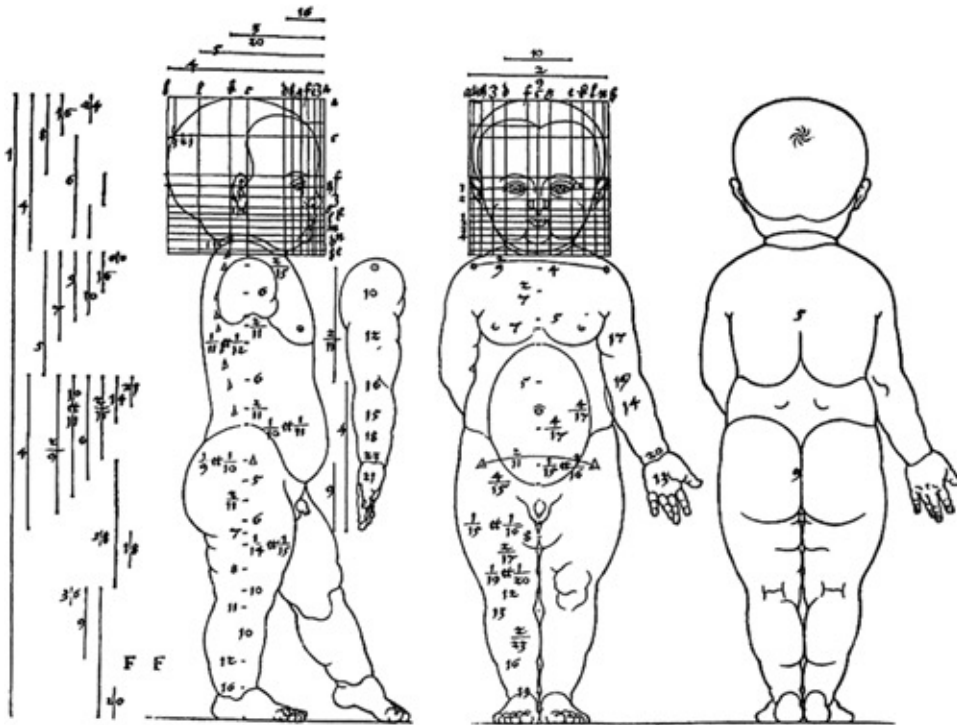


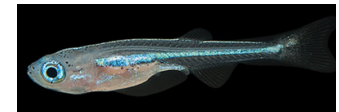
# Breaking symmetry: embryonic establishment of body axes (I)

Sept. 11, 2024

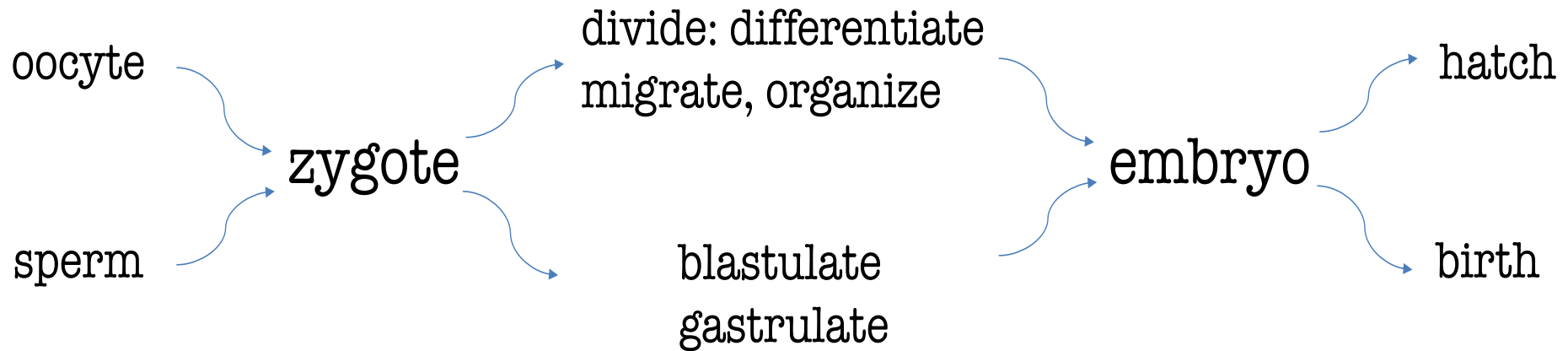
# Three body axes define asymmetries



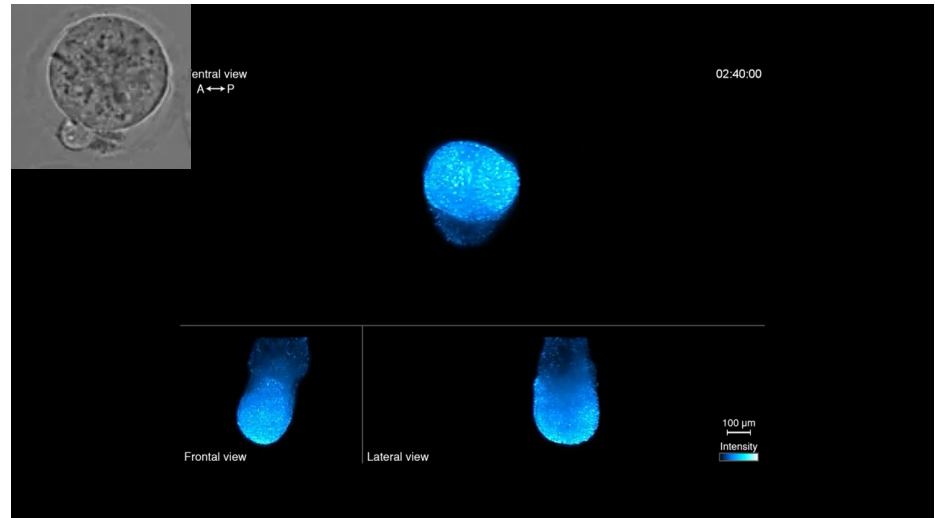
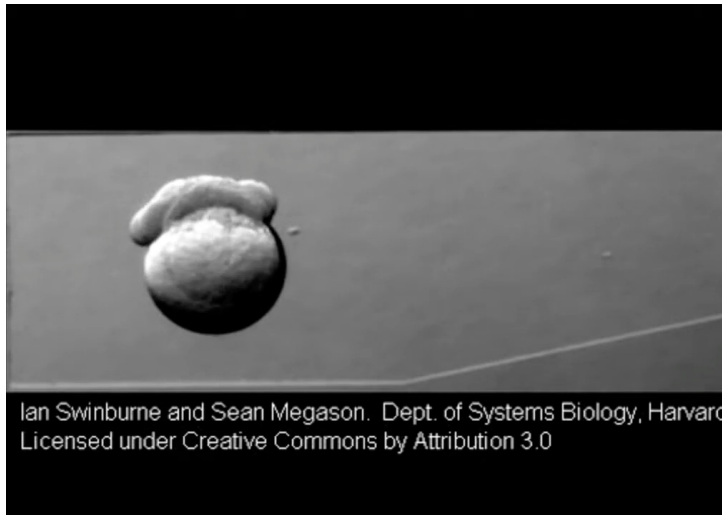
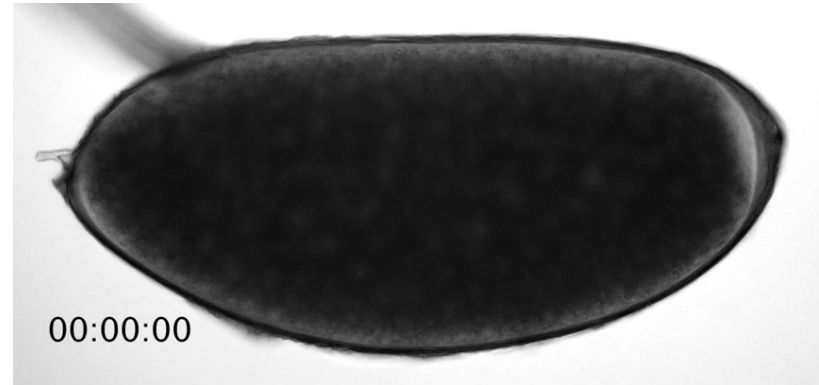
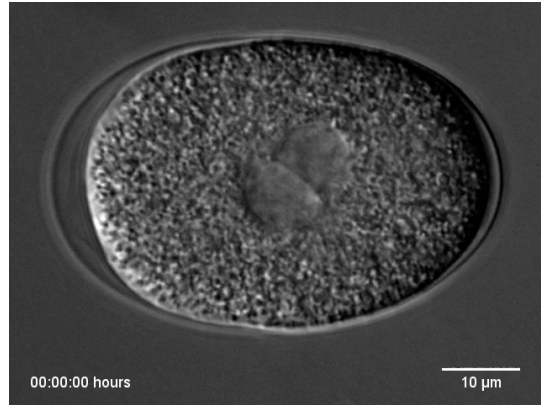
sketch: Albrecht Durer



