

Generating Textures on Arbitrary Surfaces Using Reaction-Diffusion

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Motivation

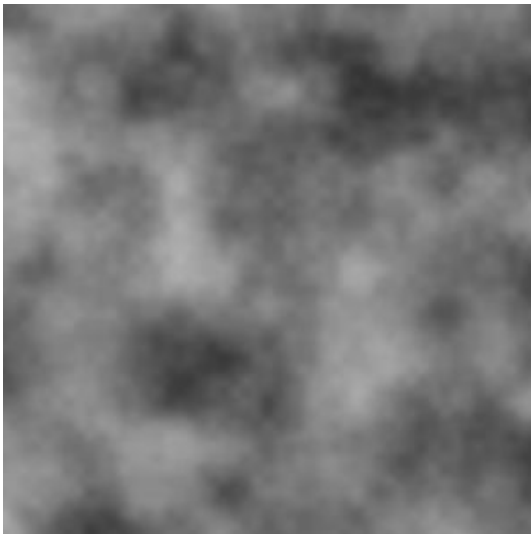
- Biological processes for creating animal coats unknown
- Reaction-Diffusion possible candidate
- How can we mimic this process for more effective texture generation and mapping to an object from 2D to 3D space?

Goals

- Two main goals
- Pattern generation with reaction-diffusion
 - Simple patterns
 - Complex with cascading R-D multiple times
- Generate mesh and synthesize a texture directly on a mesh
 - Removes need to map texture coordinates to the polygon's space

Background: Texture Generation

- Gardner – Summing sine waves to create a texture [2]
 - Similar to a Fourier series
- Perlin – “Noise” [3] and Lewis – Isotropic noise functions [4]
 - Isotropic: No orientation
- Witkin and Kass – Beginnings of Reaction-Diffusion [5]



Public Domain,
<https://en.wikipedia.org/w/index.php?curid=10409833>

Background: Texture Mapping

- “Assigning texture coordinates from a rectangle to the vertices of the polygons in a model”[1]
- Mapping depends on the model shape
 - Should match the object’s coordinate system
- Projecting texture onto an object’s surface
- Solid textures
 - Function defining color of the object at specific point in space

Reaction-Diffusion

- Chemicals diffuse and interact with each other, promoting or discouraging growth of other chemicals

$$\frac{\partial a}{\partial t} = F(a, b) + D_a \nabla^2 a \quad \text{[General form of a two-chemical reaction-diffusion]}$$

$$\frac{\partial b}{\partial t} = G(a, b) + D_b \nabla^2 b \quad \text{[1]}$$

reactions of a and b and the diffusion of a from places nearby [1]

- D_a : Constant, speed of a 's diffusion
- $\nabla^2 a$: Concentration of a compared to neighbor concentrations of a
 - Decides where a will diffuse to/away from

One-Dimensional R-D Example

$$\Delta a_i = s (16 - a_i b_i) + D_a (a_{i+1} + a_{i-1} - 2a_i)$$

$$\Delta b_i = s (a_i b_i - b_i - \beta_i) + D_b (b_{i+1} + b_{i-1} - 2b_i)$$

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Change in chemical concentration over time

Concentrations of a and b and a scaling/shape regularity constant (Reaction of a and b)

Concentrations of chemical at a point compared to its neighbors (Diffusion)

