

Phenologs

A case study of using bioinformatics to find new genes for genetic traits

BCH394P/364C Systems Biology / Bioinformatics

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Are *you* a research parasite?



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EDITORIAL

Data Sharing

Dan L. Longo, M.D., and Jeffrey M. Drazen, M.D.
N Engl J Med 2016; 374:276-277 | January 21, 2016 | DOI: 10.1056/NEJMe1516564

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The aerial view of the concept of data sharing is beautiful. What could be better than having high-quality information carefully reexamined for the possibility that new nuggets of useful data are lying there, previously unseen? The potential for leveraging existing results for even more benefit pays appropriate increased tribute to the patients who put themselves at risk to generate the data. The moral imperative to honor their collective sacrifice is the trump card that takes this trick.

However, many of us who have actually conducted clinical research, managed clinical studies and data collection and analysis, and curated data sets have concerns about the details. The first concern is that someone not involved in the generation and collection of the data may not understand the choices made in defining the parameters. Special problems arise if data are to be combined from independent studies and considered comparable. How heterogeneous were the study populations? Were the eligibility criteria the same? Can it be assumed that the differences in study populations, data collection and analysis, and treatments, both protocol-specified and unspecified, can be ignored?

“The aerial view of the concept of data sharing is beautiful.”

[but!]

A ... concern ... is that a new class of research person will emerge...the system will be taken over by ...

“research parasites.”

<http://www.nejm.org/doi/full/10.1056/NEJMe1516564>

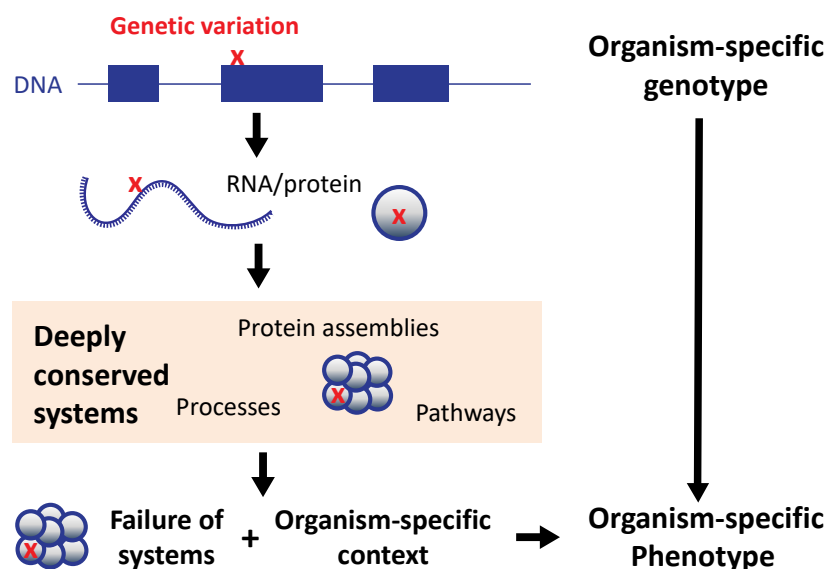
Let's think in the abstract for a moment:

How are mouse models useful for studying human disease? Are some models better than others?

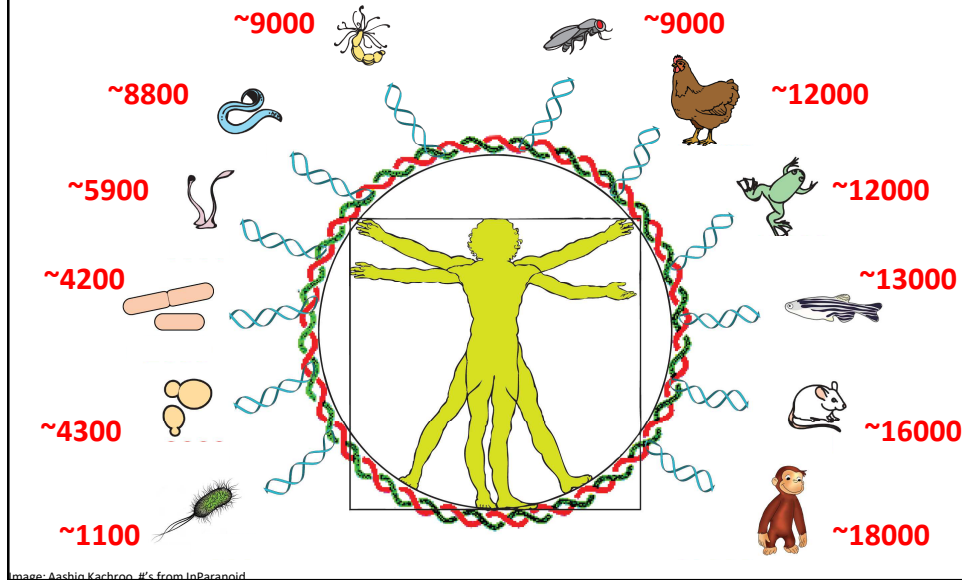
What's the worm equivalent of breast cancer?

