Up until this point we have focused on "**Classical Genetics**": Starting with a biochemical, developmental, or other process, identify the genes involved and figure out how they work together...

FROM FUNCTION TO GENES

Starting in the early 90s, we knew about a lot of genes that were emerging from genome sequencing projects, but whose function was completely unknown.

"**Reverse Genetics**" - investigating the function of known genes by targeted disruption

FROM GENES TO FUNCTION

Reverse genetics in mice

The 2007 Nobel Prize
in Physiology or Medicine
was awarded to...

Mario Capecchi

Sir Martin J. Evans

Oliver Smithies

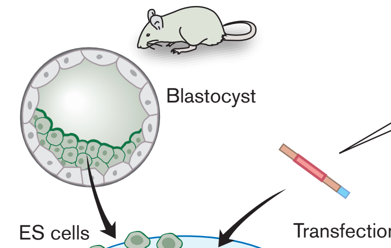
...for developing methods for gene disruption (a.k.a. gene targeting, or genetic knockouts) in mice

Gene disruption in mice is a long and laborious process... it sure would be nice to characterize genes of interest in a simpler organism before going to all this trouble!

General strategy for gene targeting in mice

Step 1 Gene targeting in ES cells

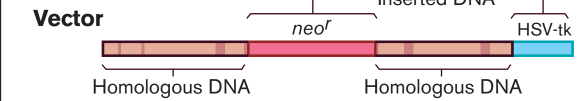
1. ES cell culture
Embryonic stem (ES) cells are cultivated from mouse pre-implantation embryos (blastocysts).



Rare cell carrying targeted gene

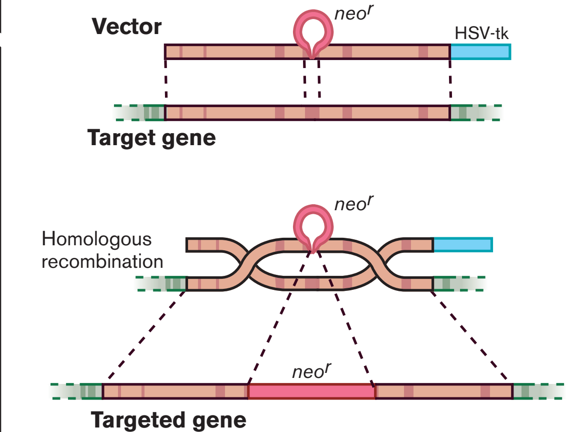
Target gene

Positive-negative selection

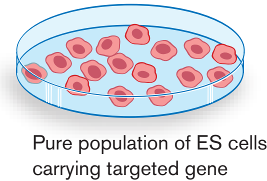


2. Construction of targeting vector
The vector contains pieces of DNA that are homologous to the target gene, as well as inserted DNA which changes the target gene and allows for positive-negative selection.

3. ES cell transfection
The cellular machinery for homologous recombination allows the targeting vector enables the target vector to find and recombine with the target gene.



4. Proliferation of targeted ES cell
Selection for presence of neo^r and absence of HSV-tk enriches targeted ES cells.



Pure population of ES cells carrying targeted gene

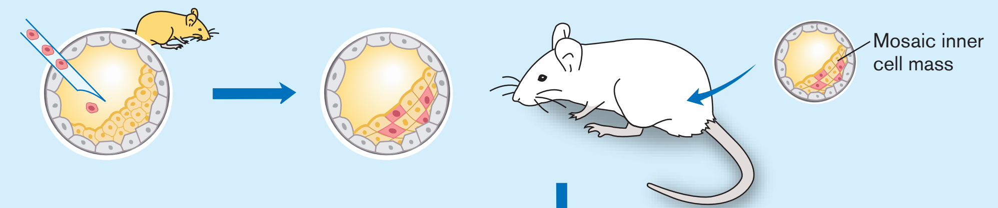
Step 2 From gene targeted ES cells to gene targeted mice

5. Injection of ES cells into blastocysts

The targeted ES cells are injected into blastocysts...

...where they mix and form a mosaic with the cells of the inner cell mass from which the embryo develops.

The injected blastocysts are implanted into a surrogate mother where they develop into mosaic embryos.

