

Discovering the Mysteries of Pha-4

In the Mango lab, we study organogenesis in the pharynx of the *Caenorhabditis elegans*. The *pha-4* gene specifies pharynx organogenesis during *C.elegans* embryogenesis.

- if there is no *pha-4* present then there will be no pharynx.
- if there is excess *pha-4*, then excess pharynx is produced.

The goal of my research is to find out how *pha-4* works and to identify its regulators and partners. We know that there must be other factors that work with *pha-4* to regulate pharyngeal growth because *pha-4* is expressed in all the pharyngeal cells. But there are 5 cell types, so there must be other factors contributing to differentiation among cells. We also know very little about how *pha-4* itself is controlled and there may be regulators of *pha-4* that are important for proper pharynx development.

We performed a suppressor screen for partners, regulators of *pha-4*. The screen introduces a series of random mutations in the worms using a chemical mutagen. We searched for mutations that allowed for the *smg-1(ts);q500* strain to survive at 20°C.

We predicted that we would isolate 5 classes of suppressors:

1. Regulators of *pha-4* (these

2. Partners of *pha-4* (these work with *pha-4* to turn other genes on or off.) c.. *pha-4* itself (e.g. making the *q500* protein more stable or more active and therefore able to function at 20°C.) d.. *smg*-mutations (we are not interested in these because the effect is global and has nothing directly to do with *pha-4*.) 5. Down stream genes involved with pharynx development. For example *ceh-22* or *myo-2* target genes that *pha-4* and partners control.

Ultimately, we hope to be able to map and eventually clone the suppressor mutation to identify factors that act on or with *pha-4* in pharynx development.



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