Spatial gene drives and pushed genetic waves

Hidenori Tanaka, Howard A. Stone, and David Nelson

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Mendelian Genetics 101

Question #1:

Microbiologists have managed to genetically engineer a mutant mosquito which does not propagate a certain disease.

Say that this mutant mosquito is AA for a particular trait and the wild type is aa for this trait. The mutant and the wild type mate and have an offspring.

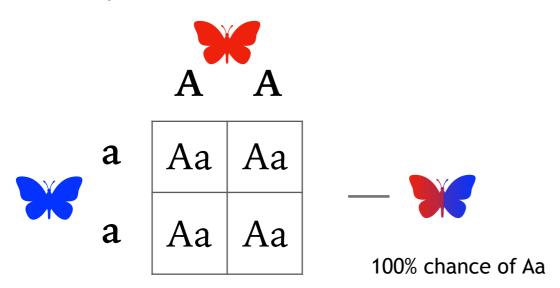
What is the probability that said offspring will propagate the desired A allele into the next generation?

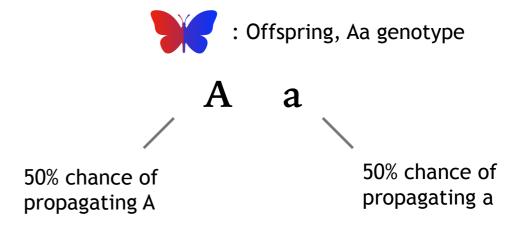
: Mutant type, AA genotype

: Wild type, aa genotype

: Offspring, ??

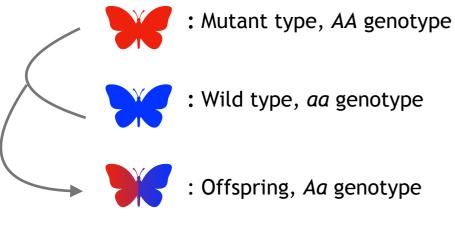
Solution: Setup an inheritance matrix





There is a 50% chance that the offspring will propagate the allele A to the next generation.

Mendelian Genetics 101



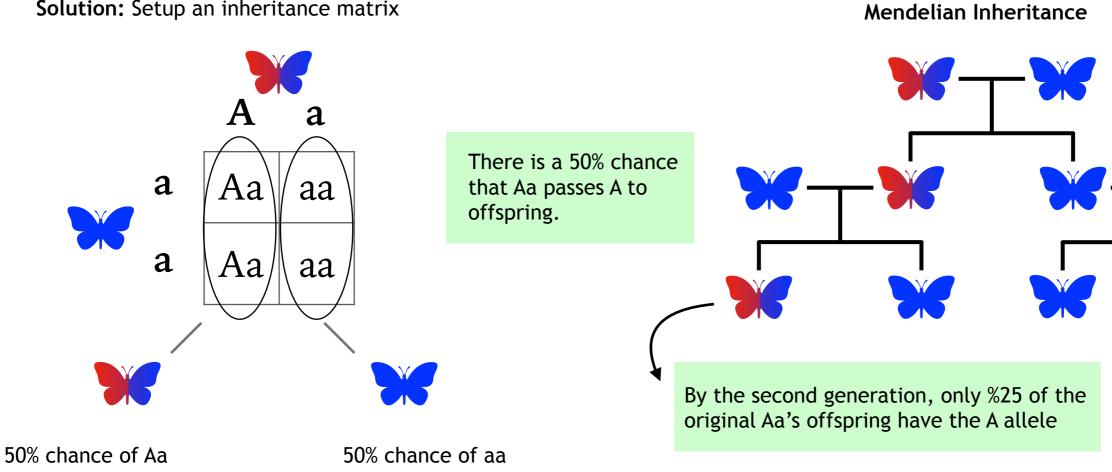
* Mixed colors refers to allele mixing, NOT phenotype mixing.

Question #2:

This mixed allele Aa offspring is placed in a population completely composed of Wild type aa organisms.

