## B. Pattern Formation

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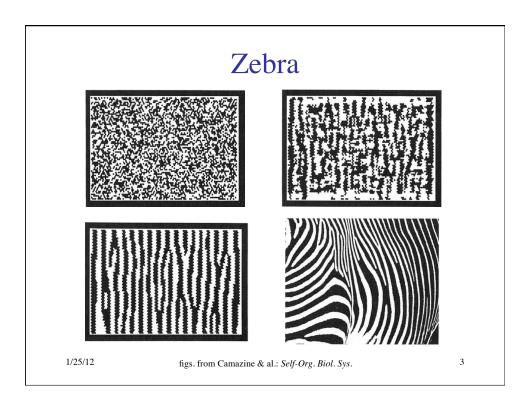


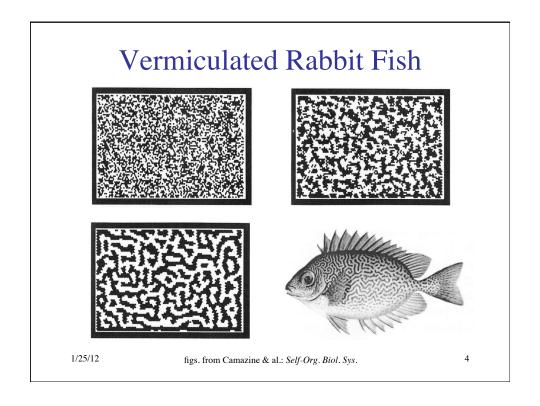


1/25/12 photos ©2000, S. Cazamine

- A central problem in development: How do cells differentiate to fulfill different purposes?
- How do complex systems generate spatial & temporal structure?
- CAs are natural models of intercellular communication

2



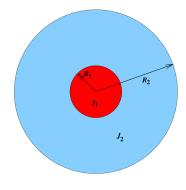


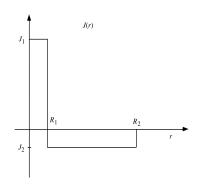
## Activation & Inhibition in Pattern Formation

- Color patterns typically have a characteristic length scale
- Independent of cell size and animal size
- Achieved by:
  - short-range activation ⇒ local uniformity
  - long-range inhibition ⇒ separation

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### **Interaction Parameters**





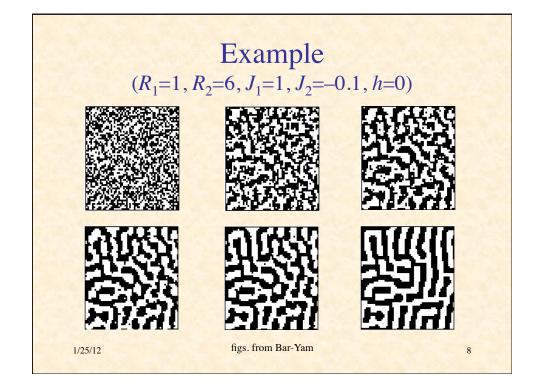
- $R_1$  and  $R_2$  are the interaction ranges
- $J_1$  and  $J_2$  are the interaction strengths

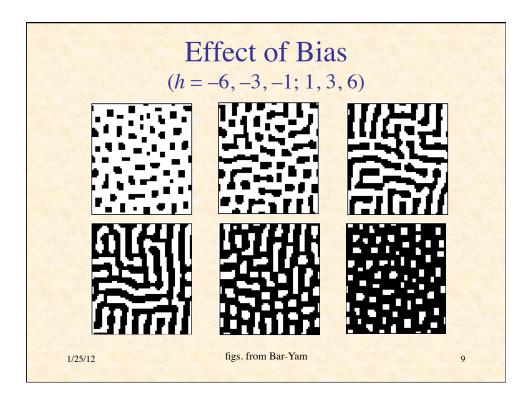
### CA Activation/Inhibition Model

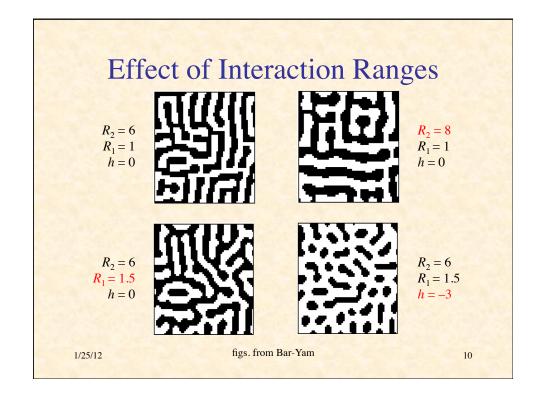
- Let states  $s_i \in \{-1, +1\}$
- and h be a bias parameter
- and  $r_{ij}$  be the distance between cells i and j
- Then the state update rule is:

$$s_i(t+1) = \text{sign}\left[h + J_1 \sum_{r_{ij} < R_1} s_j(t) + J_2 \sum_{R_1 \le r_{ij} < R_2} s_j(t)\right]$$