**Are there plant versions of human diseases?
Why would they possibly be useful?**

Conserved systems intermediate between organism-specific genotype and phenotype



We share genes with almost every known organism



**All genetic traits
and diseases
affect molecular
structures that
are evolutionarily
conserved.**

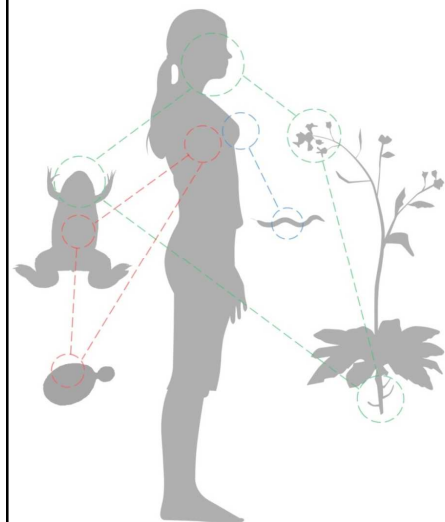
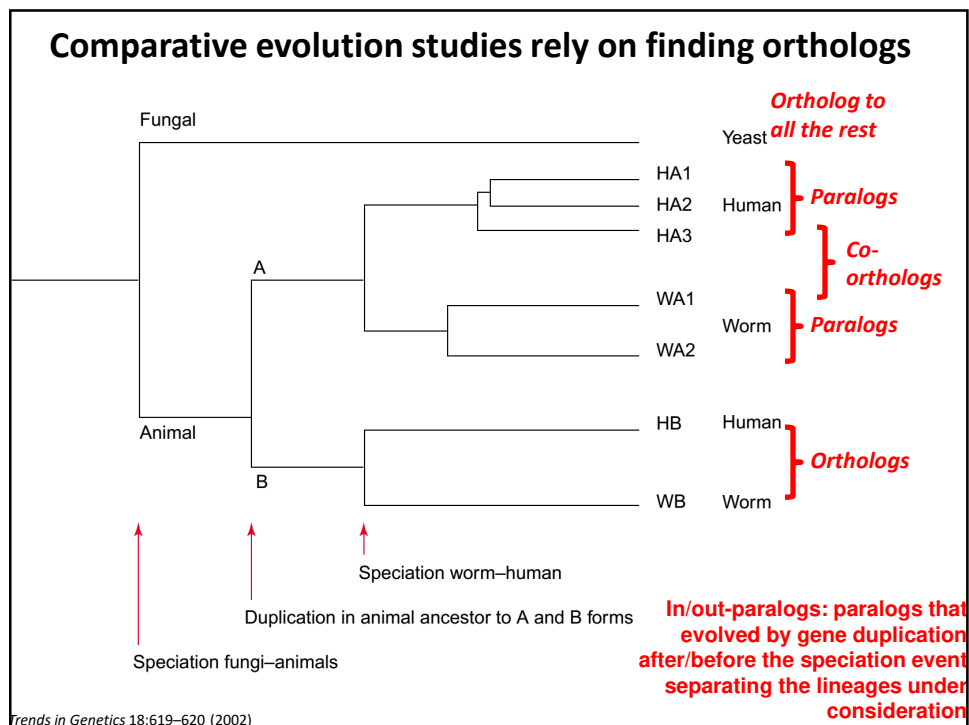


Illustration by Kathryn Weir

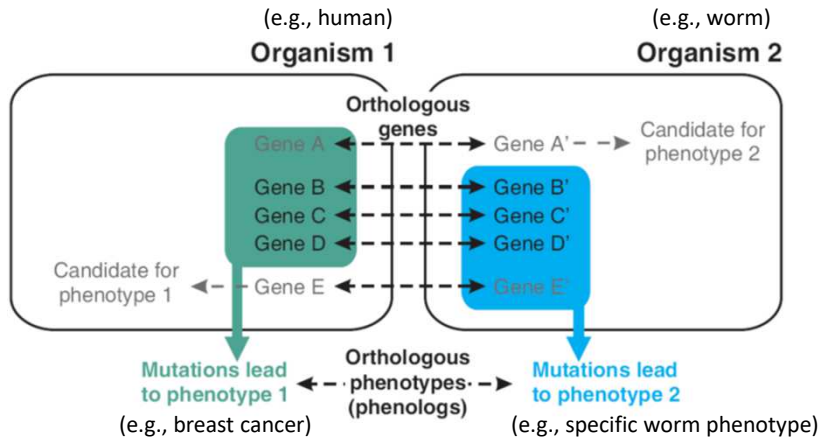
Comparative evolution studies rely on finding orthologs