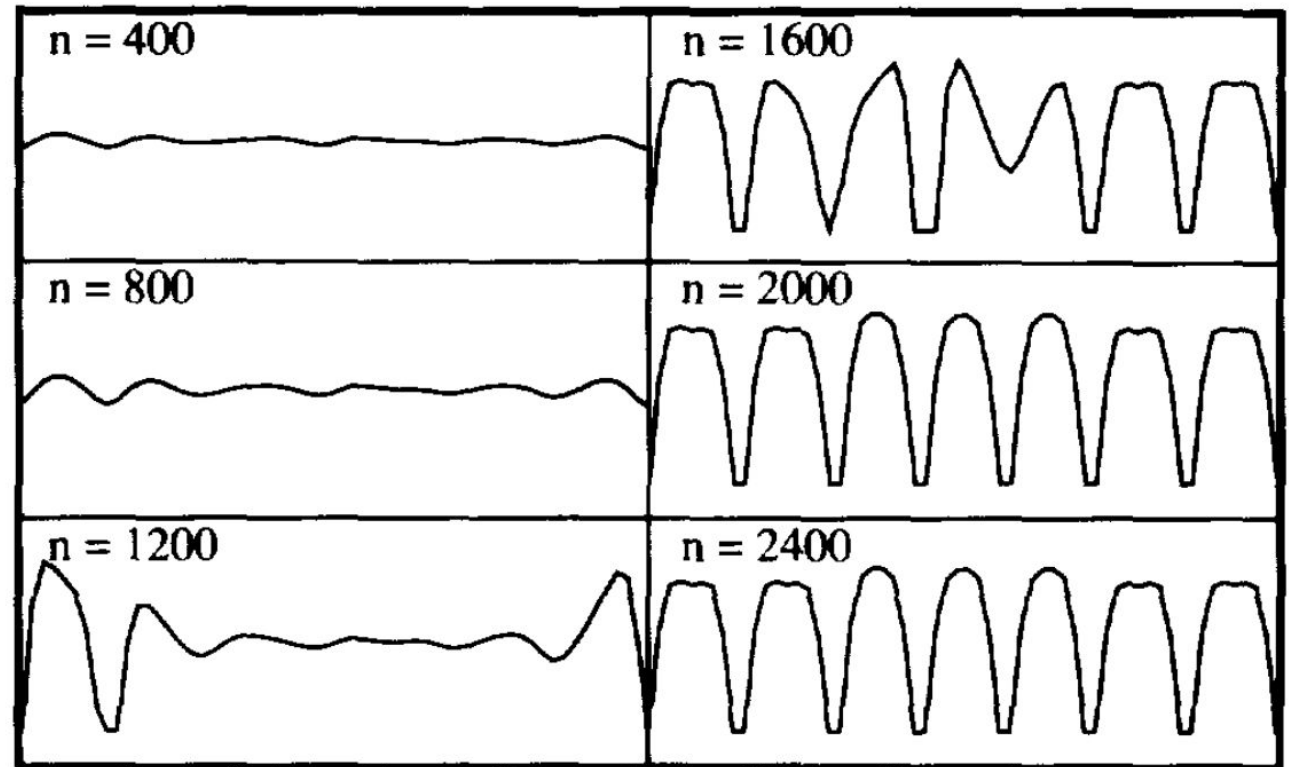
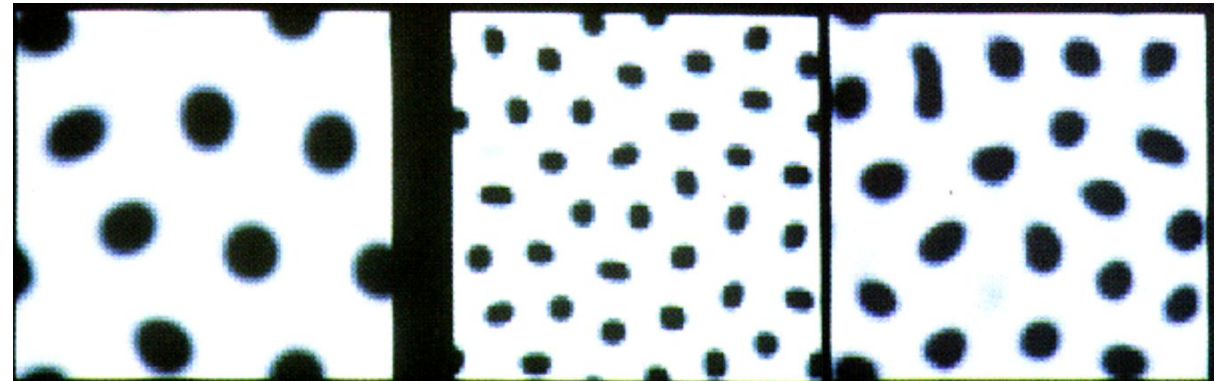


Figure 1: One-dimensional example of reaction-diffusion. [1]
Chemical concentration is shown in intervals of 400 time steps.

