Principles of Development 7.22/7.72 Fall 2005

Principles are in **boldface**, specific terms you should know are in **boldface italic**.

- 1. Development occurs in four dimensions: in three-dimensional space and over time.
- 2. Complexity of the embryo increases over time, in a stepwise, hierarchical fashion.
- 3. **Symmetry must repeatedly be broken** to allow formation of different daughter cells or functionally distinct regions.
- 4. The progression of a naïve cell towards its final function (*fate*) includes the decision to assume a fate (*determination*) and formation of the final fate (*differentiation*).
- 5. Determination and differentiation each involve multiple, sequential steps.
- 6. There are many possible *cell fates*.
- 7. Information that controls *cell type* can be distinct from information that indicates the *position* or *shape* of a cell in an embryo.
- 8. Development is controlled by spatially and temporally regulated *differential gene expression*.
- 9. Different cells express different *regulatory genes* that control fate decisions, and different *differentiation genes* that control final cell function.
- **10.** A *combinatorial code* of gene expression allows many different developmental decisions using a limited number of genes.
- 11. Gene function is conserved through evolution (homologs, orthologs).
- 12. Chromatin structure plays a crucial role in regulating gene expression (epigenetics).
- 13. Embryonic regulatory molecules may be stored in the egg (*maternal factors*) or produced by new transcription after fertilization (*zygotic factors*).
- 14. Cells can read and interpret chemical, electrical and mechanical signals originating from other cells or the environment.
- **15.** Cell communication can alter cell fate (*induction*). Thus, regulatory molecules can act between cells, that is, non-autonomously (*inducers*).
- 16. Cell fate can be controlled by inherited regulatory molecules, which act cell autonomously (*determinants*).

- 17. Cells respond differently to the same signal when it is presented at different concentrations (*morphogen*). Continuous information contained within a concentration gradient is converted to a discrete outcome through *thresholds* of activation.
- 18. Cell fate can be regulated by independent *initiation* and *maintenance* mechanisms.
- **19.** The number of cell fates that a cell has the potential to assume progressively diminishes as development proceeds (*potency*).
- 20. Cell fate decisions are generally stable.
- **21.** Cells can have restricted temporal and spatial ability to respond to signals (*competence*).
- 22. Groups of cells have the potential to assume a particular fate (*equivalence group, field*), but often only part of the group does so.
- 23. Cell fate decisions often involve both stimulatory and inhibitory signals. Stimulatory signals encourage cells to adopt a specific fate whereas inhibitory signals prevent a cell from adopting an alternate fate.
- 24. Cells may inhibit their neighbors from assuming the same fate (lateral inhibition).
- 25. Similar cells may cooperate to promote formation of their fate (community effect).
- 26. Cells that can generate replacements for a specific cell type may be present in many embryonic and adult tissues (*stem cells*).
- 27. Development requires both cell division (proliferation) and cell death (apoptosis).
- 28. Cells can form sheets (epithelia) or persist as single cells (mesenchyme).
- 29. Single cells can move (*migrate*). Direction of migration can be regulated (*guidance molecules*).
- **30.** Cells can *sort* into groups of like cells, through selective adherence to each other and to the extracellular matrix.
- 31. Groups of similar cells can become a functional unit (tissue).
- 32. Boundaries are established to keep cells where they need to be (compartments).
- 33. Groups of tissues can work together for a common function (organ).
- 34. Cells move or change shape to build three-dimensional structures (morphogenesis).
- 35. Three-dimensional organization of tissues is required for organ function (*morphogenesis*).