1/25/12







# Demonstration of NetLogo Program for Activation/Inhibition Pattern Formation: Fur

#### RunAICA.nlogo

1/25/12

### Differential Interaction Ranges

- How can a system using strictly local interactions discriminate between states at long and short range?
- E.g. cells in developing organism
- Can use two different *morphogens* diffusing at two different rates
  - activator diffuses slowly (short range)
  - inhibitor diffuses rapidly (long range)

### Digression on Diffusion

• Simple 2-D diffusion equation:

$$\dot{A}(x,y) = c\nabla^2 A(x,y)$$

• Recall the 2-D Laplacian:

$$\nabla^2 A(x,y) = \frac{\partial^2 A(x,y)}{\partial x^2} + \frac{\partial^2 A(x,y)}{\partial y^2}$$

- The Laplacian (like 2<sup>nd</sup> derivative) is:
  - positive in a local minimum
- negative in a local maximum

13

### Reaction-Diffusion System

diffusion

$$\frac{\partial A}{\partial t} = \begin{bmatrix} d_{A} \nabla^{2} A + f_{A}(A, I) \\ \frac{\partial I}{\partial t} = d_{I} \nabla^{2} I + f_{I}(A, I) \end{bmatrix}$$
 reaction

$$\frac{\partial}{\partial t} \begin{pmatrix} A \\ I \end{pmatrix} = \begin{pmatrix} d_{A} & 0 \\ 0 & d_{I} \end{pmatrix} \begin{pmatrix} \nabla^{2} A \\ \nabla^{2} I \end{pmatrix} + \begin{pmatrix} f_{A}(A, I) \\ f_{I}(A, I) \end{pmatrix}$$

$$\dot{\mathbf{c}} = \mathbf{D}\nabla^2 \mathbf{c} + \mathbf{f}(\mathbf{c}), \text{ where } \mathbf{c} = \begin{pmatrix} A \\ I \end{pmatrix}$$

1/25/12

14

## Continuous-time Activator-Inhibitor System

- Activator A and inhibitor I may diffuse at different rates in x and y directions
- Cell becomes more active if activator + bias exceeds inhibitor
- Otherwise, less active

$$\frac{\partial A}{\partial t} = d_{Ax} \frac{\partial^2 A}{\partial x^2} + d_{Ay} \frac{\partial^2 A}{\partial y^2} + k_A (A + B - I)$$

$$\frac{\partial I}{\partial t} = d_{Ix} \frac{\partial^2 I}{\partial x^2} + d_{Iy} \frac{\partial^2 I}{\partial y^2} + k_I (A + B - I)$$

1/25/12

15

## NetLogo Simulation of Reaction-Diffusion System

- 1. Diffuse activator in X and Y directions
- 2. Diffuse inhibitor in X and Y directions
- 3. Each patch performs:

  stimulation = bias + activator inhibitor + noise
  if stimulation > 0 then

set activator and inhibitor to 100 else

set activator and inhibitor to 0

## Demonstration of NetLogo Program for Activator/Inhibitor Pattern Formation

#### Run Pattern.nlogo

1/25/12

# Demonstration of NetLogo Program for Activator/Inhibitor Pattern Formation with Continuous State Change

Run Activator-Inhibitor.nlogo

### **Turing Patterns**

- Alan Turing studied the mathematics of reaction-diffusion systems
- Turing, A. (1952). The chemical basis of morphogenesis. *Philosophical Transactions of the Royal Society* **B 237**: 37–72.
- The resulting patterns are known as *Turing* patterns

1/25/12

# A Key Element of Self-Organization

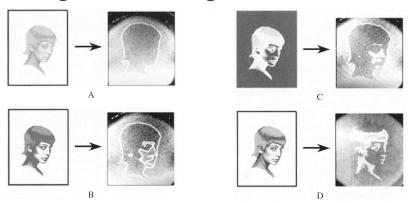
- Activation vs. Inhibition
- Cooperation vs. Competition
- Amplification vs. Stabilization
- Growth vs. Limit
- Positive Feedback vs. Negative Feedback
  - Positive feedback creates
  - Negative feedback shapes

### Reaction-Diffusion Computing

- Has been used for image processing
  - diffusion ⇒ noise filtering
  - reaction ⇒ contrast enhancement
- Depending on parameters, RD computing can:
  - restore broken contours
  - detect edges
  - improve contrast

1/25/12

### Image Processing in BZ Medium

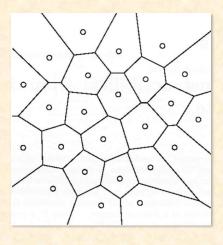


• (A) boundary detection, (B) contour enhancement, (C) shape enhancement, (D) feature enhancement

1/25/12 Image < Adamatzky, Comp. in Nonlinear Media & Autom. Coll.

