Statistical mechanics of clonal expansion in lymphocyte networks modelled with slow and fast variables

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Abstract
We use statistical mechanical techniques to model the adaptive immune system, represented by lymphocyte networks in which B cells interact with T cells and antigen. We assume that B- and T-clones evolve in different thermal noise environments and on different timescales, and derive stationary distributions and study expansion of B clones for the case where these timescales are adiabatically separated. We compute characteristics of B-clone sizes, such as average concentrations, in parameter regimes where T-clone sizes are modelled as binary variables. This analysis is independent of the network topology, and its results are qualitatively consistent with experimental observations. To obtain the full distributions of B-clone sizes we assume further that the network topologies are random and locally equivalent to trees. This allows us to compete these distributions via the Bethe–Peierls approach. As an example we calculate B-clone distributions for immune models defined on random regular networks.

Keywords: adaptive immune system, lymphocyte network, statistical physics

(Some figures may appear in colour only in the online journal)

1. Introduction
The main task of the immune system is to defend an organism from invading pathogens such as viruses, bacteria, parasites, etc. In complicated multicellular organisms, such as vertebrates, the immune system is usually divided into two subsystems: the innate immune system and the adaptive immune system [1]. The former can be seen as a first line of defence on which the
organism relies for protection in the first hours and days of infection with a new pathogen, but its immune response is not specific to this particular pathogen. The latter is a second line of defence, which is usually triggered by the innate immune system, whose immune response is more specific and also offers a more long-term protection. These properties of the adaptive immune system arise from its ability to learn and memorise a wide range of pathogens, an important part of which is learning to recognise the molecules of the organism [2], which emerges from the interactions of its cells, mediated by signalling proteins (cytokines).

Interactions between B cells and T cells are dominant in the adaptive immune response [1]. The main feature of a B cell is its B cell receptor (BCR) which is used to recognise antigen (Ag). Antigen is a (unique) protein on the surface of the pathogen. However, a BCR recognises only a specific part of the Ag, which is called epitope, so potentially a single BCR can recognise many different Ags [3, 4]. An Ag that binds to the BCR is subsequently internalised and is broken by the cell into peptides (short proteins) which are then displayed on its surface. The T cell also has its T cell receptor (TCR) to recognise Ag, however, in contrast to B cells, a T cell binds to the peptides on the surface of antigen presenting cells (APCs), such as dendritic cells, B cells, etc. The adaptive immune response is triggered when T helper cells that are attached by their TCRs to B cells, activate these B cells (by cytokine-mediated signals) that recognise the same Ag. The strength of this interaction between the BCR (TCR) and the Ag is called affinity. Upon activation, the B cells with highest affinity begin to proliferate in a process known as clonal expansion. A group of B cells (or T cells) with the same BCR (or TCR) is called a clone. While B cell clones (B clones) are expanding the T cells, activated by Ag, they are also going through the process of clonal expansion. We note that the mode of B cell activation described here is called ‘T-dependent B cell activation’, however for some ‘simple’ Ags with repetitive patterns, such as polysaccharides, the B cell can also be activated directly by Ag and without the help from a T cell (‘T-independent activation’).

Some proliferating B cells differentiate into plasma cells, which secrete large quantities of antibodies (BCRs in soluble form). The antibodies (Ab) protect the host from infection by binding to the pathogen and thus, by blocking its external parts, make the attack ineffective or ‘mark’ the pathogen for ingestion by other host cells. However, the Abs produced in this initial clonal expansion are of very low affinity, especially if the host is infected with this pathogen for the first time. The other proliferating B cells are used to seed the germinal centres (GCs). GCs are special micro-environments, formed within secondary lymphoid organs, which are divided into the light and dark zones. In the dark zone B cells undergo somatic hypermutation which introduces random mutations into the genes of BCRs. This process is followed by the interaction of B cells with the helper cells and Ag in the light zone of the GC. Here B cells with higher affinity are selected and differentiated into plasma cells and memory B cells. Presence of high affinity memory B cells in the organism leads to a more vigorous immune response on the subsequent infections by the same Ag.

Statistical physics (SP) approaches to modelling the adaptive immune system [5, 6] are predominantly based on the idea that the immune system can be viewed as a network (or graph). The nodes in such networks are lymphocyte clones (B cells and T cells), antibodies, antigens, etc and the edges model interactions such as B cells receiving ‘signals’ from T cells, antibodies binding to antigen, etc. One of the first such studies was [7], describing a network of interacting antibodies [8], i.e. an idiotypic network. Despite being very simple (binary variables assigned to the nodes represented Ab concentrations, and the interactions were assumed to be random), this model exhibited memory, which was interpreted as the (immunological) memory of the past exposure to the Ag. A more detailed model with lymphocytes and cytokines was introduced in [9]. Here the (bipartite) network was formed by interactions between effector clones, formed by B cells and killer T cells, and helper clones
(T cells), mediated by cytokines (strengths of (random) interactions in the model). The immunological memory in this model arises from the ability of T clones to memorise an extensive number of cytokine signalling ‘strategies’ (or patterns), allowing to cope with possible Ag stimulation. However, the fully connected network topology used in this model restricted the immune memory to retrieval of only one pattern at a time, limiting the abilities of the immune system to fight against multiple Ags [9]. The finitely connected network topology, which is more realistic from the biological point of view [10], allows to retrieve an extensive number of patterns in parallel.

The above studies all used the equilibrium SP framework, and assumed existence of an energy function, governing behaviour of Abs, lymphocytes, etc, from the very beginning. However, whenever it is not clear that such a function exists, or if we want to study relaxation to equilibrium, we must study dynamics. Examples of studies of the dynamics of immune idiotypic networks and bipartite lymphocyte networks are [11] and [12]. Finally, we note that SP models and concepts are also used in statistical inference of immune system data. The maximum entropy model of amino acid sequences was used to study the repertoire of memory B cell receptors in zebrafish [13]. Recently, similar approaches were applied to study the repertoires of T cells [14] and B cells [15] in humans.

In this paper we develop further the lymphocyte network model [9, 10], in both qualitative and quantitative directions. The main difference between the current and previous versions of the model is that we relax the assumption that B clones, represented by log-concentrations of B cells, and T cell clones (T clones), represented by concentrations of T cells, are subject to the same thermal noise [10]. We note that both populations of B cells and T cells are affected by various random events such as stochasticity in cell division and cell death [16], thermal fluctuations in the TCR-peptide [17–20] bond strengths [21], etc. Also, these are populations of biochemically, and possibly physically [22], distinct cells. Furthermore, in the T-dependent immune response the magnitude of fluctuations in the populations of B cells and T cells are expected to be very different due to the fact that the former are subject to the process of somatic hypermutation and the latter are not [1]. For the above reasons it is implausible for the random noise in B and T clones to be of identical strength.