# When do you start?



# Some zygotes break symmetry earlier than others



# Does the early polarity relate to final body planes?

- How might we study this question?
- How is early polarity established?

What to ask, while we learn a bit about it? e.g.

Is there an order to three axial development?

How similar or different is it among animals?

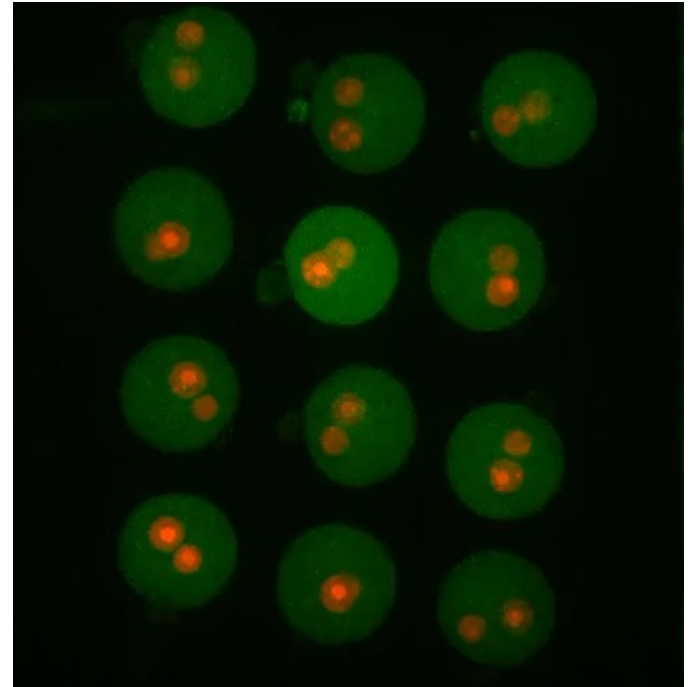
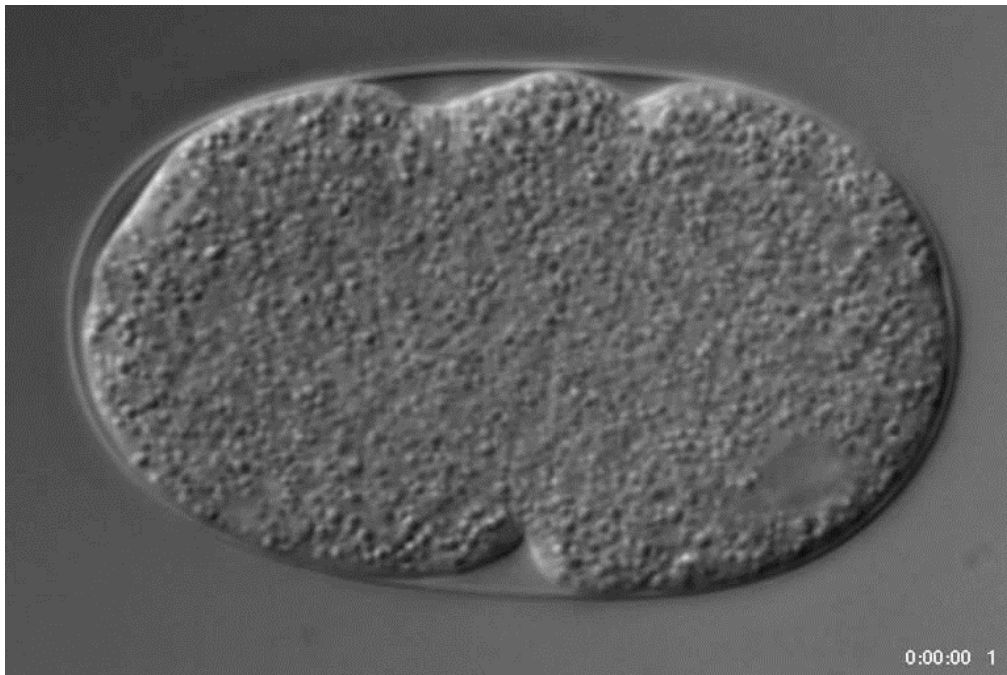
What drives the similarity and difference?

Why asymmetry at all?

