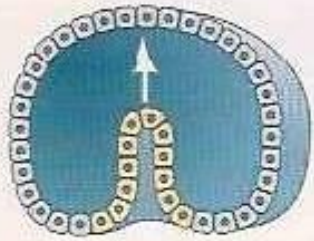


Morphogenetic movements and fate maps

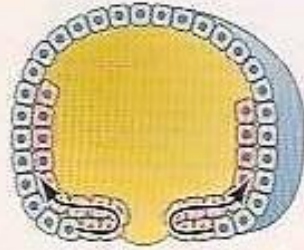
*Dr. Debnarayan Roy
Associate Professor
Department of Zoology
Jhargram Raj College*

Invagination:
Infolding of cell sheet into embryo



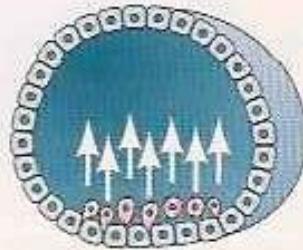
Example:
Sea urchin
endoderm

Involution:
Inturning of cell sheet over the basal surface of an outer layer



Example:
Amphibian
mesoderm

Ingression:
Migration of individual cells into the embryo



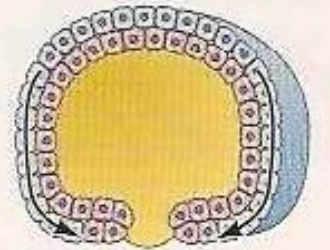
Example:
Sea urchin mesoderm,
Drosophila neuroblasts

Delamination:
Splitting or migration of one sheet into two sheets



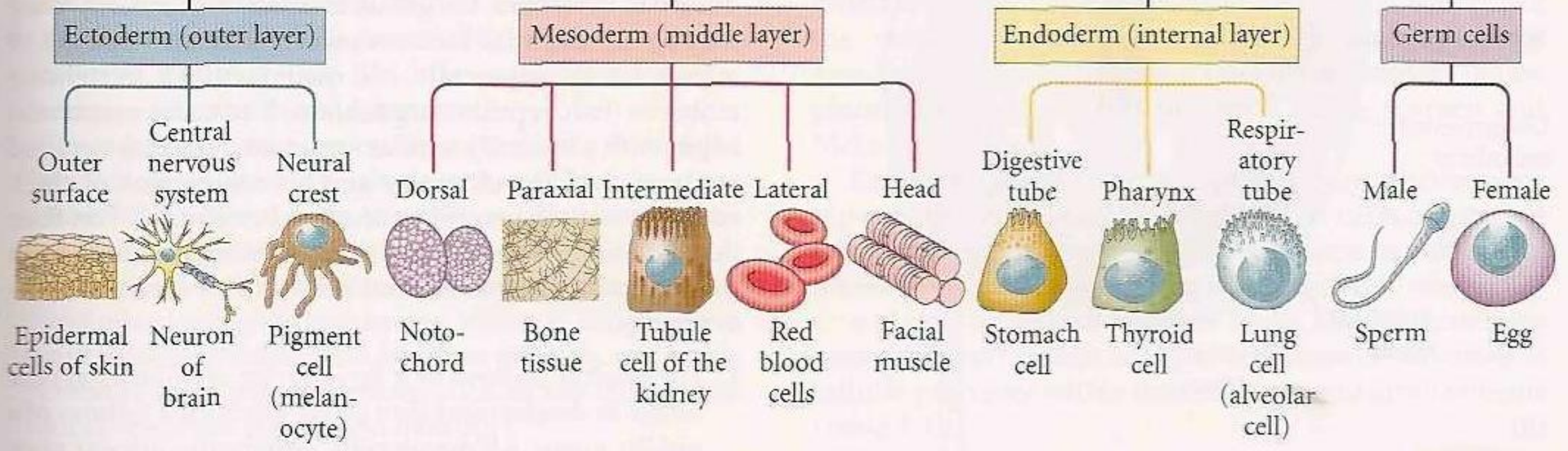
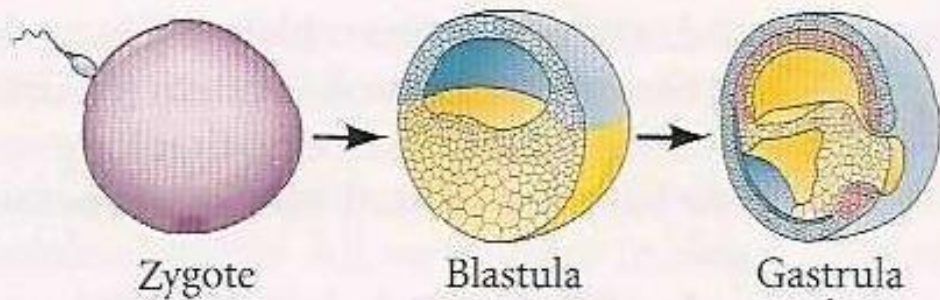
Example:
Mammalian and bird
hypoblast formation

Epiboly:
The expansion of one cell sheet over other cells



Example: Ectoderm
formation in amphibians,
sea urchins, and tunicates


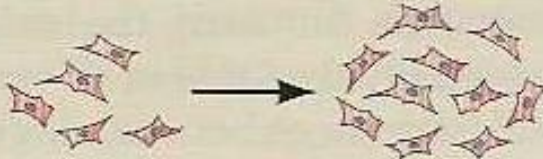
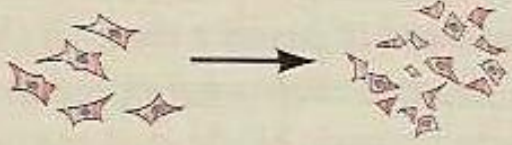
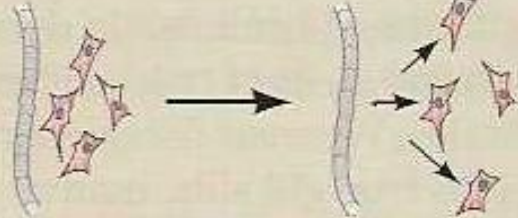
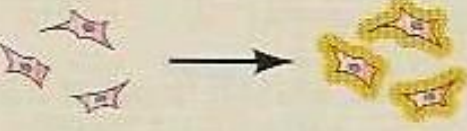
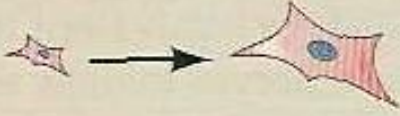
- **Invagination:** The infolding of a region of cells, much like the indenting of a soft rubber ball when it is poked.
- **Involution:** The inturning or inward movement of an expanding outer layer so that it spreads over the internal surface of the remaining external cells.
- **Ingression:** The migration of individual cells from the surface layer into the interior of the embryo. The cells become mesenchymal (i.e., they separate from one another) and migrate independently.
- **Delamination:** The splitting of one cellular sheet into two more or less parallel sheets. While on a cellular basis it resembles ingression, the result is the formation of a new sheet of cells.
- **Epiboly:** The movement of epithelial sheets (usually of ectodermal cells) that spread as a unit (rather than individually) to enclose the deeper layers of the embryo. Epiboly can occur by the cells dividing, by the cells changing their shape, or by several layers of cells intercalating into fewer layers. Often, all three mechanisms are used



Keeping Track of Moving Cells: Fate Maps and Cell Lineages

cells do not stay still in the embryo...