Having different thermal noise levels, i.e. different ‘temperatures’, in the B and T clone evolution introduces some technical difficulties in the analysis of the problem. In contrast to previous studies [9, 10], it prevents us from using the equilibrium framework directly. However, we can make progress by assuming that either B clones or T clones are ‘fast’ variables [23], i.e. they are evolving on different time-scales. The latter allows us to obtain the stationary distribution of a two-temperature system in its explicit form [24], unlike the situation with the same or comparable time-scales [25].

Furthermore, previous studies did not distinguish the T helper cells from the T regulator cells [26]. The latter play an important role in preventing autoimmunity by suppressing the growth of self-reactive B clones which could be produced during the T-dependent immune responses [27]. Also, B cells infiltrate tumours and there is an evidence for in situ immune responses [28], to tumour-associated (self) Ags, which are thought to be strongly influenced by the presence of regulator T cells [29].

Finally, we note that our main focus in this paper will be on the properties of B clone distributions and how these are affected by the parameters of the model, such as network topology, concentration of Ag, etc. Existing models of clone distributions usually disregard such details (see [30] and references therein) and do not explicitly include interactions between the B cells and T cells dominant in the T-dependent immune response. Recently, the importance of these properties in our understanding of the ageing immune system was emphasised in [31] and we also envisage that these properties are important in tumour immunology [28].
2. Dynamics

We consider $M$ B-clones interacting with $N$ T-clones on a bipartite graph $G = (V, \mathcal{F}, \mathcal{E})$, where $N = |V|$, $M = |\mathcal{F}|$ and $\mathcal{E}$ is the set of edges (see figure 1). The set of indices $\partial \mu$ contains all T-clones $i$ connected to B-clone $\mu$, and the set $\partial \theta$ defines all B-clones $\mu$ that are connected to T-clone $i$. The B-clone sizes are specified by the log-concentrations $b = (b_1, \ldots, b_M)$, and we assume that these are governed by the Langevin equation

$$\tau_b \frac{db_{\mu}}{dt} = J_{\mu} \left( \sum_{i \in \partial \mu} \xi_i^\eta \sigma_i + \theta_i \right) - \rho b_{\mu} + \chi_\mu(t)$$

(1)

where the zero-average Gaussian noise $\chi_\mu(t)$, with $\langle \chi_\mu(t) \chi_\nu(t') \rangle = 2\eta_0 \beta^{-1} \delta_{\mu \nu} \delta(t - t')$, is characterised by a ‘temperature’ parameter $\beta = \beta^{-1}$. In this dynamics the $\mu$th B-clone receives the ‘signal’ $\sum_{i \in \partial \mu} \xi_i^\eta \sigma_i$ from $|\partial \mu|$ T-clones, whose sizes are specified by the concentrations $\sigma = (\sigma_1, \ldots, \sigma_N)$. The $i$th T-clone is either formed by T-helper or by T-regulator cells. The efficacies of the T-helper and T-regulator clones are encoded, respectively, by ‘cytokine’ variables $\xi_i = 0$ and $\xi_i < 0$. The signal $\sum_{i \in \partial \mu} \xi_i^\eta \sigma_i$ from the T-clones is modulated by the interaction strength $J_{\mu}(a) = \sum_{\nu \in M} S_{\mu \nu} a_{\nu}$, which depends on the Ags, as represented by the vector of epitope ‘concentrations’ $a = (a_1, \ldots, a_M)$. Here $S_{\mu \nu} \geq 0$ is an element of an ‘affinity’ matrix which specifies how well the $\nu$th epitope is ‘matched’ by the $\mu$th B-clone. A very specialised B-clone will interact with only one epitope (i.e. $S_{\mu \nu} = \delta_{\mu \nu}$), whereas poly-reactive B-clones [3, 4] can interact with many different epitopes (e.g. $S_{\mu \nu} > 0$ for all $\nu$). The $\theta_i$ term gives the possibility of B-clone activation even in the absence of a signal (or in the presence of only a weak signal) from the T clones, i.e. it facilitates T-independent activation.

A positive (negative) ‘field’ $F_\mu(\sigma) = J_{\mu} \left( \sum_{i \in \partial \mu} \xi_i^\eta \sigma_i + \theta_i \right)$ has an excitatory (inhibitory) effect on the growth of $\mu$th B-clone. The strength of this effect is increasing with larger amounts of Ag: the interaction $J_{\mu}(a)$ is a monotonic non-decreasing function of the antigen $a$, i.e. $J_{\mu}(a) \geq J_{\mu}(\tilde{a})$ for all $\mu$ when $a_\nu \geq \tilde{a}_\nu$ for all $\nu$. Furthermore, the growth of clone $\mu$ is kept in check by the ‘apoptosis’ term $-\rho b_{\mu}$, which limits its amplitude. For $J_{\mu} = 0$, i.e. without Ag, the distribution of log-concentration $b_{\mu}$ in equilibrium takes the Gaussian form

$$p(b_{\mu}) = \left( \frac{2\pi \rho \beta}{3} \right)^{\frac{1}{2}} \exp \left( -\frac{1}{2\rho \beta} b_{\mu}^2 \right)$$

In order to derive a dynamical equation for the T-clones we will follow the ideas of [9, 10]. Firstly, we note that if we define the energy function (or Hamiltonian)

$$\mathcal{H}(b, \sigma) = -\sum_{\mu=1}^M b_{\mu} F_\mu(\sigma) + \frac{1}{2} \rho \sum_{\mu=1}^M b_{\mu}^2,$$

(2)

then equation (1) can be written in the form

$$\tau_b \frac{db_{\mu}}{dt} = -\frac{\partial}{\partial b_{\mu}} \mathcal{H}(b, \sigma) + \chi_\mu(t).$$

(3)

We note that this dynamics is invariant under the transformation $\mathcal{H}(b, \sigma) \rightarrow \mathcal{H}(b, \sigma) + V(\sigma)$ where $V(\sigma)$ is any function of $\sigma$. Secondly, we assume that

$$\tau_\sigma \frac{d\sigma_i}{dt} = -\frac{\partial}{\partial \sigma_i} \mathcal{H}(b, \sigma) + \eta_i(t),$$

(4)
where the zero-average Gaussian noise $\eta_i(t)$, with $\langle \eta_i(t)\eta_j(t') \rangle = 2\tau_i\beta^{-1}\delta_{ij}\delta(t-t')$, is characterised by a ‘temperature’ $T = \beta^{-1}$. From the above follows the equation
\[
\tau_i \frac{d\sigma_i}{dt} = \sum_{\mu\in\Omega_i} J_{\mu i} \xi_i^\mu b_\mu - \frac{\partial}{\partial \sigma_i} V(\sigma) + \eta_i(t). \quad (5)
\]
For now we will leave the function $V(\sigma)$ unspecified—it will be used later to ‘restrict’ the range of $\sigma_i$’s—while allowing us to define various thermodynamic functions. We note that the advantage of assuming that T-clones are governed by the same Hamiltonian as B-clones is that it allows us to use analytical tools from equilibrium statistical mechanics. The disadvantage is that whether this approach is correct or not, as in any other phenomenological approach, can be only established \textit{a posteriori}. One of the consequences of using the present approach, which is explicit in the T-clone equation (5), is that, in the presence of Ag, the evolution of the T-clones is governed by the B-clones. The latter can be interpreted as ‘B-cells acting as Ag presentation cells (APCs) for the T-cells’ which is a well known immunological fact \cite{1}. However, besides B-cells there are other dedicated APCs, such as dendritic cells, etc. The latter can be included via the potential $V$. Finally, we note that the relation between the time-scales of the T- and B-clone subsystems ($\tau_\sigma \gg \tau_b$ versus $\tau_\sigma \gg \tau_b$) give us two possible scenarios to analyse the equilibrium state \cite{32}.