22

### Voronoi Diagrams



- Given a set of generating points:
- Construct a polygon around each generating point of set, so all points in a polygon are closer to its generating point than to any other generating points.

1/25/12

Image < Adamatzky & al., Reaction-Diffusion Computers

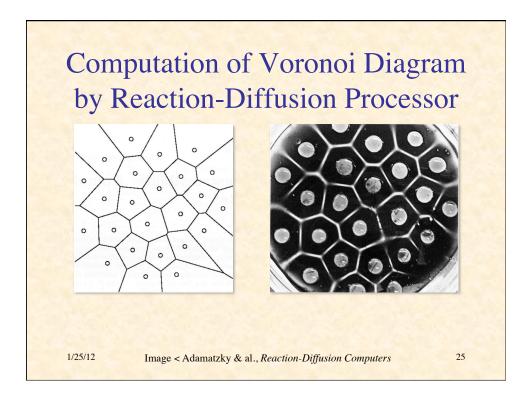
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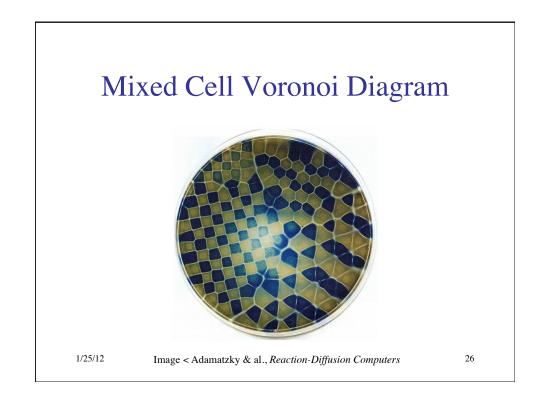
## Some Uses of Voronoi Diagrams

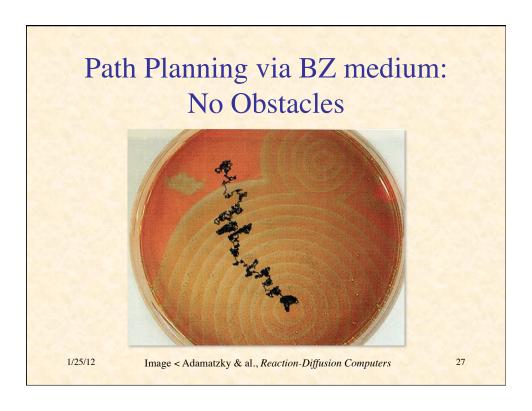
- Collision-free path planning
- Determination of service areas for power substations
- Nearest-neighbor pattern classification
- Determination of largest empty figure

1/25/12

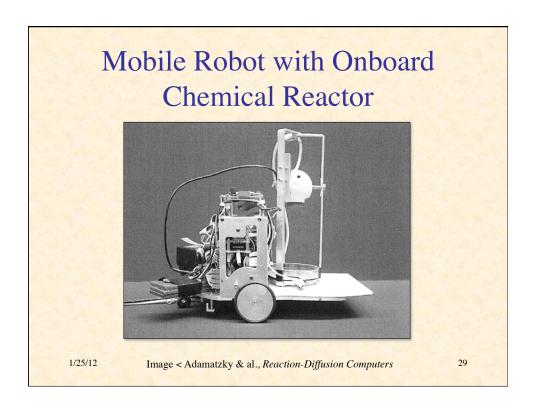
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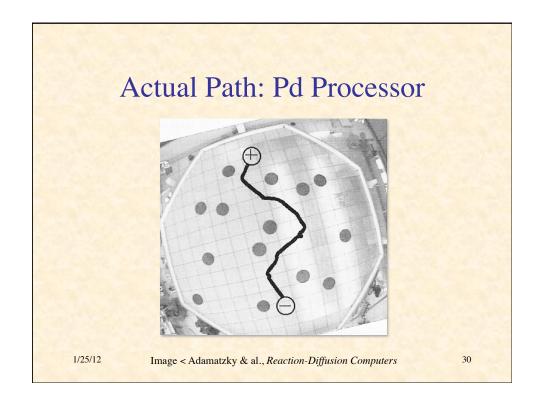


