

Extensive exploration of various ABM implementations suggested that four rules, derived from four main regulatory events, were sufficient to recreate the structuring of the blastocyst. Further, we required that all of the included genes be regulated by the known early cell fate transcription factors, by demanding significant differential transcript accumulation in all three non-hair fate mutants *wer myb23*, *gl3 egf3*, and *ttg* relative to the hair fate mutant *cpc try* in two independent labs.

## **Erzberger Group**

ABMs RD models employ relatively simple PDEs, which are sufficient for generic analyses of morphogen gradients and their interactions.

### **Testing Turing's theory of morphogenesis in chemical cells**

CHEM 116 General Chemistry I 3-3-4 Prerequisite: CHEM 115 This is the second course in the General Chemistry series. However, the system evolves sufficiently slowly that it can adiabatically exhibit the dynamical instabilities predicted by Turing for open systems , ,.

### **A Dynamical Paradigm for Molecular Cell Biology**

With respect to the root hair branching character, the composite Bayesian network identified two root-hair genes, bHLH66 and AT4G13390 encoding a proline-rich extensin-like wall protein , as the best predictors of the degree of root hair branching in an inverse correlation. For the analysis of root hair branching, 50 root hairs were examined per root in each of 9 seedlings 450 total hairs. These rather vague objectives can in retrospect be attributed to the fact that ABMs were relatively new in the field and that it was necessary to gain experience with exploring specific model features and the role of synergism among processes and rules.

### **Molecular control of cell differentiation and morphogenesis; a systematic theory.**

Quantitative experimental results obtained using this artificial cellular system establish the strengths and weaknesses of the Turing model, applicable to biology and materials science alike, and pinpoint which directions are required for improvement. Although quantifiable validation is limited, the model is capable of creating tissue-scale morphological features that depend solely on single-cell decisions in response to environmental cues.

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