In its second generation, what proportion of Aa's offspring would have the allele A?

Solution: Setup an inheritance matrix



Why do we need Gene Drives?

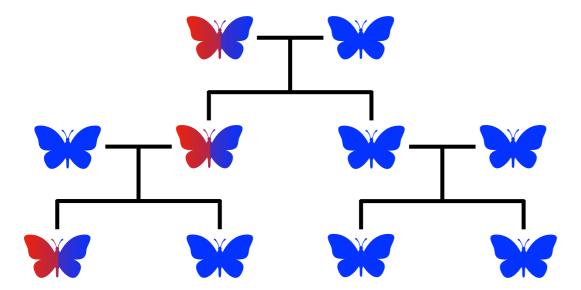
A

a

Desired mutant allele

Wild allele

Mendelian Inheritance



By Aa's second generation, only %25 of the original Aa's offspring have the A allele

There's a problem:

Our desired mutation does not propagate fast enough through the population.

Question: How can we get our desired mutation to comprise a larger proportion of the progeny?

Answer: Gene Drives!

(**Definition**) Gene drives are genes which code for their own duplication in the partner chromosome.

(Functionally) Gene drives convert heterozygous genotypes into homozygous genotypes. (This occurs between generation cycles)



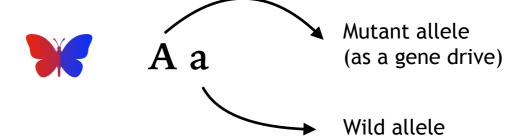
Aa

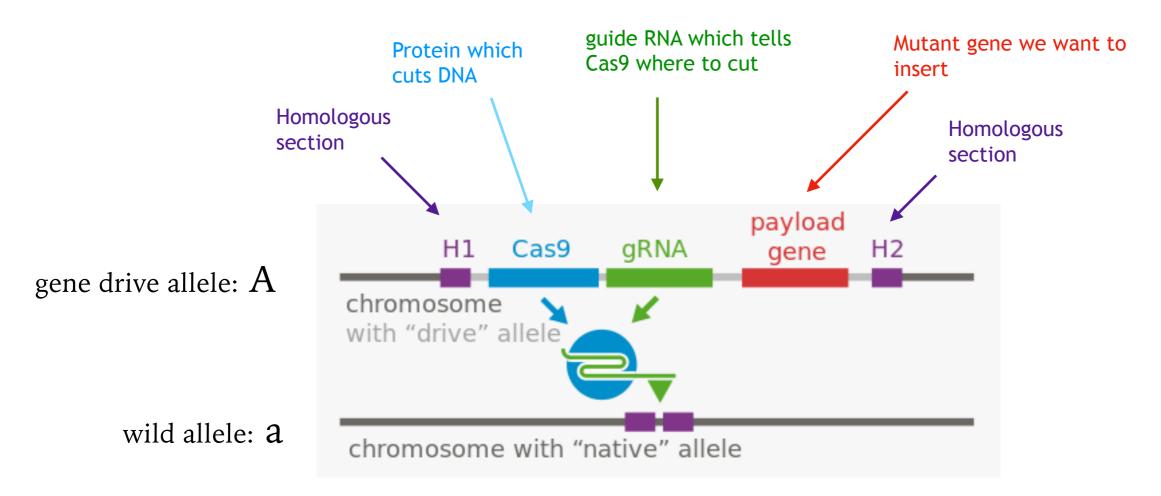
AA

How do Gene Drives work?

(**Definition**) Gene drives are genes which code for their own duplication in the partner chromosome.