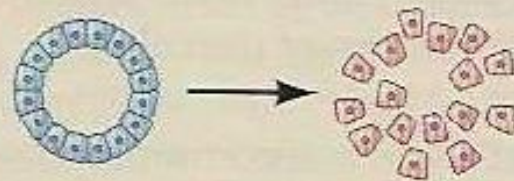
TABLE 1.1 Summary of major morphogenic processes regulated by mesenchymal and epithelial cells

Process	Action	Morphology	Example
MESENCHYMAL CELLS			
Condensation	Mesenchyme becomes epithelium		Cartilage mesenchyme
Cell division	Mitosis produces more cells (hyperplasia)		Limb mesenchyme
Cell death	Cells die		Interdigital mesenchyme
Migration	Cells move at particular times and places		Heart mesenchyme
Matrix secretion and degradation	Synthesis or removal of extracellular layer		Cartilage mesenchyme
Growth	Cells get larger (hypertrophy)		Fat cells

EPITHELIAL CELLS

Dispersal

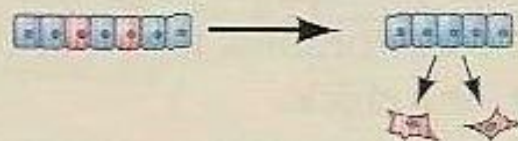
Epithelium becomes mesenchyme (entire structure)



Müllerian duct degeneration

Delamination

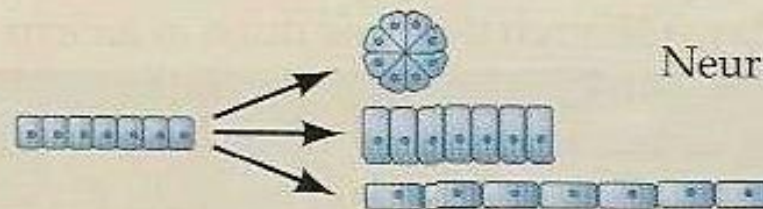
Epithelium becomes mesenchyme (part of structure)



Chick hypoblast

Shape change or growth

Cells remain attached as morphology is altered



Neurulation

Cell migration (intercalation)

Rows of epithelia merge to form fewer rows



Vertebrate gastrulation

Cell division

Mitosis within row or column



Vertebrate gastrulation

Matrix secretion and degradation

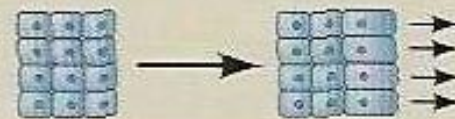
Synthesis or removal of extracellular matrix



Vertebrate organ formation

Migration

Formation of free edges



Chick ectoderm

Fate map

Diagrams that follow cell lineages from specific regions of the embryo in order to "map" larval or adult structures onto the region of the embryo from which they arose. The superimposition of a map of "what is to be" onto a structure that has yet to develop into these organs

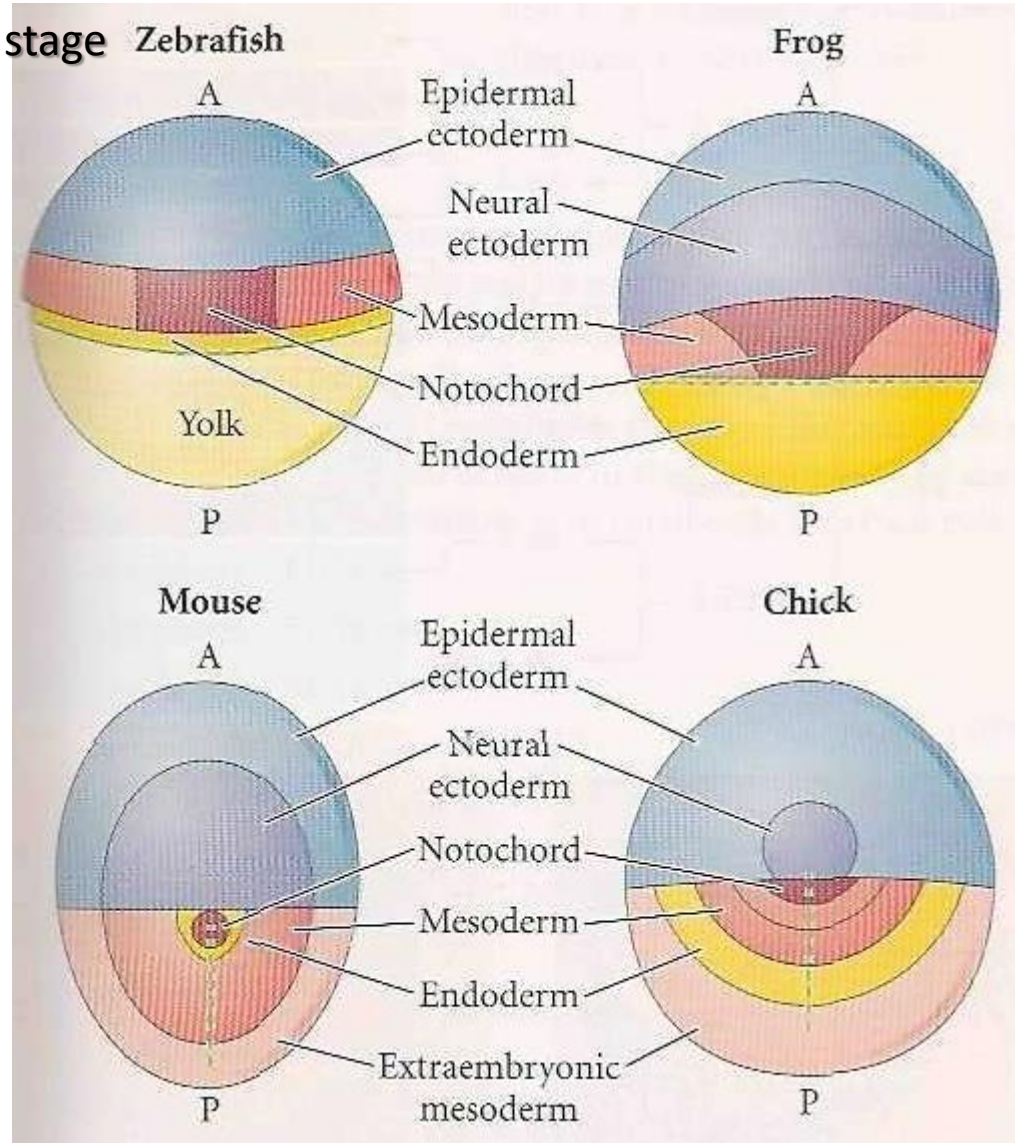
one of the most important programs of descriptive embryology became
the tracing of cell lineages:

following individual cells to see what those cells become

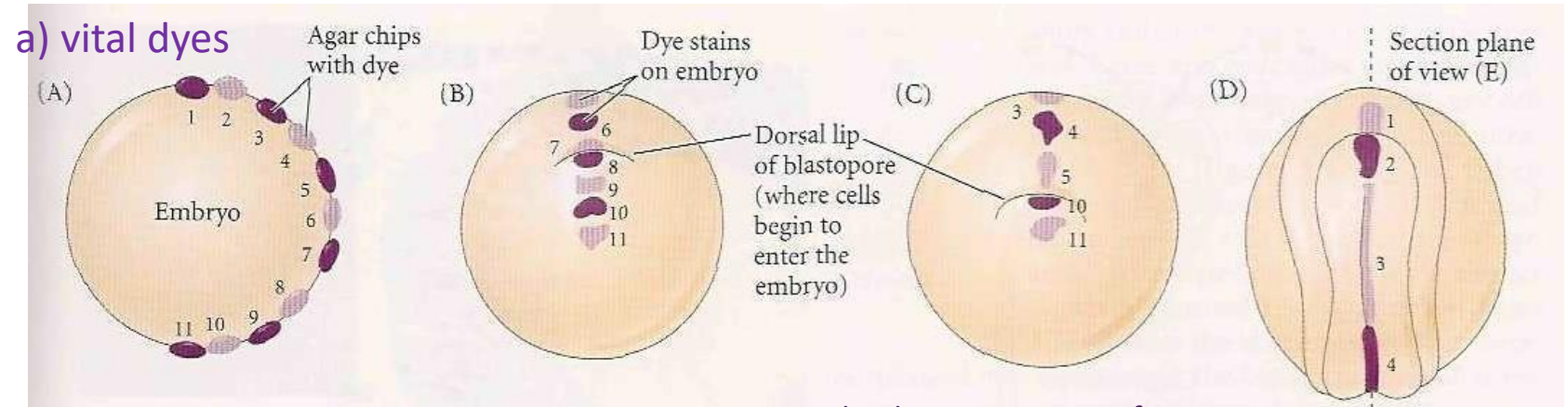
one can label *groups of embryonic* cells to see what
that area becomes in the adult organism. By bringing
such studies together, one can construct a fate map

These diagrams "map" larval or adult structures onto
the region of the embryo from which they arose.