2.1. Fast equilibration of B-clones

We first assume that the B-clone variables $b_\mu$ are ‘fast’ variables, and equilibrate on timescales much shorter than those characterising the evolution of the T-clones. In equilibrium the former will then be governed by the distribution
\[
P^\beta_\mu (b|\sigma) = \frac{1}{Z^\beta(\sigma)} e^{-\beta H_\mu(b,\sigma)}, \quad (6)
\]
where $Z^\beta(\sigma) = \int db e^{-\beta H_\mu(b,\sigma)}$ is a partition function. From this it then follows that the dynamic equation (4) becomes\footnote{We note that the ‘implicit averaging’ procedure used here is exact when $\eta_i \to 0$ \cite{23}.}
\[
\tau_\sigma \frac{d\sigma_i}{dt} = -\left\langle \frac{\partial}{\partial \sigma_i} H(b, \sigma) \right\rangle_\beta + \eta_i(t) \\
= -\frac{\partial}{\partial \sigma_i} F^\beta_\sigma (\sigma) + \eta_i(t). \quad (7)
\]
where $F^\beta_\sigma (\sigma) = -\beta^{-1} \log Z^\beta(\sigma)$ is (formally) a free energy of a state $\sigma$. The above subsequently implies that in equilibrium the T-clones are governed by the distribution
\[
P^\beta_\beta (\sigma) = \frac{1}{Z^\beta_\beta} e^{-\beta F^\beta_\beta (\sigma)}, \quad (8)
\]
where $Z^\beta_\beta = \int d\sigma e^{-\beta F^\beta_\beta (\sigma)}$.

From (6) and (8) we can construct the joint distribution
\[ P_{\beta,\beta}(b, \sigma) = P(b|\sigma)P_{\beta,\beta}(\sigma) \]
\[ = \left\{ \prod_{\mu=1}^{M} e^{\frac{-1}{2}\beta(b_{\mu} - \langle b_{\mu}\rangle_{\beta})^2} \right\} \int D\sigma e^{\frac{1}{2}\beta\sum_{\mu=1}^{M} F^2_{\mu}(\sigma)} \]
\[ \int D\sigma e^{\frac{1}{2}\beta\sum_{\mu=1}^{M} F^2_{\mu}(\sigma)} \]
\[ \left( \prod_{\mu=1}^{M} \right) e^{\frac{-1}{2}\beta(b_{\mu} - \langle b_{\mu}\rangle_{\beta})^2} \int D\sigma e^{\frac{1}{2}\beta\sum_{\mu=1}^{M} F^2_{\mu}(\sigma)} \]

which in turn allow us to construct the average density

\[ P(b) = \frac{1}{M} \sum_{\mu=1}^{M} \langle \delta(b - b_{\mu}) \rangle_{\beta,\beta} = \int dF P(F) \frac{e^{-\frac{1}{2}\beta(b - \langle b \rangle_{\beta})^2}}{\sqrt{2\pi/\beta}} , \]

where

\[ P(F) = \int D\sigma \frac{e^{\frac{1}{2}\beta\sum_{\mu=1}^{M} F^2_{\mu}(\sigma)}}{\int D\sigma e^{\frac{1}{2}\beta\sum_{\mu=1}^{M} F^2_{\mu}(\sigma)}} \frac{1}{M} \sum_{\mu=1}^{M} \delta(F - F_{\mu}(\sigma)) \]

is a distribution B-clone activation fields. It follows from the above that the number of B cells in a volume \( V \), their concentration \( c \), is governed by the distribution\(^4\)

\[ P(c) = \int dF P(F) \frac{e^{-\frac{1}{2}\beta(c - \langle c \rangle_{\beta})^2}}{\sqrt{2\pi/\beta}} . \]

We note that in equation (9) the ‘Boltzmann’ factor \( \exp \left[ -\frac{1}{2}\beta \sum_{\mu=1}^{M} F^2_{\mu}(\sigma) \right] \) can be written, up to an irrelevant constant, as \( \exp \left[ -\beta \sum_{j=0}^{j} \eta_{j} \sigma_{j} - \sum_{i=0}^{i} \theta_{i} \sigma_{i} \right] \), where \( J^1_{\mu} = (J^1_{\mu}(\eta_{j})^{\epsilon}_{j} \eta_{j}) \) and \( \theta_{i} = (J^1_{\mu}(\theta_{i})^{\epsilon}_{i} \theta_{i}) \) can be represented as a fully connected weighted graph (or clique) formed by those the T-clones that control B-clone \( \mu \). Thus the B-clone dynamics (1), given the assumptions of this section, leads us to the result that in equilibrium the T-clones are interacting via this network of cliques (see figure 2).

2.2. Fast equilibration of T-clones

Here we assume, contrary to the previous section, that the T-clones equilibrate first, and are upon equilibration governed by the distribution

\[ P_{\beta}(\sigma|b) = \frac{1}{Z_{\beta}(b)} e^{-\beta H(b, \sigma)} , \]

where \( Z_{\beta}(b) = \int d\sigma e^{-\beta H(b, \sigma)} \). The dynamics of B-clones is then given by

\[ \frac{db_{\mu}}{dt} = -\left( \frac{\partial}{\partial b_{\mu}} H((b, \sigma)) \right)_{\beta} + \chi_{b}(t) \]
\[ = -\frac{\partial}{\partial b_{\mu}} F_{\beta}(b) + \chi_{b}(t) , \]

where \( F_{\beta}(b) = -\beta^{-1}\log Z_{\beta}(b) \). The latter thus evolve towards the equilibrium state

\(^4\)For any distribution \( P(b) \) we have \( P(c) = \int P(b) \delta(c - e^{b}) db = \frac{1}{e} P(\log(c)) \).
where \( \tilde{\beta} / \beta = n \). We can use (15) and (13) to construct the joint distribution

\[
P_{\beta, \tilde{\beta}}(b) = \frac{1}{Z_{\beta, \tilde{\beta}}} e^{-\beta J(b)} = \frac{Z_{\beta}^n(b)}{\int db Z_{\tilde{\beta}}^n(b)},
\]

where \( n = \tilde{\beta} / \beta \). Let us first consider the case when \( n \in \mathbb{Z}^+ \). Here

\[
P_{\beta, \tilde{\beta}}(b, \sigma) = \frac{e^{-\frac{1}{2} \sum_{i<j} J_{ij}^\mu b_i b_j} e^{\tilde{\beta} \sum_{i=1}^M b_i F_i(\sigma)} W_{\beta}^{n-1}(b)}{\int db e^{-\frac{1}{2} \sum_{i<j} J_{ij}^\mu b_i b_j} W_{\tilde{\beta}}^n(b)},
\]

where \( W_{\beta}(b) = \int d\sigma e^{\beta \sum_{i=1}^M b_i F_i(\sigma)} \).