Say we begin with the heterozygous mutant:

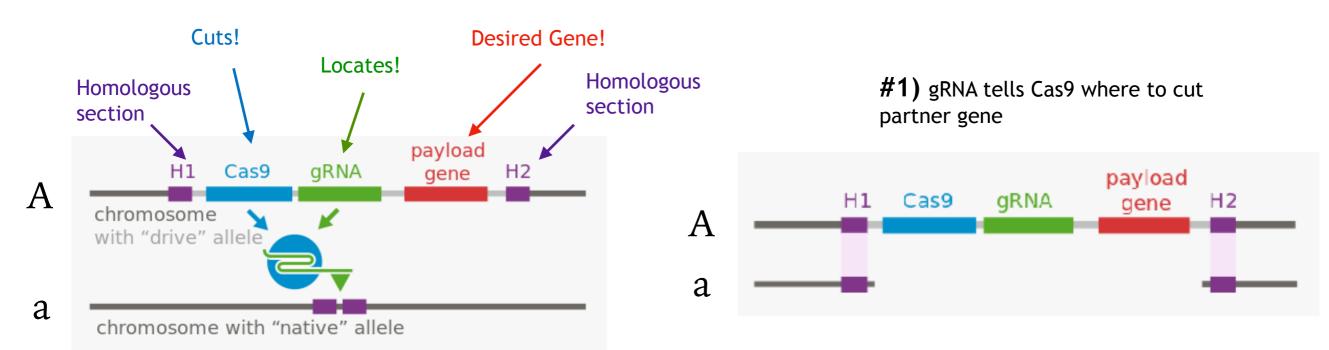




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How do Gene Drives work?

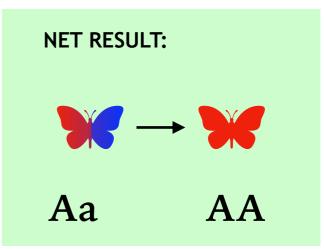
(**Definition**) Gene drives are genes which code for their own duplication in the partner chromosome.



#2) Cut is repaired by replicating* gene drive in partner chromosome



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* Replication process is prone to errors, so that it seldom occurs at 100% efficiency. (Paper assumes 100% efficiency)

"Super Mendelian" Genetics 101

Question #1:

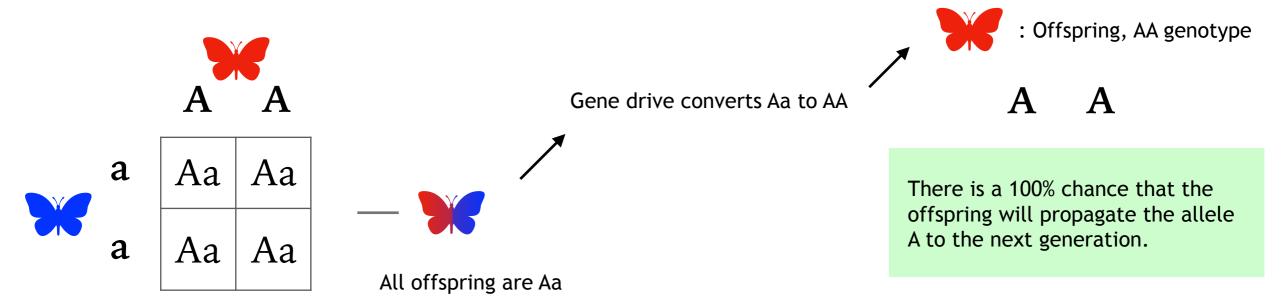
Microbiologists have managed to genetically engineer a mutant mosquito which does not propagate a certain disease.

Say that this mutant mosquito is AA (with a **GENE DRIVE**) for a particular trait and the wild type is aa for this trait. The mutant and the wild type mate and have an offspring.

What is the probability that said offspring will propagate the desired A allele into the next generation?



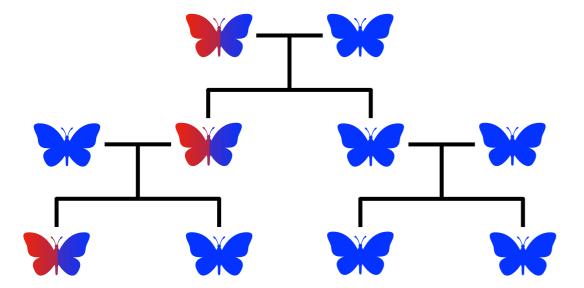
Solution: Setup an inheritance matrix



"Super Mendelian" Genetics 101

Question: How can we get our desired mutation to comprise a larger proportion of the progeny?

