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Trainable Gene Regulation Networks with Applications to \textit{Drosophila} Pattern Formation

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1 Introduction

This chapter will very briefly introduce and review some computational experiments in using trainable gene regulation network models to simulate and understand selected episodes in the development of the fruit fly, \textit{Drosophila melanogaster}. For details the reader is referred to the papers introduced below. It will then introduce a new gene regulation network model which can describe promoter-level substructure in gene regulation.

As described in chapter 2, gene regulation may be thought of as a combination of cis-acting regulation by the extended promoter of a gene (including all
regulatory sequences) by way of the transcription complex, and of trans-acting regulation by the transcription factor products of other genes. If we simplify the cis-action by using a phenomenological model which can be tuned to data, such as a unit or other small portion of an artificial neural network, then the full trans-acting interaction between multiple genes during development can be modelled as a larger network which can again be tuned or trained to data. The larger network will in general need to have recurrent (feedback) connections since at least some real gene regulation networks do. This is the basic modeling approach taken in (Mjolsness et al. 1991), which describes how a set of recurrent neural networks can be used as a modeling language for multiple developmental processes including gene regulation within a single cell, cell-cell communication, and cell division. Such network models have been called "gene circuits", "gene regulation networks", or "genetic regulatory networks", sometimes without distinguishing the models from the actual modeled systems.

In (Mjolsness et al. 1991) a number of choices were made in formulating the trainable gene regulation network models, which affect the spatial and temporal scales at which the models are likely to be useful. The dynamics was chosen to operate deterministically and continuously in time, on continuous-valued concentration-like variables, so that the dynamical equations for the network are coupled systems of ordinary differential equations (ODE's). One such form was
\[ \tau_i \frac{dv_i}{dt} = g \left( \sum_j T_{ij} v_j + h_i \right) - \lambda v_i \]

in which \( v_i \) is the continuous-valued state variable for gene product \( i \), \( T_{ij} \) is the matrix of positive, zero, or negative connections by which one transcription factor can enhance or repress another, and \( g() \) is a nonlinear monotonic sigmoidal activation function. When a particular matrix entry \( T_{ij} \) is nonzero, there is a regulatory "connection" from gene product \( j \) to gene \( i \). The regulation is enhancing if \( T \) is positive and repressing if it is negative. If \( T_{ij} \) is zero there is no connection. Figure 1 sketches the model, drawing a few representative nonzero connections as arrows between "genes" represented by open circles. The entire network is localized to a cell but communicates with other such networks in nearby cells.

[Figure 1 goes here.]

Such equations are often stiff due to the nonlinear transfer function \( g(u) \). Optimizing the unknown parameters \( T, \ldots \) and \( \ldots \) has so far proven to be computationally difficult: special versions of simulated annealing optimization (Lam and Delosme 1988a, 1988b) have been required for good results, e.g. to start from expression patterns derived from a known model and recover its parameters reliably (Reinitz and Sharp 1995). As discussed in Chapter 2, this kind of training is quite different and much slower than the usual
"backpropagation of error" training used with feed-forward (nonrecurrent) artificial neural networks. Informatics work on improving this situation could be important.

In addition to the analog circuit model, the framework of (Mjolsness et al. 1991) also proposes a dynamic grammar by which multiple biological mechanisms can be modeled by networks and then combined into a consistent overall dynamical model. The grammar/circuit combination has some similarities to hybrid models incorporating Discrete Event Systems and ODE's. In this way one can for example combine intracellular and intercellular regulation network submodels. The grammar is also suitable for implementation in object-oriented computer simulation programs.