Figure 1. Bi-partite network of T-clones and B-clones, generated by the T-dependent immune response. A red link between a (helper) T-clone \( i \) and a B-clone \( \mu \) represent that this B-clone receiving a signal to expand. A blue link between a (regulator) T-clone \( i \) and B-clone \( \mu \) represents a signal to contract.

Figure 2. Example of a local topology in the T-clone network generated by the system with fast B-clone equilibration. The evolution of B-clone \( \mu \) (large grey circle) is governed by three T helper clones (red nodes) and one T regulator clone (blue node), forming a clique of four nodes.
\[ W^\alpha_{\beta}(b) = \int D\sigma^1 \ldots \int D\sigma^n e^{\beta \sum_{\alpha=1}^n b_{\alpha} \sum_{i=1}^n F_i(\sigma^\alpha)} \] (17)

and the thermal average \( \langle f(\sigma) \rangle = \int D\sigma P_{\beta,\beta}(\sigma) f(\sigma) \) of any function \( f(\sigma) \) will be given by the following ‘replica’ relation

\[
\langle f(\sigma) \rangle = \frac{\int d\bar{b} e^{-\frac{1}{\beta} \sum_{\alpha=1}^n \sum_{i=1}^n F_i(\sigma^\alpha)} \int \{ D\sigma^n \} e^{\beta \sum_{\alpha=1}^n b_{\alpha} \sum_{i=1}^n F_i(\sigma^\alpha)} f(\sigma^\alpha) \int d\bar{b} e^{-\frac{1}{\beta} \sum_{\alpha=1}^n \sum_{i=1}^n F_i(\sigma^\alpha)} W^\alpha_{\beta}(\bar{b})}{\int \{ D\sigma^n \} e^{\frac{\beta}{\beta} \sum_{\alpha=1}^n \sum_{i=1}^n F_i(\sigma^\alpha)} }.
\] (18)

where we have defined the average

\[
\langle \ldots \rangle_n = \frac{\int \{ D\sigma^n \} e^{\frac{\beta}{\beta} \sum_{\alpha=1}^n \sum_{i=1}^n F_i(\sigma^\alpha)} }{\int \{ D\sigma^n \} e^{\frac{\beta}{\beta} \sum_{\alpha=1}^n \sum_{i=1}^n F_i(\sigma^\alpha)} } \langle \ldots \rangle.
\] (19)

The Boltzmann weight \( \exp\left[\frac{\beta}{\beta} \sum_{i=1}^n \sum_{\alpha=1}^n F_i(\sigma^\alpha)\right] \) in this expression can be written, up to an irrelevant constant, in the form \( \exp\left[-\beta \sum_{i=1}^n E_i(\{\sigma^\alpha\})\right] \). The energy function \( E_i,\{\sigma^\alpha\} = -\frac{1}{2} \sum_{\alpha,\gamma} \sum_{i,\ell=1}^n \sum_{\ell' \in \partial_\mu} (J_{i,\ell'}^\alpha) \sigma^\alpha_i \sigma^\gamma_{\ell'} - \Sigma_{\alpha=1}^n \sum_{i,\ell=1}^n \theta^\alpha_i \sigma^\alpha_i \), where \( J_{i,\ell'}^\alpha = (J_{\beta,\beta}^\alpha)^{\ell' \ell} \) and \( \theta^\alpha_i = (J_{\beta,\beta}^\alpha)^{\ell' \ell} \), can be represented as a clique constructed from those \( n \times |\partial_\mu| \) T-clones that control B-clone \( \mu \). Thus for \( n \in \mathbb{Z}^+ \) the fast T-clone equilibration leads to the enlargement of the ‘local’ T-clone system (see figure 3).

The B-clone density \( P(b) = \frac{1}{M} \sum_{\alpha=1}^n \langle \delta(b - b_\alpha) \rangle_{\beta,\beta} \) can now be computed by using the identity (17) in distribution (16), which gives us

\[
P(b) = \int dF [P(F) e^{-\frac{1}{\beta} \sum_{i=1}^n (b_i - \delta)^2} \frac{1}{\sqrt{2\pi l^\alpha (\beta)}}]
\] (20)

with the distribution of (replicated) fields.
\[ P(F) = \frac{1}{M} \sum_{\beta_i=1}^{M} \left( \delta \left( F - \sum_{\alpha=1}^{n} F_{\alpha}(\sigma^{\alpha}) \right) \right) . \]  

From this also follows the distribution of B-cell concentrations

\[ P(c) = \int dF \ P(F) \frac{e^{-1/2 \rho \ln \left( \log(c) - E \right)^{2}}}{e^{2\pi l \ln \beta}}. \]  

Let us now compute the distribution (16) for the more general case where \( n \in \mathbb{R}^{+} \), not necessarily integer. In order to do this we assume that \( V(\sigma) = \sum_{i} V(\sigma_{i}) \), and use the short-hand \( D\sigma = e^{-2\beta \sum_{i}^{M} V(\sigma_{i})} d\sigma \). We then consider the integral

\[ W_{b}(b) = \int D\sigma \ e^{iM_{\alpha=1} b_{\alpha} F_{\alpha}(\sigma)} = e^{iM_{\alpha=1} b_{\alpha} \delta_{\alpha} + \sum_{i=1}^{n} \log \int D\sigma \ e^{i\rho \sum_{\alpha=1}^{n} b_{\alpha} \delta_{\alpha}}}, \]

where \( D\sigma = e^{-2\beta \sum_{i}^{M} V(\sigma_{i})} d\sigma \). Upon inserting this result into (16) we can extract the marginal distributions

\[ P_{\beta,\delta}(b) = \frac{1}{Z_{\beta,\delta}} e^{-\frac{1}{2} M \beta \sum_{\alpha=1}^{n} \left( b_{\alpha} - \delta_{\alpha} \right)^{2}} + n \sum_{i=1}^{n} \log \int D\sigma \ e^{i\rho \sum_{\alpha=1}^{n} b_{\alpha} \delta_{\alpha}} \]

and

\[ P_{\beta,\delta}(\sigma) = \int dB P_{\beta,\delta}(\sigma) \left\{ \prod_{i=1}^{N} \frac{e^{i\rho \sum_{\alpha=1}^{n} b_{\alpha} \delta_{\alpha}}}{\int D\bar{\sigma} \ e^{i\rho \sum_{\alpha=1}^{n} b_{\alpha} \delta_{\alpha}}} \right\}. \]

