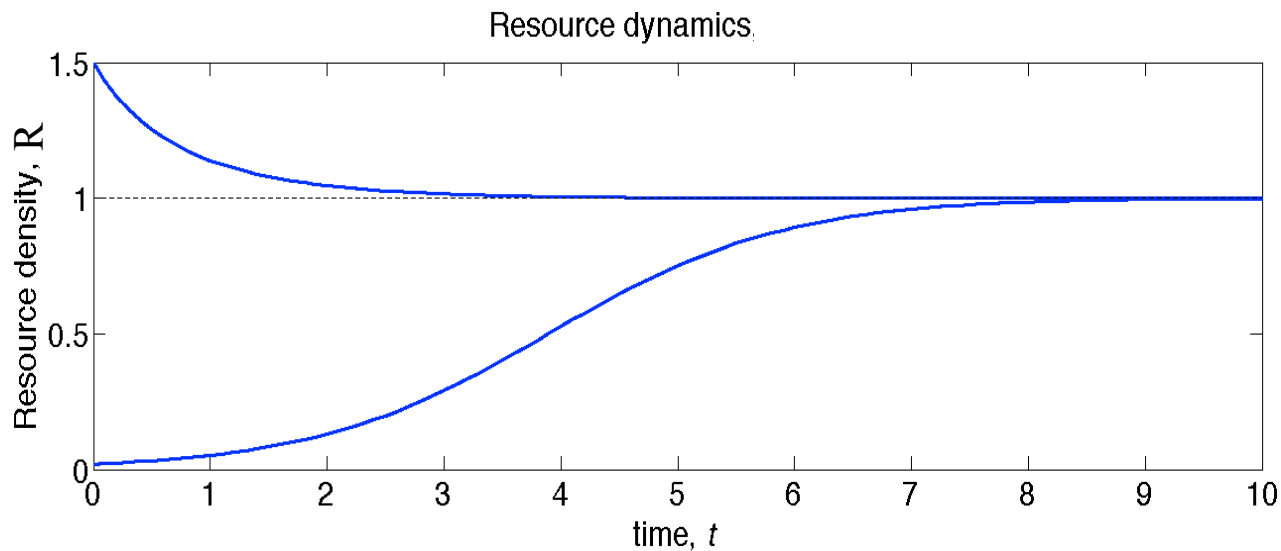
# Adding prey density dependence

Resource dynamics:

$$\frac{dR}{dt} = rR \left( \frac{K - R}{K} \right)$$

density dependence

$R$  = resource density  
 $r$  = intrinsic growth rate  
 $K$  = carrying capacity



# Modelling predation

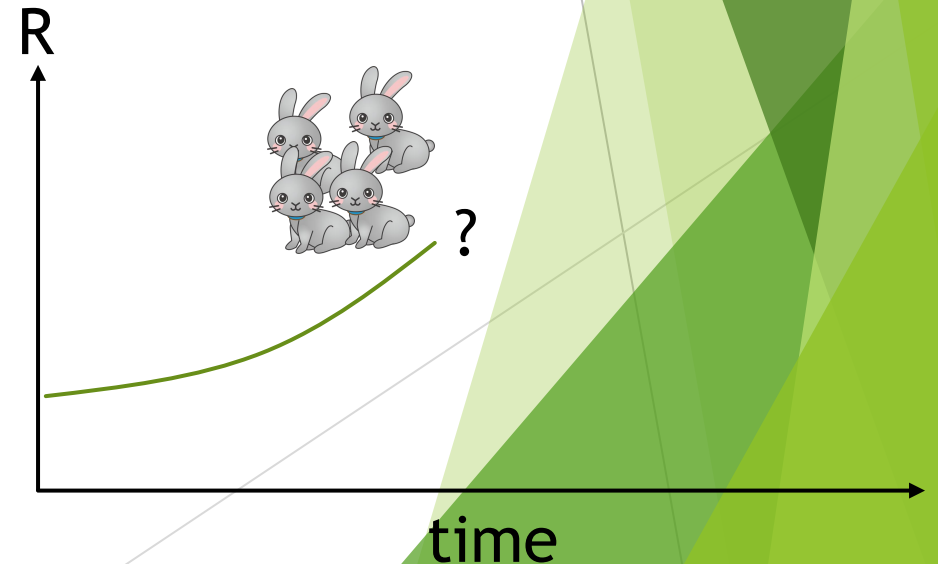
## Prey population growth

Prey population growth = prey births - prey deaths

Prey population growth = prey birth - prey background mortality - predation

$$\frac{dR}{dt} = rR \left( \frac{K-R}{K} \right) - ?$$

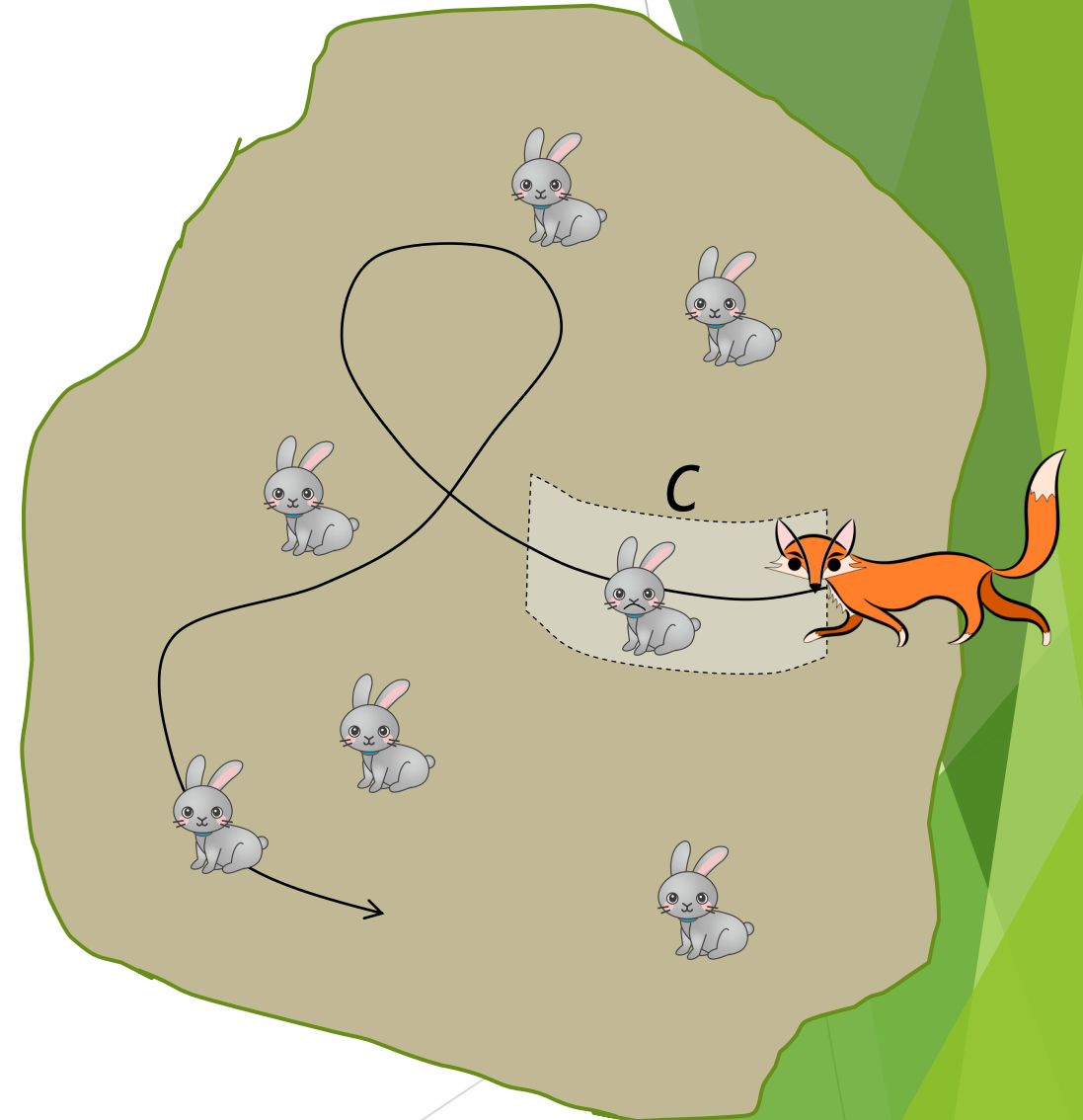
$r$  = prey intrinsic growth rate ( $b-d$ )