# How might we study this question?

- Watch and describe processes

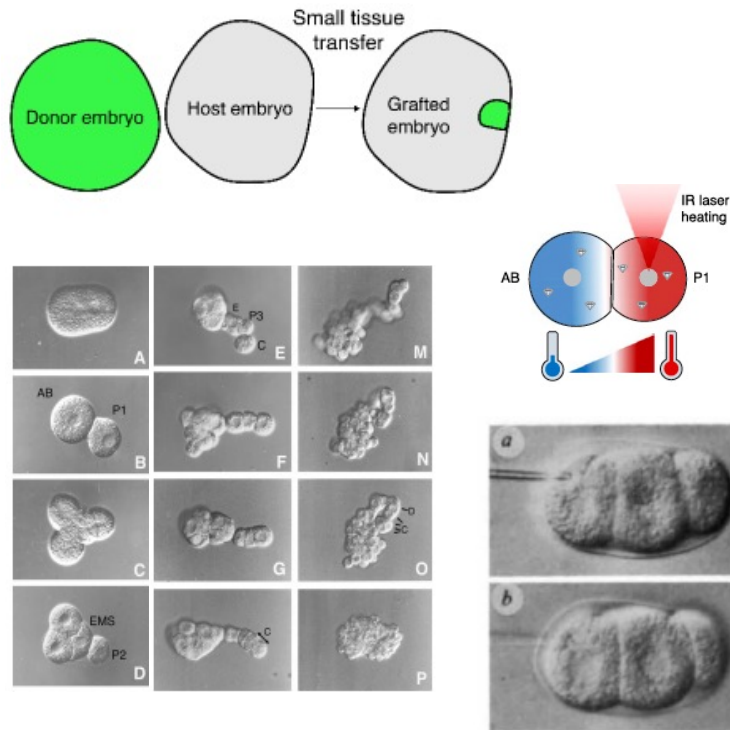
live imaging



# How might we study this question?

- Perturb and sort effects

## cell-manipulation



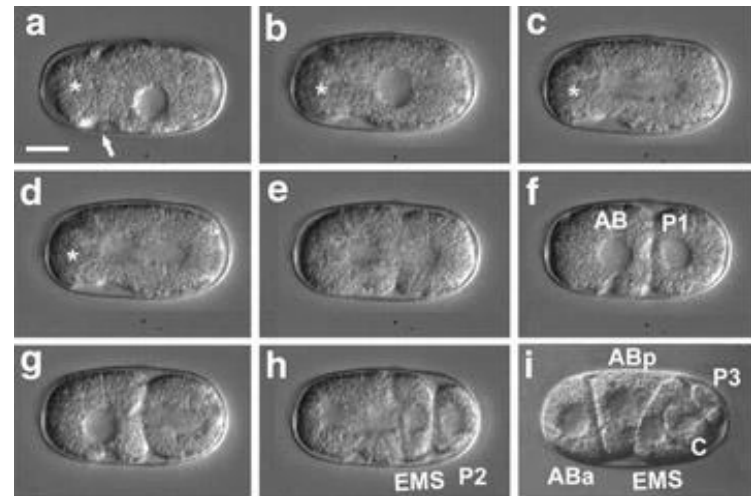
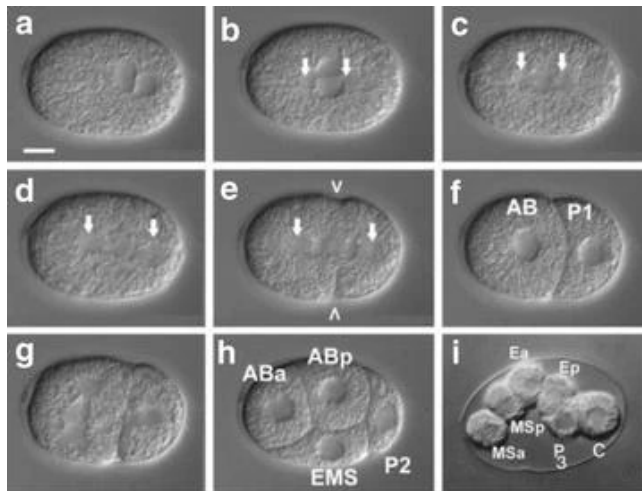
## gene-manipulation

- Mutagenesis  
(forward genetic screens)
- Mutagenesis  
(reverse genetic or  
'genome-wide' screens)

# How might we study this question?

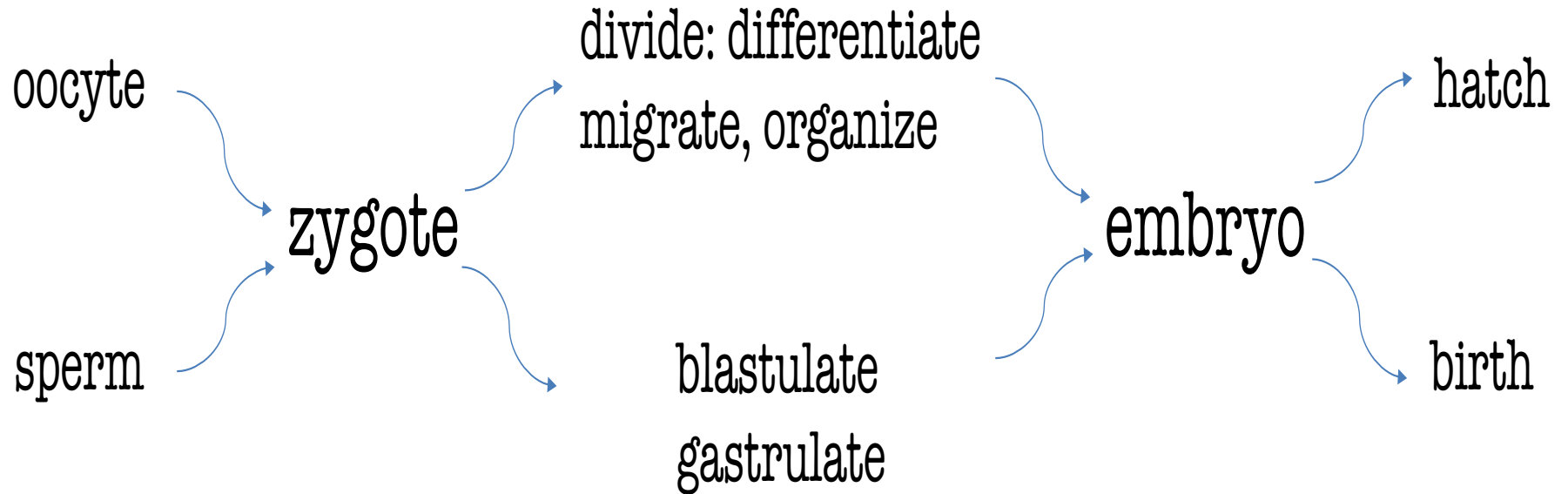
- Compare among sub-species

Correlate genomic-cellular variations





# When do you start?



How early is 'early'?