The distributions (24) and (25) can be used to construct the densities \( P(b) = M^{-1} \sum_{i=1}^{M} P_{\beta,\delta}(b) \) and \( P(\sigma) = N^{-1} \sum_{i=1}^{N} P_{\beta,\delta}(\sigma) \) in terms of the marginal distribution \( P(\sigma) \) is given by

\[ P(\sigma) = \int dB P_{\beta,\delta}(\sigma) \frac{\int D\tau \ e^{i\rho \sum_{\alpha=1}^{n} b_{\alpha} \delta_{\alpha}}}{\int D\bar{\tau} \ e^{i\rho \sum_{\alpha=1}^{n} b_{\alpha} \delta_{\alpha}}} \delta(\sigma - \sigma_{i}), \]

respectively. In a similar manner, using the definition

\[ P_{e}(F) = \int D\sigma P_{\beta,\delta}(\sigma) \delta(F - F_{\mu}(\sigma)), \]

we can derive the distribution of fields

\[ P_{\mu}(F) = \int dB P_{\beta,\delta}(b) \delta(F - F_{\mu}(\sigma)) \prod_{i \in \mu} \frac{\int D\tau_{i} e^{i\rho \sum_{\alpha=1}^{n} b_{\alpha} \delta_{\alpha}}}{\int D\bar{\tau}_{i} e^{i\rho \sum_{\alpha=1}^{n} b_{\alpha} \delta_{\alpha}}} \delta(F - F_{\alpha}(\sigma)) \]

Similarly to a B-clone, the \( i \)th T-clone experiences the field \( \sum_{\alpha=1}^{n} b_{\alpha} \delta_{\alpha} \); see equation (26).
3. Equilibrium analysis

In the remainder of this article we consider the simplest case when the T clones (or single T cells) are modelled by the (binary) Ising variables $\sigma_i \in \{-1, 1\}$, or by the binary variables $\sigma_i \in \{0, 1\}$. The motivation for this choice of variables is as follows. A single T helper cell can be either active or inactive, i.e. ‘on’ or ‘off’, and is activated by the Ag presenting cell which could be a dendritic cell, a B cell, etc. Then this T helper cell can activate a B cell if it receives the ‘right’ Ag related signal from it. For the T regulator cells we assume that a similar mechanism is at work [27]. On the level of T-clone we could say that the T-clone is active (inactive) if majority of its cells are active (inactive). Thus the use of binary variables can be seen either as a crude approximation of T-clones, where we only retain information about the state of a clone but disregard its size, or all $N$ variables are simply treated as single T cells and we no longer distinguish the T clones to which they belong.

Within the analytical framework of the previous section the choices $\sigma_i \in \{-1, 1\}$ and $\sigma_i \in \{0, 1\}$ can be obtained by using the double well potentials $V(\mathbf{\sigma}) = \Delta \sum_{i=1}^{N}(\sigma_i^2 - 1)^2$ and $V(\mathbf{\sigma}) = \Delta \sum_{i=1}^{N}(\sigma_i - 1)^2 + \frac{1}{2} \omega \sum_{i=1}^{N} \sigma_i^2$ respectively. Here the ‘chemical’ potential $\omega$ allows us to control the number of activated T cells $\sum_{i=1}^{N} \delta_{\sigma_i, 1}$, via the integral measure $D\mathbf{\sigma}$. Taking the limit $\Delta \to \infty$ converts the N-dimensional integral $\int D\mathbf{\sigma}^f(\mathbf{\sigma})$, for example in the equation (18), into a sum $\sum_{\mathbf{\sigma}} f(\mathbf{\sigma})$ over binary variables. Furthermore, the choice $\sigma_i \in \{-1, 1\}$, combined with $\xi_i^0 = 1$, gives us the scenario when each active T cell is either a regulator cell or a helper cell. The fact that the $i$th T cell can change from being a regulator, $\sigma_i = -1$, to being a helper, $\sigma_i = 1$ (or vice versa), must then be seen as an assumption that changes in T cell function occur on the same time scale as the immune response to the Ag. However, in experiments [27, 33] one usually has access only to the number of T cells of either type, and such an assumption may be acceptable to capture the observed phenomena. For the alternative choice of active or inactive T cells where $\sigma_i \in \{0, 1\}$ for all $i$, with $\xi_i^0 \in \{-1, 1\}$ (in this case $\xi_i^0 = \xi_i$), the number of regulating T cells $\sum_{i=1}^{N} \delta_{\xi_i, -1}$ is independent of the immune response.

The consequence of our choice for the measure $V(\mathbf{\sigma})$ describing the variables $\sigma_i$, when T cells are fast variables with $n \in \mathbb{Z}^+$ (see section 2.2), is that the T cells are governed by the distribution

$$P(\mathbf{\sigma}^1, \ldots, \mathbf{\sigma}^n) = \frac{e^{\beta \sum_{i=1}^{n} \sum_{\mathbf{\sigma}} F(\mathbf{\sigma}) \xi_i^0}}{\sum_{\mathbf{\sigma}} e^{\beta \sum_{i=1}^{n} \sum_{\mathbf{\sigma}} F(\mathbf{\sigma}) \xi_i^0}}, \quad (28)$$

where $F(\mathbf{\sigma}) = J \sum_{i} \sum_{j} \xi_i \xi_j \sigma_i^0 \sigma_j^0$, which is equivalent to an Ising spin model for $\sigma_i \in \{-1, 1\}$ or a ‘lattice gas’ for $\sigma_i \in \{0, 1\}$. For the latter we need to add the term $-\frac{1}{2} \beta \omega \sum_{i=1}^{N} \sum_{\mathbf{\sigma}} \sigma_i^0$ to the ‘energy’ function in the exponential of (28). We note that $F(a) = J \sum_{i,j} S_{ij} a_i a_j$, where $J \geq 0$, allows us to control the ‘amount’ of Ag $a$ by increasing or decreasing $J$. Then the quantity $\beta J^2$ controls either the level of noise in the T clone system for fixed $J$ or the amount of Ag for fixed $\beta$. Also the distribution of T-clones (9) in the fast B clone equilibration regime can be obtained from (28) by setting $n = 1$, so for T-clones the case of integer $n$ covers both equilibration scenarios. Let us for now discuss the case of $n = 1$ and $\theta_\mu = 0$.