Two-Dimensional R-D

- Inhibitor: High values of the inhibitor prevent other chemicals from concentrating
- Chemical $a \rightarrow$ Spots \rightarrow Prevent other spots from forming nearby
- Spots form in low concentrations of b
- s parameter affects spot size
- β affects consistency in shape

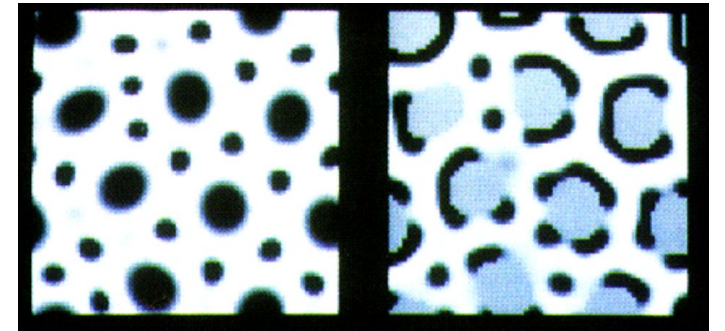
$$\Delta a_{ij} = s(16 - a_{ij}b_{ij}) + D_a(a_{i+1,j} + a_{i-1,j} + a_{i,j+1} + a_{i,j-1} - 4a_{ij})$$
$$\Delta b_{ij} = s(a_{ij}b_{ij} - b_{ij} - \beta_{ij}) + D_b(b_{i+1,j} + b_{i-1,j} + b_{i,j+1} + b_{i,j-1} - 4b_{ij})$$



Complex Patterns

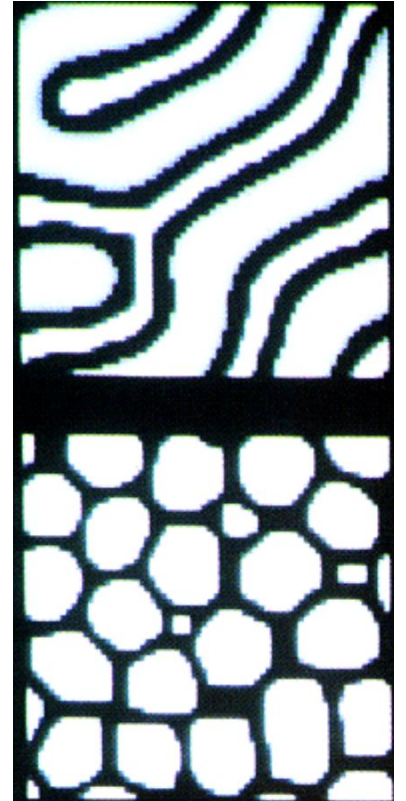
- Cascade Process
 - First system lays out a pattern
 - Second system modifies the pattern further
- Left figure (Spots)
 - Large spots first
 - Cells are “frozen”
 - Small spots synthesized in the non-frozen region
- Right figure (Leopard)
 - Large spots first
 - Cells are frozen and a and b concentrations $\rightarrow 4$
 - Smaller spots tend to form AROUND large spots due to concentration values

[1]



Complex Patterns (Continued)

- Additional examples include Lionfish and Giraffe textures
- Lionfish
 - Large stripes formed first
 - Small stripes form in-between “frozen” large stripes
- Giraffe
 - Irregularly shaped spots initially created
 - Stripes form between the spots to create the “web-like pattern called reticulation” [1]



[1]

[1]