*C. elegans* is quite early:

sperm entry + first 3 rounds of divisions

5 asymmetric divisions at 3 planes:

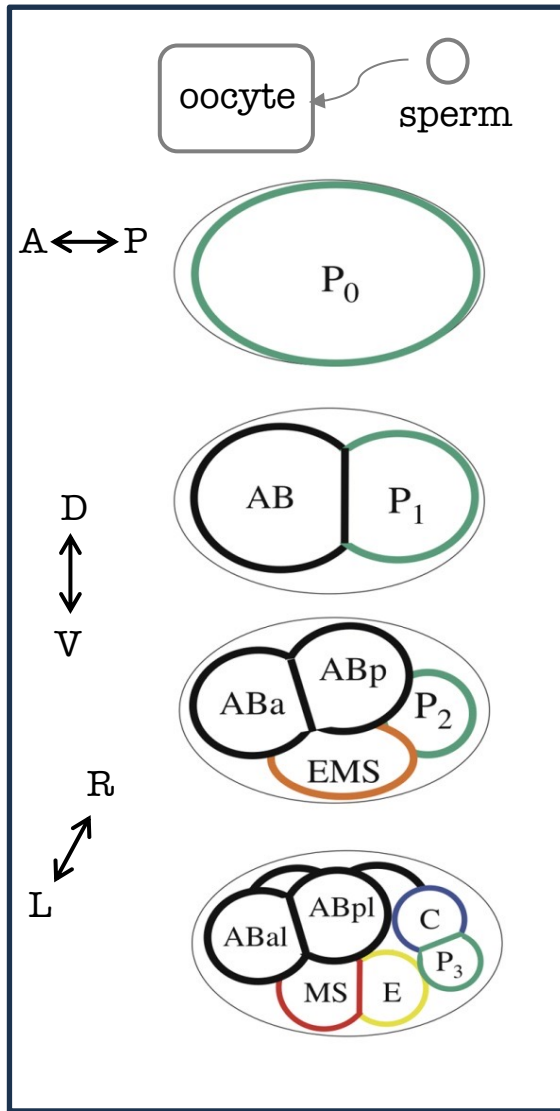
rough alignment with the body axes

6 founder cells (non-exchangeable fate):

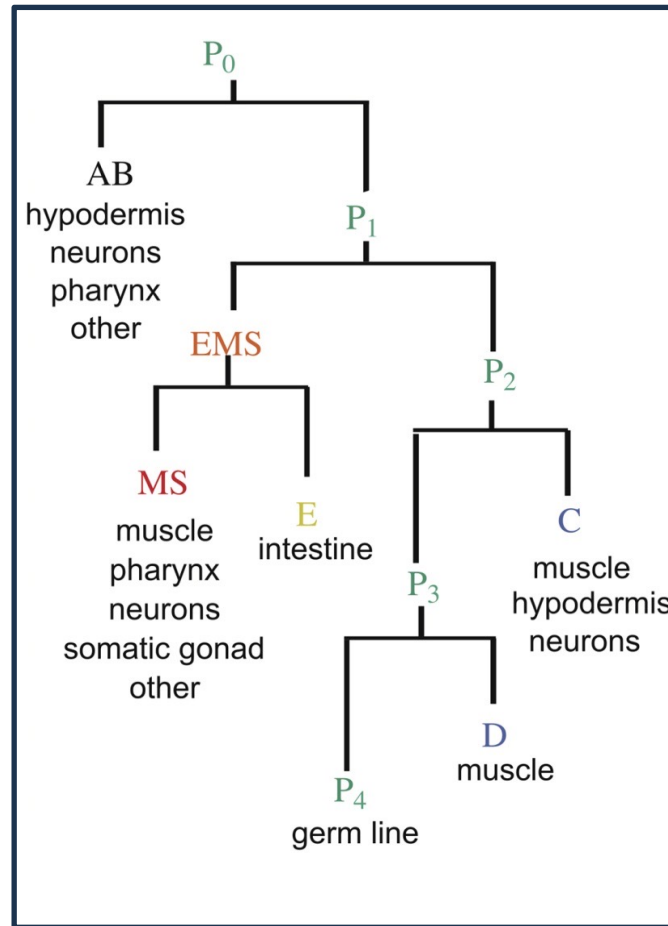
induction-determination

Gastrulation: the final body plane layout

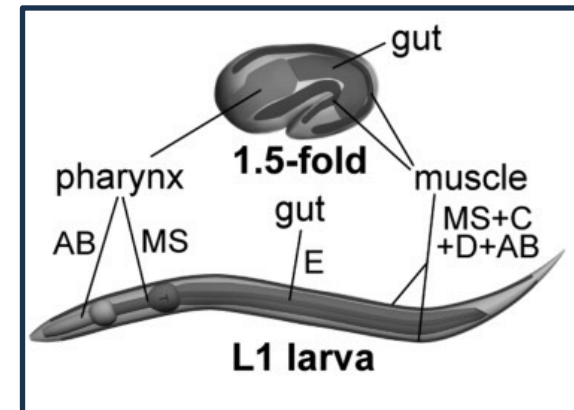
# 5 asymmetric divisions



## 6 founder cells



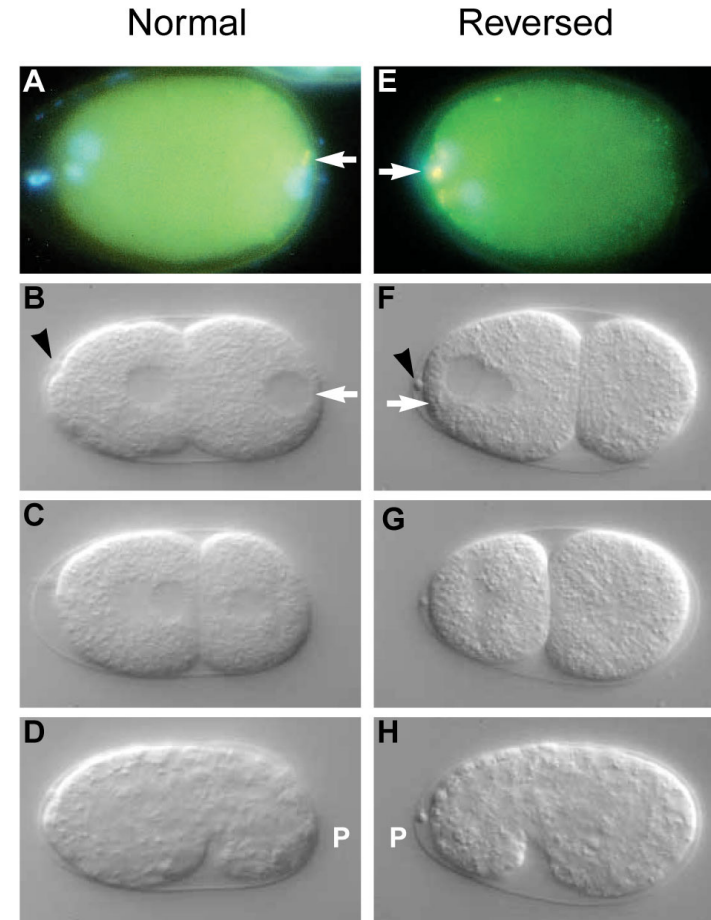
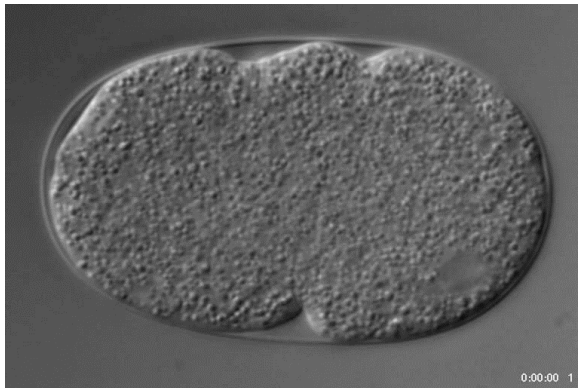
## gastrulate & unfold



# Let's review key evidences

## 1) A-P axis:

Sperm polarizes zygote, the  
and A-P asymmetry in the  
first division.