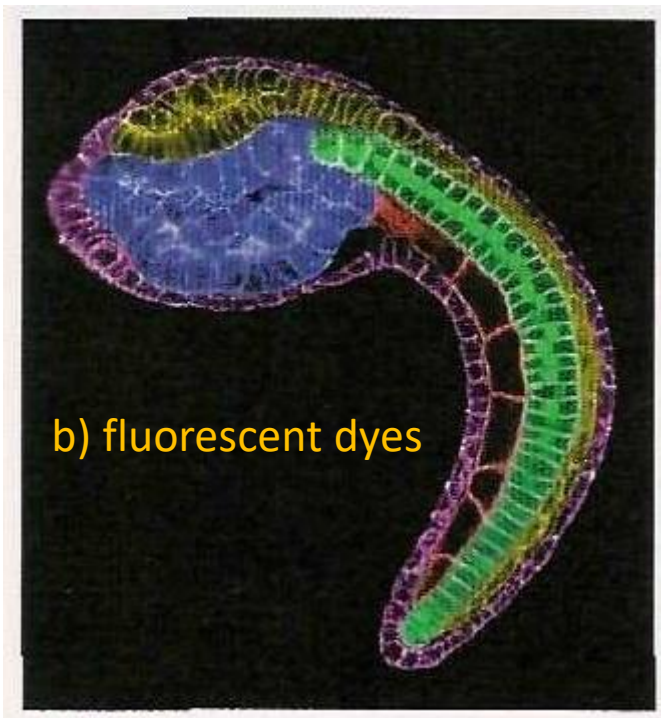
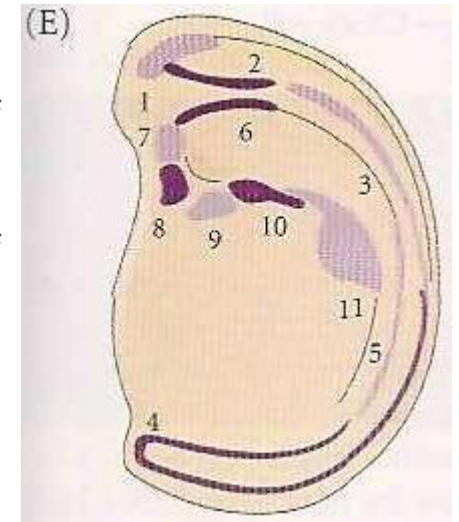
Fate maps of vertebrates at the early gastrula stage



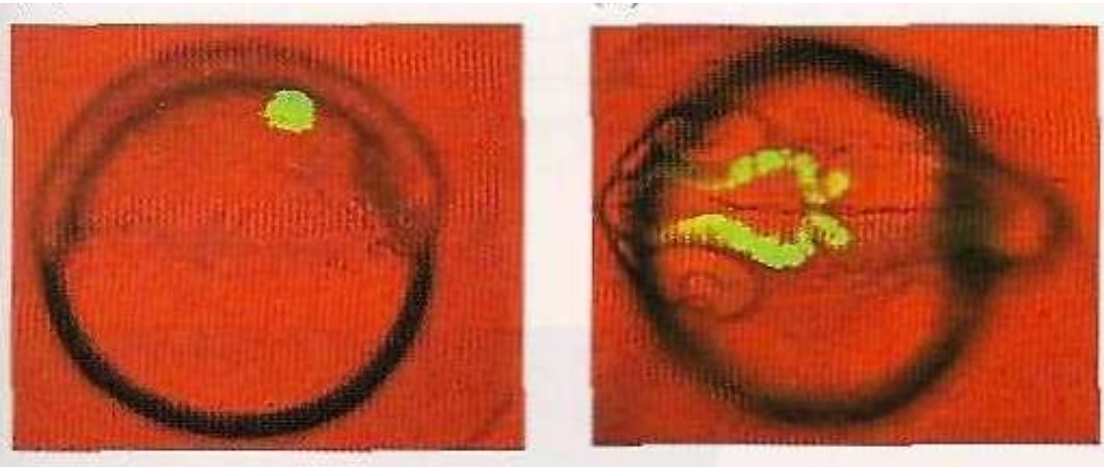
Dye marking (Vogt (1929))



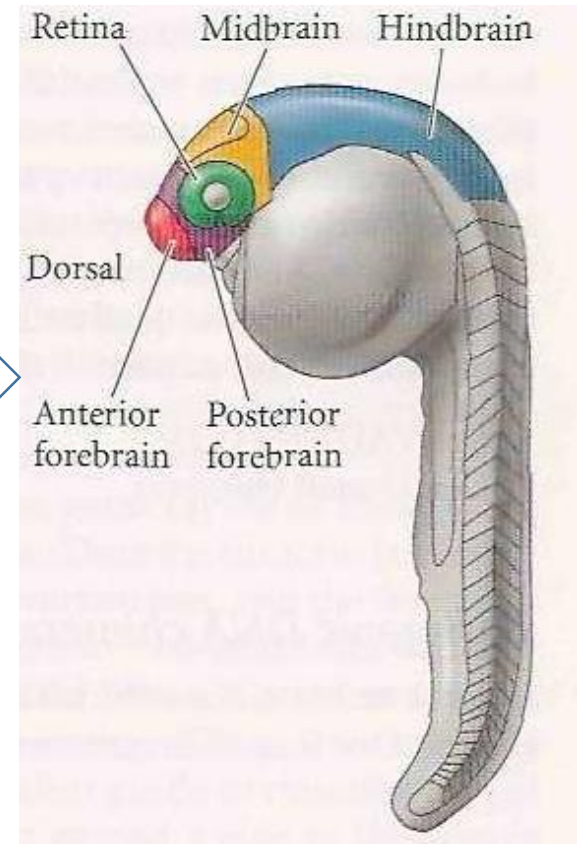
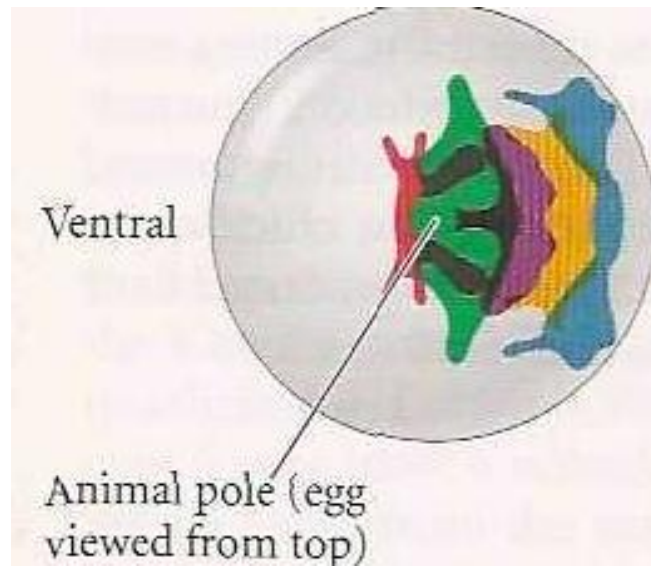
Vital dye staining of amphibian embryos. (A) Vogt's method for marking specific cells of the embryonic surface with vital dyes. (B-D) Dorsal surface views of stain on successively later embryos. (E) Newt embryo dissected in a medial sagittal section to show the stained cells in the interior. (After Vogt 1929.)