Mendelian Inheritance

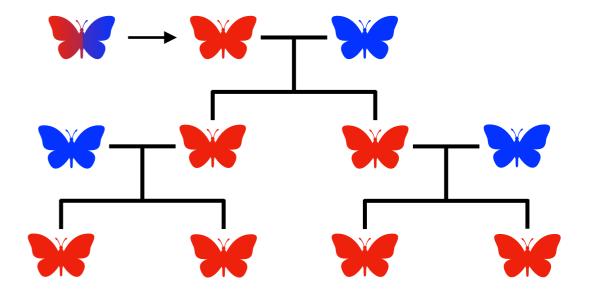


By the second generation, only %25 of the original Aa's offspring have the A allele

A a

Desired mutant allele Wild allele

"Super Mendelian"/Gene Drive Inheritance



By the second generation, 100% of the original Aa's offspring have the A allele.

Problem: Gene drive overcorrects Mendelian genetics!

Previously (Mendelian Genetics):

Engineered mutation does not propagate fast enough through the population.

Now (Super Mendelian Genetics):

Engineered mutation (with gene drive) completely overtakes the population.

"Super Mendelian" Genetics 101

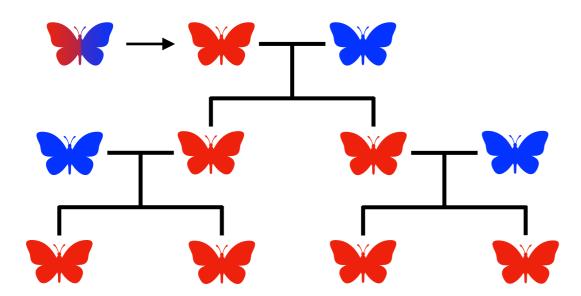
A

a

Desired mutant allele

Wild allele

"Super Mendelian"/Gene Drive Inheritance



By the second generation, 100% of the original Aa's offspring have the A allele.

AND 0% have the a allele

Paper's Main Question:

How can we control whether a genedrive mutation overtakes a population?

Partial Answer:

The gene drive mutant has a selective disadvantage relative to the wild type.

$$w_{\rm drive} = w_0(1-s)$$

Main Question (Rephrased): For what selection coefficients can an organism with a gene drive be controlled so that it neither overtakes nor is eliminated from a population?

Paper's Main Question: For what selection coefficients can an organism with a gene drive be controlled so that it neither overtakes nor is eliminated from a population?

Main Strategy: Use Hardy-Weinberg Principles to derive an equation for allele frequency as a function of time.

$$w_{\text{drive}} = w_0(1-s)$$

What is s for a controllable gene drive?

$$q$$
 : frequency of gene drive allele
$$\frac{dq}{dt} + U'(q,s) = 0 \label{eq:qp}$$

Hardy Weinberg (HW) Principle: The allele frequencies in a population (without selection and mutations) remains constant throughout time.

Example 1: HW with NO SELECTION and NO GENE DRIVE

q: frequency of "A" (mutant allele) at time t

p: frequency of "a" (wild allele) at time t

*p + q = 1 (frequencies sum to 1)

Genotype	Frequency	Fitness
AA	q^2	w_0
Aa	2pq	w_0
aa	p^2	w_0

How does the allele frequency of A change for an incremental time step?

$$egin{aligned} q(t+\Delta t) &= rac{ ext{Mean fitness for allele A}}{ ext{Mean fitness overall}} \ &= rac{w_0 q^2 + rac{1}{2} w_0 2 p q}{w_0 q^2 + w_0 2 p q + w_0 q^2} \ &= q^2 + q p \ &= q \end{aligned}$$

As we expect, the frequency of allele A remains the same for all time changes.