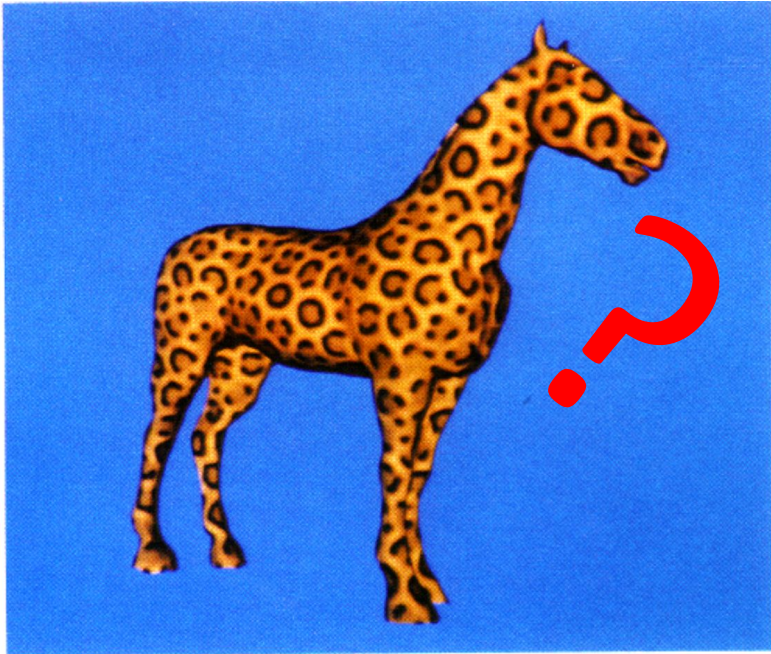


Figure 4: Leopard-Horse

[1]

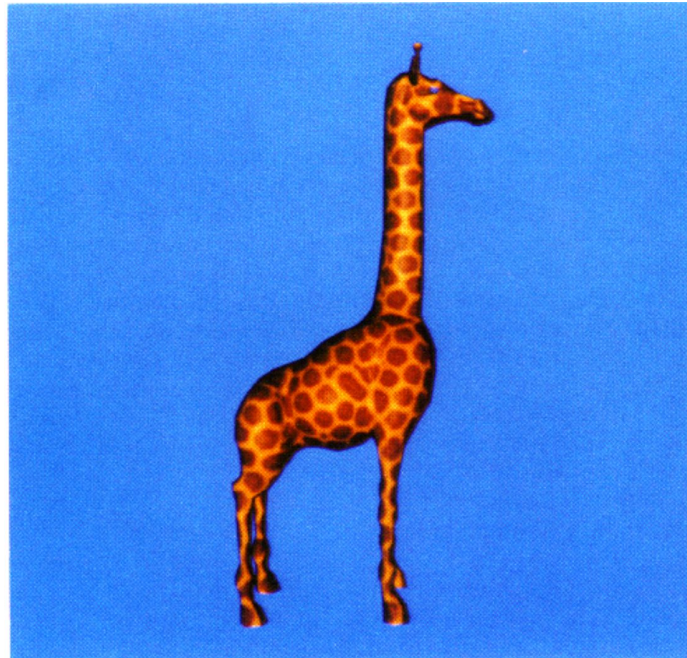


Figure 5: Giraffe

[1]