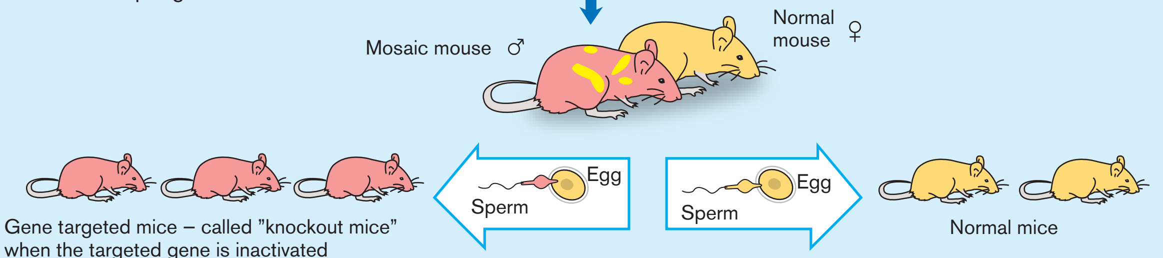


6. Birth and breeding of mosaic mice
The mosaic mice mate with normal mice to produce both gene targeted and normal offspring.

Born mosaic mice

Mosaic mouse ♂

Normal mouse ♀



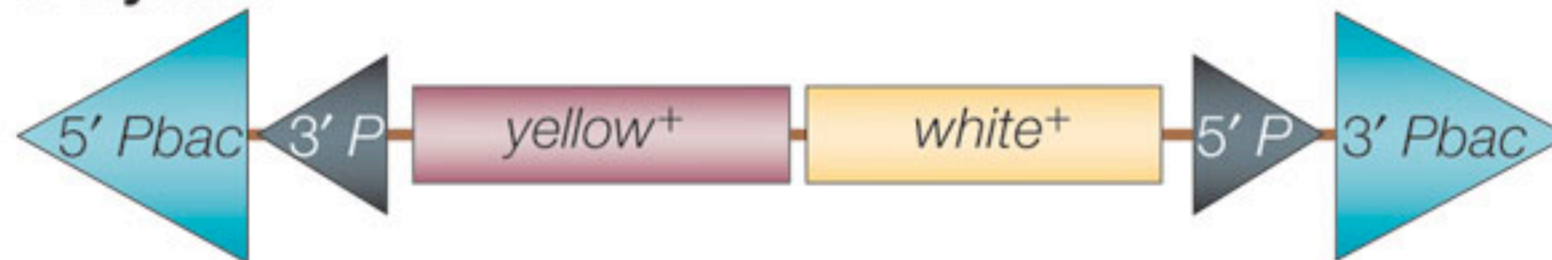
Gene targeted mice – called "knockout mice" when the targeted gene is inactivated

Normal mice

© The Nobel Committee for Physiology or Medicine Illustration: Annika Röhl

Reverse genetics in Drosophila

In *Drosophila*, it is possible (but not trivial) to generate mutations in specific genes by "hopping" transposable elements around the genome and then sifting through the collection of resulting flies for individuals that have a transposon in the gene of interest.



A few different transposons are used to generate these insertion collections, because individual transposons have "hotspots" where they like to jump, and may never land in certain genes.

A transposon insertion can create a loss-of-function mutation, but sometimes it doesn't (for example, transposons have a tendency to jump into introns rather than exons, in which case they can get spliced out of the messenger RNA).

In these cases, you have to get the transposon to hop *out* of the gene and hope for an *imprecise excision* that deletes some of the gene.

In *C. elegans*, Craig Mello's lab reported in 1997 that injection of "antisense" RNA could apparently reduce the function of a gene of interest...

It was *thought* that this effect occurred through a mechanism that blocks translation:

Antisense DNA oligonucleotide

mRNA

Transcription

DNA

Translation

Ribosome

Amino acids

Protein

...but, there was some serious weirdness. They noticed that the "control" sense RNA could induce the same effect.