Example 2: HW with SELECTION and NO GENE DRIVE

q: frequency of "A" (mutant allele) at time t

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How does the allele frequency of A change for an incremental time step?

$$egin{aligned} q(t+\Delta t) &= rac{ ext{Mean fitness for allele A}}{ ext{Mean fitness overall}} \ &= rac{w_0(1-s)q^2 + rac{1}{2}w_02pq}{w_0(1-s)q^2 + w_02pq + w_0q^2} \ &= rac{q(1-sq)}{1-sq^2} \end{aligned}$$

We find that $q(t+\Delta t) < q$. Thus when the mutant genotype is selected against, its frequency decreases

Example3: HW with SELECTION and GENE DRIVE (case in the paper)

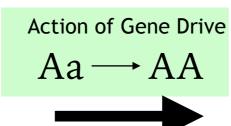
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Genotype	Frequency	Fitness
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How does the allele frequency of A change for an incremental time step?

Genotype	Frequency	Fitness
AA	q^2	$w_0(1-s)$
Λa	2pq	w_0
Aa	p^2	w_0

Action of Gene Drive
$$Aa \longrightarrow AA$$

Genotype	Frequency	Fitness
AA	q^2+2pq	$w_0(1-s)$
aa	p^2	w_0

$$egin{aligned} q(t+\Delta t) &= rac{ ext{Mean fitness for allele A}}{ ext{Mean fitness overall}} \ &= rac{w_0(1-s)(q^2+2pq)}{p^2+w_0(1-s)(q^2+2pq)} \ &= rac{(1-s)q(q+2p)}{1-sq(2-q)} \end{aligned}$$

It is unclear how $q(t+\Delta t)$ compares to q. We require a more involved analysis.



Define:

$$\lim_{\Delta t \to 0} q(t + \Delta t) - q \equiv \tau_g \frac{dq}{dt}$$

We then obtain

 τ_g : generation time

$$\tau_g \frac{dq}{dt} = \frac{sq(1-q)(q-q^*)}{1-sq(2-q)}$$

where
$$q^* \equiv \frac{2s-1}{s}$$

HW with SELECTION and GENE DRIVE (case in the paper)

$$\tau_g \frac{dq}{dt} = \frac{sq(1-q)(q-q^*)}{1-sq(2-q)}$$

where
$$q^* \equiv \frac{2s-1}{s}$$

This equation determines how the mutant (gene drive) frequency evolves under selection

Limiting Cases: No Selection and Full Selection

– No Selection; s=0(Mutant has wild type fitness) $\tau_g \frac{dq}{dt} = q(1-q)$

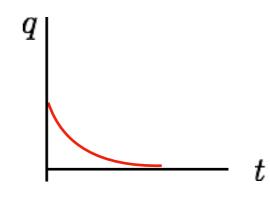
$$\tau_q \frac{dq}{dt} = q(1-q)$$

- Full Selection; s = 1 (Mutant has no fitness)

$$au_g rac{dq}{dt} = -q$$



The mutant (gene drive) allele completely overtakes the population



The mutant (gene drive) allele completely dies out

Reality is somewhere in between.

How do we analyze the long time behavior of the mutant allele for intermediate values of s?

Potential Energy Landscape for Long-Time Behavior

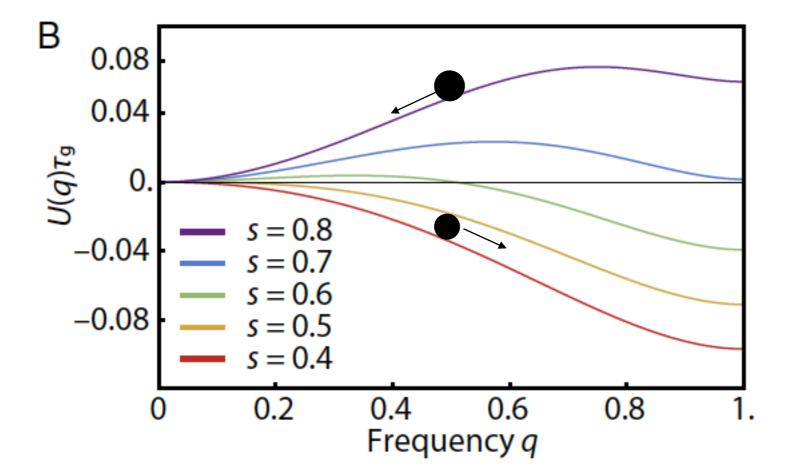
How do we analyze the long-time behavior of q(t) for arbitrary selection coefficient s?