2. Three Case Studies

In this section, some of the literature on trainable gene circuit models which have been fit to *Drosophila* gene expression patterns is reviewed. Three applications to pattern formation are shown to demonstrate the generality of the methods. First, a model of gap gene expression patterns along the anterior-posterior axis will be described. Second, the extension of this model to incorporate the important pair-rule gene *eve* and a number of other improvements will be introduced. Finally, a
gene circuit model of neurogenesis incorporating nonlinear signaling between nearby cells through the Notch receptor and the Delta ligand will be briefly described.
2.1 Gap Gene Expression

Such gene regulation network models can be tuned or "trained" with real gene expression data, and then used to make robust and at least qualitatively correct experimental predictions, as was shown in (Reinitz et al. 1992). In that study the goal was to understand the network of gap genes expressed in bands (domains) along the anterior-posterior (A-P) axis of the very early embryo (the syncytial blastoderm) of *Drosophila*. This experimental system has the advantage that there are no cell membranes between adjacent cell nuclei, so elaborate cell-cell signalling mechanisms do not need to be modeled. Also *Drosophila* is an easy species to manipulate genetically, as for example "saturation mutagenesis" - finding all the genes affecting a particular process - is possible.

Positional information along the A-P axis of the syncytial blastoderm is encoded in a succession of different ways during development. At first the main encoding is a roughly exponential gradient of *bicoid (bcd)* protein imposed by the mother fly, along with maternal *hunchback (hb)* expression. These provide gene regulation network inputs to the gap genes: *Kruppel (Kr)*, *knirps (kni)*, *giant (gt)*, *tailless (tll)*, and *hunchback (hb)* again. These each establish one or two broad domains of expression along the A-P axis. The gap genes then serve as network inputs to the pair-rule genes including *even-skipped (eve)* and *fushi tarazu (ftz)*, which establish narrow, precise stripes of expression and precise positional
coding. These in turn provide input to segment-polarity genes such as *engrailed* and *wingless* which are the first to retain their expression pattern into adulthood. For example, *engrailed* is expressed in bands just one cell wide which define the anterior borders of the parasegments. Introductions to the relevant *Drosophila* developmental biology may be found in (Lawrence 1992) and (Alberts et al. 1994).

An example of a spatial gene expression pattern along the A-P axis of a triple-stained embryo is shown in Figure 2. Here, fluorescently labelled antibodies simultaneously label those nuclei in the syncytial blastoderm expressing *Kruppel*, *giant*, and *even-skipped*.

[Figure 2 goes here.]

The first computer experiments with fitting such analog gene regulation nets to real expression data concerned the establishment of the broad gap gene domains (excluding the extreme ends of the A-P axis) from maternally supplied initial conditions, by a gene regulation network in which all gap genes interact with all others and *bcd* provides input to, but does not receive any input from, the gap genes.
Figure 3 shows the experimentally observed and model-fitted curves for gap gene expression. They are in qualitative agreement, which is the most that can be expected from the expression data that was available at the time. The extra dip in $gt$ expression could not be predicted by the model, which can be interpreted as an indication of the role of circuit components not included in the model.

The most important predictions of the model concerned the anomalous dose-response observed by (Driever and Nusslein-Volhard 1988). Figure 4 shows the prediction in detail; it may be summarized by saying that positional information for the gap gene system is specified cooperatively by maternal $bcd$ and $hb$. This qualitative behavior was observed to be robust over many runs of the simulated annealing parameter-fitting procedure, and therefore taken to be a prediction of the model. Essential features of the cooperative control of positional information by maternal $bcd$ and $hb$ were verified experimentally in (Simpson-Brose et al. '1994). The gap gene model prediction and the experiment occurred independently of one another.
2.2 Eve Stripe Expression

Following the gap gene computer experiments, (Reinitz and Sharp 1995) went on to perform a detailed study of the gap gene circuit as extended to include the first of the pair-rule genes, \textit{eve}. The further observations which could be included in this model allowed an important milestone to be reached: not only qualitative behaviors, but also the circuit parameter signs and rough magnitudes became reproducible from one optimization run to another, and some parameters such as connections to \textit{eve} were still more reproducible. Hence, far more could be predicted. For example the diffusion constant for \textit{eve} was much lower than for other transcription factors in successful runs. This has an experimental interpretation: \textit{eve} mRNA is expressed in the outer part of each future cell just as the cell membranes are invaginating into the blastoderm embryo, providing an apical obstruction to diffusion.

More importantly, each of the eight boundaries of the four central stripes of \textit{eve} expression could be assigned a particular gap gene as the essential controller of that boundary. This picture is in agreement with experimental results with the possible exception of the posterior border of \textit{eve} stripe 3, the interpretation of which is an interesting point of disagreement (Small et al. 1996, Reinitiz and
Sharp 1995, Frasch and Levine 1987) and a possible focal point for further laboratory and/or computer experiments.