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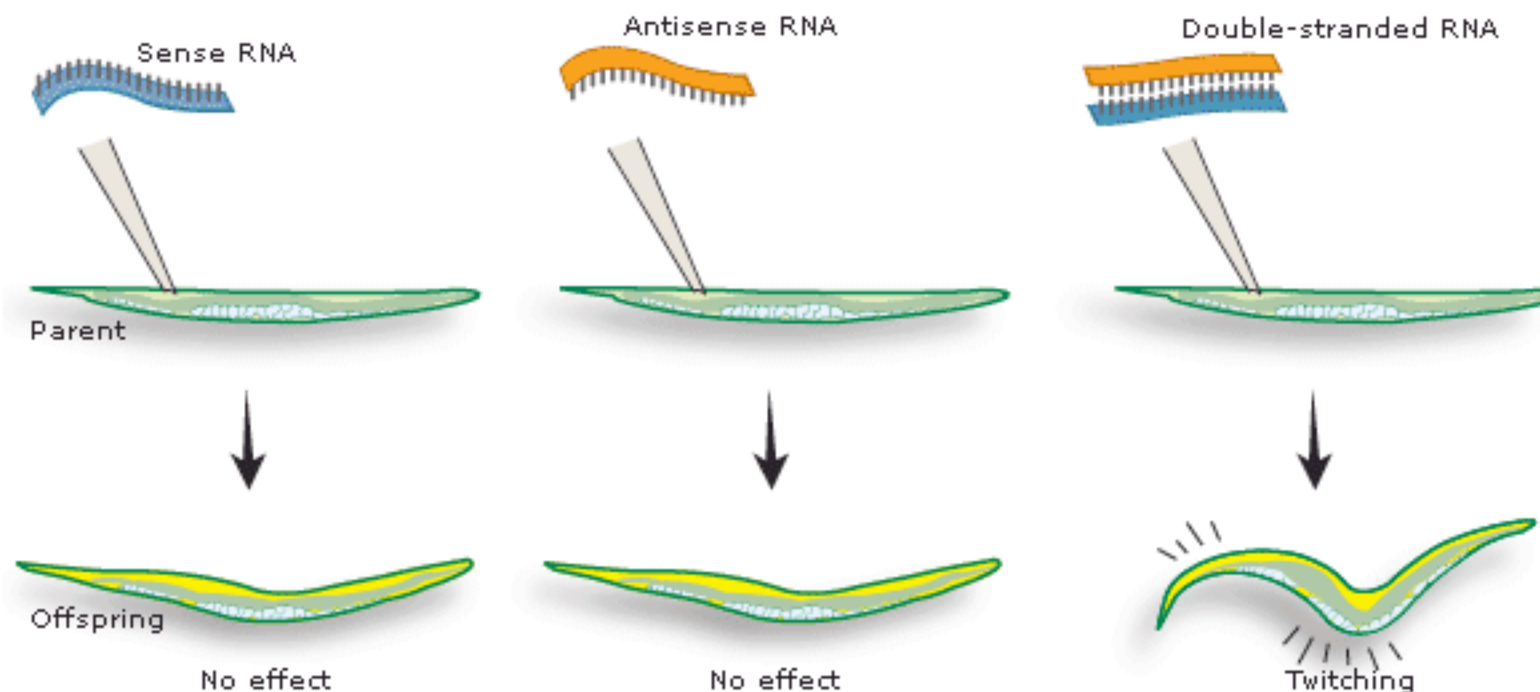
Potent and specific genetic interference by double-stranded RNA in *Caenorhabditis elegans*

Andrew Fire*, SiQun Xu*, Mary K. Montgomery*, Steven A. Kostas*†, Samuel E. Driver‡ & Craig C. Mello‡

Andy Fire and Craig Mello figured out that the interference was due to small amounts of *double-stranded* RNA in the "sense" and "antisense" preparations.



Injection of purified sense or antisense RNA from the *unc-22* gene into wild-type worms did not produce a mutant phenotype, but mixing the two strands did.

For this discovery, they were awarded the 2006 Nobel Prize in Physiology or Medicine