$$au_g rac{dq}{dt} = rac{sq(1-q)(q-q^*)}{1-sq(2-q)}$$
 where $q^* \equiv rac{2s-1}{s}$

Introduce a Potential Function

$$\begin{split} \frac{dq}{dt} &= -U'(q,s) \\ U(q,s) &= -\frac{1}{\tau_g} \int_0^q dq' \, \frac{sq(1-q)(q-q^*)}{1-sq(2-q)} \end{split}$$

Potential Landscape



Long Time Behavior:

For a given initial q, we can find the long time behavior by letting q "roll" down the potential landscape

The system evolves to either q = 0 or q = 1.

Say
$$q(t = 0) = 0.5$$

- High selection coefficient (s = 0.8)
$$q(t \to \infty) = 0$$

- Low selection
$$q(t \to \infty) = 1$$

Gene Drive Dynamics with Diffusion

How do the dynamics change when the alleles are allowed to diffuse through space?

Qualitatively, the authors find three regimes

Explosion (s<0.5):

Gene drive allele ALWAYS TAKES OVER

Tunable (0.5<s<0.697):

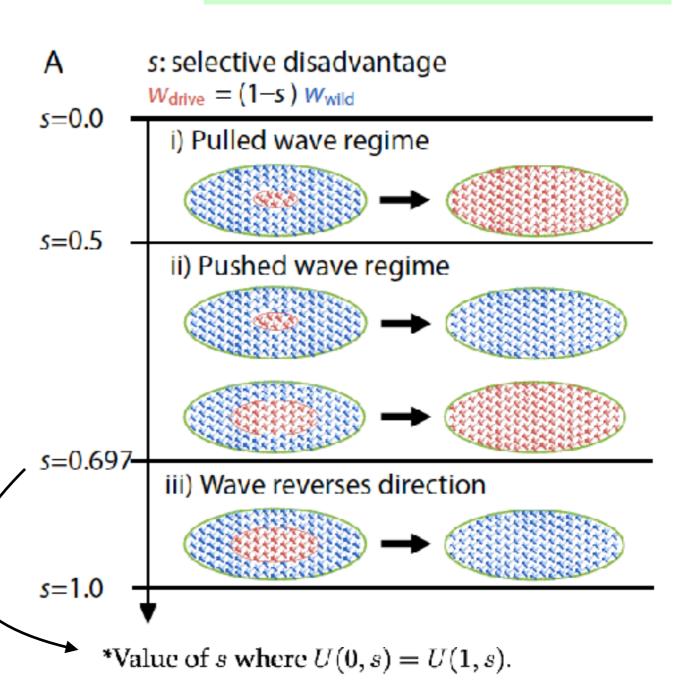
Gene drive allele only takes over for certain conditions

Failure (s>0.697):

Gene drive allele ALWAYS COLLAPSES

Answer found by solving the 1D Diffusion equation

$$\tau_g \frac{\partial q}{\partial t} = D \frac{\partial^2 q}{\partial x^2} + \frac{sq(1-q)(q-q^*)}{1-sq(2-q)}$$



Specific Solutions to Gene Drive Diffusion Equation

How does the nature of solutions to

$$\tau_g \frac{\partial q}{\partial t} = D \frac{\partial^2 q}{\partial x^2} + \frac{sq(1-q)(q-q^*)}{1-sq(2-q)}$$