Further experimental understanding of the gap genes' influence on eve expression is obtained in (Reinitz et al. 1998), where it is shown that the fact that eve is unregulated by other pair-rule genes can be understood by the phase of its periodic spatial pattern: no other phase offset pattern of pair-rule expression (e.g. the phase-shifted patterns of hairy or fushi-tarazu) can be produced from gap gene input alone.

[Figure 5 goes here.]

Related work on modeling the gap gene and eve system of A-P axis positional information in Drosophila includes (Hamahashi and Kitano 1998).

2.3 Neurogenesis and Cell-Cell Signaling

The syncytial blastoderm is very favorable, but also very unusual, as morphogenetic systems go because there is no cell membrane interposed between nearby cell nuclei and therefore the elaborate mechanisms of cell-cell signaling do not come into play. But if we are to model development in its generality it is essential to include signaling along with gene regulation.
networks. As a first attempt in this direction, we have modeled the selection of particular cells in an epithelial sheet (later in *Drosophila* development) to become neuroblasts. Virtually the same gene network is thought to be involved in the selection of particular cells in wing imaginal disks to be sensor organ precursors. The essential molecule to add is the Notch receptor, a membrane-bound receptor protein responsible for receiving the intercellular signals which mediate this selection process. It binds to a ligand molecule ("Delta" for this system) on neighboring cells. Recent experiments (Schroeter et al. 1998) indicate that it acts on the nucleus (following activation by a ligand on another cell) by having an intracellular domain cleaved off and transported there. Variants of the Notch receptor occur in many developmental subsystems where a subpopulation of cells must be picked out, in *Drosophila* and homologously across many species.

In (Marnellos 1997) and (Marnellos and Mjolsness 1998a, 1998b) are reported computer experiments incorporating both intracellular and intercellular components in a gene regulation network model of neurogenesis. A minimal gene circuit model with lateral inhibition (such as depicted in Figure 6) was not quite sufficient to produce the observed patterns of selection robustly. Incorporating a denser intracellular connection matrix and/or the dynamic effects of delamination on the geometry of cell/cell contact area produced better results. However, the "data" to which the fits were made was highly abstracted from real gene expression data so it is premature to draw a unique biological
hypothesis from the model. Figure 7 shows the resulting model behavior in the case of dense interconnections.

[Figure 6 goes here.]

Related work on Notch-mediated signaling in *Drosophila* developmental models includes the appearance of Notch and Delta in the ommatidia model of (Morohashi and Kitano 1998).

[Figure 7 goes here.]

3 Extending the Modeling Framework to Include Promoter Substructure

A very important scientific problem is to understand the influence of promoter substructure on *eve* stripe formation. The *eve* promoter has many transcription factor binding sites, some of which are grouped more or less tightly into promoter elements such as the stripe 2 “minimal stripe element” (MSE 2) (Small et al. 1992), or a similar less tightly clustered element for stripes 3 and 7 (Small et al. 1996). As an example of the scientific problems that are raised, if is *hb* an enhancer for MSE 2 but an inhibitor for MSE 3, what is its net effect on *eve* and can it change sign (Reinitz et al. 1998)? And how are we to understand the action of “silencer” elements such as the one apparently responsible for long-range
repression of *zen* by *dorsal* (Gray et al. 1995)? Such questions point to the need for at least one additional level of complication in the phenomenological models of gene networks whose application is described above, to describe the substructure of promoters: binding sites, their interactions, and promoter elements. Otherwise the relevant experiments cannot even be described, let alone predicted, with network models.