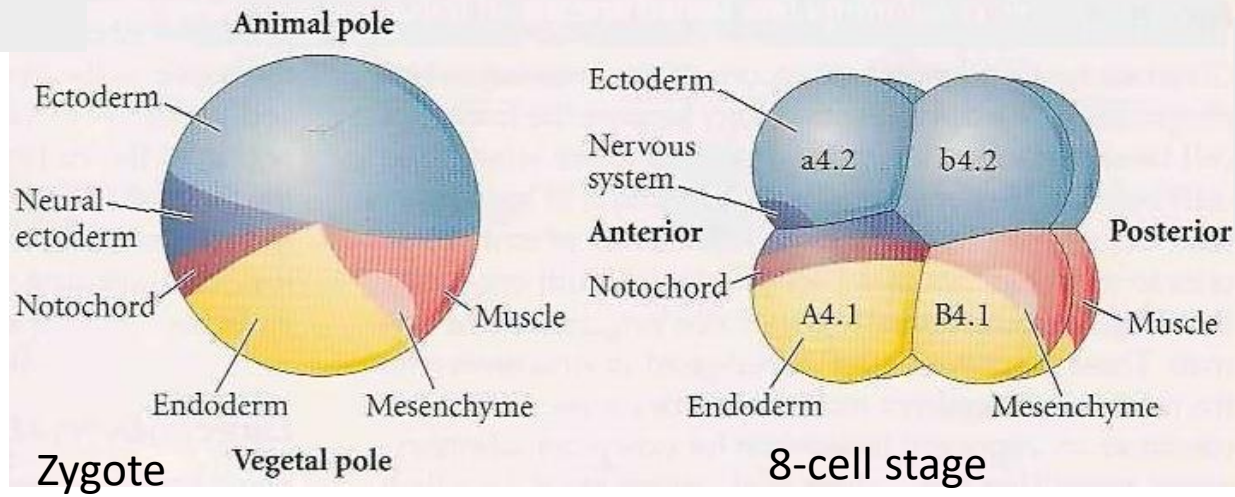


Fate map of the tunicate embryo: Confocal section through a larva of the tunicate *Ciona savignyi*. The **notochord** cells are stained green; the cell boundaries are stained white. The **endoderm** is blue, the **muscles** red, the neural tube yellow, and the epidermis magenta.