Orthologs = genes from different species that derive from a single gene in the last common ancestor of the species

Paralogs = genes that derive from a single gene that was duplicated within a genome

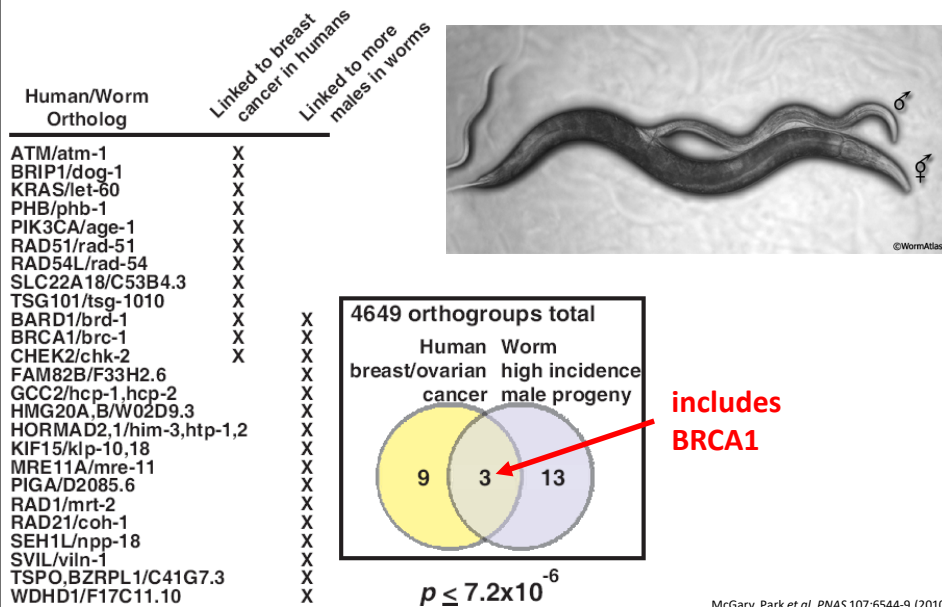


Phenologs = significantly overlapping sets of orthologous genes, such that each gene in a given set gives rise to the same phenotype in that organism



McGary, Park et al. PNAS 107:6544-9 (2010)

E.g., 'high incidence of male' *C. elegans* genes predict human breast/ovarian cancer genes



Building & searching a collection of phenotypes

Mining available databases +
manual collection from the primary literature



gene-phenotype
associations

<u>Organism</u>	<u># gene-phenotype associations</u>
human	1,923
mouse	74,250
worm	27,065
yeast	86,383
<i>Arabidopsis</i>	22,921

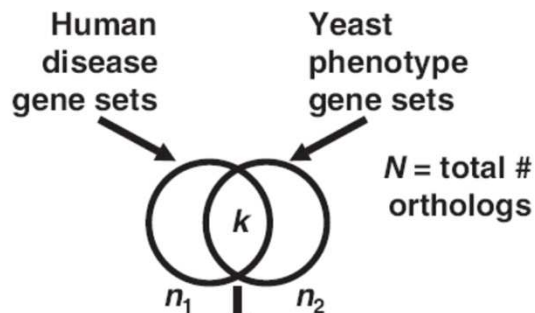
Spanning ~300 human diseases,
>7,000 model organism mutational phenotypes



Computational scan phenotypes for novel models of a disease of interest,
identify significant phenologs using permutation tests

McGary, Park et al. PNAS 107:6544-9 (2010)

Discovering phenologs



Measure p ($\text{overlap} \geq k \mid n_1, n_2, N$) for each
disease-phenotype pair,
considering only human-yeast orthologs



Identify all significant phenologs
by permutations or reciprocal best hits

McGary, Park et al. PNAS 107:6544-9 (2010)

**Computationally, we find many genes shared
between human diseases and
mouse, yeast, worm, and even plant traits**

McGary, Park *et al.* *PNAS* 107:6544-9 (2010)
Woods, Blom *et al.* *BMC Bioinformatics*, 14:203 (2013)



Michael Murphy, M.D.