**Table 1** Maternal Loci in *C. elegans*: Gene Names and Molecular Identities (See Text for References)

Gene	Name	Molecular identity
Par Group Genes		
<i>let-99</i>	<i>Lethal</i>	?
<i>par-1</i>	<i>Partitioning-defective</i>	Ser-Thr kinase; binds a nonmuscle myosin
<i>par-2</i>	Same	Novel; ATP-binding site
<i>par-3</i>	Same	Novel; two PDZ domains
<i>par-4</i>	Same	Ser-Thr kinase
<i>par-5</i>	Same	?
<i>par-6</i>	Same	?
<i>mes-1</i>	<i>Maternal-effect sterile</i>	?
Blastomere Identify Group Genes		
P <sub>1</sub> subgroup		
<i>pal-1</i>	<i>Posterior alae defective</i>	Homeodomain protein; putative transcription factor
<i>pie-1</i>	<i>Pharynx and intestine excess</i>	TIS-11-like Zn <sup>2+</sup> finger ptn
<i>skn-1</i>	<i>Skin excess</i>	bZIP-like putative transcription factor; lacks a leucine zipper
<i>pop-1</i>	<i>Posterior pharynx defective</i>	HMG domain protein; putative transcription factor
<i>mom-1</i>	<i>More mesoderm</i>	Porcupine homologue; ER protein required for Wnt secretion
<i>mom-2</i>	Same	Wingless/Wnt homologue; putative secreted glycoprotein ligand
<i>mom-3</i>	Same	?
<i>mom-4</i>	Same	?
<i>mom-5</i>	Same	Frizzled homologue; putative receptor for Wnt ligands
AB subgroup		
<i>aph-2</i>	<i>Anterior pharynx defective</i>	Novel membrane-associated extracellular protein
<i>apx-1</i>	<i>Anterior pharynx excess</i>	Delta-like transmembrane protein; putative GLP-1 ligand
<i>glp-1</i>	<i>Germline proliferation defective</i>	Notchlike transmembrane protein; putative receptor
Intermediate Group Genes		
<i>mex-1</i>	<i>Muscle excess</i>	TIS-11-like Zn <sup>2+</sup> finger ptn
<i>mex-3</i>	Same	Two KH domains; putative RNA-binding protein
<i>pos-1</i>	<i>Posterior localized mRNA</i>	TIS-11-like Zn <sup>2+</sup> finger ptn

# What might be the sperm's polarity cue?

Its pronucleus or DNA?      **No!**

embryos show normal polarity when oocytes are fertilized by anucleate sperm (Sadler and Shakes, 2000. Development 127: 355-366.)

Its centrosomes?      **Yes!**

embryo fails to polarize when its centrosome was destroyed by a laser (Cowan and Hyman, 2004. Nature 431: 92-96.)

**Requirement for initiating embryo polarity, from the centrosome:**

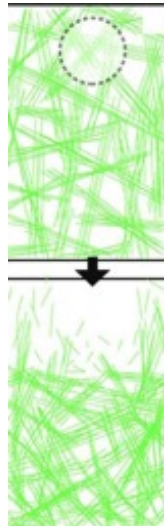
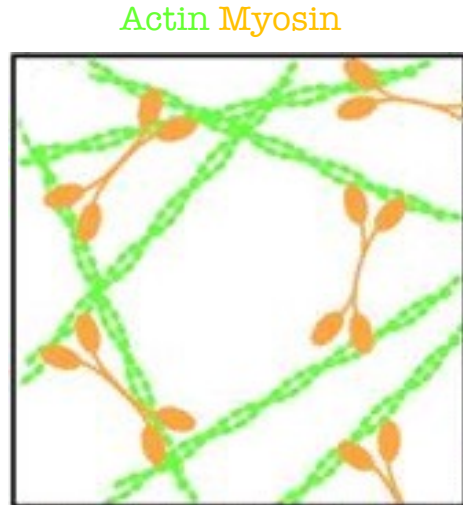
its microtubule extension to cortex

its close association with cortex

its component: e.g. Aurora kinase AIR-1

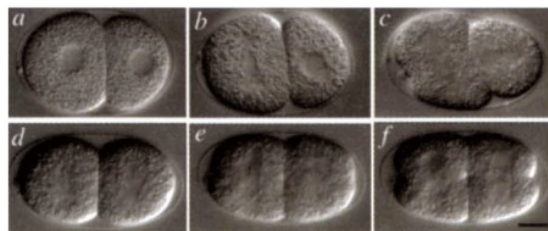
# How might the centrosome polarize the zygote?

*asymmetric cortical actomyosin contraction*

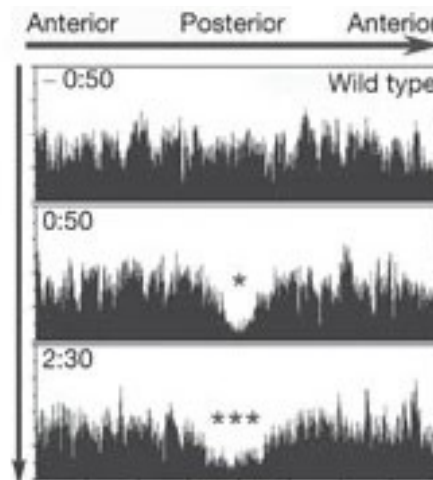


- Cytochalasin D blocks all division asymmetry
- Removing non-muscle myosin blocks polarity
- Posterior cortex RhoGEF cortex is excluded, which requires centrosome proteins