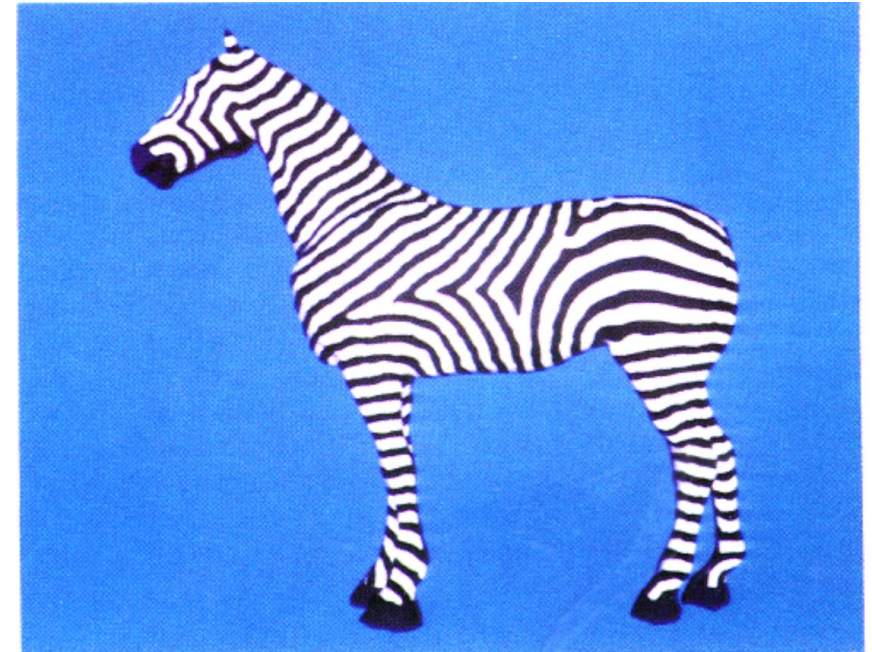


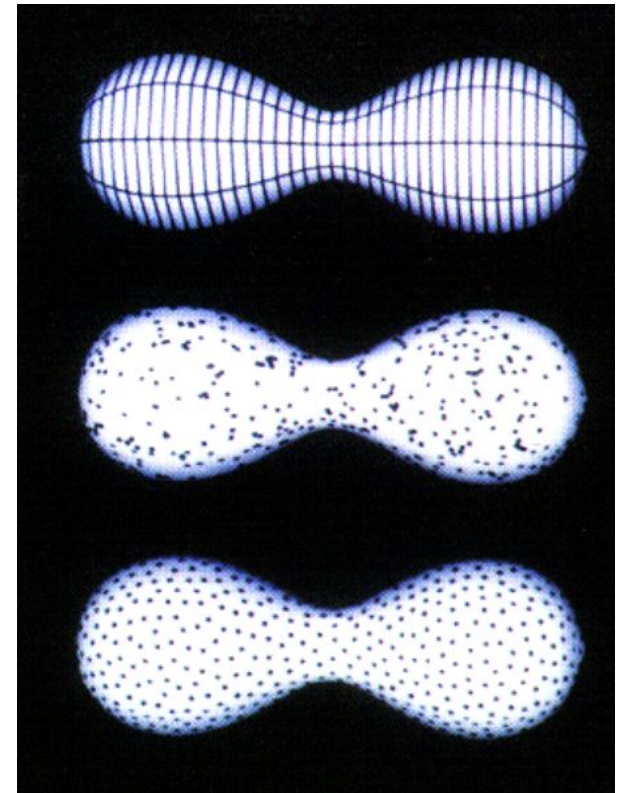
Figure 6: Pseudo-Zebra

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Varied R-D parameters across the model so that the stripes matched its animal counterpart at every location of the body

Mesh Generation

- Mesh must be created that matches model's shape
 - Mesh is made for texture synthesis
- Points created across mesh
- Relaxation sets these points to be evenly spaced
- Points pushed away at distance of repulsion radius
- Points can be pushed off of a polygon
 - Rotated about polygon's edge it crossed until it lies on the surface of a polygon



$$r = 2 \sqrt{a / n}$$

a = area of surface

n = number of points on surface

Voronoi Regions

- “Given a set of points S in a plane, the Voronoi region of a particular point P is that region on the plane where P is the closest point of all points in S ” [1]
- Helps decide what cells are adjacent for R-D
- Voronoi Regions can be found in $O(n)$ time
 - Points within distance $2r$ of a point are in the Voronoi Region
- R-D is then run on the Voronoi Regions of the mesh
- Chemical concentrations found by R-D must be rendered into texture

Rendering

- A function is used to convert chemical concentrations to texture color
- When mapping colors, repulsion radius and scaling can cause texture aliasing
 - Blurring texture colors reduces effect efficiently
 - Supersampling (rendering texture with higher resolution and scaling down) not efficient
- Bump mapping possible by using the gradient of the cell colors
 - Perturbation vector added to “surface normal” to give lighting effect of bumpy or rough texture [3]
 - Normal used “to determine a surface's orientation toward a light source” [8]

