Redundancy and error resilience in Boolean Networks

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We consider the effect of noise in sparse Boolean Networks with redundant functions. We show that they always exhibit a non-zero error level, and the dynamics undergoes a phase transition from non-ergodicity to ergodicity, as a function of noise, after which the system is no longer capable of preserving a memory of its initial state. We obtain upper-bounds on the critical value of noise for networks of different sparsity.

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Introduction— Biological systems are unavoidably noisy in their nature, but often need to function in a predictable fashion \[1\]. In such a situation, strategies to diminish the harmful effect of noise will significantly impact the fitness of a given organism. The most fundamental protection mechanism a system can adopt is the redundancy of its underlying components, since the resulting coincidences necessary to impact the proper function of the system can drastically diminish the probability of error. In this letter we are concerned with the effect of redundancy in gene regulation; in particular in a simple Boolean Network (BN) model. We assume that each component in the system is arbitrarily redundant, with the only restriction that the number of inputs per component is fixed and finite. In a general manner, we are able to show that redundancy can always guarantee reliable dynamics, up to a given critical value of noise, above which the system is incapable of maintaining any memory of its past states. From simple considerations, we are able to obtain upper bounds on the maximum resilience attainable. This provides an important frame of reference to determine the reliability of a system with a given sparsity.

We begin by defining the model, and how noise is introduced. A Boolean Network (BN) \[2\] is a directed graph with \( \mathcal{N} \) nodes, representing the genes, which have an associated Boolean state \( \sigma_i \in \{0,1\} \), corresponding to the transcription state, and a function \( f_i(\{\sigma_j\}) \), which determines the state of node \( i \) given the states of its input nodes \( \{\sigma_j\} \). The number of inputs of a given node is \( k_i \), or simply \( k \) if its the same for all nodes. This system is usually updated in parallel, such that at each time step \( t \), we have \( \sigma_i(t+1) = f_i(\{\sigma_j(t)\}) \). Starting from an initial configuration, the system will evolve, and eventually settle on an attractor. In a real system, the expression level of a particular gene can fluctuate, despite the stability of its input states \[3\]. This characteristic can be incorporated qualitatively in the BN model as uniform noise \[4\]—\[10\], defined as a probability \( p \) that, at each time step, the value of a given input \( \sigma_j \in \{\sigma_j\} \), of a node \( i \) is flipped, prior to the evaluation of the function \( f_i \). The value of \( p \) plays the role of a temperature in the system. If \( p = 0 \) the original deterministic model is recovered, and if \( p = 1/2 \) the system becomes effectively decoupled, with entirely stochastic dynamics.

In the model above, it is known that error resilience does not spontaneously emerge, since Random Boolean Networks (BNs with random topology and functions \[11\]), and simple functional elements such as loops always exhibit ergodic behaviour in the presence of noise \( (p > 0) \) \[10\]. To obtain resilience, some level of functional redundancy must be introduced in the network. In the following we describe how this can be done, and analyse the optimal situation where all functions are arbitrarily redundant. From this situation we obtain upper bounds on the maximum reliability attainable, which is characterized by a transition from non-ergodic to ergodic behaviour at a critical noise value.

Redundancy in sparse networks— Given a finite number of inputs per node \( k \ll \mathcal{N} \), for any given (non-constant) function, the probability that noise will change the output of the function will always be above zero, independent of the size of the network. Therefore, in average there will always be a non-vanishing fraction of the nodes which will be at the wrong state at any given time. The most that can be expected is that this fraction be as small as possible, and remain small as the dynamics evolve. The issue of achieving the first goal was first approached by von Neumann \[12, 13\], who described a general mechanism of optimal redundancy, which is capable of reducing the propagation of errors in a BN. We will briefly outline this mechanism, and then show how it can be used to construct a dynamical model of error propagation in resilient Boolean Networks.

The mechanism proposed in \[12\] consists of locally replicating a given function, such that the replicated input and output edges will form bundles which will all carry the same information, in the absence of noise (see Fig 1). The edges of

![FIG. 1: Redundancy construction method. Left: The original function in the network; Middle: Equivalent redundant function, composed of the executive and restoring organs, with an edge bundle of size four. The grey rectangle corresponds to a random rewiring of the edges; Right: The resulting “pseudo-function”, with incoming and outgoing edge bundles.](image-url)
the output bundle are then randomized and fed into appropriate restoration functions which will independently query the majority state carried in them. The number of edges in the bundle can be arbitrarily large, but the number of inputs per node must remain fixed. The output bundle of the last functions will then propagate the information to the rest of the network, which is also modified in the same manner. The first stage was dubbed in [12] the “executive organ”, and the second stage, the “restoring organ”. We note that while this method outlines a specific construction, it has a general nature, since it incorporates the two the most necessary features to be resilient against noise: replication and restoration of majority values. It does so piecewise for all functions in a given network, depicting an alternative version with an optimal level of redundancy. This robust version will then function exactly as the system shows ergodic dynamics (as previously discussed), of the dynamics, with different initial states where