Andrew Fire

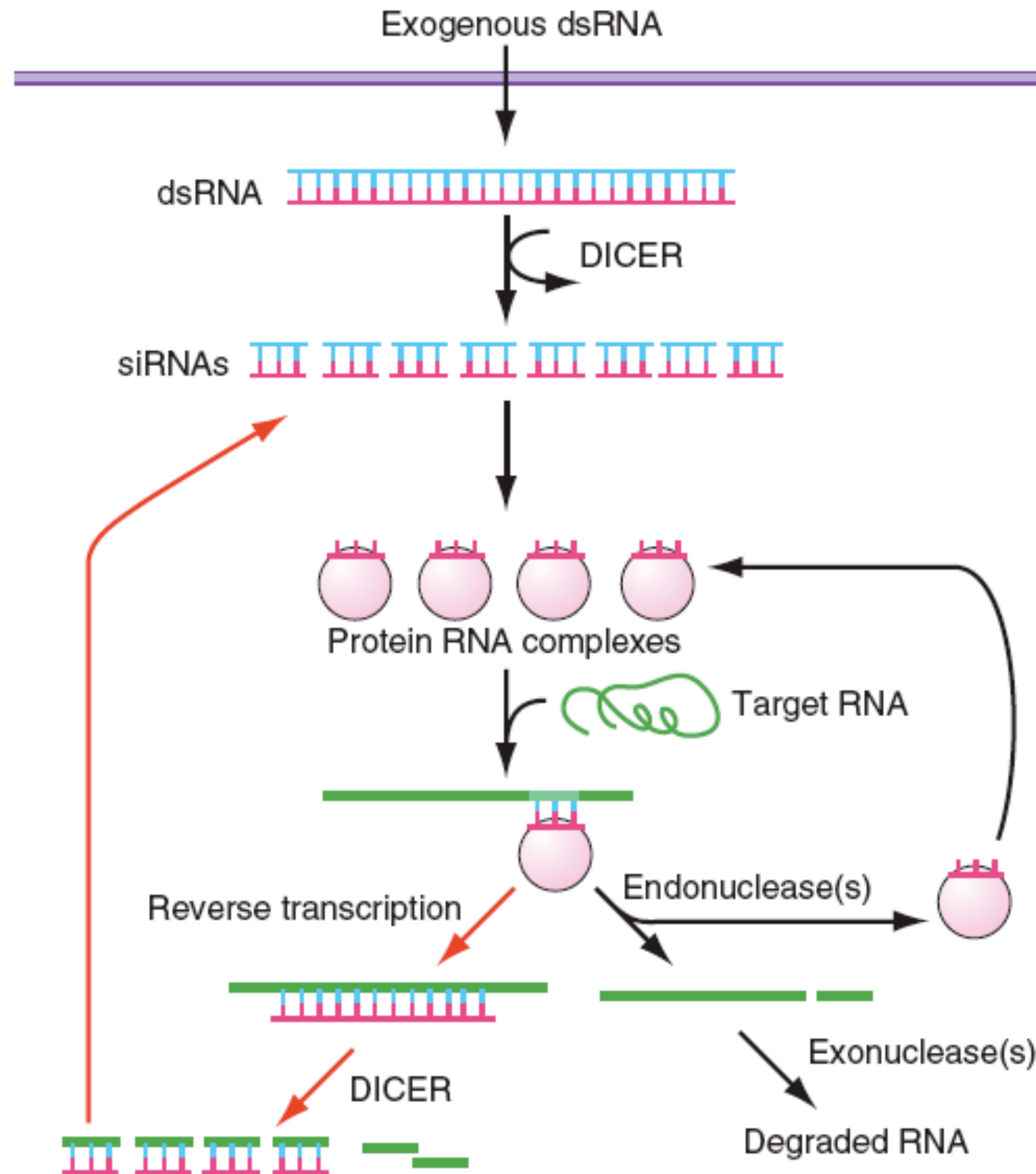
Craig Mello

Why was this simple finding so revolutionary?

Their experiments, along with follow-up work by their labs and others, uncovered the existence of an unknown mechanism in plants, animals, and many fungi (but not budding yeast) called "double-stranded RNA-mediated interference," or RNAi.

This knowledge has radically changed experimental biology, and led to the possibility of RNAi-based therapeutics