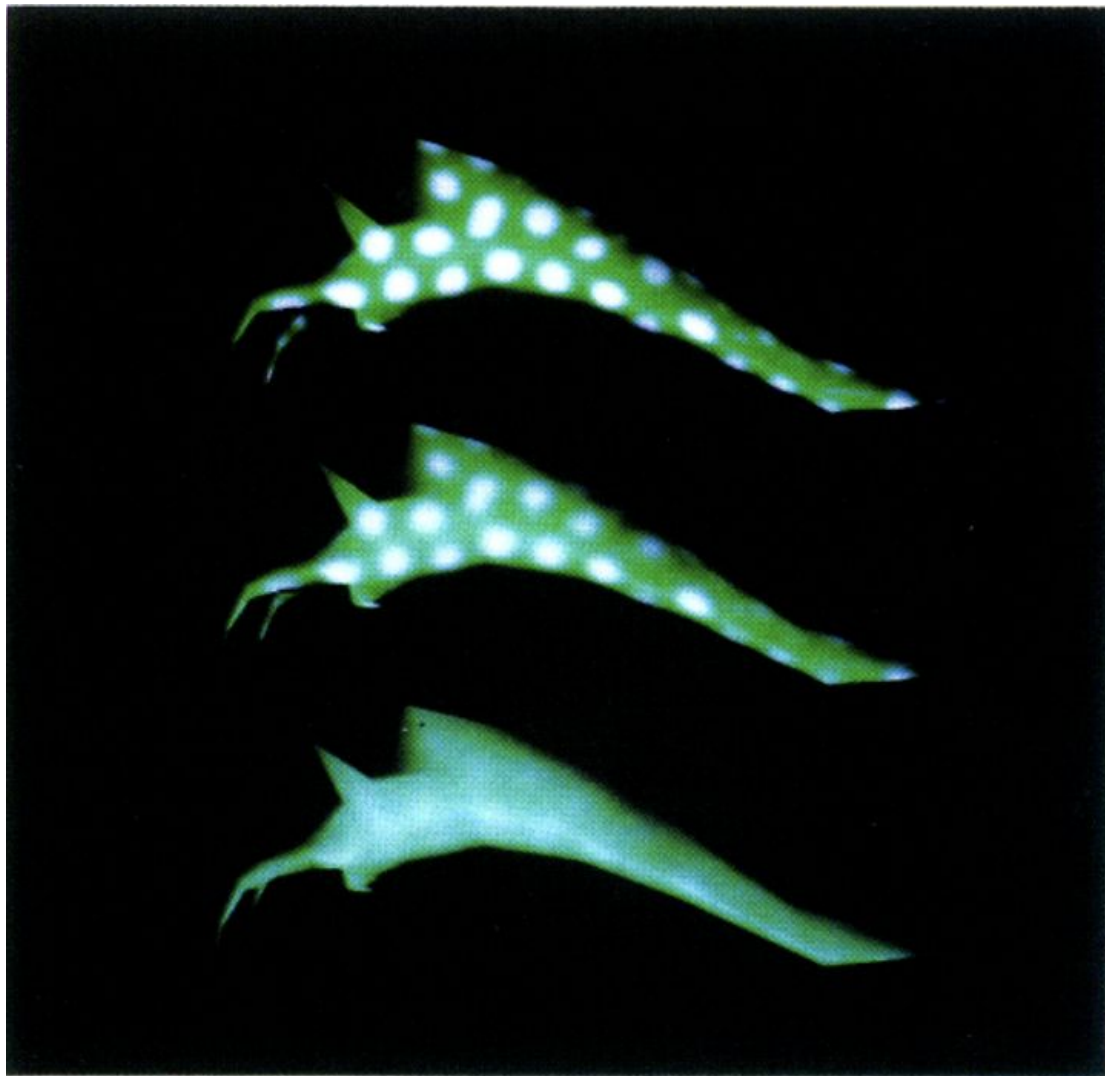
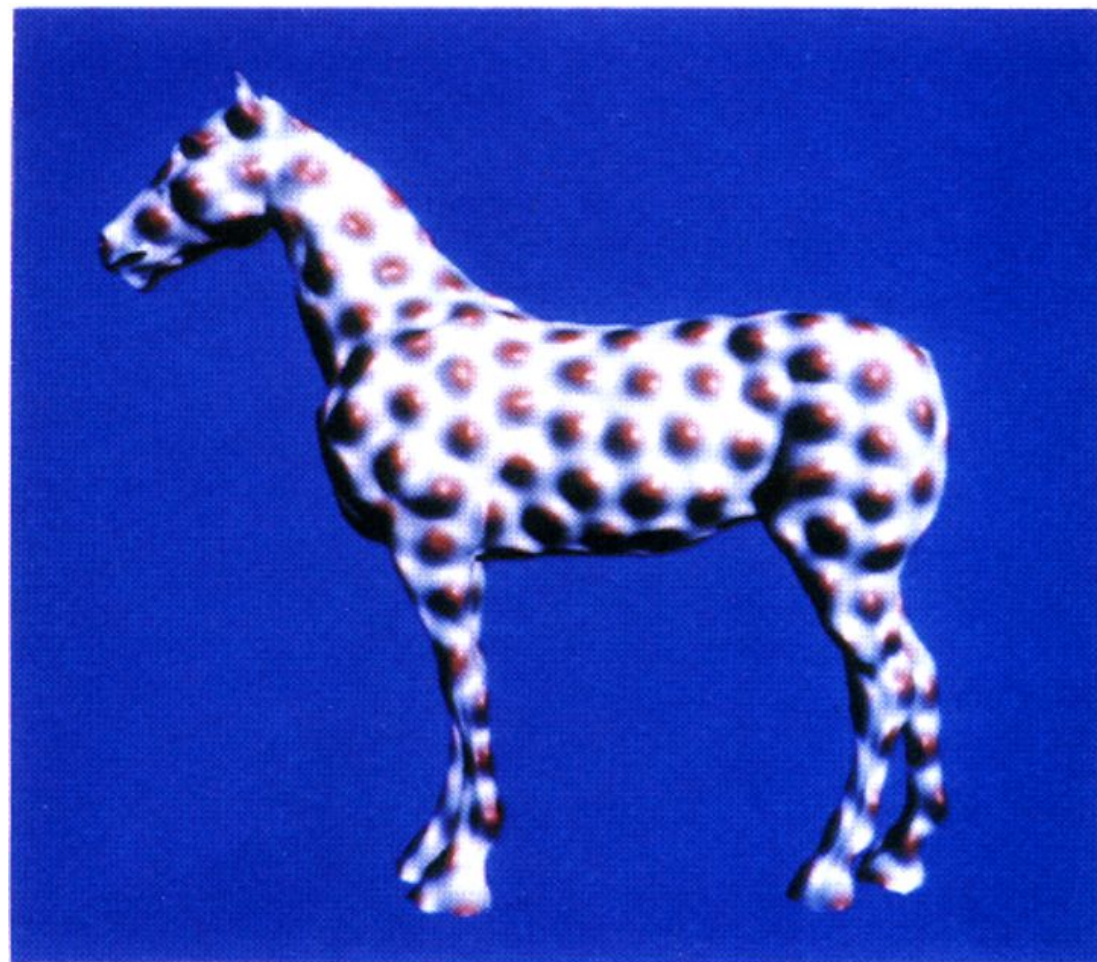