How many prey does a single predator catch, per time unit?

- Prey density, prey per area unit:  $R$
- Search area per predator, per time unit:  $a$  (predator *attack rate*)
- Prey caught per time unit, per predator:  $aR$
- Predator density:  $C$
- Total prey caught:  $aRC$
- Total prey growth:

$$\frac{dR}{dt} = rR \left( \frac{K-R}{K} \right) - aRC$$



# Modelling predation

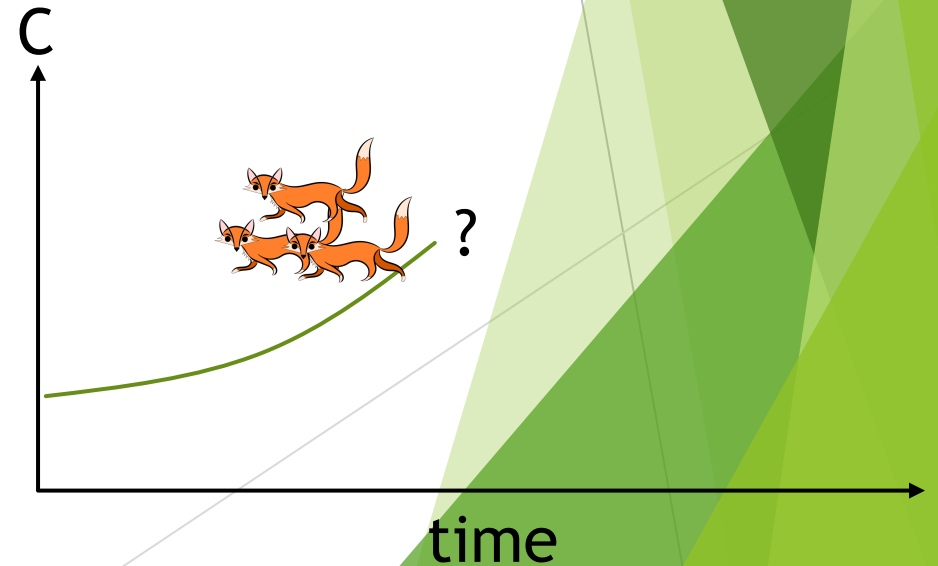
## Predator population growth

Predator population growth =  
predator growth due to predation - background predator mortality

$$\frac{dC}{dt} = \varepsilon aRC - \mu C$$

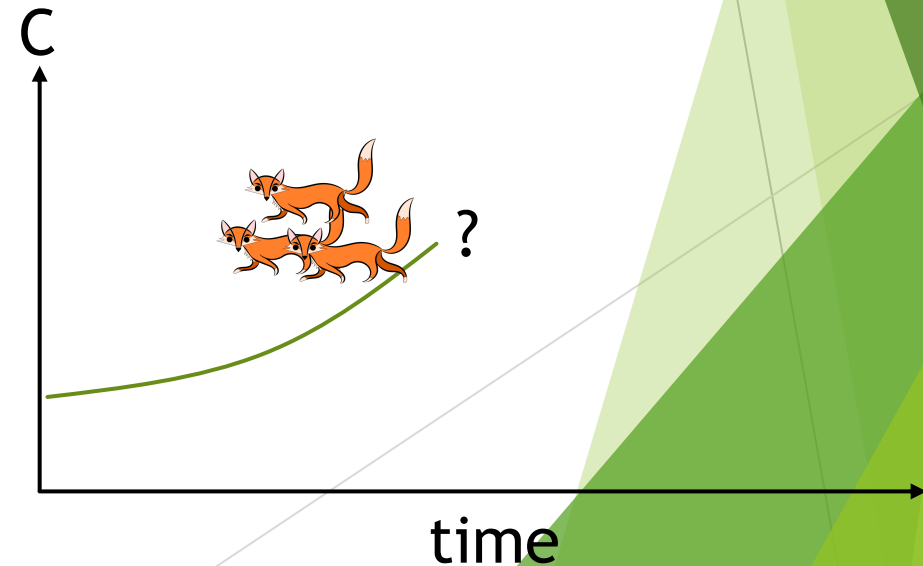
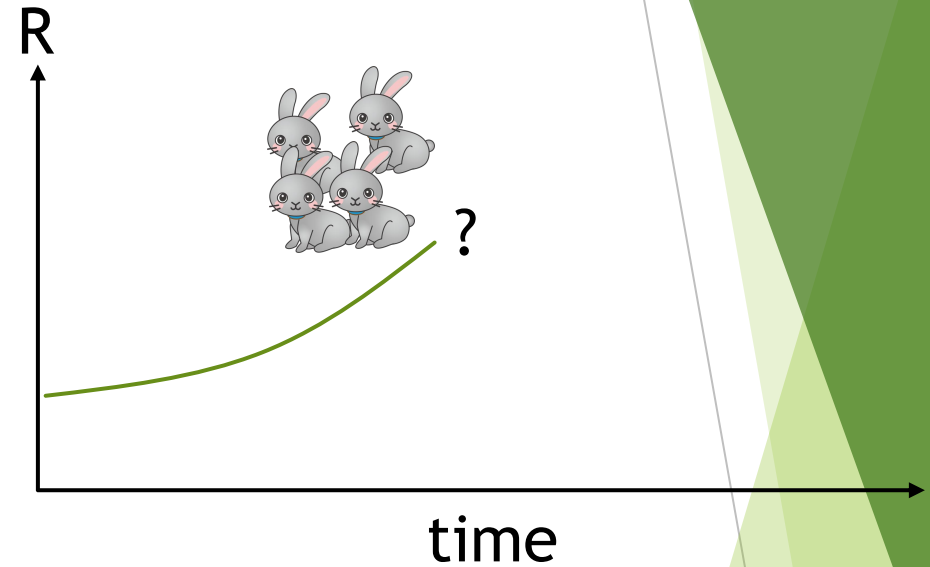
$\varepsilon$  = conversion factor from prey to predators