wildtype

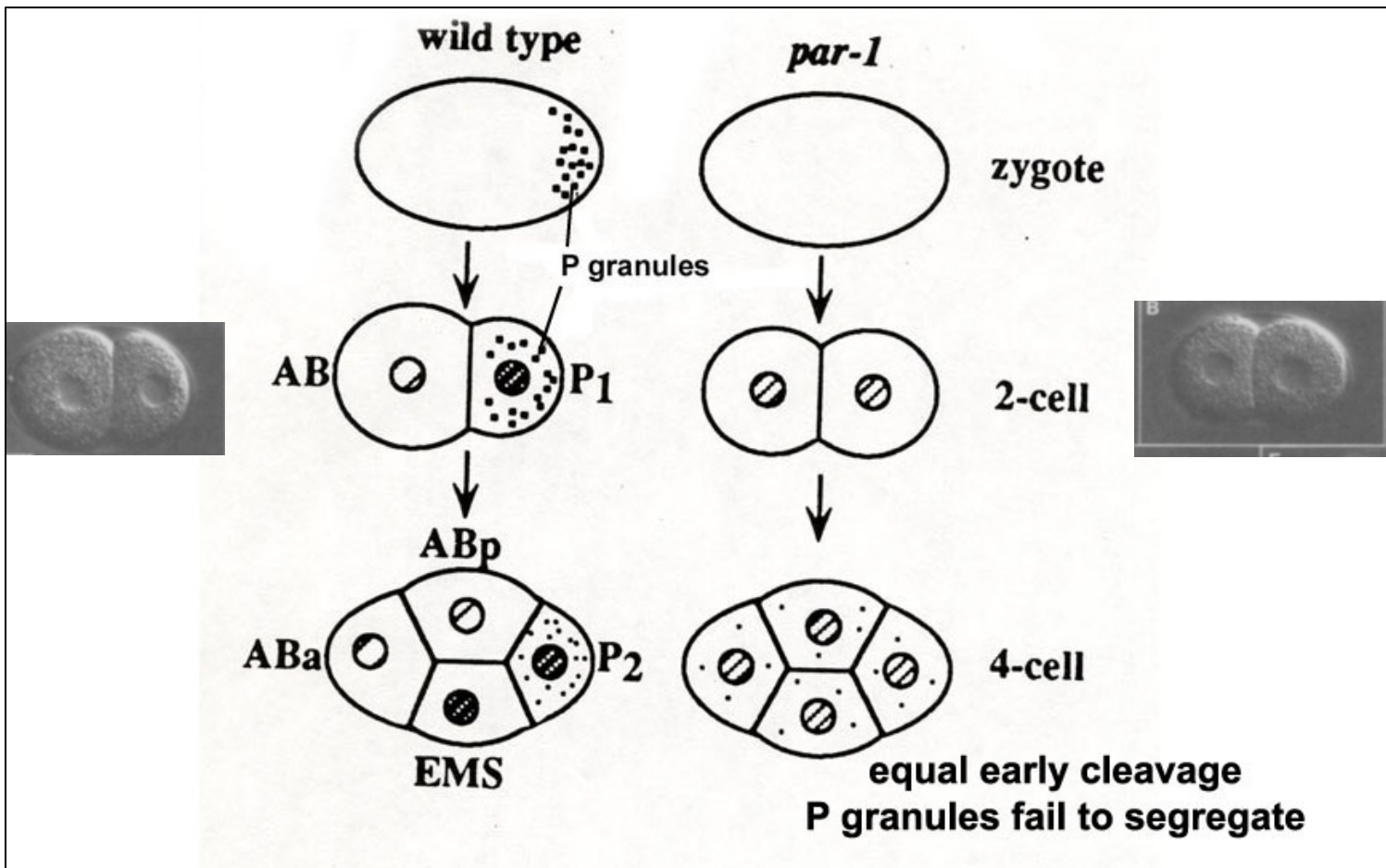


nmy-2  
RNAi



# How might cortical contraction polarize the zygote?

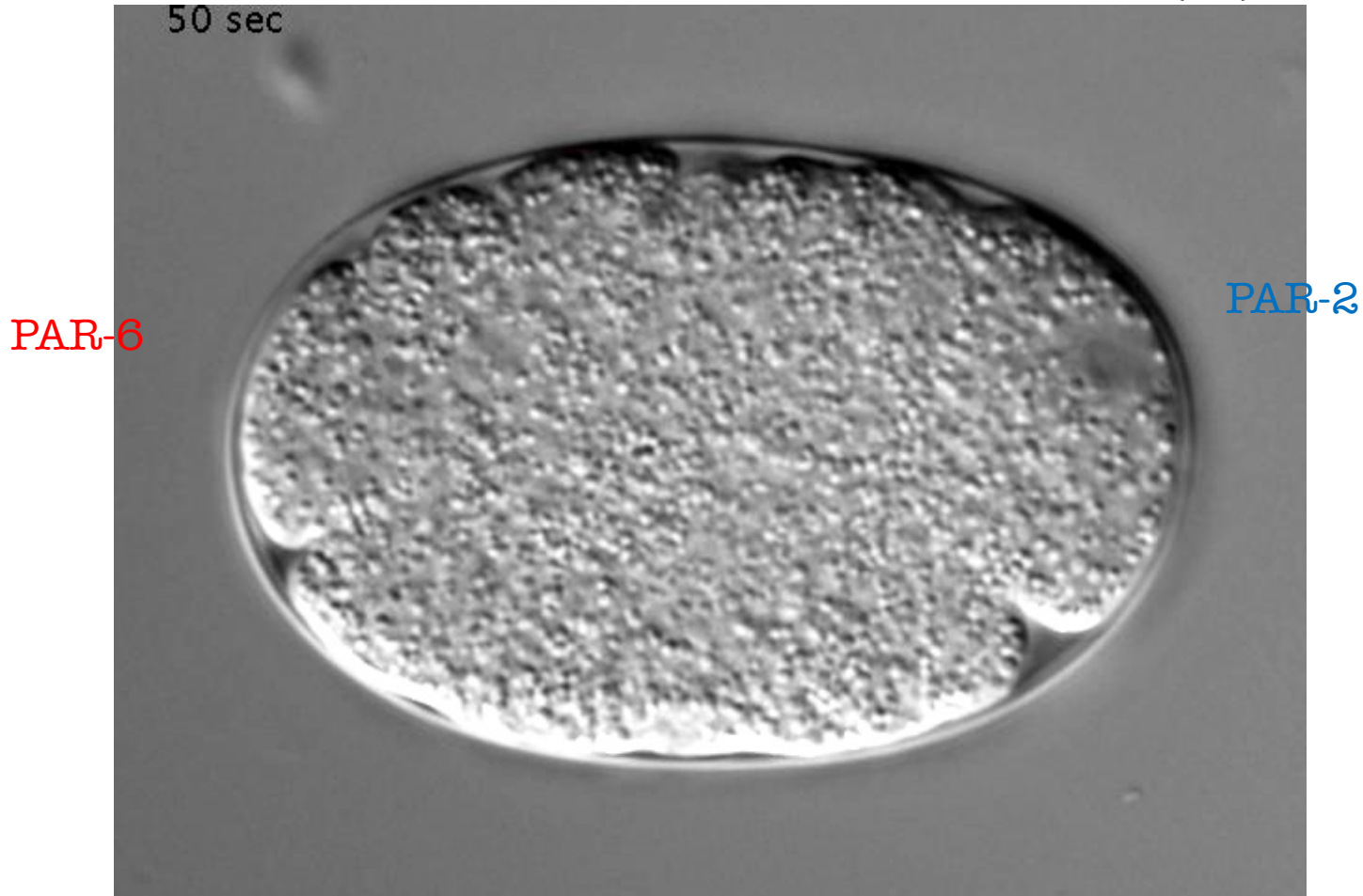
*polarized cortical par location (I)*





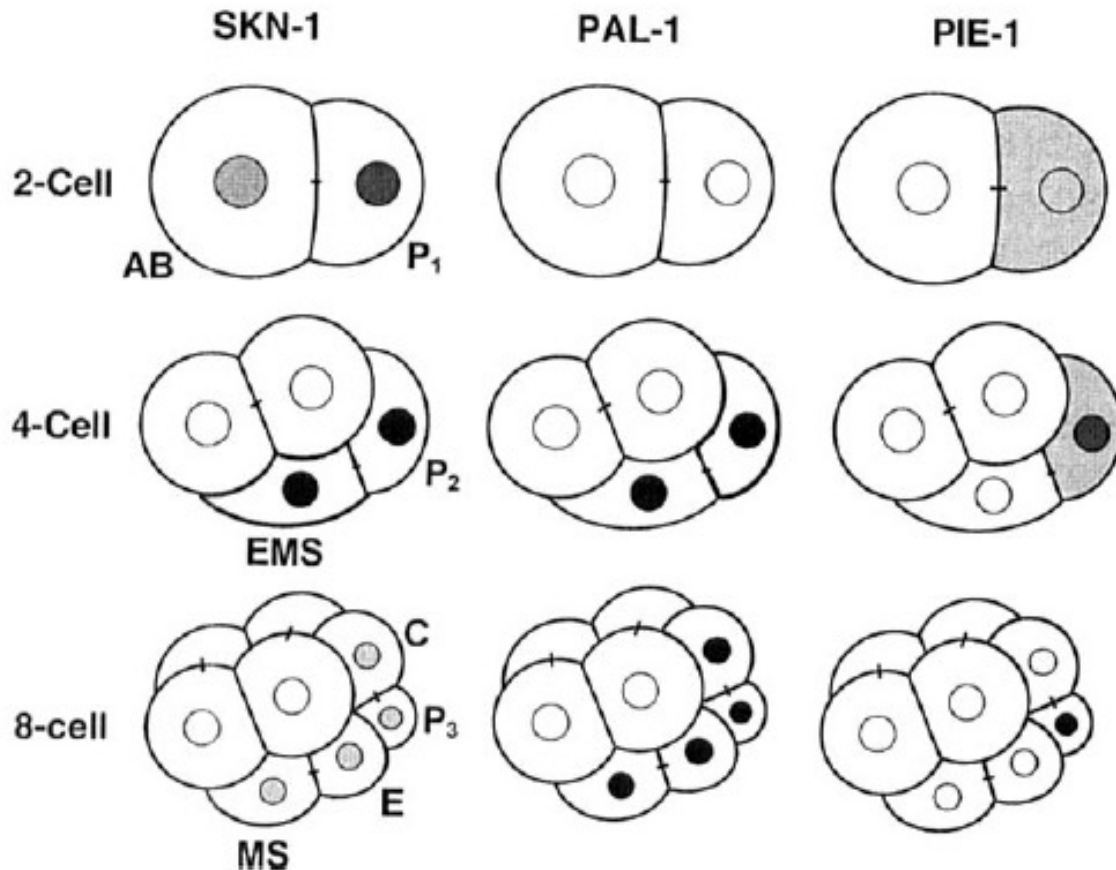
# How might cortical contraction polarize the zygote?

*polarized cortical par location (II)*



# How might the blastomere cell fate be specified?

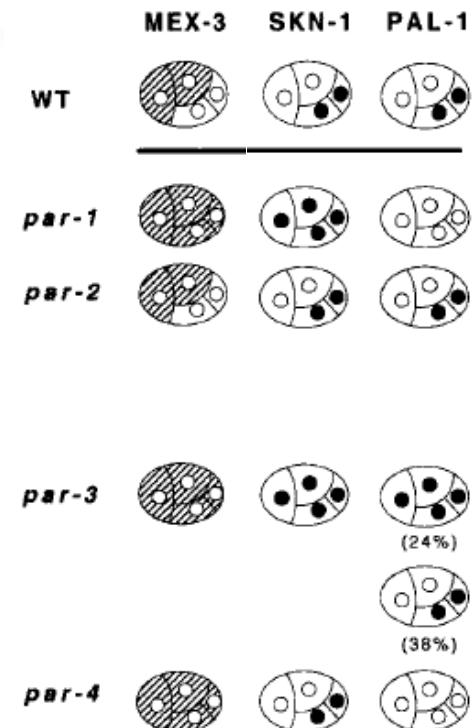