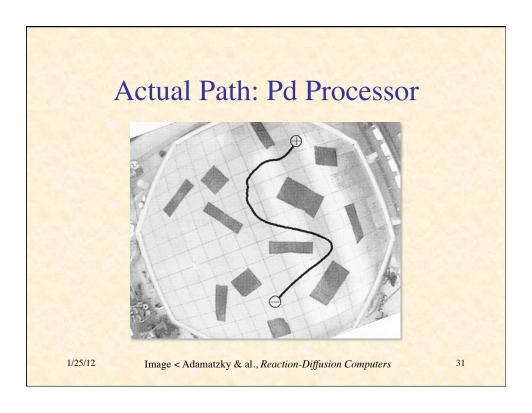


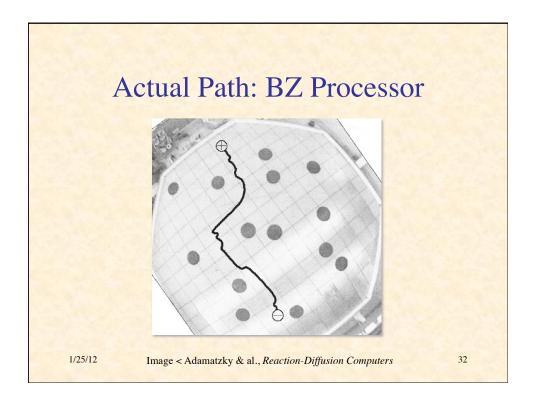












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- 2. Adamatzky, Adam, De Lacy Costello, Ben, & Asai, Tetsuya. *Reaction Diffusion Computers*. Amsterdam: Elsevier, 2005.

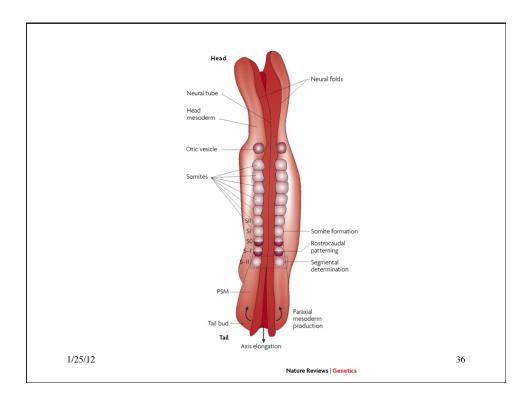
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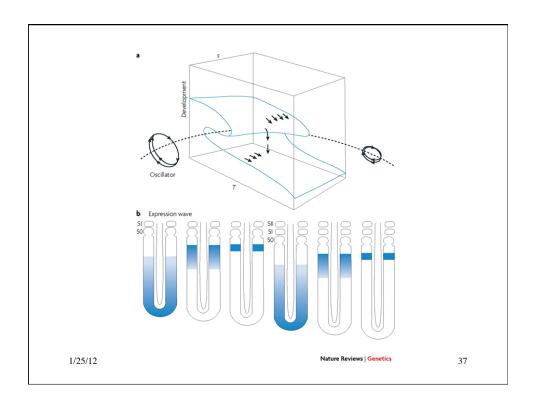
### Segmentation

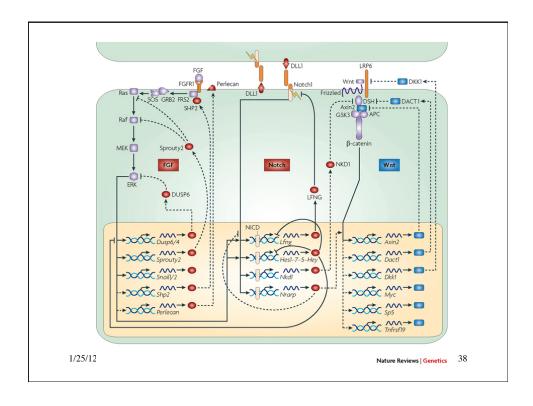
(in embryological development)