**Waardenburg syndrome
accounts for ~2-5% of
cases of deafness**



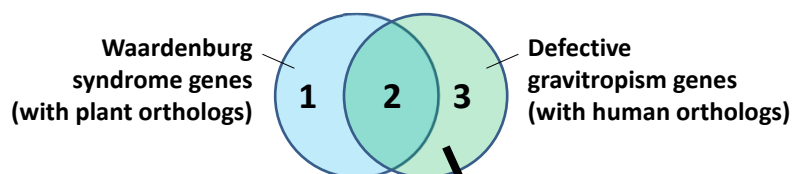
Associated websites: <http://www.verywell.com/waardenburg-syndrome-1048892>, <http://tebb.animalsandpeople.blogspot.com/>

Plants sense and respond to gravity → gravitropism



Fukaki et al., *The Plant Journal*
14, 425-430 (1998)

Plant gravitropism genes predict Waardenburg syndrome, a human congenital deafness syndrome



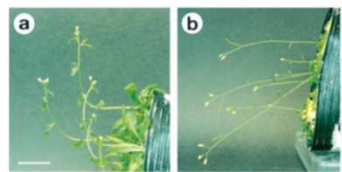
Michael Murphy, M.D.



Waardenburg syndrome

Human versions of these plant genes are candidate Waardenburg genes

≈



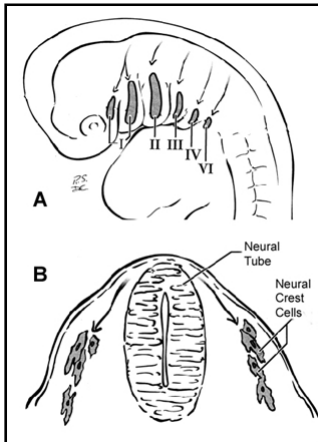
Gravitropism defects

Fukaki et al., *The Plant Journal*
14, 425-430 (1998)

McGary et al., *PNAS* 107:6544-9 (2010)

Waardenburg syndrome is a defect of neural crest cells

Neural crest cells migrate during embryonic development



Heike & Hing, *Gene Reviews* (2009)

Some WS correlates in other animals:
Deafness in Dalmatian dogs (22% unilaterally deaf)



www.petplanet.co.uk

Variations in the Blenheim spot
Cavalier King Charles Spaniels

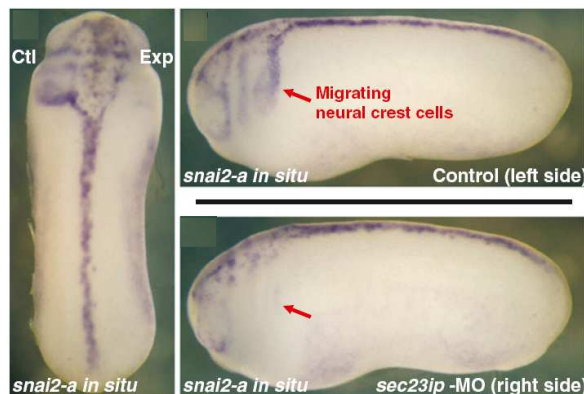
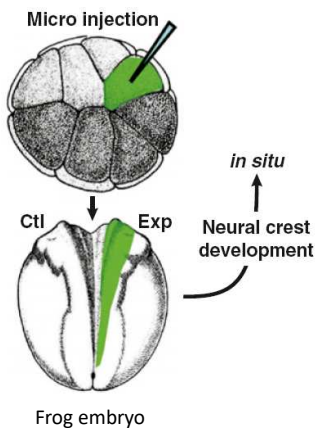