\textsuperscript{6}In immune response experiments, as in [27], such fractions can be computed from the number of activated T helper, T regulator cells and the total number of activated T cells, N.
For the Ising case with $\xi_i = \beta_1 i$, the average magnetization $\langle m \rangle = \frac{1}{N} \sum_{i=1}^{N} \langle \sigma_i \rangle$, which is related to the fraction of activated helper (regulator) T-cells via the identity $m_+ = 1 - m_- = \frac{1}{2} \left( 1 - m \right)$, has a phase transition at $\beta_c = \beta$ from the disordered paramagnetic (PM) $m = 0$ phase to the ordered ferromagnetic (FM) phase $m \neq 0$ (see figure 4) when $N \to \infty$ [34]. We note that in this framework the PM and FM phases can be interpreted respectively, as the ‘low-dose tolerance’ and ‘vigorous immune response’ phases of a real immune system. The former is consistent with its insensitivity to ‘small’ amounts of Ag and the latter is consistent with its strong reaction to a larger amounts of Ag [1]. The noise parameter $\beta$ controls this ‘sensitivity’ to the Ag: for small (large) $\beta$, i.e. in a high (low) noise regime, a larger (smaller) amount of Ag is needed to trigger a vigorous immune response.

There are many possible topologies of a bipartite graph $G$ for which the T cell system (28) has a finite critical inverse noise level $\beta_c$. Suppose we choose one such topology (see

Figure 4. Fraction of regulator T cells (lower branch) $m_- = \frac{1}{2} \left( 1 - m \right)$, fraction of helper T cells (upper branch) $m_+ = \frac{1}{2} \left( 1 + m \right)$, and average B-clone size $\langle c \rangle$, in the fast B-clone equilibration regime, as a function of the inverse temperature $\beta$ in the T-clone system modelled with the binary variables $\sigma_i \in \{-1, 1\}$.

Figure 5. Two possible topologies for bipartite immunological graphs (left and centre), to illustrate the consequences of eliminating ‘fast’ B-clones. T and B clones are represented by circles and squares respectively. Upon integrating out the B-clone variables, the bipartite graph on the left gives rise to an effective T-clone system shown on the right, in which the T-clones (circles) interact on the classical square lattice (drawn as thick solid lines). Also the bipartite graph in the centre gives rise to an effective T-clone system on the square lattice shown on the right, but now with extended interaction range (drawn as thin solid lines).

For the Ising case with $\xi_i = 1$, the average ‘magnetization’ $m = \frac{1}{N} \sum_{i=1}^{N} \langle \sigma_i \rangle$, which is related to the fraction of activated helper (regulator) T-cells via the identity $m_+ = 1 - m_-$, has a phase transition at $\beta_c = \beta$ from the disordered paramagnetic (PM) $m = 0$ phase to the ordered ferromagnetic (FM) phase $m \neq 0$ (see figure 4) when $N \to \infty$ [34]. We note that in this framework the PM and FM phases can be interpreted respectively, as the ‘low-dose tolerance’ and ‘vigorous immune response’ phases of a real immune system. The former is consistent with its insensitivity to ‘small’ amounts of Ag and the latter is consistent with its strong reaction to a larger amounts of Ag [1]. The noise parameter $\beta$ controls this ‘sensitivity’ to the Ag: for small (large) $\beta$, i.e. in a high (low) noise regime, a larger (smaller) amount of Ag is needed to trigger a vigorous immune response.
for the $\beta \in (\beta_c, \infty)$ the fraction of helper (regulator) T cells $m_+$ ($m_-$) is either a monotonic non-decreasing or a monotonic non-increasing function of the noise parameter $\beta$ (see figure 4) and of interaction strength $\langle J^2 \rangle / \rho$. This follows from the Griffiths–Kelly–Sherman (GKS) theorem [35], which holds for any ferromagnetic Ising spin system. Furthermore, if we know the average magnetization $m \equiv m(\langle J^2 \rangle / \rho)$ for the T cell system with uniform interactions, i.e. with $J_0 = J$ for all $\mu$, then by the same theorem $m(\langle J^2 \rangle / \rho) \leq m(\langle J^2 \rangle / \rho) \leq m(\langle J^2 \rangle / \rho)$, where $J_0 = \inf J_\mu$ and $J_1 = \sup J_\mu$, from which it also follows that $\beta \equiv \beta(\langle J^2 \rangle / \rho)$ obeys: $\beta(\langle J^2 \rangle / \rho) \leq \beta(\langle J^2 \rangle / \rho)$. Figure 5)

We note that the ratio of the number of regulator T cells to the total number of T cells was observed, during a normal immune response to the Ag stimulation, to be in the range $0.1 - 0.25$ in experiments on mice [27]. In the phase diagram (see figure 4) this corresponds to the lower branch of the $m_-$ plot. Also in this regime, the fraction $m_+$ of helper T-cells is a monotonic non-decreasing function of Ag concentration, which is consistent with the experimental data [33]. The case of a ‘lattice gas’, i.e. $\sigma \in [0, 1]$ and $\xi_0 = 1$, which is the scenario where we have only T helper cells which are either active ($\sigma = 1$) or inactive ($\sigma = 0$), can be mapped similarly into the ferromagnetic Ising model with (positive) external field [34]. For example, by writing $\sigma_i = \frac{1}{2}(1 + s_i)$, where $s_i \in \{-1, 1\}$. As a consequence, also here the fraction of helper T clones $m_+ = 1 - m_-$ is a continuous and monotonic function of $\beta$ and $\langle J^2 \rangle / \rho$ [35].

We now turn to the analysis of B-clone properties in the fast B-clone and fast T-clone ($n \in \mathbb{Z}^+$) equilibration regimes. For the former the average B-clone size (or average B cell concentration) $\langle c \rangle$ can be computed from (12), and for the latter the same average can be computed from (22). In order to simplify the analysis of both equilibration regimes, we define the distribution

$$P(c) = \int dF P(F) \frac{e^{-\frac{1}{2}c \log(c) - \frac{1}{2}c^2}}{c \sqrt{2\pi} \beta}.$$

(29)

$$P(F) = \sum_{\sigma^1, \ldots, \sigma^n} P(\sigma^1, \ldots, \sigma^n) \frac{1}{M} \sum_{\nu=1}^M \delta \left( F - \sum_{\alpha=1}^n F_\nu(\sigma^\nu) \right).$$

(30)

Formula (29) gives the distribution of B-clone sizes for the fast B-clone and fast T-clone equilibration regimes when $n = 1$ and $\beta = n\beta$, respectively. This log-normal distribution can be interpreted as the (asymptotic) distribution of the ‘size’ $c_i$ of an element in a growth process, which changes its size at rate $w(c \rightarrow c') = \lambda_b(c' - (1 + g)c)$, in which $\lambda$ is a (mean) growth rate and $g$ is a growth factor [36]. The growth factor $g$ is related to the apoptosis parameter $\rho$ via $\log^2(1 + g) = \rho^{-1}$ and the noise parameter $\beta$ is related to the (rescaled) time $\lambda t$, where $\lambda$ is a (mean) growth rate, via $\lambda t = \beta^{-1}$. Furthermore, when the initial size $c_i = c_0$ then $c_0 = \exp[(F ln - \sqrt{\rho} \beta)/\rho]$ and the distribution (29) represents an average over random initial conditions. Since $c$ is a clone size, the growth process picture is consistent with the adaptive immune response: those B-cells which ‘survived’ interactions with the T cells, of which there are $c_0$, serve to ‘seed’ the B cell proliferation process.