$\mu$  = predator mortality, *per capita*



# The complete population dynamics

$$\begin{cases} \frac{dR}{dt} = rR \left( \frac{K - R}{K} \right) - aRC \\ \frac{dC}{dt} = \varepsilon aRC - \mu C \end{cases}$$



# The complete population dynamics

$R$  : Resource abundance (e.g. kg/ha)

$C$  : Consumer abundance (e.g. kg/ ha)

$a$  : Consumer attack rate (e.g. ha/year)

$\epsilon$  : conversion coefficient (e.g. kg<sup>-1</sup>)

$\mu$  : background mortality (e.g. year<sup>-1</sup>)

Fitness = per capita growth rate (why?)

Resource population growth:  $\frac{dR}{dt} = fR = rR(1 - R) - aRC$

Consumer population growth:  $\frac{dC}{dt} = fC = (\epsilon aR - \mu)C$

Resource fitness:  $f_R = r(1 - R) - aC$

Consumer fitness:  $f_C = \epsilon aR - \mu$

Resources consumed per time unit, per consumer.

What if the attack rate  $a$  evolves (the consumers become more efficient)?



# The complete population dynamics

Resource dynamics:

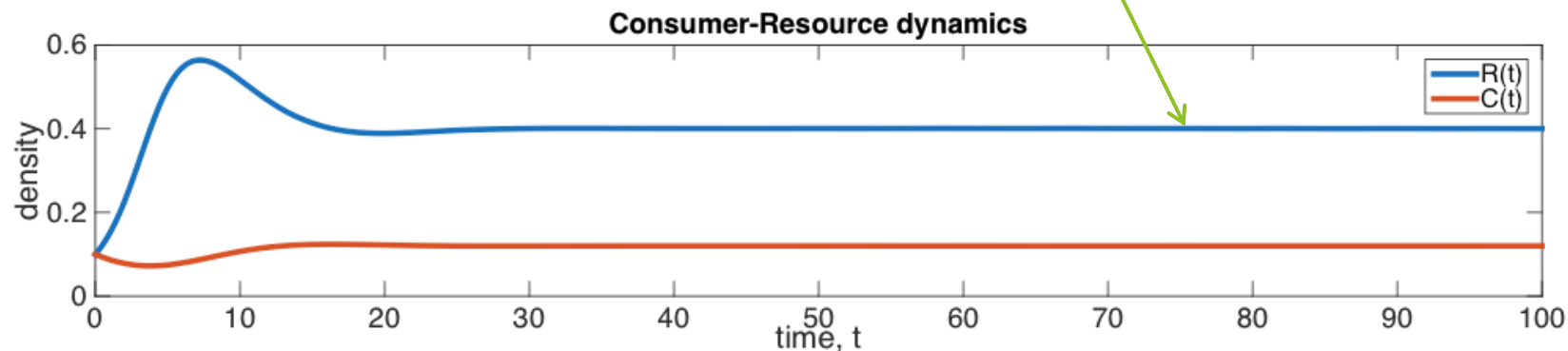
$$\frac{dR}{dt} = rR \left( \frac{K - R}{K} \right) - aRC$$

Consumer dynamics:

$$\frac{dC}{dt} = \varepsilon aRC - \mu C$$

Consumer per capita growth (fitness!):

$$\frac{dC}{dt} \frac{1}{C} = \varepsilon aR - \mu \quad \square \quad R^* = \frac{\mu}{\varepsilon a}$$



# The complete population dynamics

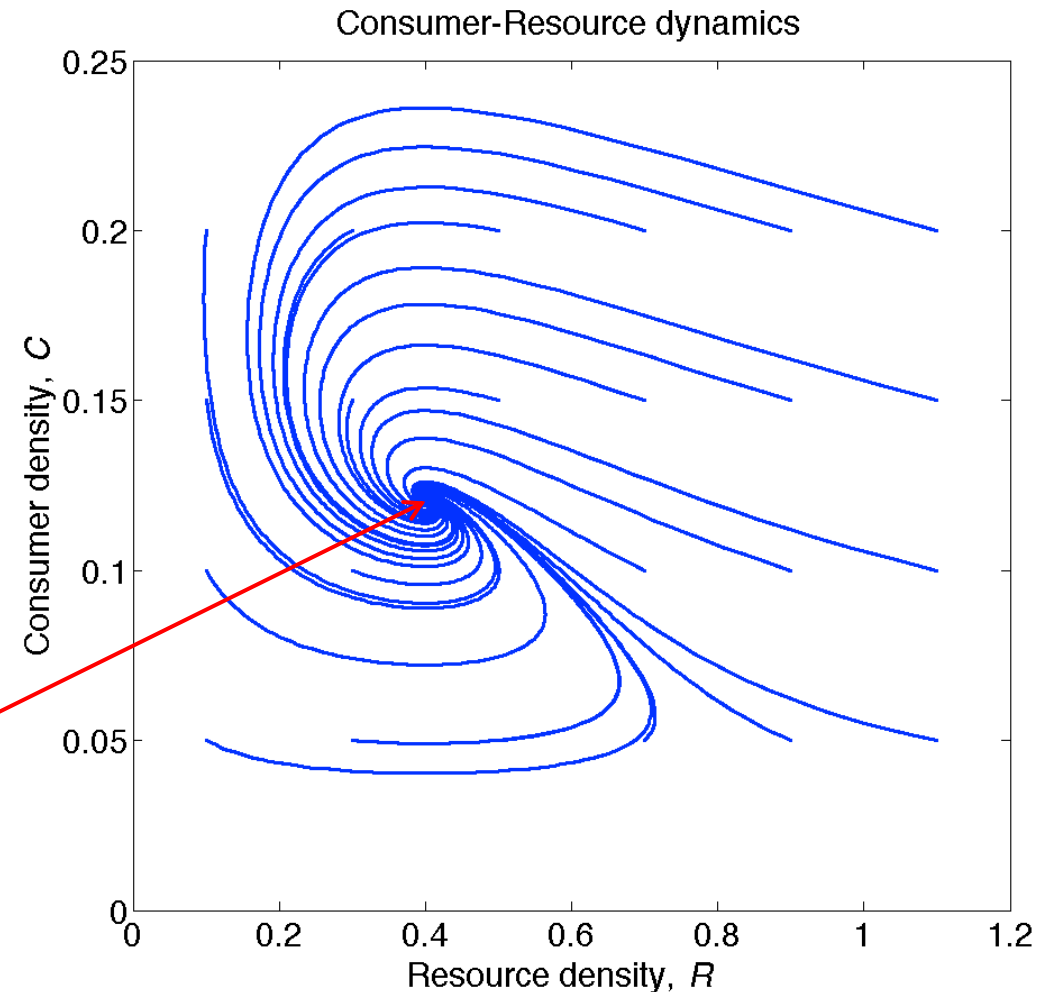
A dynamically stable, fixed point equilibrium