www.silvarcea.co.uk

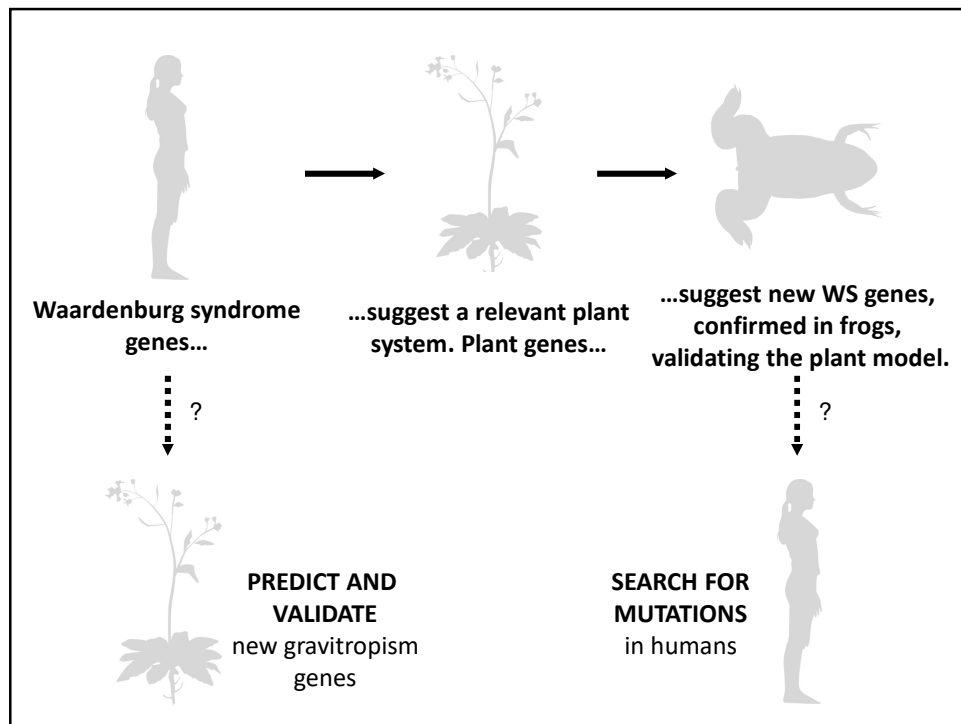
Association between white blue-eyed cats and deafness (noted by Darwin in 1859)

White forelock and deafness/bowel blockage in foals & many more...

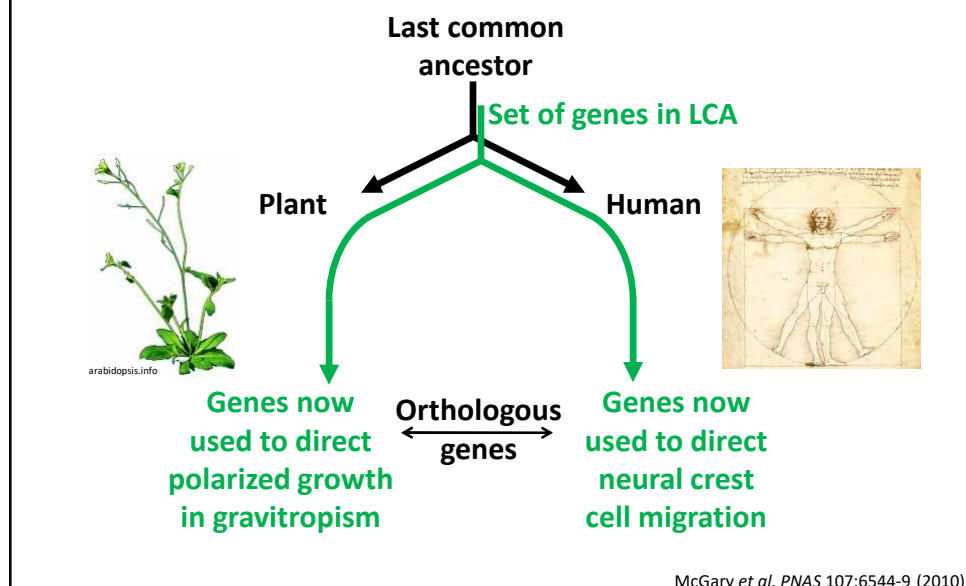
Sure enough, inactivating one of the genes—predicted from plants—in a tadpole disrupts neural crest cells, consistent with Waardenburg syndrome



McGary, Park et al. *PNAS* 107:6544-9 (2010)



Conservation and “repurposing” of core cell machinery across deep evolutionary time



Let's talk about how such projects play out in practice.

**How are discoveries made?
How do you computationally explore ideas?**

Let's step through this particular discovery process:

- 1. We had an idea, based on a puzzling observation:**

Why do mutations in worm retinoblastoma genes induce ectopic vulva while a mutation in the human ortholog causes eye cancer?