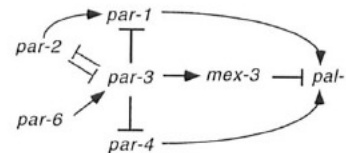
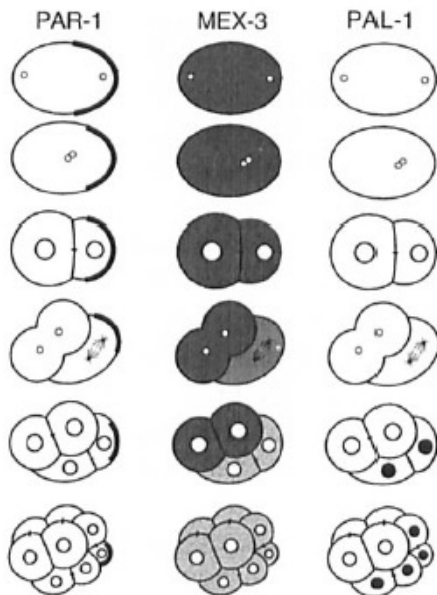
*founder cell transcription factors*



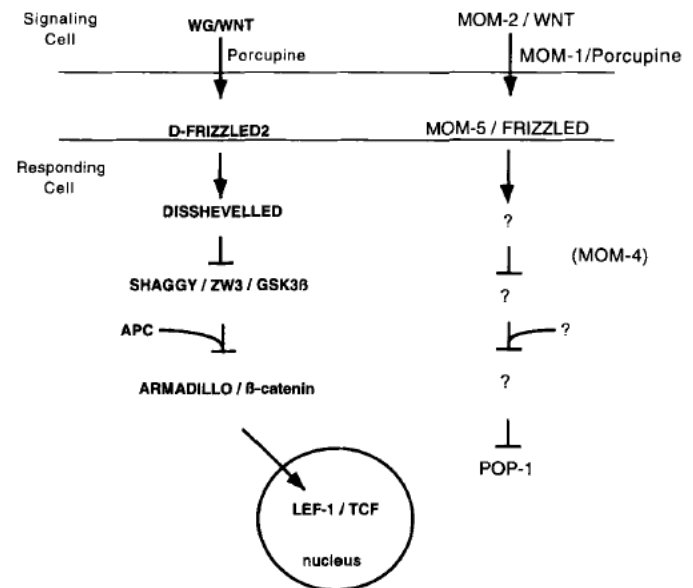
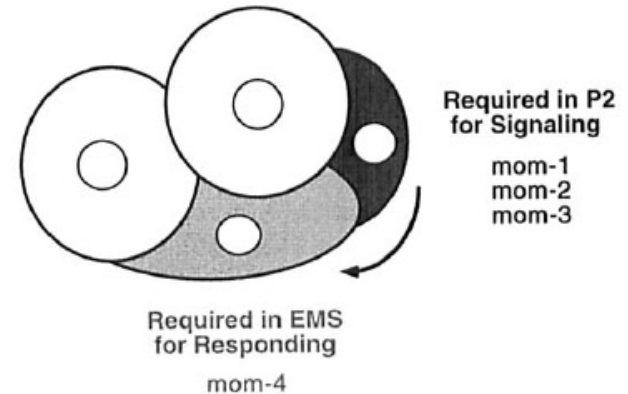
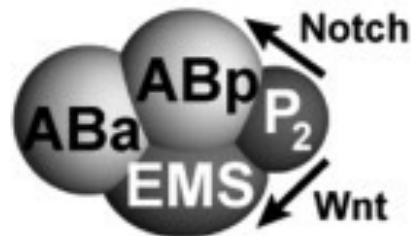
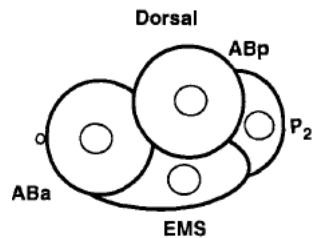
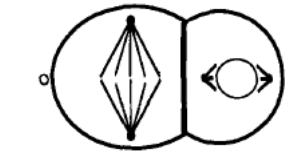
# How might the blastomere cell fate be specified?