$$\begin{cases} \frac{dR}{dt} = rR \left( \frac{K - R}{K} \right) - aRC \\ \frac{dC}{dt} = \varepsilon aRC - \mu C \end{cases}$$

Substitute  $C^*$  into the resource equation and solve

Scaling the system by setting  $K=1$  (commonly done to simplify the system) leads to

$$(R^*, C^*) = \begin{bmatrix} \frac{\mu}{\varepsilon a} \\ \frac{r}{a} \left( 1 - \frac{\mu}{\varepsilon a} \right) \end{bmatrix}$$



What if the  
attack rate  $a$   
evolves (the  
consumers  
become more  
efficient)?

Consumer fitness:  $f_c = \epsilon a R - \mu$

**First**, sort out the ecological dynamics (we already did).

**Second**, we need a theoretical evolutionary foundation (e.g. Game Theory)

**Third**, we need a evolutionary theory and eco-evolutionary model analyses (e.g. Adaptive Dynamics)

# Evolving attack rate

Assume the consumer attack rate  $a$  depends on some heritable trait  $x$ .

In other words, assume there can be several types of consumers present in the population, each with its own trait value  $x$  and attack rate  $a(x)$ :

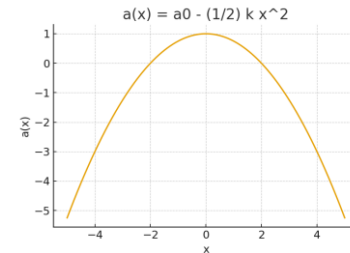
A consumer with trait  $x$  thus has *per capita growth rate (fitness)*

Trait-based fitness

$$f(x) = \epsilon a(x)R - \mu$$

Trait-based attack rate

$$a(x) = a_0 - \frac{1}{2}kx^2$$



The trait-based resource dynamics becomes:

$$\frac{dR}{dt} = rR(1 - R) - \sum_{i=1}^C a(x_i)R \quad C = \text{the number of consumers}$$

# Adaptive Dynamics assumptions

Assume mutations are rare, such that:

1. There is most of the time only one type (the resident) present in the population
2. An ecological equilibrium is established before any new mutant occurs

If a resident type  $x$  has established an ecological equilibrium

$$(R^*, C^*) = \left( \frac{\mu}{\varepsilon a(x)}, \frac{r}{a(x)} \left( 1 - \frac{\mu}{\varepsilon a(x)} \right) \right),$$

what is then fitness of a rare invading mutant type  $x'$ ?

$$f(x', x) = \varepsilon a(x') R^*(x) - \mu = \mu \frac{a(x')}{a(x)} - 1$$

This is the *invasion fitness*.

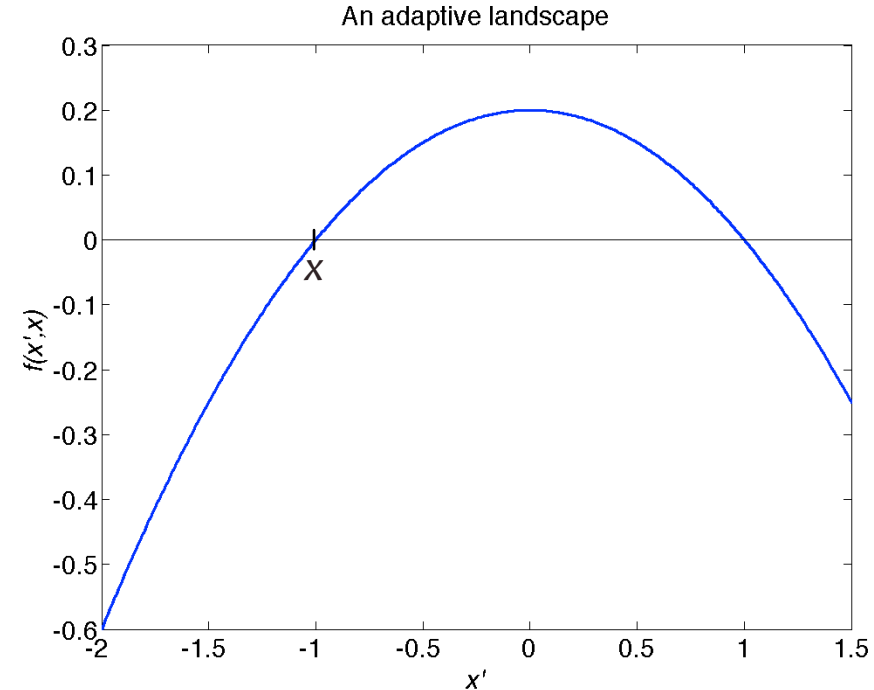
# Invasion fitness

General definition:

$f(x', x)$  = per capita growth rate of rare type  $x'$  in the equilibrium environment set by resident type  $x$ .

By assumption:  $f(x, x) = 0$

Keeping resident type  $x$  fixed, but varying the mutant type  $x'$  generates an *adaptive landscape*.