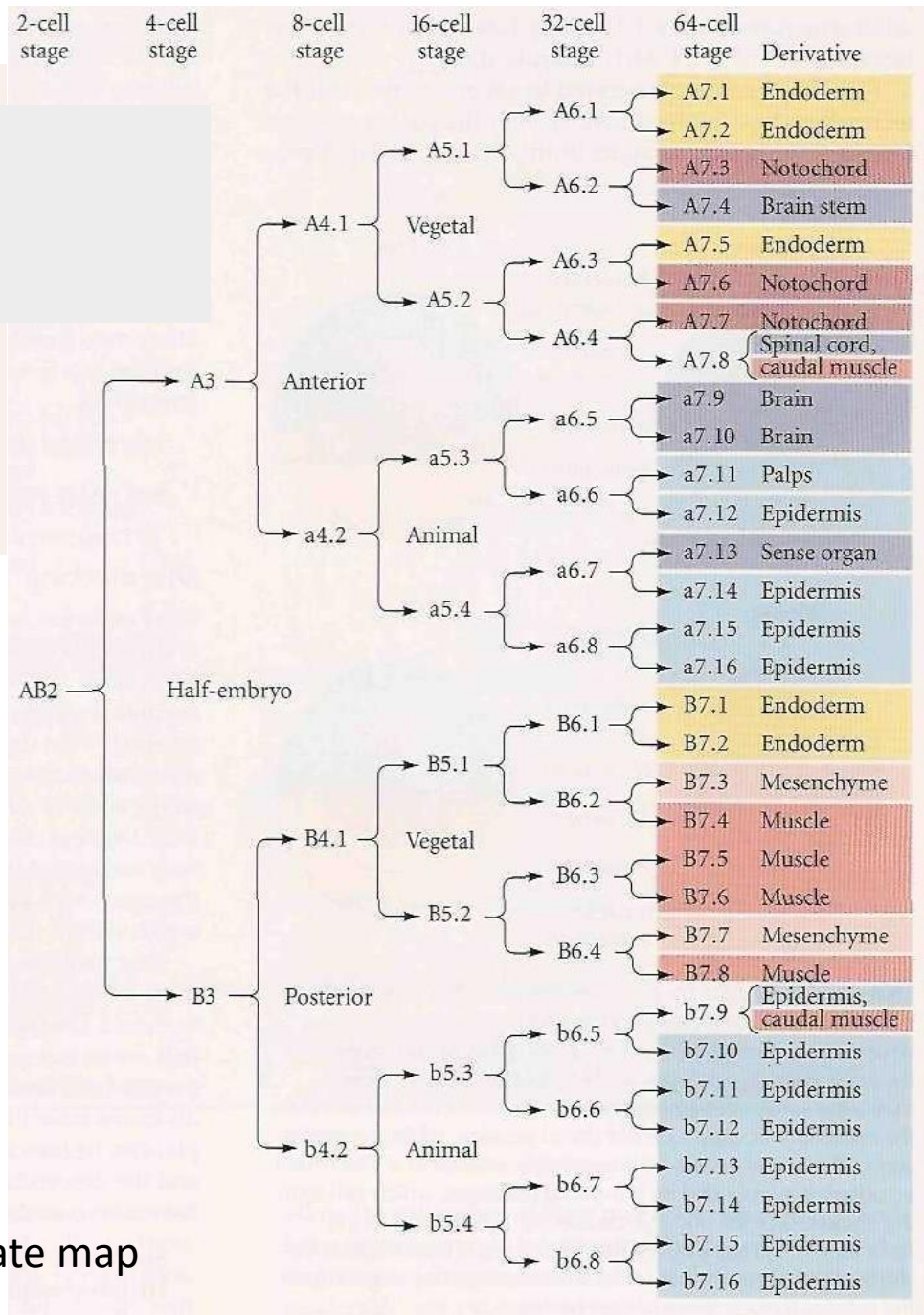


Zebra fish





Styela partita



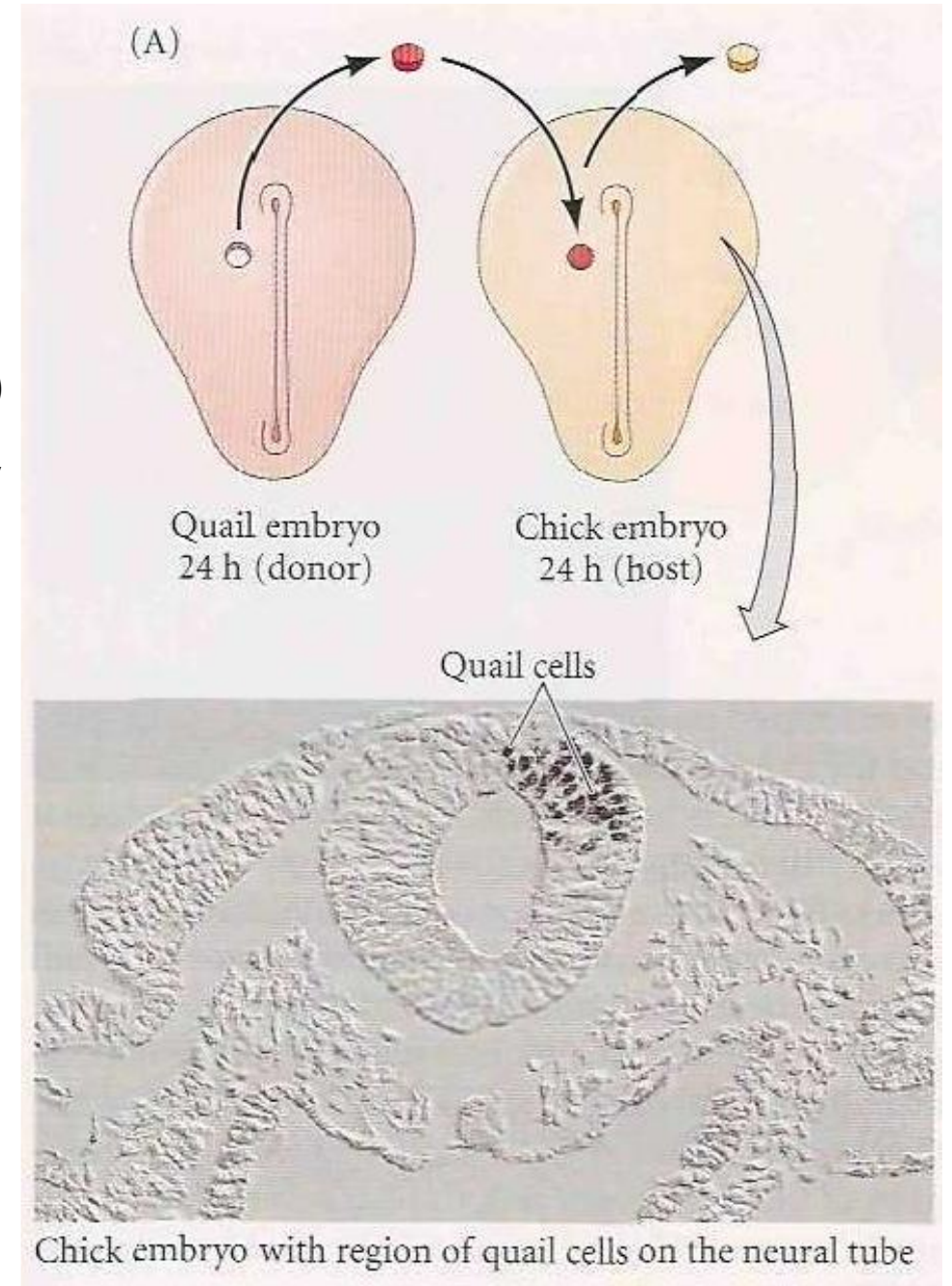
Linear fate map

Chimera

Quail cells differ from chick cells in two important ways.

First, the quail nucleus has condensed DNA (*heterochromatin*) concentrated around the nucleoli, making quail nuclei easily distinguishable from chick nuclei.

Second, cell-specific antigens that are quail-specific can be used to find individual quail cells, even if they are "hidden" within a large population of chick cells.

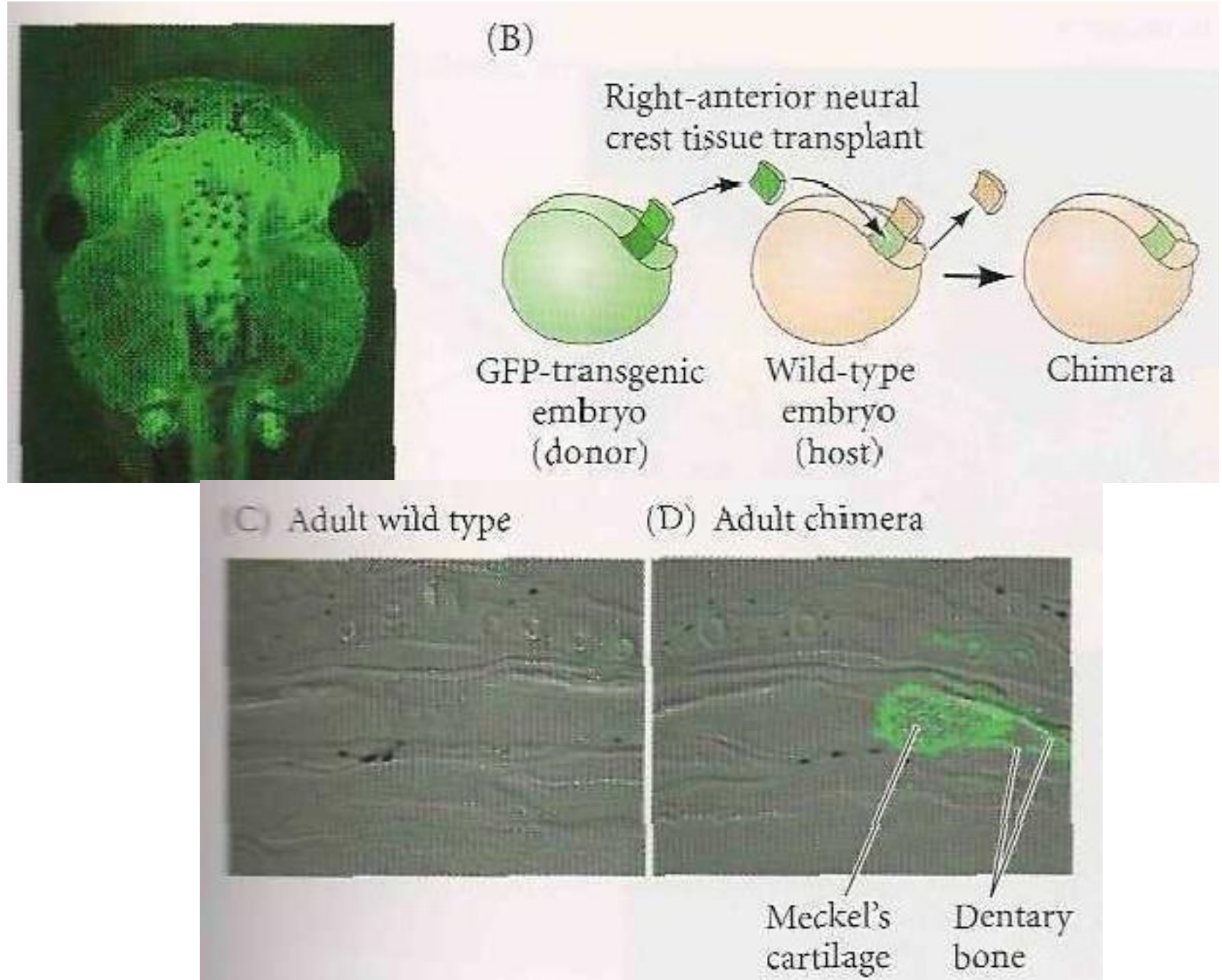


Chick resulting from transplantation of a trunk neural crest region from an embryo of a pigmented strain of chickens into the same region of an embryo of an unpigmented strain. The neural crest cells that gave rise to the pigment migrated into the wing epidermis and feathers.



Transgenic DNA chimera

Fate mapping with transgenic DNA shows that the neural crest is critical in making the bones of the frog jaw.



SPECIFICATION

*Introducing Cell Commitment and
Early Embryonic Development*

“the greatest of all wonders of the material universe: the existence, in a simple, unorganized egg, of a power to produce a definite adult animal.”