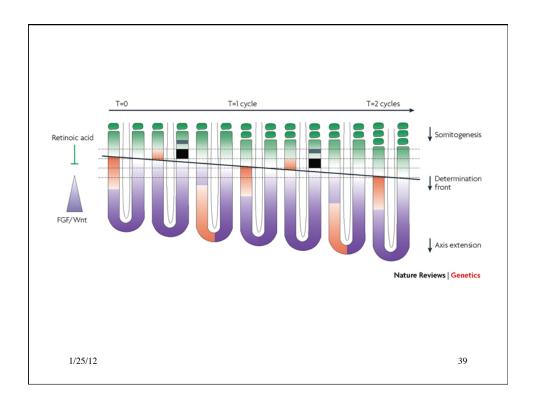
### Vertebrae

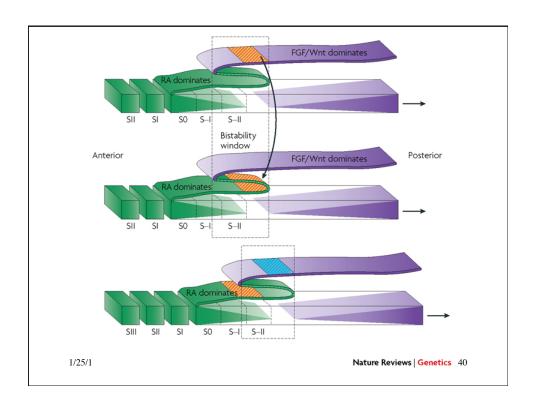
- Humans: 33, chickens: 55, mice: 65, corn snake: 315
- Characteristic of species
- How does an embryo "count" them?
- "Clock and wavefront model" of Cooke & Zeeman (1976).

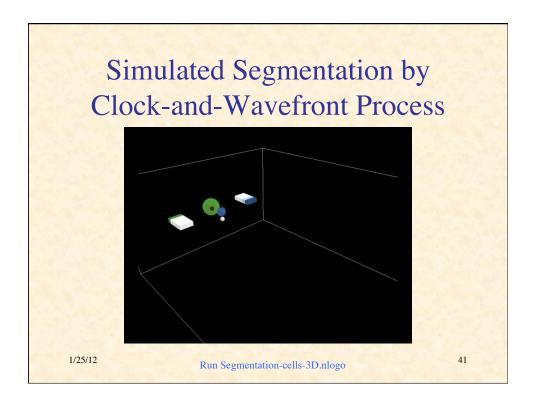


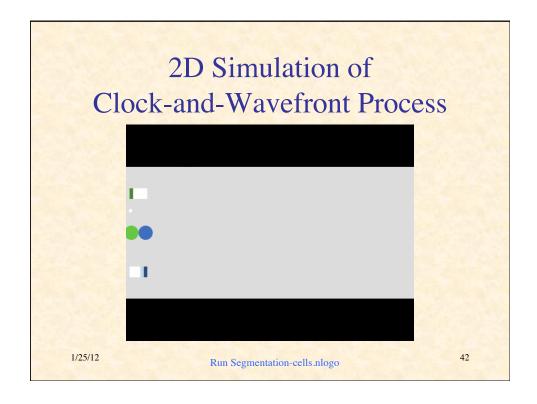


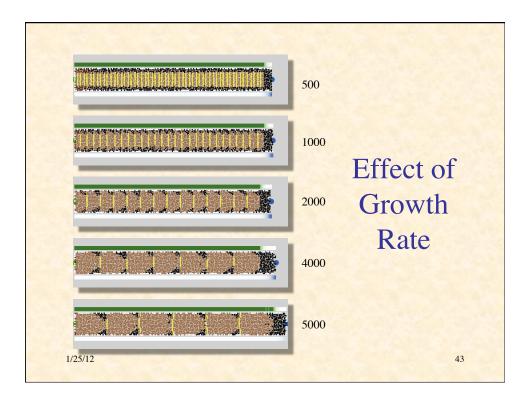


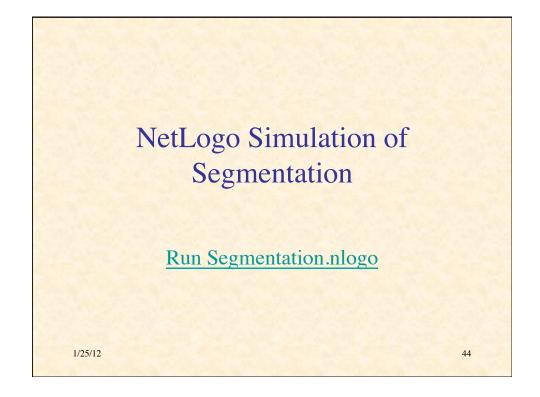












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1/25/12 45

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- 6. Solé, R., & Goodwin, B. Signs of Life: How Complexity Pervades Biology. Basic Books, 2000.

1/25/12 continue to "Part 2C" 46