\[ b_{i}(t) \text{ is the value of } b_i \text{ for pseudo-node } i \text{ at time } t, \text{ with a starting state } b_i(0|\sigma) = \sigma. \text{ The average } \langle \ldots \rangle \text{ is taken over the whole network, and several independent realizations of the dynamics, with different initial states } \{\sigma_i^0\} \text{ and } \{a_i^0\}. \text{ If the system shows ergodic dynamics (as previously discussed), the value of } h \text{ should converge to zero, corresponding to only one possible fixed point in the values of } b_i. \text{ Otherwise, it should decrease with the noise strength } p, \text{ as the effects of noise bring the system closer to the ergodic phase [17].} \]

We will consider separately the case of networks with functions \( k = 2 \), and later the case \( k > 2 \). The case \( k = 1 \) will not be analysed since it does not allow for the construction of a restoring organ.

\[ k = 2 \text{— The most crucial part in the procedure outlined above is the selection of the function to be used in the restoring organ. The function must be able to transform the values of the majority of the edges in the input bundle, into a even greater majority in the output bundle. However, no } k = 2 \text{ function can act as a simple majority function. There are however some functions which behave as a majority function for certain input combinations, but not for others. These functions are the AND (8), NAND (7), OR (14) and NOR (1) which react only to two simultaneous input changes, if the original inputs are all 1 or 0, but react to any input flip if the original inputs are in the opposite state. Therefore those functions would be able to correct either value passing on the bundle, but not both. The solution proposed in [12] is to construct the restoring organ with two modules connected in sequence, both with either NAND or NOR functions [18], as can be seen in Fig. 2. The first tier will correct one of the values if it can and intrinsically flip the majority value of the bundle, and the second tier will then have its chance of correcting, now that the majority value has changed. We can verify the actual response of this scheme to noise, by defining two maps. First, the actual noise on the bundle, \]

\[ \epsilon(\eta) = (1 - 2p)\eta + p, \] (3)

where \( \epsilon \) is the fraction of the edges in the bundle with a given value, given the original fraction of same value \( \eta \). Second, the response of the NOR and NAND functions,

\[ a_{\text{nor}}(\eta) = (1 - \eta)^2, \quad a_{\text{nand}}(\eta) = 1 - \eta^2, \] (4)

where \( a \) is the fraction of edges in the output bundle with value 1, and \( \eta \) the fraction of inputs with the same value. The full restoration map is then given by

\[ b^r(\eta) = (a_{\text{nor/nand}} \circ \epsilon)^2(\eta), \] (5)
which is plotted in Fig. 3, for some values of $p$. We can see that the majority value on the bundle is preserved, even for non-zero values of $p$. However, the question remains if this restoration will be enough to maintain trajectories of a network from diverging. For that, we need to couple the restoration map above with the functions present on the network and iterate the system. However, there is one specific situation which represents the limiting case of maximum resilience, namely when after each pseudo-function there are infinitely many restoring organs in sequence. In this case, the response to noise of the function in the executive organ can be neglected, and the (infinitely long) restoring organ alone will determine the resilience of the network. Conveniently, this can be done with successive iterations Eq. 6 which should eventually reach a fixed point, corresponding to the roots of the equation \((b^{\text{meanhand}} \circ e)^2(\eta) = \eta\). This is a fourth order polynomial in $\eta$, and the roots can be obtained analytically. The system exhibits a typical pitchfork bifurcation, with three solutions of any starting state. In order to characterize this transition more precisely, we can write the expression for the previously defined order parameter in Eq. 2 as

$$h = \lim_{t \to \infty} [b(t) - b(t)] = \frac{[8(p - p_c)(p - p^*_c)]^{1/2}}{(2p - 1)^2} [p \leq p_c]$$

where $b(t|\sigma)$ is the value of $b(t)$ with the starting point $b(0) = \sigma$, and $p^*_c = (3 + \sqrt{7})/4$. The values of the order parameter are plotted in Fig. 4. From Eq. 6 we also see easily that the critical exponent is 1/2 (mean-field universality class).

![FIG. 3: Fixed points, or period-2 points, $b(\infty)$ (top) and long-term hamming distance $h$ (bottom) for the NAND (black) and NOR (gray) maps, as a function of noise. Black curves correspond to NAND restoration and grey ones to NOR.](image)

existence of this critical value of noise points to a direct up-

per bound on the reliability attainable by $k = 2$ networks, since it represents the maximum limit of error correction of the restoring organ. Additionally, this critical value of noise corresponds exactly to the upper bound found rigorously by Evans and Pippinger [14] for reliable computation of Boolean formulas composed of noisy NAND gates.