-William Keith Brooks, 1883

Differentiation The process by which an unspecialized cell becomes specialized into one of the many cell types that make up the body.

Commitment Describes a state in which a cell's developmental fate has become restricted even though it is not yet displaying overt changes in cellular biochemistry and function.

Simply...

The generation of specialized cell types is called *differentiation*

and the process is

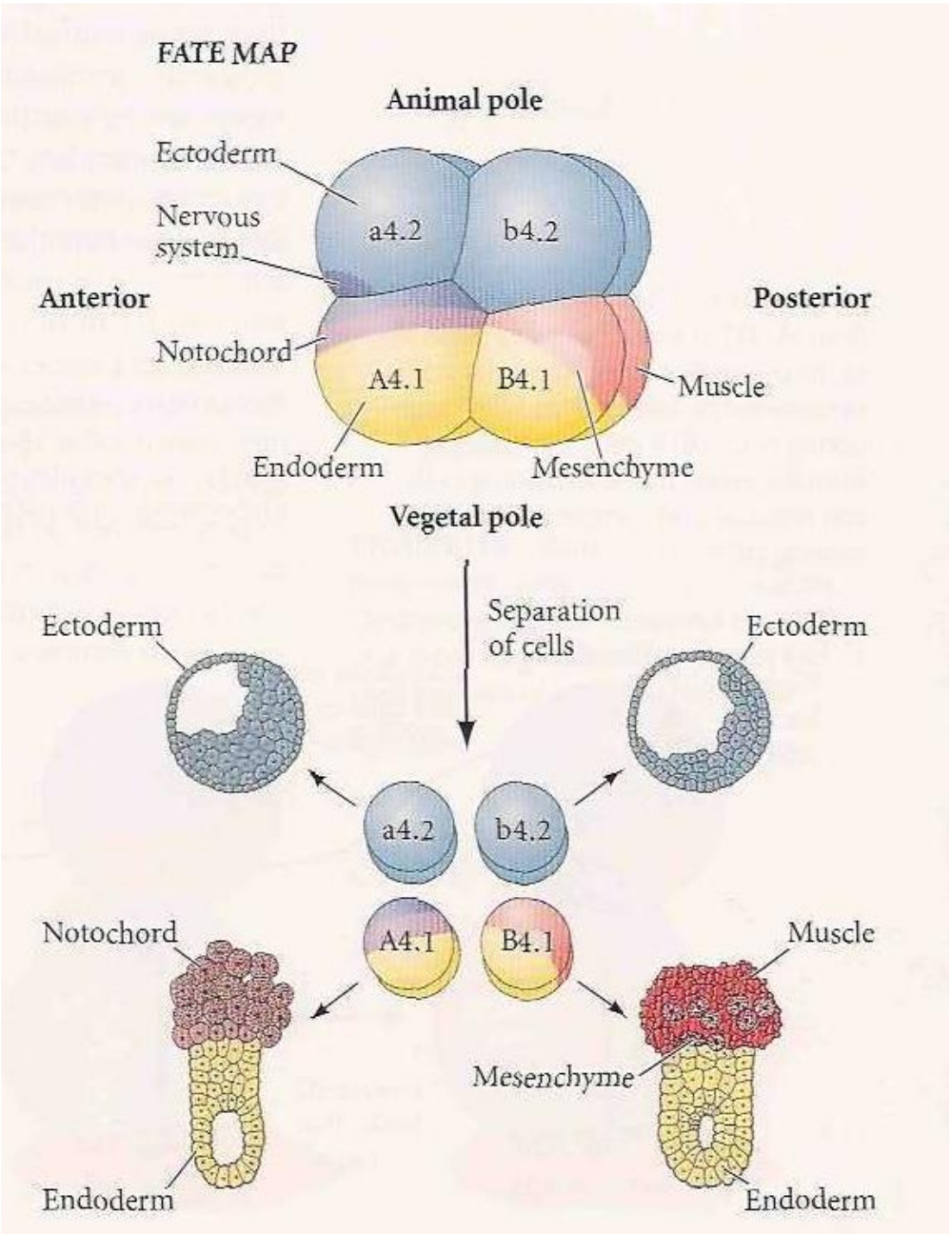
commitment

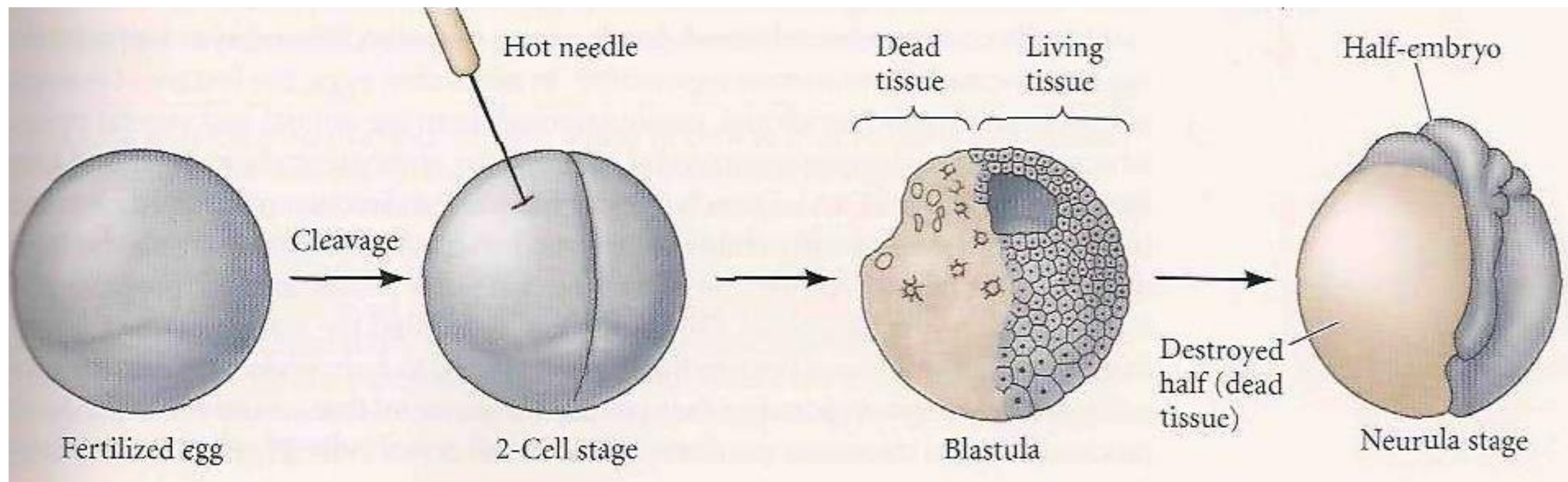
Commitment have two stages...