# Trait substitution sequence

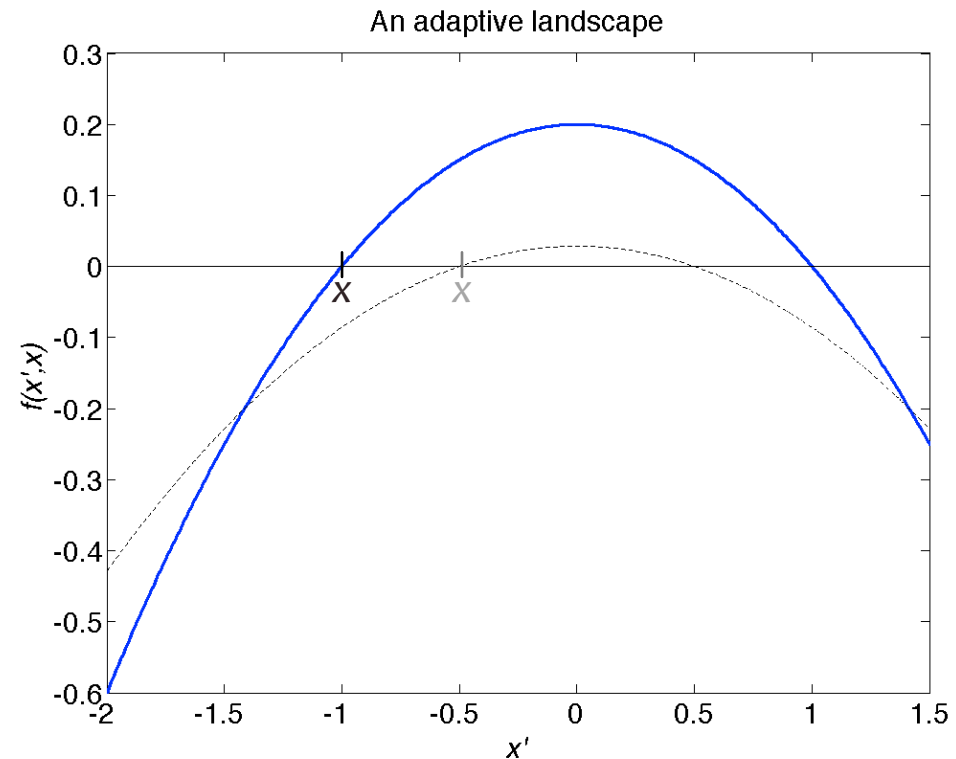
A mutant with positive invasion fitness can invade and replace the resident type, establishing a new equilibrium.

Evolution will in this way proceed in a ‘trait substitution sequence’.

The adaptive landscape shows which mutants can invade, but the landscape will change when the resident trait changes.

Always,  $f(x, x) = 0$ .

Evolution is in this way  
‘climbing a sinking landscape’.



# Genotype-Phenotype Mapping in Adaptive Models



# Overview

- ▶ Multilocus diploid genotypes dictate traits
- ▶ Multiple additive alleles effects traits
- ▶ Traits of particular interest: ecological and mating traits
- ▶ Adaptation emerges from genetic variation, GP mapping and selection

# Example: Genotype Architecture in evolutionary models

- ▶ Each evolving trait is encoded by:
  - ▶ Multiple diploid loci (20-60 per trait)
  - ▶ Diallelic alleles, each with some numeric value
  - ▶ Additive allele effects (sum across loci)
  - ▶ After allele summation, traits are rescaled into continuous trait values
  - ▶ Recombination leads to new variants on which selection can act
  - ▶ Mutations lead to new variations on which selection can act

# How Phenotypes Determine Fitness & Mating

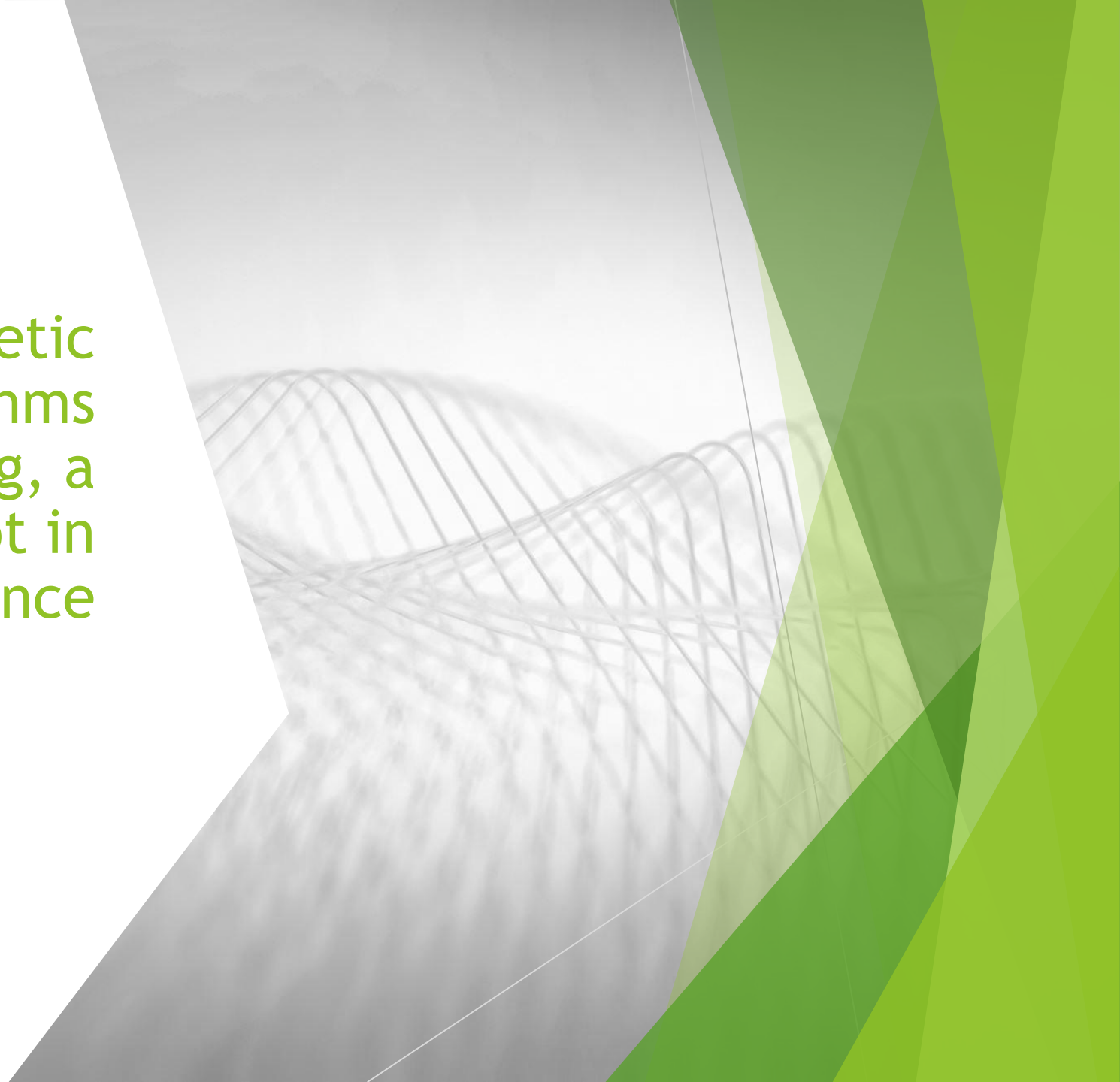
- ▶ Phenotype values affect:
  - ▶ Resource uptake
  - ▶ Habitat survival
  - ▶ Assortative mating via trait similarity
  - ▶ Sexual selection