$k > 2$— If the functions have $k > 2$, the choice of the restoring organ becomes more obvious, and the most natural choice is the majority function, which returns simply the majority value of its inputs. Since it will work equally well if the value on the bundle is either 0 or 1, the majority function is capable of performing restoration with only one tier of functions, without accumulating noise in an intermediate step, which provides it with superior characteristics. The restoration map of the majority function is given by

$$b^m(\eta) = 1 - \sum_{i=0}^{[k/2]-1} \binom{k}{i} \eta^i (1 - \eta)^{k-i} + \frac{\delta_{k/2,[k/2]}}{2} \binom{k}{k/2} \eta^{k/2} (1 - \eta)^{k/2}. \quad (8)$$

The last term is added only for functions with even $k$, which have an indeterminate majority state. In this case, it is assumed that half the restoring functions output 1 and the other half 0. It is also clear that majority functions with even $k > 2$ will perform just as well as a $k - 1$ odd function, and therefore the extra input is, for this purpose, wasted. We can analyse the quality of this restoration by iterating Eqs. 8 and 9 in sequence, like it was done for $k = 2$. In the absence of noise, this will lead to one of two fixed points, depending on the starting condition. It can be seen in Fig. 4 that those fixed points also merge into one at a critical value of noise, and the associate order parameter $h$ also indicates a second order transition, with the same critical exponent, but different critical noise values. As expected, the value of $h$ is larger for larger $k$, for the same value $p$, and the critical noise is also larger. The critical values $p_c$ match exactly the upper bounds for Boolean formulas using noisy majority functions of $k$ inputs found by Evans and Schulman [15], given by,

$$p_c = \frac{1}{2} - \frac{2^{k-2}}{k \left(\frac{k-1}{2}\right)}.$$  

for odd $k$. This scales with \((1/2 - p_c) \sim 1/\sqrt{k}\) (see Fig. 5), leading to a strictly resilient situation when $k \to \infty$.

One can test the effectiveness of the majority functions, by considering other functions as the executive organ. Instead of systematically analysing all $2^{2k}$ functions, we can consider functions which are pathological in their stability to noise. Here we will consider a function with maximum sensitivity, which output 0 or 1 if all inputs are 1 or 0, respectively, but otherwise the function is uniformly distributed, and the outputs will be 0 or 1 with equal probability for each input combination. The corresponding map can be written as

$$b^{\text{min}}(\eta) = \frac{1}{2} (1 + (1 - \eta)^k - \eta^k). \quad (10)$$
Its properties can be seen in 

We see that indeed it becomes progressively difficult to stabilize for larger $k$, and the critical point now scales as $p_c \sim 1/k$, and the transition becomes first-order. On the other hand, the mere existence of the critical point confirms some level of resilience, which is not present in the system without redundancy, where the critical point is always $p_c = 0$.

![Figure 4: Fixed points and long-term hamming distance, for the majority restoring organ (left), and the maximum sensitivity executive organ with the majority restoring organ (right).](image)

![Figure 5: Critical noise value $p_c$ as a function of $k$, for the majority restoring organ, and $k = 2$ NAND/NOR restoration (left), and the maximum sensitivity executive organ with the majority restoring organ (right). The dashed lines correspond to $1/\sqrt{k}$ (left) and $1/k$ (right).](image)

**Conclusion**— We have shown that sparse networks, while they cannot be arbitrarily resilient, they can have stable dynamics in the presence of noise, if redundancy is correctly introduced. The stability is marked by second or first-order transitions, from non-ergodic to ergodic behaviour. We obtain upper-bounds on the error resilience attainable by redundant networks with a given $k$. This is in stark contrast to what is observed in Random Boolean Networks [10], which never exhibit memory of its past states when noise is introduced, either in its frozen or “chaotic” phases.

We have shown that the stabilization through redundancy is successful even with the most pathologically sensitive functions, such as the function with maximum sensitivity discussed. We note also that redundancy provides additional benefits, such as robustness against damage and mutations, as was shown in [16]. We stipulate that due to these robust features, redundancy must be present in some extent in real gene regulatory networks; if not in the entire network, at least in its more dynamically relevant modules. On the other hand, arbitrary redundancy close to the optimal bound is very unlikely due to its high putative cost to the organism, which would favor instead a genetic circuit composed of fewer elements, with only enough resilience sufficient for survival. It remains to be seen to what extent is redundancy desirable, and how it may be connected with other topological and functional restrictions of gene regulation.

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