1)Specification_the labile one

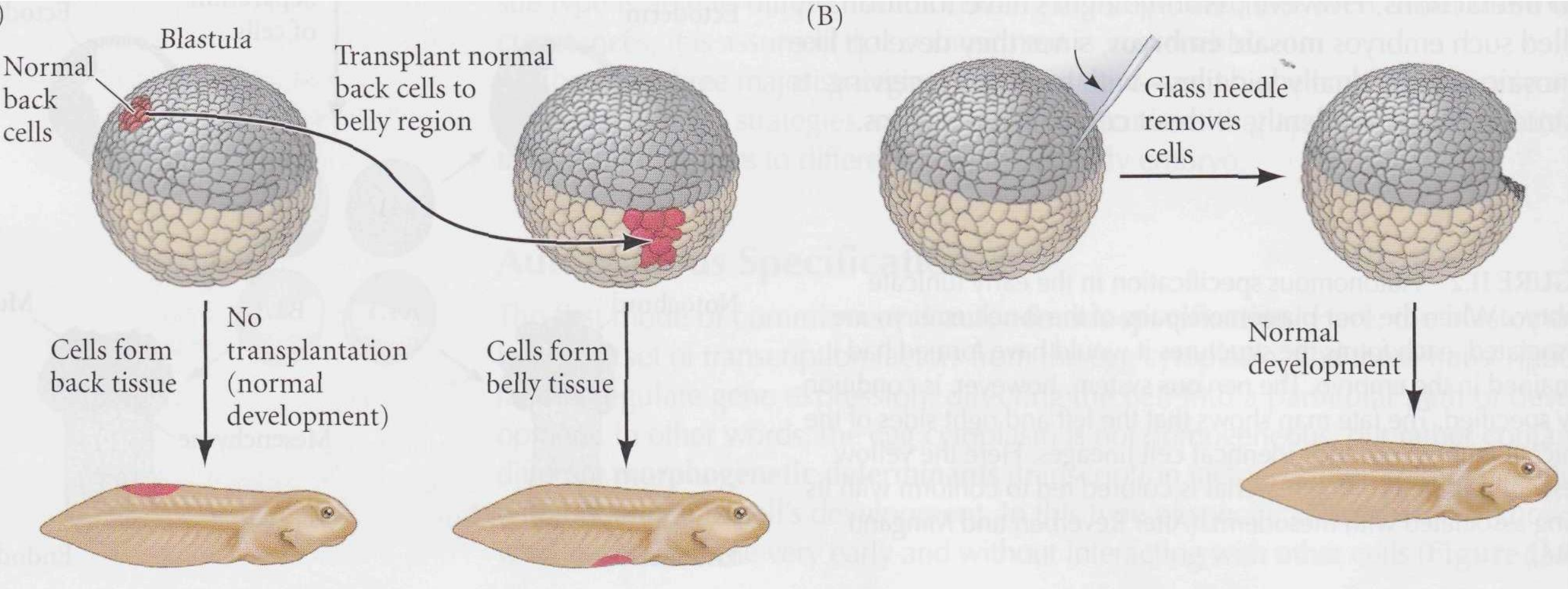
2)Determination_the irreversible one

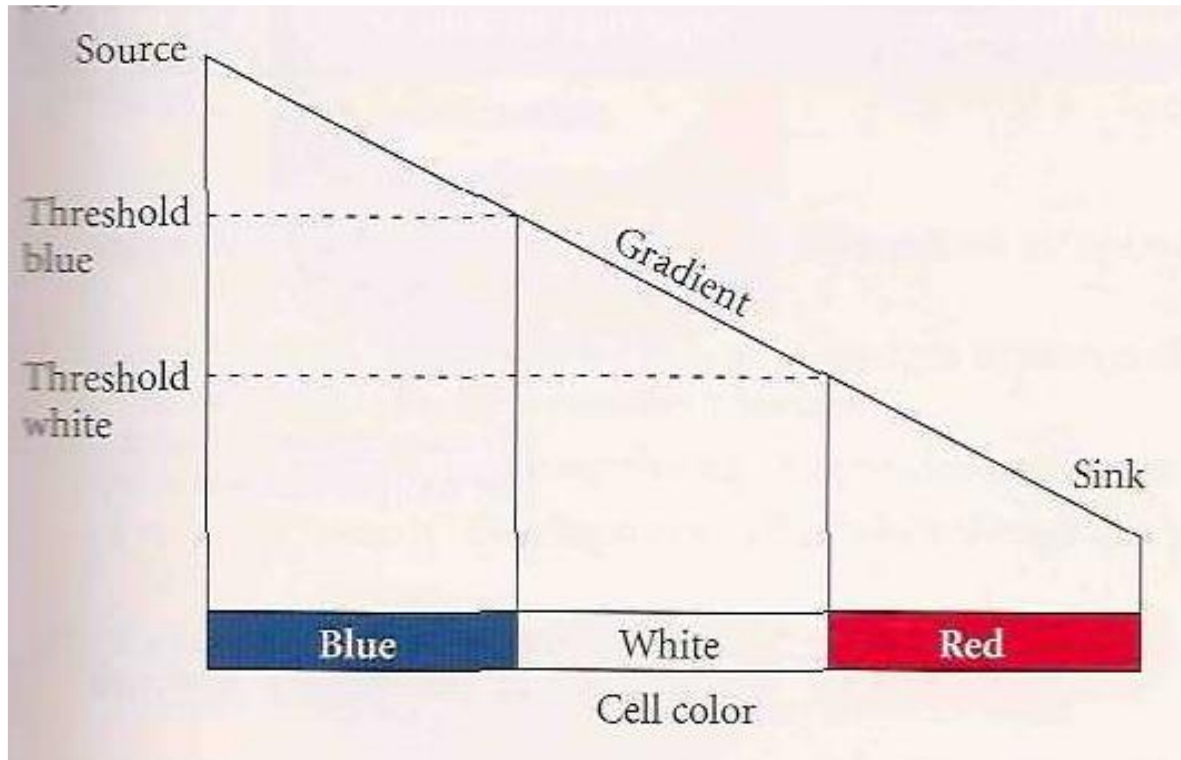
Autonomous specification



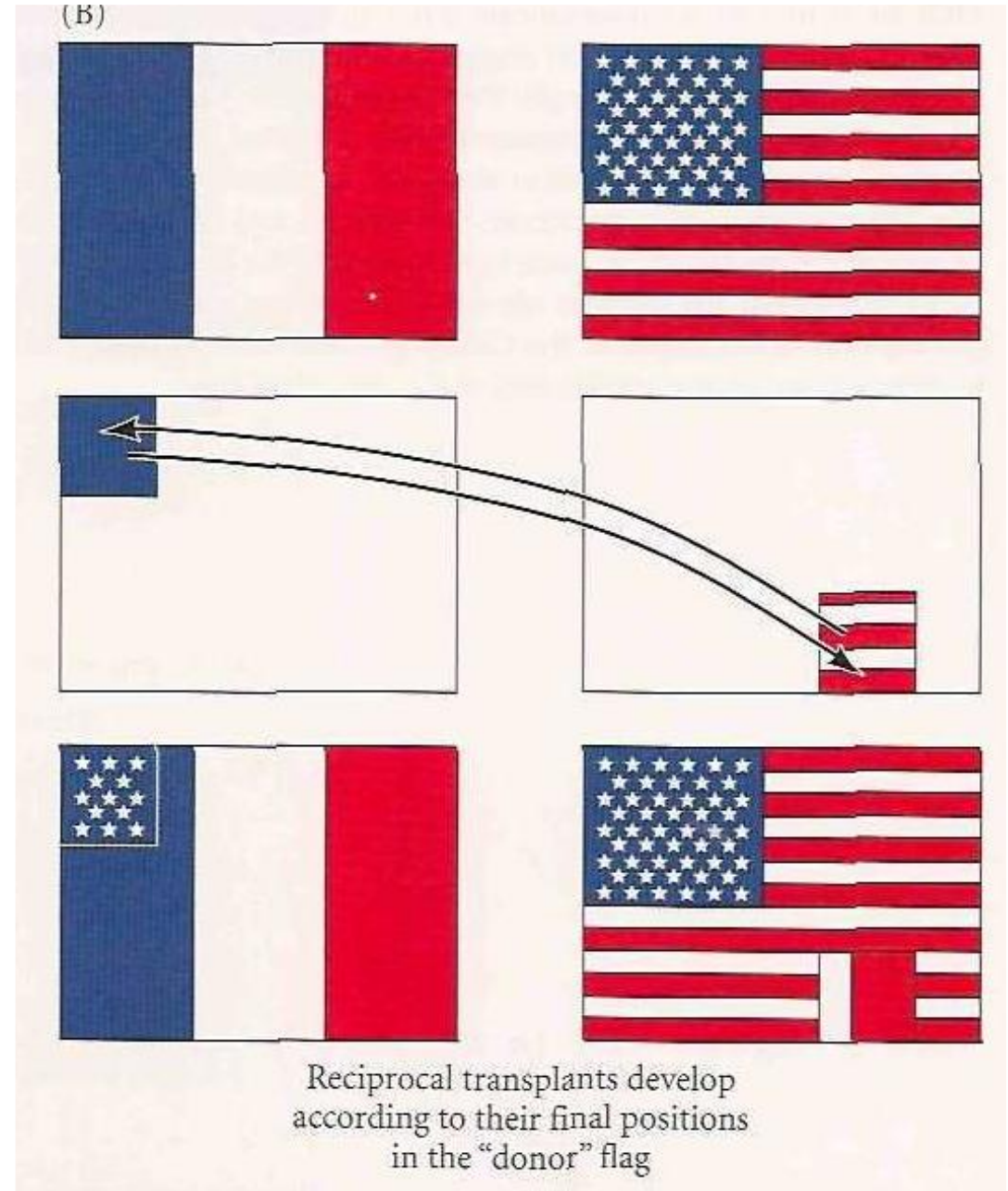