# How Adaptation Emerges

- ▶ Adaptive dynamics in the model arise from:
  - ▶ Ecological selection traits
  - ▶ Sexual & assortative mating selection on traits
  - ▶ Mutation introducing variation
  - ▶ Recombination mixing genotypes

# Evolution and Genetic Algorithms

- Evolutionary computing, a general concept in computational science



# Evolutionary computing

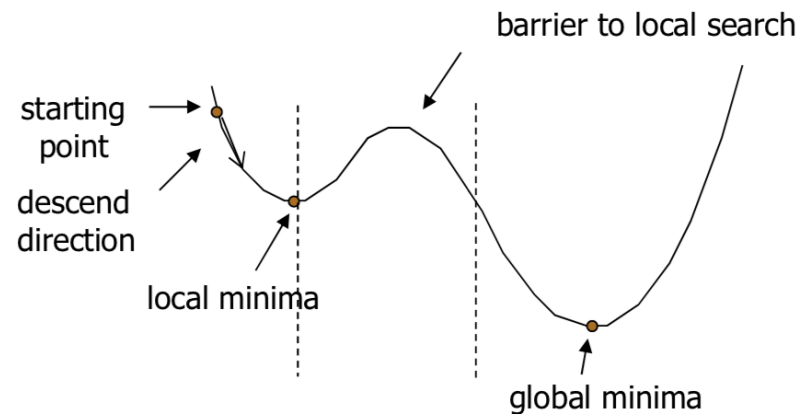
- ▶ Algorithmic models that use Darwinian-like evolution
- ▶ Iterative optimization
- ▶ E.g. Evolutionary strategy models
- ▶ E.g. Genetic algorithms
- ▶ E.g. Genetic programming

# Evolutionary computing

- ▶ Algorithmic models that use Darwinian-like evolution
  - ▶ Iterative optimization
  - ▶ E.g. Evolutionary strategy models
  - ▶ E.g. Genetic algorithms
  - ▶ E.g. Genetic programming
- 
- ▶ Key components:
    - ▶ Selection (better solutions survive)
    - ▶ Mutation (random change)
    - ▶ Crossover/ recombination (combining parts)
    - ▶ Inheritance (solutions pass on structure to next iteration)

# Genetic algorithms, GA

- ▶ Robust and global compared to calculus and enumerative solutions
- ▶ Easier to use when little is known about the problem
- ▶ Can search a large solution space and find a global optimum
- ▶ Use only for complex problem (expensive)





# GA, step by step

1. Represent your problem as genes in a binary value  
chromosome (= solution)

Example:

Find the maximum of a 10 digits base 2 number:

```
MyChromosome <- c(1, 0, 0, 0, 1, 0, 1, 1, 0, 0)
```

# Maximum Value of a 10- Digit Binary Number, known solution

- ▶ A 10-digit base-2 number has 10 bits, each being either 0 or 1.
  - ▶ The largest value is achieved when all bits are set to 1:  $1111111111_2$
  - ▶ Each bit represents a power of 2:  $2^9, 2^8, \dots, 2^1, 2^0$
  - ▶ Therefore:  $1111111111_2 = 2^9 + 2^8 + \dots + 2^1 + 2^0$
  - ▶ This is a geometric series:  $\text{sum} = 2^{10} - 1 = 1023$
  - ▶ A 10-bit number can represent a total of  $2^{10} = 1024$  distinct values (0 to 1023).

# GA, step by step

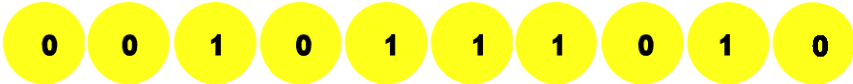
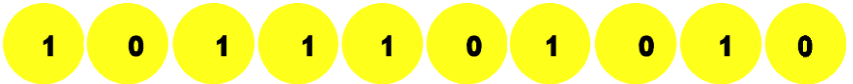
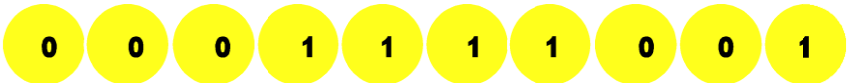

## 2. Random generation of a "population" of chromosomes

individual	representation									
1	0	0	1	0	1	1	1	0	1	0
2	1	0	1	1	1	0	1	0	1	0
3	0	0	0	1	1	1	1	0	0	1
4	1	0	0	0	0	1	1	1	1	0

In reality start with 50 chromosomes (=solutions)

# GA, step by step

## 3. Evaluate "fitness" of individual chromosomes

individual	representation	fitness
1		186
2		746
3		121
4		532

Fitness = how good is a chromosome as solution

# GA, step by step

## 4. Sort solutions after rank

individual	representation	fitness
1	1 0 1 1 1 0 1 0 1 0	746
2	1 0 0 0 0 1 1 1 1 0	532
3	0 0 1 0 1 1 1 0 1 0	186
4	0 0 0 1 1 1 1 0 0 1	121

# GA, step by step

## 5. Pairing, recombination

individual	representation	fitness
1	1 0 1 1 1 0 1 0 1 0	746
2	1 0 0 0 0 1 1 1 1 0	532
3	0 0 1 0 1 1 1 0 1 0	186
4	0 0 0 1 1 1 1 0 0 1	121

Elitistic pairing, random cut

# GA, step by step

## 5. Pairing, recombination → offspring

individual	representation	fitness
1	1 0 1 1 1 0 1 0 1 0	746
2	1 0 0 0 0 1 1 1 1 0	532
3	0 0 1 0 1 1 1 0 1 0	186
4	0 0 0 1 1 1 1 0 0 1	121
<hr/>		
5	1 0 1 1 1 1 1 1 1 0	767

High quality offspring = good solution

# GA, step by step

## 6. Insert offspring in population, discard worst solutions

individual	representation	fitness
1	1 0 1 1 1 1 1 1 1 0	767
2	1 0 1 1 1 0 1 0 1 0	746
3	1 0 0 0 0 1 1 1 1 0	532
4	0 0 1 0 1 1 1 0 1 0	186
<hr/>		
5	0 0 0 1 1 1 1 0 0 1	121