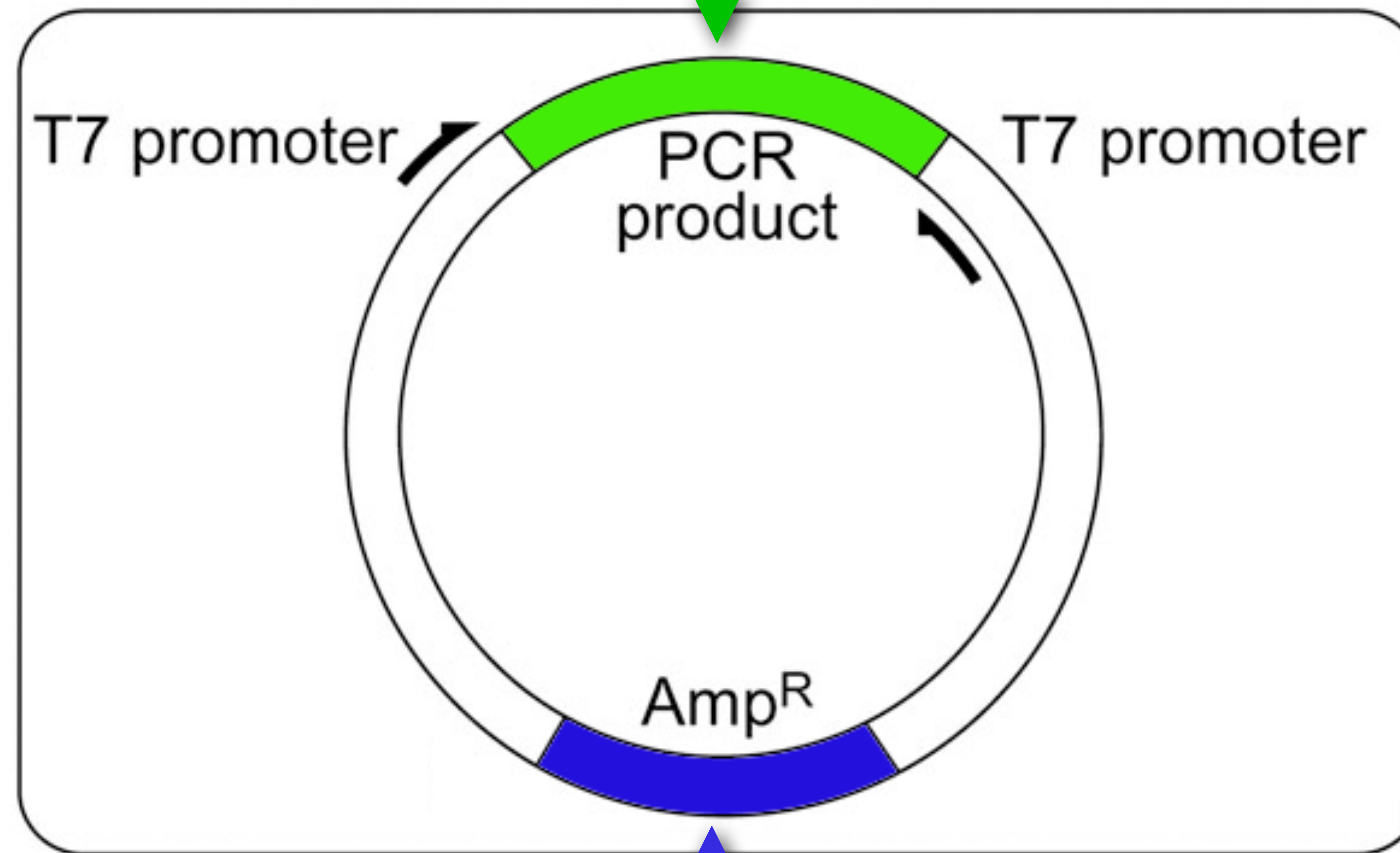
RNAi probably evolved because double-stranded RNA is viewed as "toxic" by eukaryotic cells. A special RNAse enzyme called Dicer chops up dsRNA into small fragments. The resulting *siRNAs* (small interfering RNAs) are then bound by a protein complex (the RISC) complex, which leads to destruction of any complementary mRNA



Worms that eat bacteria expressing dsRNA will undergo silencing of the corresponding gene

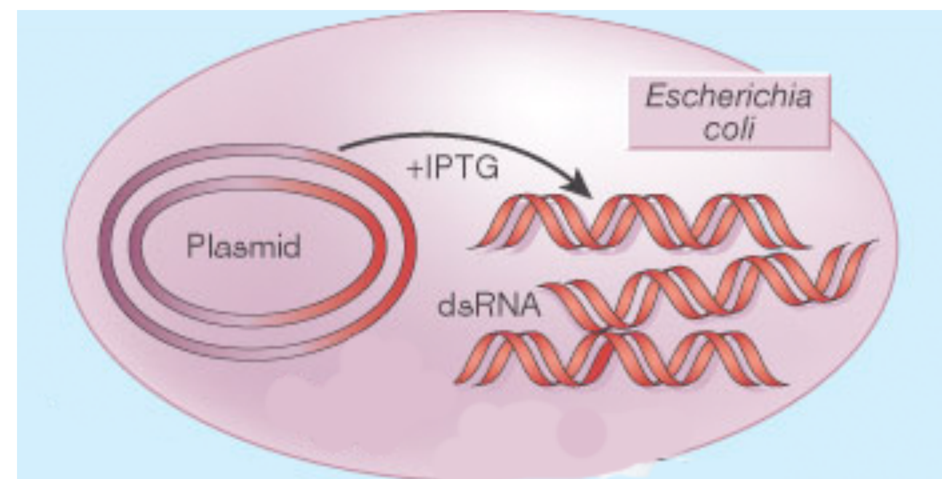
piece of worm gene coding sequence

bacterial cell (*E. coli*)