*founder cell transcription factors*

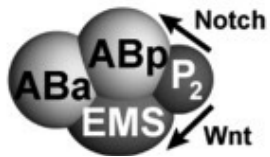
$par-1 \rightarrow mex-3 \rightarrow pal-1$



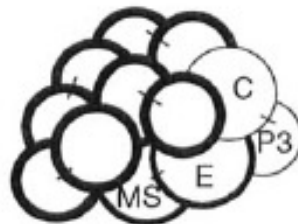
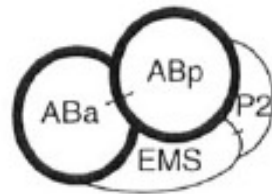
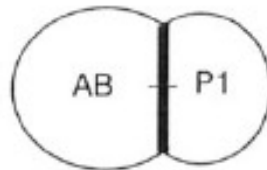
# The second round of divisions: D-V axis and fate induction



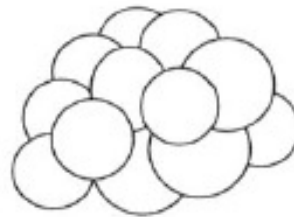
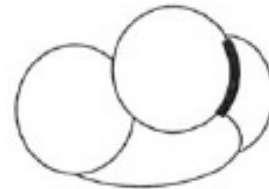
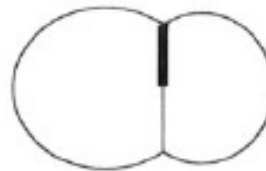
# The second round of divisions: DV axis and fate induction



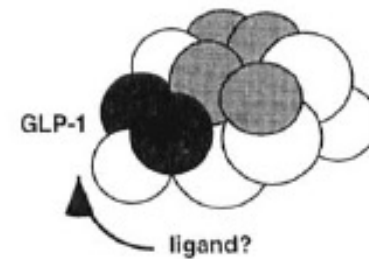
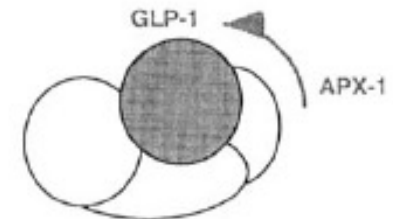
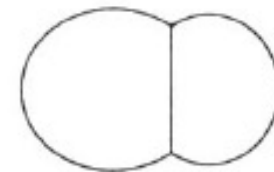
GLP-1



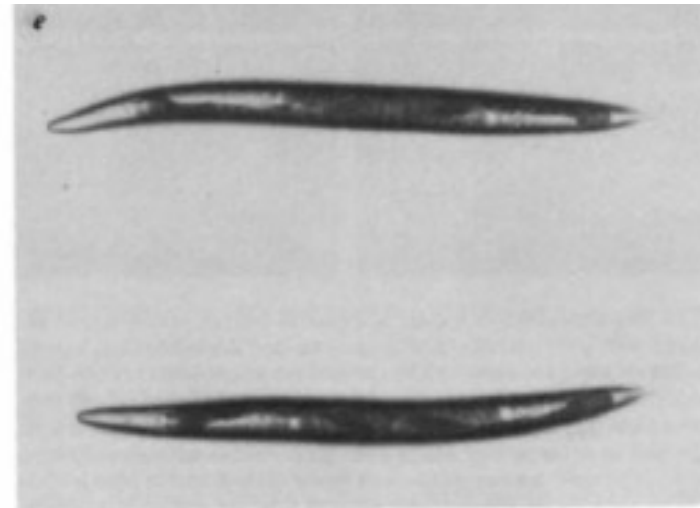
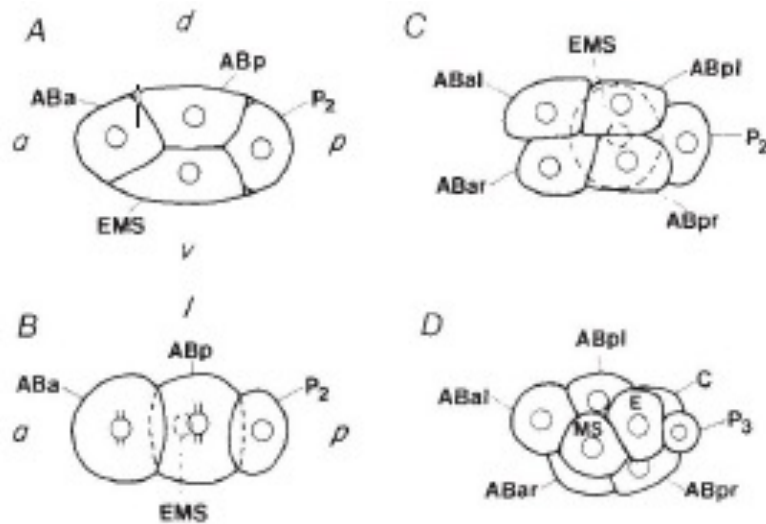
APX-1



Two Signals



# The third-round division: L-R axis



# A brief recap:

*C. elegans* axis patterning starts early: sperm entry

The first 3 rounds of zygote divisions generate 6 founder cells with asymmetry in the division plane, size or morphology, and unique fate.

The establishment of A-P axis is followed by D-V, and by (or in parallel) L-R

# A brief recap:

## Steps and purposes of blastomere patterning:

- Initiate polarized actomyosin contraction
- Partition anterior-posterior PAR complex
- Establish intrinsic or induced founder cell's transcription factors

Cell fate maps: roughly aligned with future body planes

Final body planes laid out in gastrulation



Is this early?