We weren't interested in specific mechanism here, but rather the impact of organismal context on conserved systems. In particular, *how do ever-more distant evolutionary models inform us about human disease?*

Let's step through this particular discovery process:

- 2. We thought about how this might be part of a large trend—does it illustrate a general principle? Could we look for new cases systematically?**
- 3. We thought about other examples, mentally assembling what could serve as positive and negative control cases. i.e. how to we decide if a systematic approach is working?**

Let's step through this particular discovery process:

- 4. A grad student (Kris McGary) started assembling relevant datasets. We took heavy advantage of existing resources: model organism databases that had already painstakingly curated relevant data, large-scale screens reporting easy-to-process data.**

Let's step through this particular discovery process:

5. We started inventing/evaluating statistical models/algorithms, exploring the data and thinking about how to search for the relevant trends. We iterated these steps until we thought we understood the problem better.

6. At some point, the lab bet a 6 pack of beer on the outcome:

Can we discover plant models of human disease?

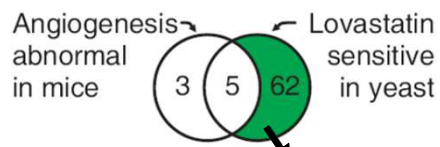
Let's step through this particular discovery process:

7. The algorithms predicted some remarkable and crazy results. We had no option but to test or reject the new predictions, so began testing, thanks to collaborators in the Wallingford lab willing to sink a few weeks into high-risk experiments.

8. Some tests worked, some didn't. We went back & thought about the ones that didn't and refined how we prioritized the results.

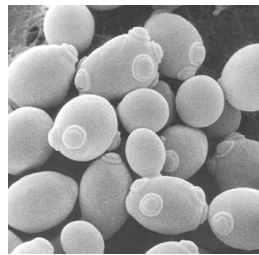
9. Iterate, iterate. Jackpot! A plant model of deafness! Shouting in the halls...

Last example: Yeast genes linked to statin sensitivity predict blood vessel defects

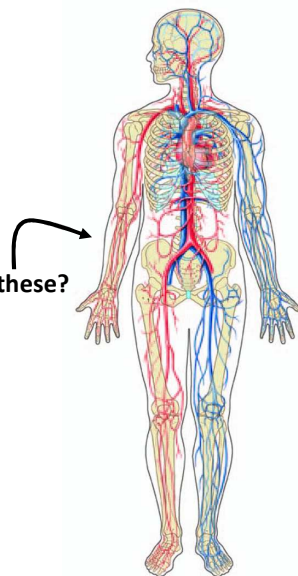


The human versions of these yeast genes are candidate angiogenesis genes

Can these really tell us about these?



www.chemistryland.com



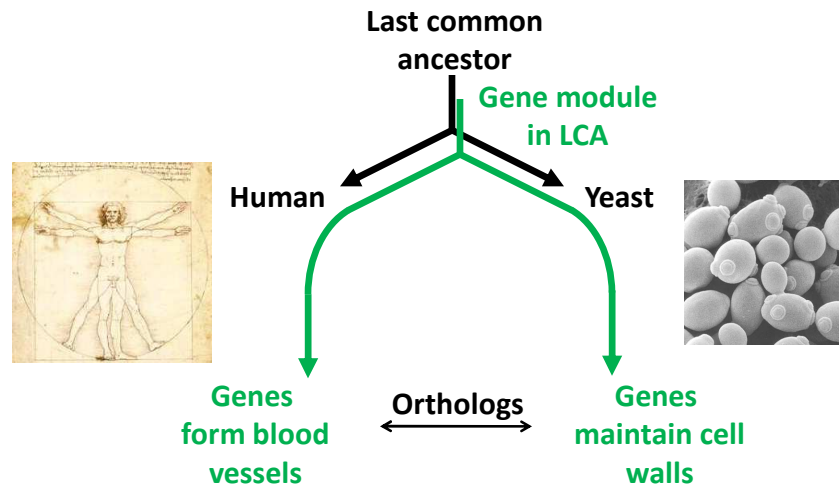
McGary, Park et al.
Dorling Kindersley PNAS 107:6544-9 (2010)

Disrupting the SOX13 gene causes strong blood vessel defects



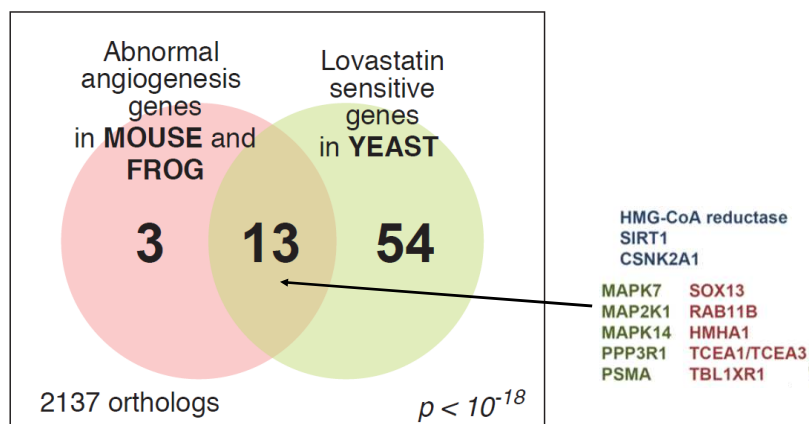
McGary, Park et al.
PNAS 107:6544-9 (2010)