"killed"! (keep the number of chromosomes at 50)



# GA, step by step

Why cannot the GA solve this problem?

individual	representation	fitness
1	1 0 1 1 1 1 1 1 1 0	767
2	1 0 1 1 1 0 1 0 1 0	746
3	1 0 0 0 0 1 1 1 1 0	532
4	0 0 1 0 1 1 1 0 1 0	186

# GA, step by step

## 7. Mutation

individual	representation	fitness
1	1 0 1 1 1 1 1 1 1 0	767
2	1 1 1 1 1 0 1 0 1 0	874
3	1 0 0 0 0 1 1 1 1 0	532
4	0 0 1 0 1 1 1 0 1 0	186

# GA, step by step

## 7. Mutation

individual	representation	fitness
1	1 0 1 1 1 1 1 1 1 0	767
2	1 1 1 1 1 0 1 0 1 0	874
3	1 0 0 0 0 1 1 1 1 0	532
4	0 0 1 0 1 1 1 0 1 0	186

Now you can make a genetic algorithm!

# GA, step by step

## 8. Cycle until termination

1. Represent your problem as genes in a chromosome
2. Random generation of a population of chromosomes
- 3. Evaluate fitness of individual chromosomes
4. Sort after rank
5. Pairing, recombination
6. Insert offspring
7. Mutation

# GA, step by step

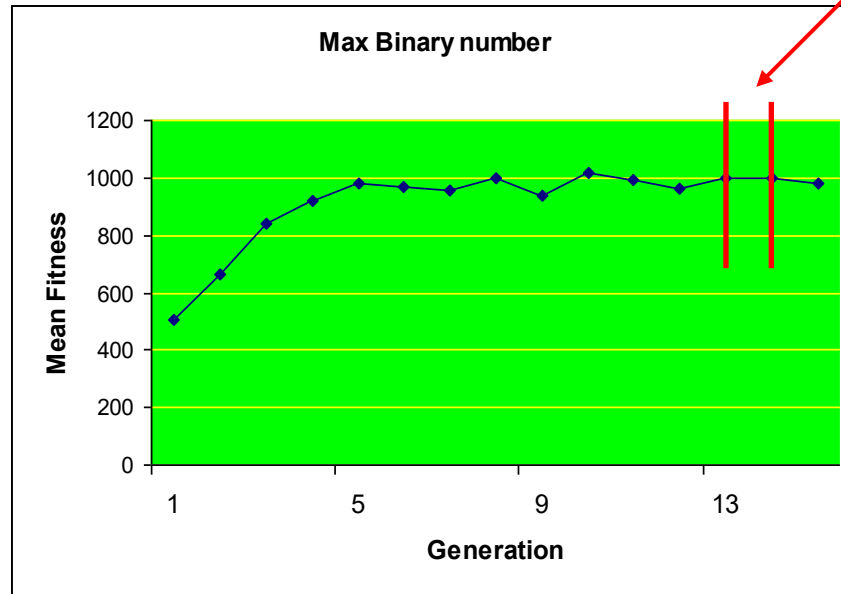
## 8. Cycle until termination

1. Represent your problem as genes in a chromosome
2. Random generation of a population of chromosomes
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4. Sort after rank
5. Pairing, recombination
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7. Mutation

Termination?

## 8. Cycle until termination

$$\text{diff} = y_1 - y_2$$

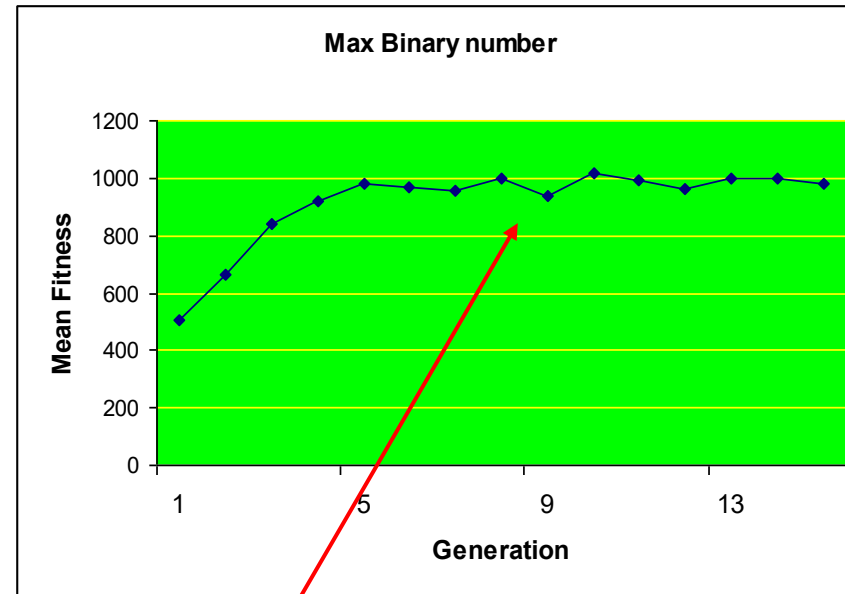


```
for (g in 1:15)  
    {cycle algorithm}
```

g = generation

```
while (diff > 10)  
    {diff =  $y_1 - y_2$ ;  
    cycle algorithm  
    }
```

## 8. Cycle until termination



Why not a straight line? It is a very simple problem.

## Genetic algorithm

Example 1, binary (= discrete) GA:

Find maximum of a 10 digits base 2 number:

```
MyChromosome <- c(1, 0, 0, 0, 1, 0, 1, 1, 0, 1)
```

Example 2, find best solution of an equation with two unknowns

$$f(x, y) = x \sin(4x) + y \sin(2y)$$

```
MyChromosome <- c( 3.43537, -1.0002345)
```



# Alternative Ways to Solve $f(x, y)$ Optimization

- ▶ Equation:  $f(x, y) = x \cdot \sin(4x) + y \cdot \sin(2y)$ 
  - ▶ Gradient-based optimization (e.g., BFGS, Newton, Conjugate Gradient)
  - ▶ Solve analytically via partial derivatives:  $\partial f / \partial x = 0$  and  $\partial f / \partial y = 0$
  - ▶ Multi-start local optimization to avoid local minima
  - ▶ Global optimization alternatives: simulated annealing, basin hopping
  - ▶ For bounded domains: grid search or fine 2D sampling

## Continuous genetic algorithms:

- Real values can be used instead of binary strings
- Decide range for gene values (e.g.  $-5 \leq x \leq 5$  )

Best solution of equation with two unknowns:

$$f(x, y) = x \sin(4x) + y \sin(2y)$$

Individual values (start with randomly created numbers)

1	[ 3.435, -1.003]
2	[ -2.331, - 4.234]
3	[ -0.877, - 4.954]
4	[ 3.537, 1.645]
5	[ 2.479, - 1.541]

# Continuous genetic algorithms:

- problems with recombination:

ind	values
1	[ 3.435, -1.003]
2	[ -2.331, - 4.234]
3	[ -0.877, - 4.954]
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Question1: Why is direct crossover not good?