Conditional specification

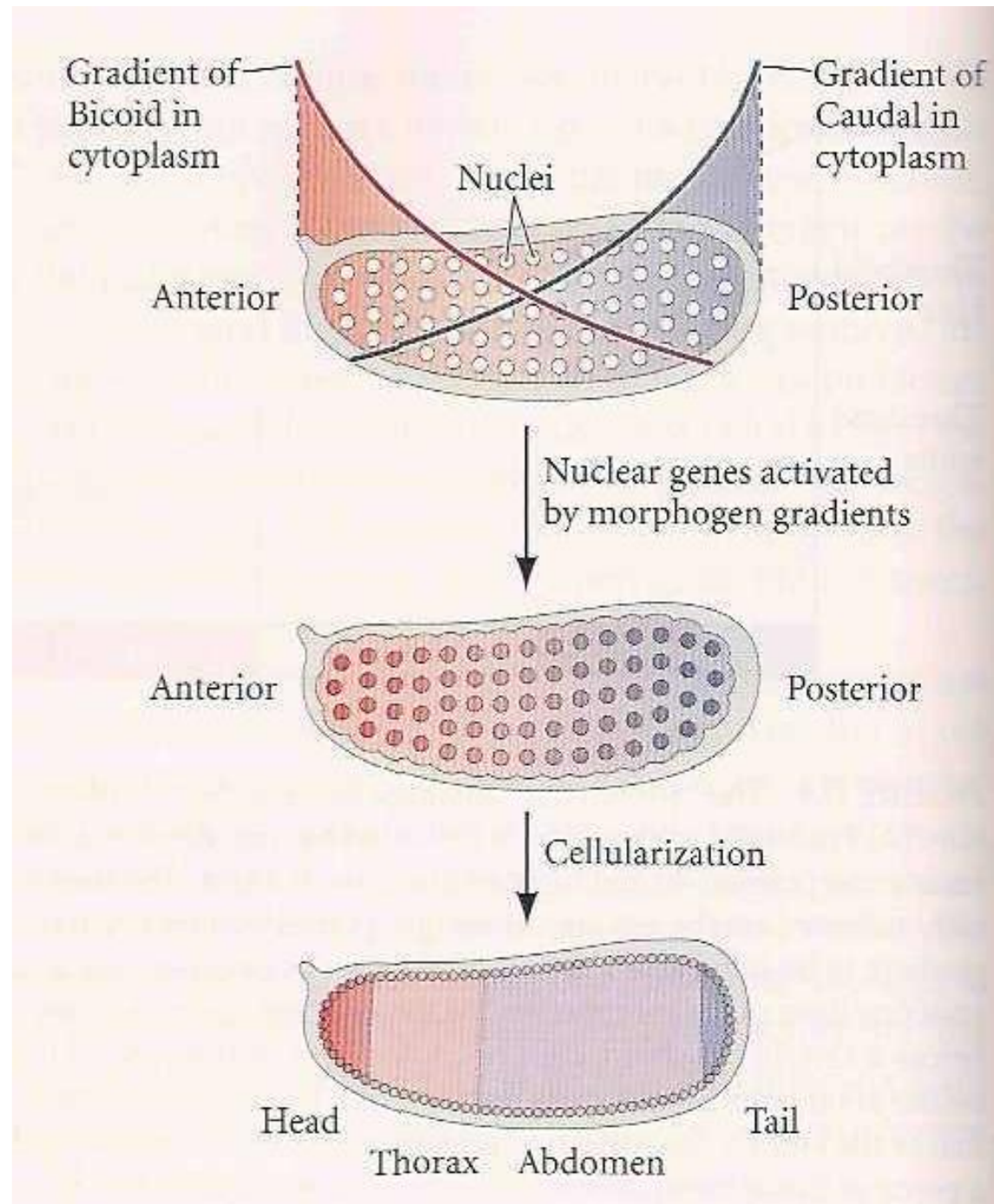




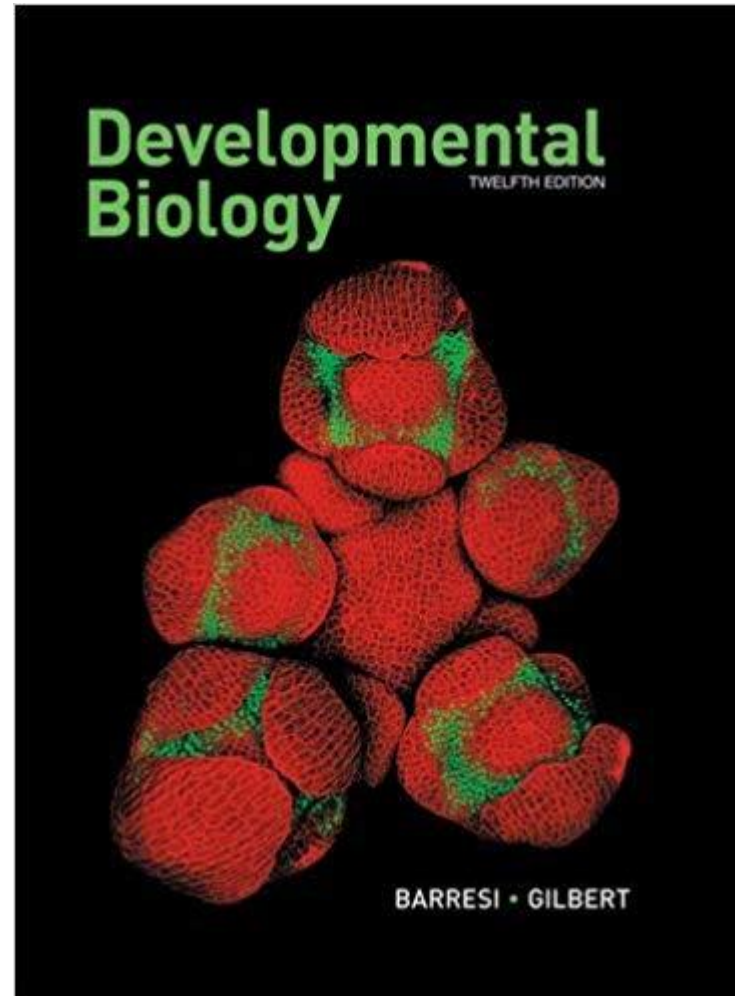
French flag analogy of conditional specification



Syncytial specification



References:



That's all...