A yeast model of angiogenesis = example of a deeply conserved, but “repurposed” gene module

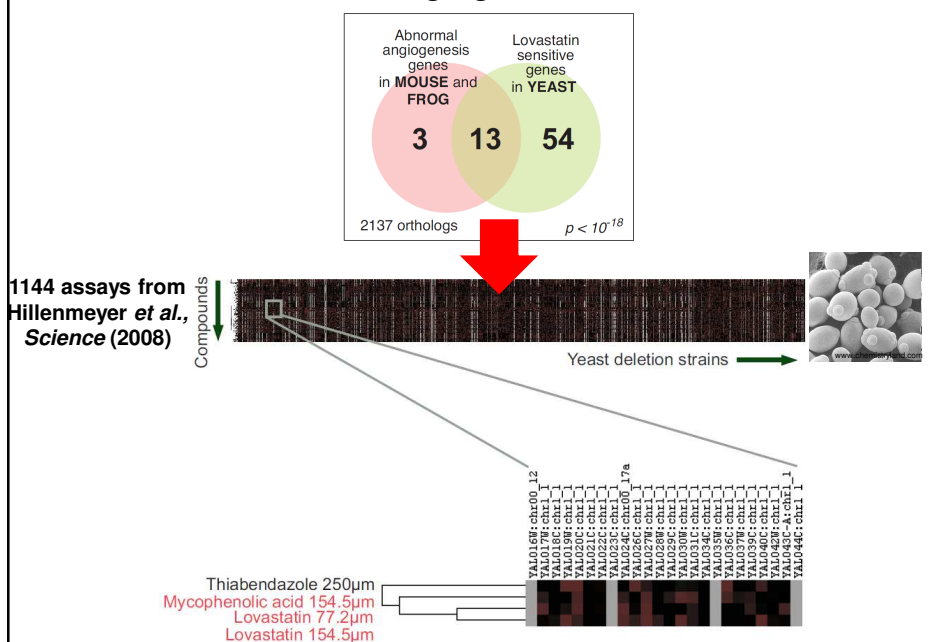


McGary, Park *et al.* PNAS (2010)

The yeast/angiogenesis gene module



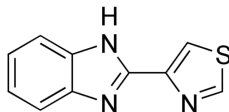
Chemicals that interact genetically with this module are candidate angiogenesis inhibitors



Screening for drugs that interact genetically with this yeast module led us to identify a new angiogenesis inhibitor

TBZ = thiabendazole

FDA-approved antifungal drug with 40 years of safety data



- Approved by U.S. Food and Drug Administration in 1967

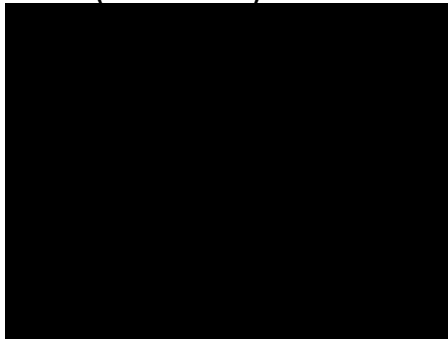
- Fungicide and parasiticide
- Not mutagenic or carcinogenic; 2 year dog safety trials
- Off-patent, marketed as a generic

Imaging the blood vessels of a living, transgenic tadpole in a dish of water

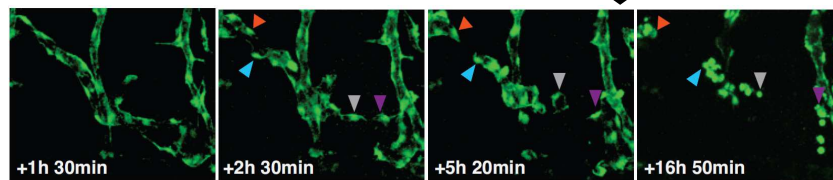
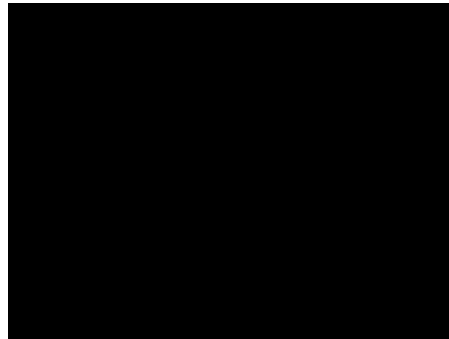