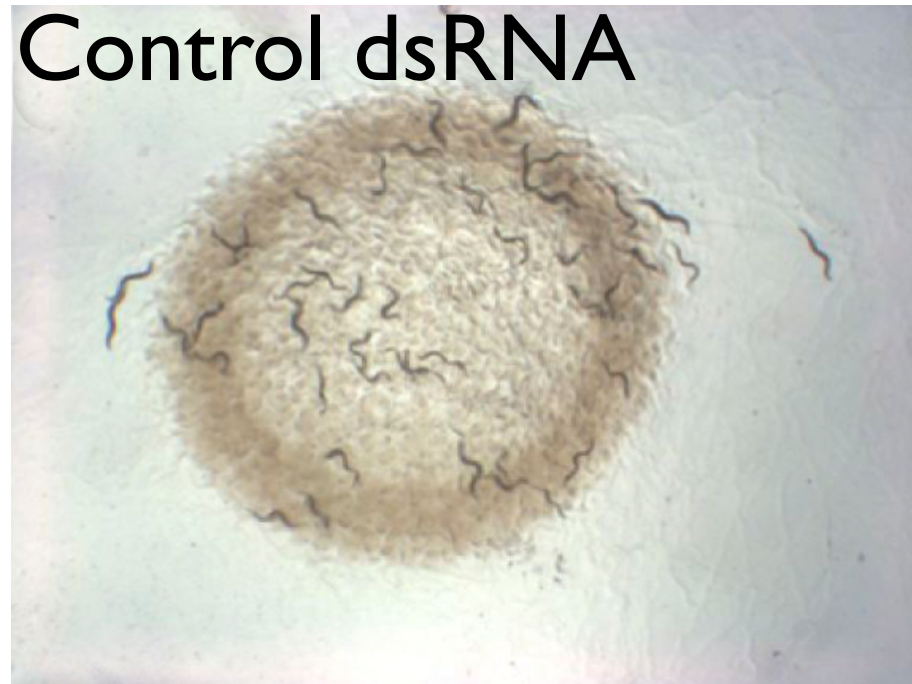
T7 promoters are turned on when bacteria are treated with IPTG

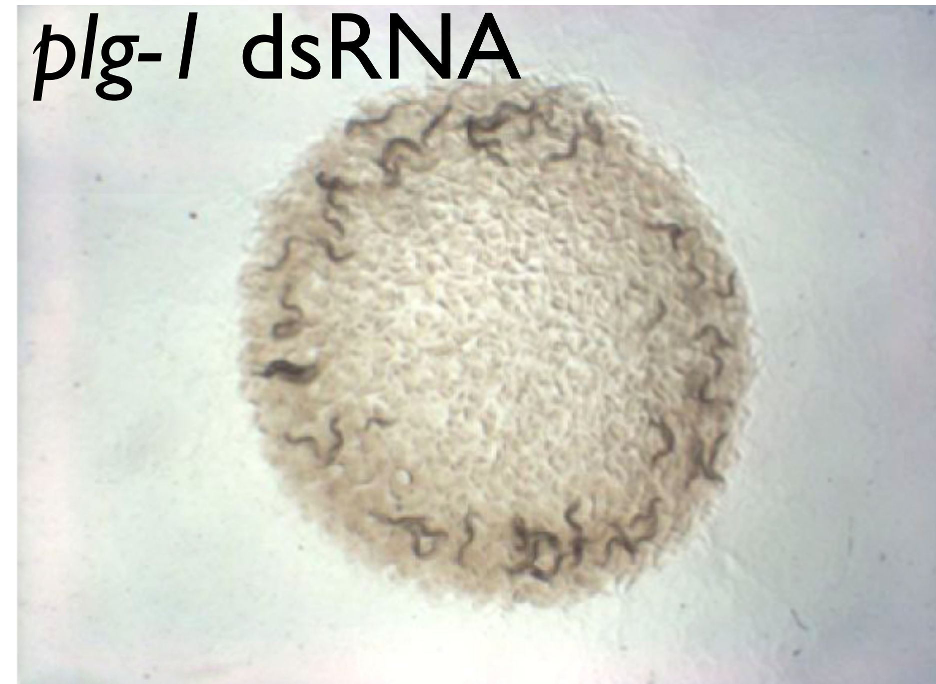
selectable marker (to maintain plasmid)