In (Small et al. 1992) an informal model for activation of MSE 2 is suggested: it is activated by *bcd* and *hb* "in concert", and repressed by *gt* anteriorly and *Kr* posteriorly. A simple "analog logic" expression for the activation of MSE 2 in terms of variables taking values in [0,1] might then be (GRN 1998):

\[
\begin{align*}
U_{MSE2} &= (bcd + \gamma \times hb)(1 - gt)(1 - Kr) \\
V_{MSE2} &= g(U_{MSE2})
\end{align*}
\]

where _ is a weight on the relative contribution of *hb* vs *bcd*. A similar simplified formula for the model of (Small et al. 1996, figure 8) for MSE 3 could be for example:

\[
\begin{align*}
U_{MSE3} &= Dstat(1 - hb)(1 - kni) \\
V_{MSE3} &= g(U_{MSE3})
\end{align*}
\]
(We omit direct activation of MSE3 by tailless (tll) since tll represses kni (Pankratz et al. 1989) which represses MSE3.) The rate of eve transcription would be approximated by a further analog logic formula including a weighted “or” of the MSE activations $v_{MSE2}$ and $v_{MSE3}$.

The validation or invalidation of such formulae and their interpretation in terms of more detailed models will require a quantitative treatment of the relevant expression data which is not yet available. It may also lead to fitting the parameters in quantitative network models of promoter-level substructure within a gene regulation network.

3.1 An Example: Hierarchical Cooperative Activation

As an example of such a gene network model incorporating promoter level substructure, I introduce here a “Hierarchical Cooperative Activation” (HCA) model for the degree of activation of a transcription complex. It at least seems more descriptive of known mechanisms than a previous attempt to derive phenomenological recurrent neural network equations as an approximation to gene regulation dynamics (Mjolsness et al. 1991). An earlier suggestion for including promoter-level substructure in gene regulation networks is described in (Sharp et al. 1993). The present HCA model is more detailed but has not been
fit to any experimental data yet and is therefore quite speculative: perhaps a next stage of successful modeling will include some of the following ingredients.

The basic idea of the model is to use an equilibrium statistical mechanics model (complete with partition functions valid for dilute solutions (Hill 1985)) of "cooperative activation" in activating a protein complex. Such a model can be constructed from the following partition function, which is essentially the Monod-Wyman-Changeux (MWC) model for a concerted state change among subunits (Hill 1995):

$$Z = K \prod_b (1 + K_b v_j(b)) + \prod_b (1 + K_b v_j(b))$$

in which the probability of activation of some complex is determined by relative binding constants for each component $b$ of the complex in the active and inactive states, but there are no other interactions. As before, $v_j$ represents the concentration of gene product $j$ of a gene circuit. In this formula, $j$ is a function of $b$ so that each binding site is specialized to receive only one particular transcription factor. To remove this assumption one could write instead

$$Z = K \prod_b (1 + \sum_j A_{b_j} K_{b_j} v_j) + \prod_b (1 + \sum_j A_{b_j} K_{b_j} v_j)$$

where $A = 0$ or $1$ specifies which transcription factors may bind to which sites by its sparse nonzero elements. For either expression, $K$ is the relevant binding
constant for a binding site when the complex is in its “active” state and $\hat{K}$ is the binding constant when the complex is inactive.

For this partition function, given a global active or inactive state, all binding sites are independent of one another. For example the components could be the occupants of all the binding sites $b$ within a particular regulatory region of a eukaryotic promoter. This conditional independence leads to the products over the binding sites in the expression for $Z$. There are two such products because there is one additional bit of global state which can be “active” or “inactive”.

For this model the probability of activation of the complex under consideration can be calculated and it is:

$$P = g(u) = \frac{Ku}{1 + Ku}$$

$$u = \prod_b \left( \frac{1 + K_b v_j(b)}{1 + \hat{K}_b v_j(b)} \right)$$

so (if $Ku \ll 1$)

$$u = 1 + \sum_b (K_b - \hat{K}_b)^{v_j(b)}.$$  

(Further simplifications result if the binding constants $K_b$ specific for a given transcription factor $j(b)$ are all roughly equal to a common value $K_j$. The final line above suggests a neural-network like approximation for $u$, although in that
regime \( g \) could be linearized also.) We will use this model as a building block to construct a more detailed one.