TBZ disrupts vascular integrity, making vascular endothelial cells retract & round up

Control (DMSO carrier)

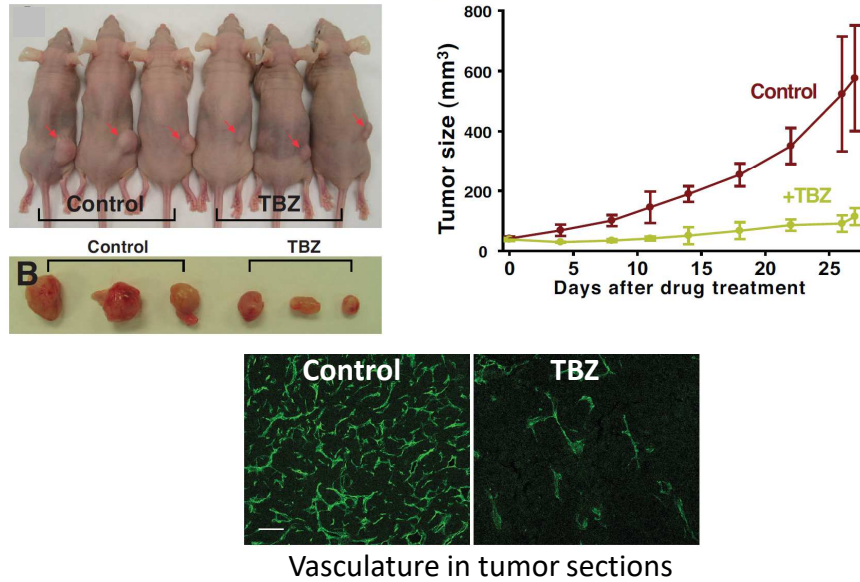


+ TBZ



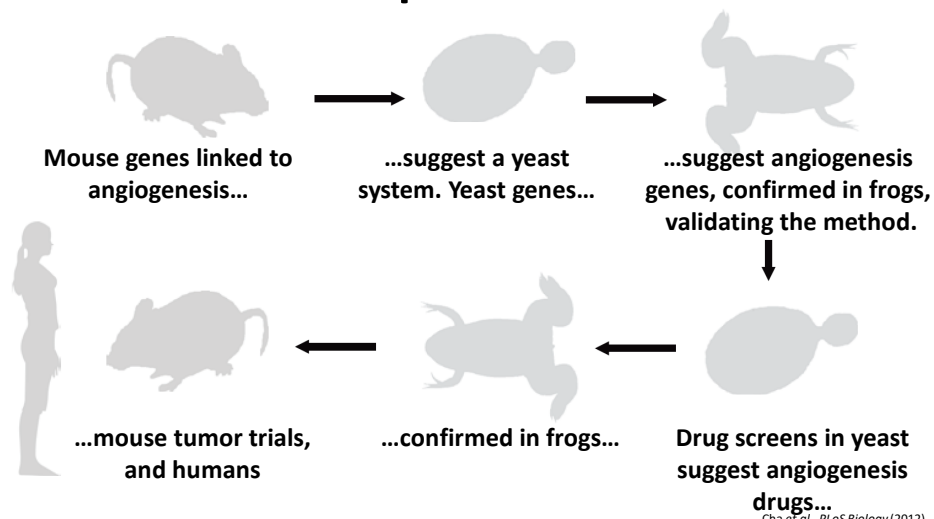
Cha et al., *PLoS Biology* (2012)

TBZ slows human fibrosarcoma tumors transplanted into immune-compromised mice



Cha et al., *PLoS Biology* (2012)

“Road map” to a new vascular disrupting agent, by mapping phenotypes across species



Cha et al., *PLoS Biology* (2012)

Try it out yourself!
<http://www.phenologs.org>

You can start by rediscovering the plant model of
Waardenburg syndrome:

Search known diseases for "Waardenburg", or enter the
human genes linked to Waardenburg (Entrez gene IDs 4286,
5077, 6591, 7299) to start.

Tools for finding orthologs are linked on the class website