Average and variance of (29) are given by

$$\langle c \rangle = e^{1/2\beta} \int dF P(F) e^{F ln \rho}.$$

(31)
\[ \langle (c - \langle c \rangle)^2 \rangle = e^{1/2 \beta} (e^{1/2 \beta} - 1) \int \! \! dF \ P(F)e^{2F}\langle p_{\text{inp}} \rangle. \] (32)

Let us next define the function \( \langle e^{F_{\text{inp}}} \rangle_\beta = \int \! \! dF \ P(F)e^{F_{\text{inp}}} \) and consider its properties. For \( \beta \to 0 \) this function can be expanded around \( \beta = 0 \) which gives \( \langle e^{F_{\text{inp}}} \rangle_\beta = M^{-1} e^{\sum_{\mu < M} b_j \sigma / \beta} + O(\beta) \), while in the opposite limit \( \beta \to \infty \) it is dominated by the ferromagnetic ground state \( \sigma = (1, \ldots, 1) \) which gives \( \langle e^{F_{\text{inp}}} \rangle_\beta = M^{-1} e^{\sum_{\mu < M} b_j \sigma / \beta} \) (note that for \( \theta_j = 0 \) the dynamics (5) can also ‘select’ the (1, ..., 1) state as \( N \to \infty \)), so \( \langle e^{F_{\text{inp}}} \rangle_\infty = (\langle F_{\text{inp}} \rangle_0)^N \). Furthermore, \( \langle e^{F_{\text{inp}}} \rangle_\beta \) is monotonic non-decreasing function of \( \beta \) and \( J(J_0 = J) \sum_{\nu < M} S_{\nu} a_{\nu} \), for which \( N < \infty \) is smooth everywhere except at \( \beta_0 \), when \( N \to \infty \) with \( \theta_j = 0 \) (see appendix A for details).

From the above analysis of \( \langle e^{F_{\text{inp}}} \rangle_\beta \) we infer that in the fast B-clone equilibration regime \( n = 1 \) the average number of B-cells \( \langle c \rangle \) is a monotonic non-decreasing function of \( \beta \) and \( J \) (see figure 4 for one of the possible behaviours). Combining the two plots in figure 4 then shows that \( \langle c \rangle \) must be a monotonic non-decreasing (non-increasing) function of the fraction of helper T-cells \( m_+ \) (regulator T-cells \( m_- \) and Ag, which is consistent with what was observed in in vivo [27, 33]. We note that the variance \( \langle (c - \langle c \rangle)^2 \rangle \) has a similar behaviour in this regime (the function \( \langle e^{2F_{\text{inp}}} \rangle_\beta = \int \! \! dF \ P(F)e^{2F_{\text{inp}}} \) used in (32) has the same behaviour as the function \( \langle e^{F_{\text{inp}}} \rangle_\beta \)).

The behaviour of the average \( \langle c \rangle \) in this fast T-clone equilibration regime with \( n \in \mathbb{Z}^+ \) is not so clear, due to the relation \( \beta = n \beta \). Here \( \langle c \rangle = e^{1/2 \rho \beta} \langle e^{F_{\text{inp}}} \rangle_\beta \) diverges for \( \beta \to 0 \) (it is finite when \( \beta \to \infty \)), and the derivative \( \partial (c) / \partial \beta = e^{1/2 \rho \beta} \partial (\langle e^{F_{\text{inp}}} \rangle_\beta) / \partial \beta = \langle e^{F_{\text{inp}}} \rangle_\beta / 2 \rho \beta \) is negative when \( \beta \to 0 \) (in this limit \( \partial (\langle e^{F_{\text{inp}}} \rangle_\beta) / \partial \beta < \infty \), and \( \langle e^{F_{\text{inp}}} \rangle_\beta / 2 \rho \beta \) is \( O(1/\beta) \)). For \( \beta \to \infty \) the derivative \( \partial (c) / \partial \beta \) could be positive or negative; now \( \langle e^{F_{\text{inp}}} \rangle_\beta / 2 \rho \beta \) is \( O(1/\beta) \), and one expects the convergence to zero of \( \partial (\langle e^{F_{\text{inp}}} \rangle_\beta) / \partial \beta \) to be strongly influenced by the topology of the effective T-clone network. However, if \( \beta \) is fixed and we vary only \( J \) then behaviours summarised in the figure 4 are also observed (due to monotonicity of the average \( \langle e^{F_{\text{inp}}} \rangle_\beta \) with respect to \( J \) which is shown in the appendix A) in the \( n \in \mathbb{Z}^+ \) fast T-clone equilibration regime.

The fast T-clone equilibration regime with \( n \in \mathbb{R}^+ \) is much more difficult to analyse, but in the low B-clone noise \( \beta \to \infty \) limit we expect at least in some regimes the same phase diagram as in the figure 4. To show this we first note that the marginal distribution (26) can be written as the integral

\[
P(\sigma) = \frac{\int db \ \theta(b)}{\int db \ e^{J b}} \frac{\int d\sigma \ e^{\beta \sum_{\mu < M} b_j \sigma / \beta} \delta(\sigma - \sigma_i)}{\int d\sigma \ e^{\beta \sum_{\mu < M} b_j \sigma / \beta}}, \tag{33}
\]

where

\[
\phi(b) = -\frac{1}{2} \rho \sum_{\mu = 1}^M b_\mu \left( \frac{J_\mu \theta_\mu}{\rho} \right)^2 + \frac{1}{\beta} \sum_{i=1}^N \log \int d\sigma \ e^{\beta \sum_{\mu < M} b_j \sigma / \beta}. \tag{34}
\]

This integral can be computed exactly by the Laplace method [37], which gives us

\[
P(\sigma) = \frac{\int d\sigma \ e^{\beta \sum_{\mu < M} b_j \sigma / \beta} \delta(\sigma - \sigma_i)}{\int d\sigma \ e^{\beta \sum_{\mu < M} b_j \sigma / \beta}}, \tag{35}
\]

where \( b_\mu^* \) is a solution of the following system of equations
corresponding to a maximum of the function $\phi(b)$. From the first moment

$$
\langle \sigma \rangle = \frac{\int \mathcal{D}\sigma \, e^{i\beta\sum_{i<j}b_{ij}\xi_{ij}^\mu\sigma_j} \delta(\sigma - \sigma_\text{eq})}{\int \mathcal{D}\sigma \, e^{i\beta\sum_{i<j}b_{ij}\xi_{ij}^\mu\sigma_j}}
$$