Figure 9: Blur levels for anti-aliasing [1]



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Figure 10: Bump mapping

[1]

Conclusion

- Took several hours to render each texture with reaction-diffusion
- Rendering models was much quicker
- Leopard Horse took 70 seconds to render to the model at 512x512
- Model without texture took 16 seconds
- “Such texture synthesis times would seem to prohibit much experimenting with reaction-diffusion textures” [1]

Strength 1 – In-Depth Background on Alternative Methods

- Provides the reader with a solid background on texture synthesis and rendering methods
- Shows how the problem domain is important
- Validates their work as an effective alternative with performant results
- Their research addresses a growing area (developmental biology) with texture generation

Strength 2 – Slowly Developed Complexity

- Initially show small reaction-diffusion demos affecting size/shape
- Add cascading process to create more complex dualities between chemicals
- Add multiple (5) chemicals to system for additional intricacy
- Include mesh generation for practical applications
- Shows each step individually to see how they all contribute to a complex system

Strength 3 – Many Applications for Research

- Inherent applications in developmental biology
 - Used reaction-diffusion to accurately generate popular animal coats
 - Validates the process as possible candidate in embryo development
- Computer graphics/Videogames, Movies, Etc.
 - Procedural content generation, especially with dynamic meshes
 - Creating unique textures time consuming. Mapping them to dynamic shapes even more difficult.