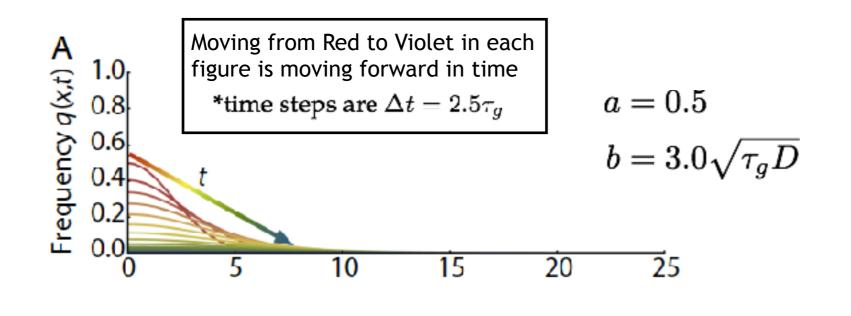
depend on solution parameters and selection coefficient?

0.0L

Defining initial condition and selection coefficient:

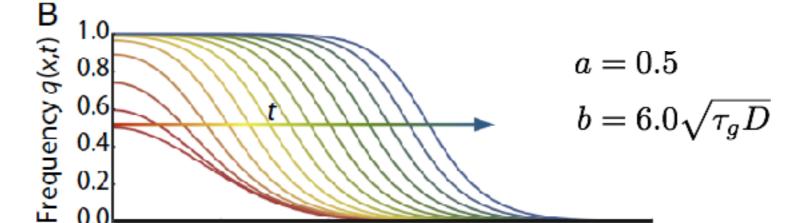
$$q(x,0) = a e^{-(x/b)^2}; s = 0.58$$

*(Tunable Regime) 0.5<0.58<0.697



Narrow distribution:

Gene drive allele eventually collapses



Distance $x/\sqrt{\tau_9D}$

15

20

10

25

Broad distribution:

Gene drive allele eventually takes over.

Gene Drive Diffusion Equation with Selection Barriers

What is the nature of these solutions when we incorporate a **selection boundary**?

$$\tau_g \frac{\partial q}{\partial t} = D \frac{\partial^2 q}{\partial x^2} + \frac{sq(1-q)(q-q^*)}{1-sq(2-q)}$$

Defining initial condition and function parameters

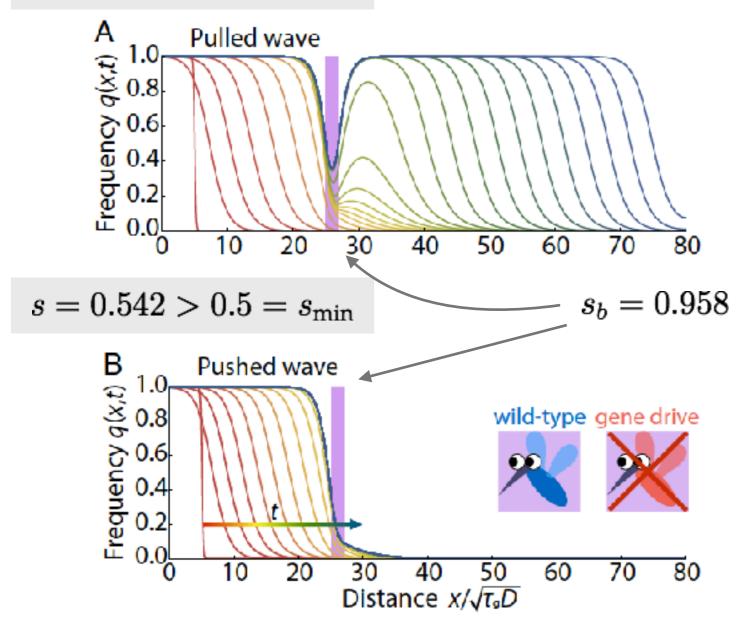
$$q(x,0) = rac{q_0}{1 + e^{10(x-x_0)/\sqrt{ au_g D}}}$$

where

$$q_0 = 1.0$$

$$x_0 = 5.0\sqrt{\tau_g D}$$

$$s = 0.479 < 0.5 = s_{\min}$$



Moving from Red to Violet in each figure is moving forward in time *time steps are $\Delta t = 2.5 au_g$