Given the MWC-style model of "cooperative activation", we'd like to use it hierarchically: to describe the activation of promoter "modules" or "elements" in terms of transcription factor concentrations, and then again to describe the activation of the whole transcription complex in terms of the "concentrations" of active promoter elements, which are proportional to their activities. An additional wrinkle is to allow either monomeric or homodimeric transcription factor binding. (Heterodimers will be introduced with appropriate notation later.) The resulting bare-bones hierarchical model would replace the neural-net activation dynamics

\[
\tau_i \frac{dv_i}{dt} = [\text{transcribing}]_i - \lambda v_i
\]

\[
[\text{transcribing}]_i = g(u_i)
\]

\[
u_i = \sum_j T_{ij}v_j + h_i
\]

with the two-level model

\[
\tau_i \frac{dv_i}{dt} = [\text{transcribing}]_i - \lambda v_i
\]

\[
[\text{transcribing}]_i = g(u_i) = \frac{Ju_i}{1 + Ju_i}
\]

\[
u_i = \prod_{a \in i} \left( \frac{1 + J \alpha P_\alpha}{1 + J \alpha P_\alpha} \right)
\]
(the product is taken over enhancer elements which regulate genei) and

\[ P_a = g_a(\tilde{u}_a) = \frac{\tilde{K}_a \tilde{u}_a}{1 + \tilde{K}_a \tilde{u}_a} \]

\[ \tilde{u}_a = \prod_{b \in \alpha} \left( 1 + \frac{K_b \nu_{j(b)}^{n(b)}}{1 + \hat{K}_b \nu_{j(b)}^{n(b)}} \right) \]

Here \( n(b) = 1 \) for monomers and 2 for homodimers. Note that for this simple feed-forward version of the model, the parameters \( K_b \) and \( \hat{K}_b \) are related to observables

\[ f_{ab} = \frac{K_b \nu_{j(b)}^{n(b)}}{1 + K_b \nu_{j(b)}^{n(b)}} \quad \hat{f}_{ab} = \frac{\hat{K}_b \nu_{j(b)}^{n(b)}}{1 + \hat{K}_b \nu_{j(b)}^{n(b)}} \]

where \( f_{ab} \) is the probability that site \( b \in \alpha \) is occupied if \( \alpha \) is active, and \( \hat{f}_{ab} \) is the probability that site \( b \) is occupied if \( \alpha \) is inactive. In principle these quantities could be observed by in vivo footprinting. Such observations could be used to evaluate the parameters \( K_b \) to use in the first expression for \( u_a \) for arbitrary inputs \( \nu_j \).

If we are modeling a network rather than a single gene, then some of the quantities listed above require an additional \( i \) index.
We have the opportunity to include a few more important biological mechanisms at this point. One is the possibility that, as in the Endo16 model of (Yuh et al. 98), the hierarchy could go much deeper than two levels - especially if transcription complex formation is a sequential process. Another significant mechanism is competitive binding within a promoter element. This could arise if several transcription factors bind to a single site, as we have formulated earlier, or if binding at one site eliminates the possibility of binding at a nearby site and vice versa. In this case the 4-term product of two 2-term binding-site partition functions is replaced with one three-term function by excluding the configuration in which both competing sites are occupied:

\[
Z_{bb'} = (1 + \sum_j A_{bj} K_{bj} v_j + \sum_k A_{bk} K_{bk} v_k)
\]

\[
\hat{Z}_{bb'} = (1 + \sum_j A_{bj} \hat{K}_{bj} v_j + \sum_k A_{bk} \hat{K}_{bk} v_k)
\]

(where again \(A = 0\) or \(1\) describes which transcription factors bind to which sites by its sparse nonzero elements) with corresponding modifications to the update equations. Also homodimeric and heterodimeric transcription factor binding are easy to accommodate with appropriate concentration products in more general one-site and two-site partition functions:
Transcription factor trimers and higher order subcomplexes at adjacent binding sites could be described by suitable generalizations of these expressions, at the cost of introducing more parameters.