Worms that eat bacteria expressing dsRNA will undergo silencing of the corresponding gene

Worms eating bacteria





Cell-specific or Tissue-specific RNAi

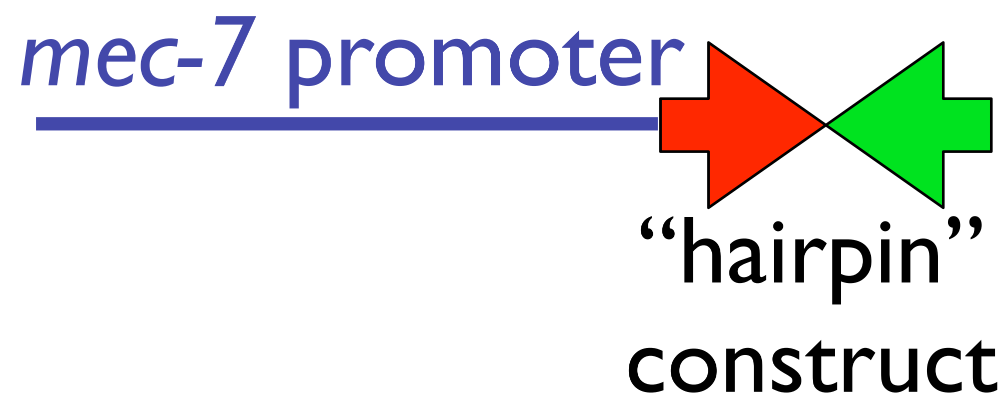
We previously discussed the idea of making a "transgene" that expresses a gene of interest from a tissue- or cell-type-specific promoter

The mec-7 gene is normally expressed in the ALM neurons

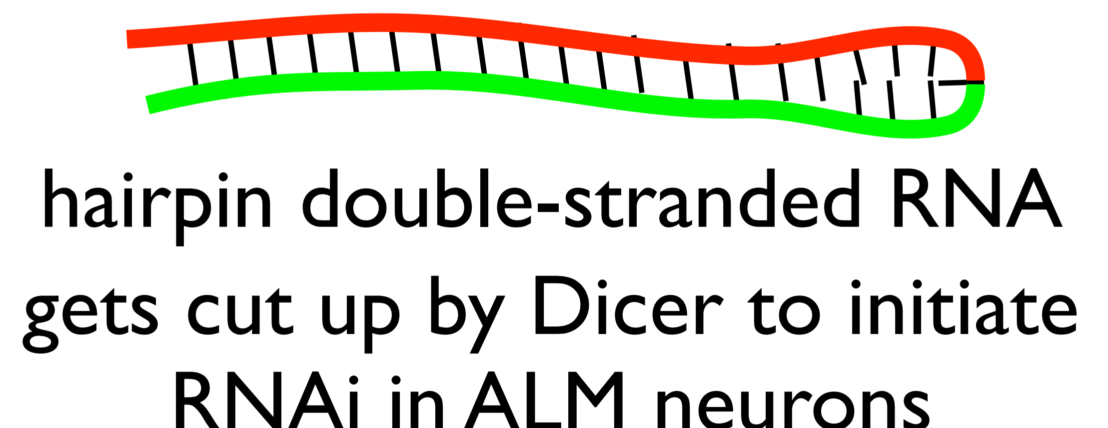
$P_{mec\text{-}7}$::*ced-3*

A different gene placed under the control of the *mec-7* promoter will also be expressed at high levels in those neurons