Low (s<0.5) Selection:

Gene drive allele overcomes selection barrier

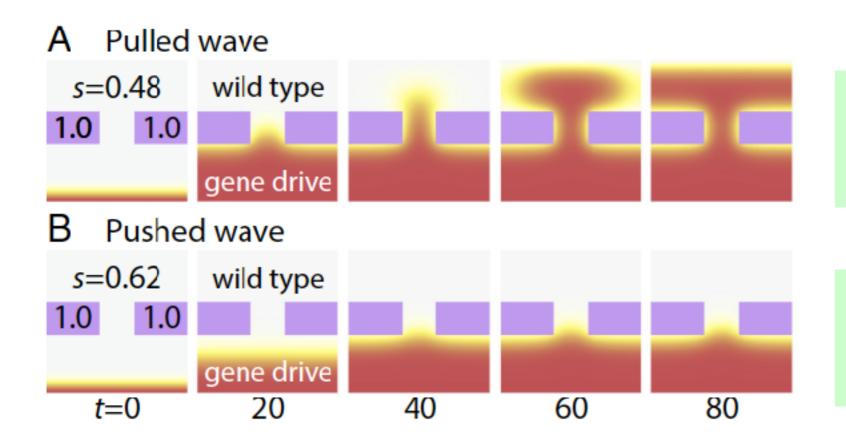
Tunable (0.5< s< 0.697) Selection:

Gene drive allele cannot get past selection barrier

Selection Barriers in Two-Dimensions

How do selection barriers affect the propagation of alleles for two dimensional systems?

$$\tau_{g} \frac{\partial q}{\partial t} = D \left(\frac{\partial^{2} q}{\partial x^{2}} + \frac{\partial^{2} q}{\partial y^{2}} \right) + \frac{sq(1-q)(q-q^{*})}{1 - sq(2-q)}$$



Low (s<0.5) selection:

Allele is able to diffuse through gap in the barrier.

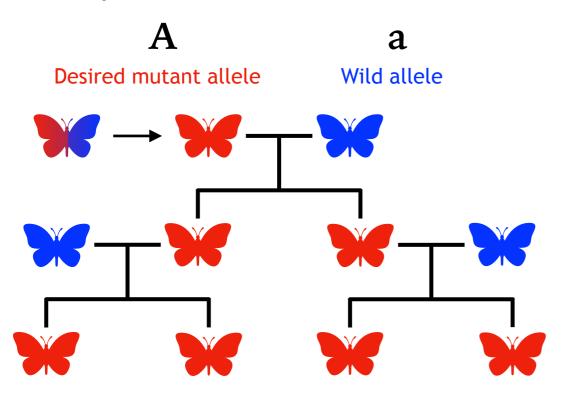
Tunable(0.5<s<0.697) selection:

Allele cannot diffuse through gap in the barrier

Even incomplete barriers in 2D can limit the spread of an allele

Concluding Summary

"Super Mendelian"/Gene Drive Inheritance



Some other questions

- How does inefficient gene-drive conversion change the results? (Answered in appendix)
- How would one specifically engineer a desired selection coefficient?

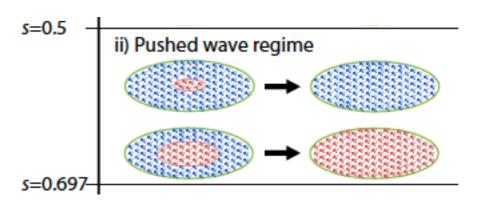
Paper's Main Question:

How can we control whether a gene-drive mutation overtakes a population?

General Answer:

We can limit the spread of the gene-drive mutation by engineering its selection coefficient to be within a specific range of values

*For the model in the paper the range is 0.5 <s < 0.697



END