Similarly, constitutive transcription factor binding with activation by phosphorylation or dephosphorylation can be described with minor modifications of the appropriate one-site or two-site partition functions. For example one could use Michaelis-Menton kinetics in steady-state for phosphorylation and dephosphorylation, and the one-site dimeric partition functions would become

\[
Z_b^{(1)} = (1 + \sum_{j} A_{b_j K_{b_j}} v_j + \sum_{j} A_{b_j k} K_{b_j k} v_j v_k)
\]

\[
\hat{Z}_b^{(1)} = (1 + \sum_{j} A_{b_j K_{b_j}} v_j + \sum_{j} A_{b_j k} K_{b_j k} v_j v_k)
\]

\[
Z_{bb'}^{(2)} = (1 + \sum_{j} A_{b_j K_{b_j}} v_j + \sum_{j} A_{b_j K_{b_j}} v_j v_k + \sum_{l m} A_{b'_l K_{b'_l}} v_j + \sum_{l m} A_{b'_l m} K_{b'_l m} v_j v_m)
\]

\[
\hat{Z}_{bb'}^{(2)} = (1 + \sum_{j} A_{b_j K_{b_j}} v_j + \sum_{j} A_{b_j K_{b_j}} v_j v_k + \sum_{l m} A_{b'_l K_{b'_l}} v_j + \sum_{l m} A_{b'_l m} K_{b'_l m} v_j v_m)
\]

where \(x_i\) is proportional to the concentration of a kinase for the bound \(j/k\) dimer (with proportionality constants depending on the catalytic reaction rates) and \(y_m\)
is proportional to a corresponding phosphatase concentration. Also $A_{hjk,lm} \leq A_{hjk}$, so that the extra indices $l$ and $m$ just specify the relevant kinase(s) and phosphatase(s) from a kinase network. For example MAP kinase mediated signaling could be modeled as activating a gene regulation network by this mechanism.

In this model formulation we have omitted lateral interactions other than competitive binding between activation of nearby binding sites. Such interactions could be modeled in the manner of an Ising model. For simplicity we just use a tree topology of states and partition functions here.

Given such one-site and two-site partition functions, the overall partition function for a promoter element in terms of its binding sites is:

$$Z_\alpha = K_\alpha \left( \prod_{b|C=0} Z_b^{(1)} \prod_{bb'|C=1} Z_{bb'}^{(2)} \right) + \left( \prod_{b|C=0} \tilde{Z}_b^{(1)} \prod_{bb'|C=1} \tilde{Z}_{bb'}^{(2)} \right).$$

Here each binding site competes with at most one other one as determined by the 0/1-valued parameters $C_b, C_{bb'}$.

In this picture, silencers are just particular promoter elements with sufficiently strong negative regulation of transcription to veto any other elements.
The Hierarchical Cooperative Activation (HCA) dynamics then become

\[ \tau_i \frac{dv_i}{dt} = \text{[transcribing]}_i - \lambda_i v_i \]

\[ \text{[transcribing]}_i = g(u_i) = \frac{J u_i}{1 + J u_i} \]

\[ u_i = \prod_{a \in i} \left( \frac{1 + J_a P_a}{1 + J_a P_a} \right) \]

and

\[ P_a = g_a(u_a) = \frac{\tilde{K}_a \tilde{u}_a}{1 + \tilde{K}_a \tilde{u}_a} \]

\[ \tilde{u}_a = \prod_{b \neq a \in C} \left( \frac{Z^{(1)}_{b}}{\tilde{Z}^{(1)}_{b}} \right) \prod_{b' \neq b \in C} \left( \frac{Z^{(2)}_{b,b'}}{\tilde{Z}^{(2)}_{b,b'}} \right) \]

with Z's as before:

\[ Z^{(1)}_b = (1 + \sum_j A_{bj} K_{bj} v_j + \sum_{jk} A_{bk} K_{bj} v_j v_k) \]

\[ Z^{(2)}_{bb'} = (1 + \sum_j A_{bj} K_{bj} v_j + \sum_{jk} A_{bk} K_{bj} v_j v_k + \sum_{im} A_{b'i} K_{b'i} v_i + \sum_{im} A_{b'lm} K_{b'lm} v_i v_m) \]

and likewise for inactive-module (hatted) Z's and K's. These partition functions encode monomeric, homodimeric and heterodimeric protein-DNA binding using the various A parameters.
The resulting HCA model (Figure 8) can describe promoter elements, silencer regions, dimeric and competitive binding, and constitutive transcription factor binding, among other mechanisms. The price is that there are considerably more unknown parameters in the model than in the previous recurrent neural network models - not exponentially many as in the general N-binding site partition function, but enough to pose a challenge to model-fitting procedures and data sets.