# Continuous genetic algorithms:

- problems with recombination:

ind	values
-----	--------

1	[ 3.435, -1.003]
---	------------------

2	[ -2.331, - 4.234]
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3	[ -0.877, - 4.954]
---	--------------------

4	[ 3.537, 1.645]
---	-----------------

5	[ 2.479, - 1.541]
---	-------------------

- Key problems with direct crossover:
  - Breaks good (x, y) combinations
  - Creates poor-quality offspring
  - Does not fit continuous-valued search spaces
  - Offspring may fall in bad regions
  - Better operators exist (arithmetic, BLX- $\alpha$ , SBX)

Question1: Why is direct crossover not good?

# Continuous genetic algorithms:

- problems with recombination:

ind	values
1	[ 3.435, -1.003]
2	[ -2.331, - 4.234]
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Question 1: Why is direct crossover not good?

Question 2: Why is averaging better but not good?

# Continuous genetic algorithms:

- problems with recombination:

ind      values

1      [ 3.435, -1.003]

2      [ -2.331, - 4.234]

3      [ -0.877, - 4.954]

4      [ 3.537, 1.645]

5      [ 2.479, - 1.541]

Averaging (arithmetic crossover) improves over direct crossover, because:

- Offspring stay between parents (more stable than direct mixing)
- Less disruption of good (x, y) combinations
- Reduces the chance of creating extreme or invalid values

But it is still not ideal because:

- Averaging reduces diversity too quickly
- Cannot explore beyond parent region (no global exploration)
- May get stuck in local optima

Question 1: Why is direct crossover not good?

Question 2: Why is averaging better but not good?

## Continuous genetic algorithms:

- Blending (recombination operator in continuous GA)

ind      values

$p_1$       [ 3.435, -1.003]

$p_2$       [ -2.331, - 4.234]

Three solutions:

i)      proportional blending:  $b * p_1 + (1 - b) * p_2$       where  $0 \leq b \leq 1$

ii) linear blending gives three offspring:

$$\text{offsp1} = 0.5p_1 + 0.5p_2$$

$$\text{offsp2} = 1.5p_1 - 0.5p_2$$

$$\text{offsp3} = 1.5p_2 - 0.5p_1$$

iii) combination between crossover and blending

# Continuous genetic algorithms:

Question: What problem may this blending process create?  
(Think about the min and max values  $-5, 5$ )

Suggest a solution to this problem



# Continuous genetic algorithms:

- Individuals can be "out of range"

$$\text{offsp2} = 1.5p_1 - 0.5p_2$$

$p_1 = 4.952$ ,  $p_2 = -4.213$  gives offspring 9.535

Three solutions:

- i) chop at limit
- ii) "gene repair"
- iii) give low fitness

# Continuous vs discrete genetic algorithms:

Discrete (0 1 1 0 1)

binary

fitness function

simple crossing over

traditional, original

Continuous (0.654, 3.564...)

real numbers

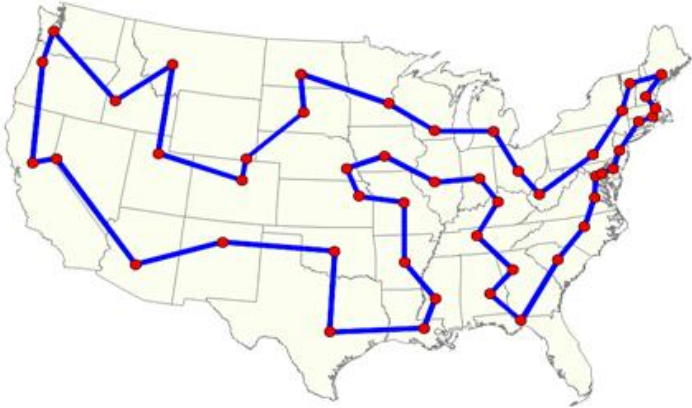
direct use of numbers

complex mating

newer

# Traveling salesman problem:

You have to visit a number of geographic positions, calculate the shortest route



## Traveling salesman problem:

You have to visit: Göteborg, Borås, Norrköping, Kristianstad, start in Lund



## Traveling salesman problem:

You have to visit: Göteborg, Borås, Norrköping, Kristianstad, start in Lund

Chromosome 1	Lund	Göteborg	Borås	N-köping	K-stad
Chromosome 2	Lund	Göteborg	N-köping	K-stad	Borås
Chromosome 3	Lund	K-stad	Göteborg	Borås	N-köping

Etc.

## Traveling salesman problem:

You have to visit: Göteborg, Borås, Norrköping, Kristianstad, start in Lund

Chromosome 1	Lund	Göteborg	Borås	N-köping	K-stad
Chromosome 2	Lund	Göteborg	N-köping	K-stad	Borås
Chromosome 3	Lund	K-stad	Göteborg	Borås	N-köping

Etc.

What is fitness?

## Traveling salesman problem:

You have to visit: Göteborg, Borås, Norrköping, Kristianstad, start in Lund

	Lund	Göteborg	Borås	N-köping	K-stad
Lund	0	262	275	431	77
Göteborg	262	0	63	311	264
Borås	275	63	0	250	264
N-köping	431	311	250	0	392
K-stad	77	264	264	392	0

# Traveling salesman problem:

You have to visit: Göteborg, Borås, Norrköping, Kristianstad, start in Lund

- ▶ TSP chromosome and fitness evaluation:
  - ▶ Chromosome = a permutation of cities (route order)
  - ▶ Example: [1, 5, 2, 3, 4] means visiting cities in that sequence
  - ▶ Distance matrix provides distance between each city pair
  - ▶ Total route distance = sum of distances between consecutive cities
  - ▶ Return to start city is included in route length
  - ▶ Fitness =  $1 / \text{total distance}$  (shorter route  $\rightarrow$  higher fitness)



# Traveling salesman problem:

You have to visit: Göteborg, Borås, Norrköping, Kristianstad, start in Lund

- **GA works well for the Traveling Salesman Problem:**
  - TSP search space grows factorially ( $n!$  routes), too large for brute force
  - GA can efficiently explore huge combinatorial spaces
  - Works well without problem-specific assumptions
  - Maintains a population of routes → avoids getting stuck in bad local optima
  - Specialized crossover/mutation operators preserve valid permutations
  - Produces good near-optimal solutions quickly, even when exact solution is hard
  - Robust against noisy or changing distance data