[1]

Weakness 1 – Relatively Simple Models Used

- Good introduction into texture/mesh generation
- For practical uses, 3D models in simulations and games are much more complex
 - Polygons continuously added to modern models
- On a workstation at the time of publishing, each texture “took several hours to generate” [1]
- Might not be as useful in real world
- Does the runtime linearly scale with model detail? (Voronei region detection is $O(n)$)

Weakness 2 – Liberties Taken in Code

- Paper is focused on biological systems
- Made algorithm choices that would not mimic biology
- To get appealing stripe patterns...
 - Run specific “Meinhardt” stripe algorithm [1]
 - Freeze cells that belong to a white stripe region
 - Run another system with altered parameters to create thinner stripes
- Too many specific changes so that their desired behavior would emerge
- Perhaps explain why this liberty was taken
 - Example of this “freezing” occurring in nature

Weakness 3 – Too Focused Biologically

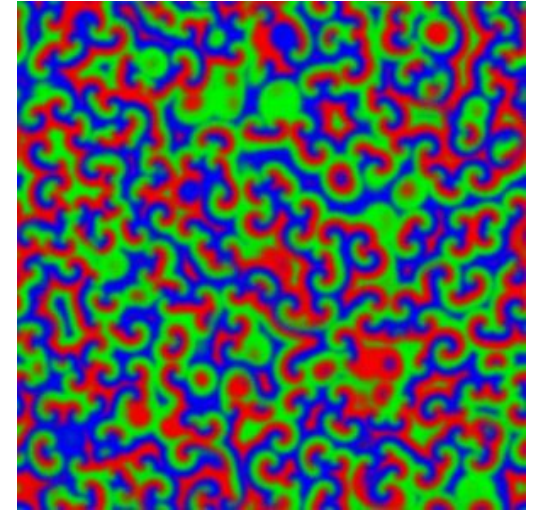
- Every pattern they discussed was deeply rooted in animal coloring
- Might have suppressed other aesthetically pleasing results by focusing on biology
- Texture generation could be important in other areas
 - Games as discussed previously
 - Design (Graphics or abstract interior design)
- What NEW patterns can we find, rather than replicate?
- If this paper focused more on the texture synthesis than developmental biology, more fields could benefit from their work
 - Put themselves into a "niche"

Extension 1– Branching Out of Biology

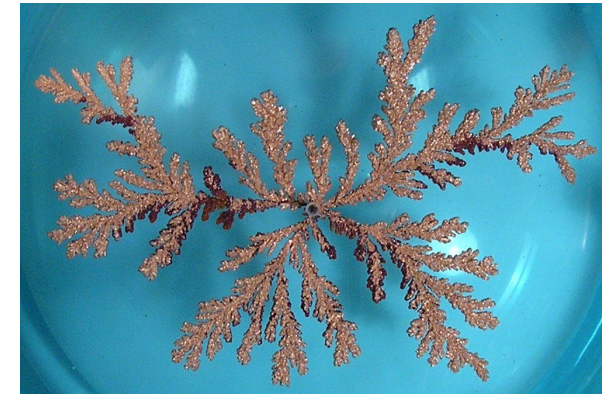
- This study was a good introduction in reaction-diffusion
- Recreating animal textures can validate the system's performance
- Create a more exploratory study in the future
- What texture synthesis results are possible that are unique and can be applied in other fields or systems?
- Is reaction-diffusion's performance acceptable for computer graphics work? Could it change the field of procedurally generated content (PCG)

Extension 2 – Additional Texture Generation Methods

- Borrow from Cellular Automata and reality
- Belousov–Zhabotinsky (BZ) reaction
 - Theoretical Chemistry reaction without equilibrium
 - Evolve “chaotically” [6]
- Diffusion-Limited Aggregation
 - Seen in some organic systems, like mineral deposits
 - Studied through computer simulations [7]



[6]



[7]

References

- [1] Turk, Greg. "Generating textures on arbitrary surfaces using reaction-diffusion." *ACM SIGGRAPH Computer Graphics*. Vol. 25. No. 4. ACM, 1991.
- [2] Gardner, Geoffrey Y. "Visual simulation of clouds." *Acm Siggraph Computer Graphics*. Vol. 19. No. 3. ACM, 1985.
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