The same strategy can be used to induce RNAi in specific cells



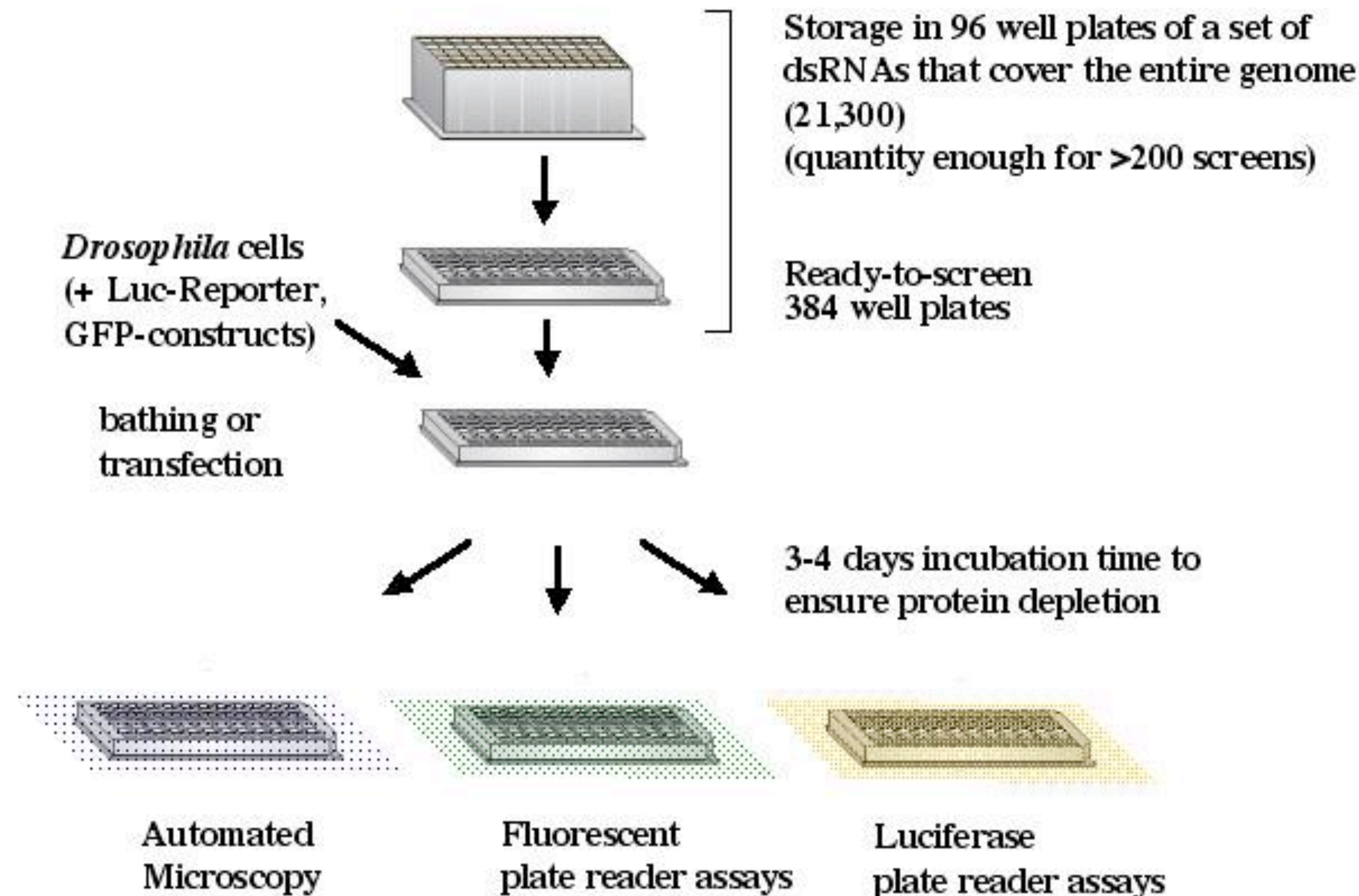
"hairpin" construct



hairpin double-stranded RNA gets cut up by Dicer to initiate RNAi in ALM neurons

Cells from other organisms (e.g., *Drosophila*) will undergo RNAi-mediated gene silencing if they are treated with dsRNA

RNAi by feeding or soaking has enabled many high-throughput (genome-wide) screens



Advantages of RNAi-based screens

Every known gene in the genome can be tested

There is no need to clone a gene that gives an interesting phenotype - you already know what it is!

Hypomorphic (reduction-of-function) phenotypes can be identified for essential genes, since RNAi gene silencing is often incomplete

This makes RNAi particularly useful to identify genetic ENHANCERS of a particular mutation, since hypomorphic alleles are frequently good enhancers