4 Conclusion

Gene regulation networks have been applied to model several episodes in the development of *Drosophila*, successfully making contact with experimental results. A variety of biological mechanisms including intercellular signaling can now be included in such models. We proposed a new version of gene regulation network models for use in describing experiments which involve promoter substructure, such as transcription factor binding sites or promoter regulatory elements.
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References


**Figure 1.** Sketch of recurrent analog neural network model for gene regulation networks. A set of analog-valued units $v$ are connected in a continuous-time, recurrent circuit by a connection matrix $T$. Communication with circuits in other cells may require additional connections, e.g. as formulated in [Mjolsness et al '91].

**Figure 2.** Spatial pattern of gene expression in a *Drosophila* syncytial blastoderm for two gap genes and one pair-rule gene. Immunofluorescent staining of nuclei for *Kruppel* (green), *giant* (blue), and *even-skipped* (red). Overlap areas of *Kr* and *eve* appear yellow, and overlaps of *gt* and *eve* appear purple. Image courtesy of John Reinitz.

**Figure 3.** Data and model for gap gene circuit. Horizontal axes are nuclei along lateral midline from anterior to posterior. Vertical axes are relative concentrations. Left: data estimated from immunofluorescence images similar to Figure 2 for pairs of gap genes. Right: output of a circuit model fit to expression data using a nonlinear least squares criterion and simulated annealing optimization. From [Reinitz et al. '92].

**Figure 4.** Predictions of the model as *bicoid* dosage is increased: location of selected landmarks along A-P (horizontal) axis vs. number of bcd copies (vertical axis). (a) Displacement of a landmark (anterior margin of the *Kr* domain)
expected if it were determined by reading off a fixed concentration value of maternal Bicoid protein alone. (b-c) Smaller displacement of the same landmark (anterior margin of the Kr domain) predicted by model. (A retrodiction.) (d) Observed anomalously small displacement of a related landmark: the first eve stripe, not available in the gap gene model but expected to be offset anteriorly from the Kr landmark. Note anomalously high slope compared to a, but as in b,c. (e) Prediction: return to the behavior of (a) if maternal hunchback is set equal to zero. From [Reinitz et al. '92].

Figure 5. Drosophila eve stripe expression in model (right) and data (left). Green: eve expression, red: kni expression. From [Reinitz and Sharp '95]. Courtesy J. Reinitz and D. H. Sharp.

Figure 6. A hypothesized minimal gene regulation circuit for lateral inhibition mediated by Notch and Delta. Redrawn from [Heitzler et al '96, Figure 6]. Two neighboring cells express Notch (N) and Delta (Dl) at their surfaces. Notch positively regulates transcription of genes of the Enhancer-of-split complex E(spl)-C, which negatively regulate transcription of genes of the achaete-scute complex (AS-C), which positively regulate transcription of Delta. Curved boundaries are the cell membranes between two neighboring cells. Related circuit diagrams have been suggested elsewhere e.g. [Lewis '96].
Figure 7. Cluster resolution. A circuit "trained" to resolve simple proneural cluster configurations into individual neuroblasts (or sensory organ precursor cells) is tested on more complex and irregular configurations. In this case each cluster was successfully resolved into a single neuroblast, but the large clusters resolve more slowly. Times: t=1 (top left), t=76 (top right), t=106 (bottom left), t=476 (bottom right). Similar to [Marnellos and Mjolsness '98a]; courtesy George Marnellos.

Figure 8. Hierarchical Cooperative Activation (HCA) model for promoter substructure within a gene "node" in a gene regulation network. Different layers of sub-nodes have different forms of dynamics. This network could be used to selectively expand some or all of the nodes in Figure 1, for example just the "eve" gene in a network for the gap genes and eve.
Figure 1:
Figure 3a:
Figure 3b:
Figure 4:
Figure 5:
Figure 6:
Figure 7:
Figure 8:

- Transcription output
- Promoter element activation
- Binding site activation
- Binding site occupation - dimerization, competitive binding
- Transcription factor inputs