(37)

of the distribution (35), and the extremum condition (36), it follows that

$$
b_{ij} = J_\mu \left( \frac{1}{\rho} \sum_{j \in \partial_i} \xi_j^\mu \langle \sigma_j \rangle + \theta_\mu \right)
$$

(38)

which gives us the equation

$$
\langle \sigma \rangle = \frac{\int \mathcal{D}\sigma \, e^{i\beta\sum_{i<j}b_{ij}\xi_{ij}^\mu\sigma_j} \delta(\sigma - \sigma_\text{eq})}{\int \mathcal{D}\sigma \, e^{i\beta\sum_{i<j}b_{ij}\xi_{ij}^\mu\sigma_j}}
$$

(39)

Solutions of this equation can be used to compute the marginal T-clone distribution (35) via the equation

$$
P(\sigma) = \frac{\int \mathcal{D}\sigma \, e^{i\beta\sum_{i<j}b_{ij}\xi_{ij}^\mu\sigma_j} \delta(\sigma - \sigma_\text{eq})}{\int \mathcal{D}\sigma \, e^{i\beta\sum_{i<j}b_{ij}\xi_{ij}^\mu\sigma_j}}
$$

(40)

and the distribution (27) via the equation

$$
P_f(F) = \prod_{i \in \partial \mu} \frac{\int \mathcal{D}\sigma \, e^{i\beta\sum_{i<j}b_{ij}\xi_{ij}^\mu\sigma_j} \delta(\sigma - \sigma_\text{eq})}{\int \mathcal{D}\sigma \, e^{i\beta\sum_{i<j}b_{ij}\xi_{ij}^\mu\sigma_j}} \times \delta(F - F_\mu(\sigma)).
$$

(41)

The average B-cell concentration $\langle \sigma \rangle = \int_0^{\infty} dc \, c P(c)$ can be also computed from the integral

$$
\langle \sigma \rangle = M^{-1} \sum_{\mu=1}^{M} \lim_{\beta \rightarrow \infty} \int \mathcal{D}b \, P_{\beta/\beta}(b) e^{b_\mu} \text{ which gives us the equation}
$$

$$
\langle \sigma \rangle = \frac{1}{M} \sum_{\mu=1}^{M} e^{b_\mu} \left( \prod_{i \in \partial \mu} \delta(b_{ij}) \right).
$$

(42)

In the case of Ising variables $\sigma \in \{-1, 1\}$ and $\xi_{ij}^\mu \in \{-1, 1\}$, which also includes the binary case $\sigma \in \{0, 1\}$, the above framework leads to

$$
\langle \sigma \rangle = \text{tanh} \left( \beta \frac{1}{\rho} \sum_{j \in \partial i} \xi_j^\mu \langle \sigma_j \rangle + \theta_\mu \right).
$$

(43)

The simplest nontrivial clonal interaction structure is a regular network topology, as in figure 5, with uniform interactions\(^7\) $J_\mu = J$, $\xi_{ij}^\mu = 1$ for all $(i, \mu)$, and no self-activation,

\(^7\)We expect that having non-uniform interactions $J_\mu$ would lead only to quantitative but not qualitative differences with the uniform $J_\mu = J$ case studied here, i.e. we expect the same phase diagram in both cases, provided that non-uniform $J_\mu$ are well behaved.
\[ \theta_i = \theta_0. \] Here equation (43) simplifies considerably. From (38) we infer that \( b_i = \frac{L}{\rho} K \langle \sigma_i \rangle, \)
with \( |\partial\mu| = K, \) and if we define the average magnetization \( m = \langle \sigma_i \rangle \) we find that
\[ m = \tanh (J^2 L K m / \rho), \tag{44} \]
with \( |\partial| = L. \) The average B cell concentration \( \langle c \rangle = e^{k \mu m} \) follows from (42). The distribution of fields (41) takes the following simple form, with \( P(F) = P_\rho(F): \)
\[ P(F) = \sum_{\sigma_i} \frac{e^{\beta_0 L K \sum_i \sigma_i}}{2^K \cosh^K \left( \frac{\beta_0 L K m}{\rho} \right)} \delta \left( F - J \sum_{j=1}^K \sigma_j \right). \tag{45} \]

Figure 6. Behaviour of B- and T-clones in the model with fast T-clone equilibration in the noiseless B-clone regime (\( \beta \rightarrow \infty \)). The system, here defined on a graph with connectivity \( L = K = 4, \) was studied for the high \( \beta < \beta_c (\beta_c = 0.0625) \) and low \( \beta > \beta_c \) T clone noise levels with \( J = 1 \) and for the low \( J < J_c (J_c = 0.25) \) and high \( J > J_c \) Ag levels with \( \beta = 1. \) Top left: The average B-clone size, \( \langle c \rangle, \) as a function of the fraction of T-regulator cells, \( m = \frac{1}{2} (1 - m), \) for \( J = 1 \) and \( \beta \in [0, 0.5]. \) Inset: \( \langle c \rangle \) as a function of \( \beta. \) Top right: \( \langle c \rangle \) as a function of \( m \) for \( \beta = 1 \) and \( J \in [0, 1]. \) Inset: \( \langle c \rangle \) as a function of \( J. \) Bottom: The distribution \( P(F) \) for \( \beta = 0.0525 \) with \( m = \frac{1}{2} \) (left), for \( \beta = 0.0855 \) with \( m = 0.1 \) (centre) and for \( \beta = 0.0855 \) with \( m = 0.9 \) (right).
Equation (44) has $m = 0$ as its solution for any $\beta$, but this solution becomes unstable for $\beta > \beta_c$, where $\beta_c = \frac{\mu}{\rho \mu}$, and two stable solutions $|m| 
eq 0$ emerge. Furthermore, the average number of B cells, $\langle c \rangle$, is a monotonic non-increasing function of the fraction of T regulator cells $m_− = \frac{1}{2}(1 − m)$ (the representative case of $L = K = 4$ and $\rho = 1$ is studied in the figure 6). The B-clones are receiving predominantly excitatory signals from the T clones, described by $P(F)$, when $m_− < \frac{1}{2}$, and predominantly inhibitory signals when $m_− > \frac{1}{2}$, as can be seen in figure 6. For $\beta < \beta_c$ the point $m = 0$ is the only solution, and the B cells are no longer controlled by the T cells: $\langle c \rangle = 1$ for all $\beta$ and $J$. For $\beta > \beta_c$ and $m_− < \frac{1}{2}$ the average number of B cells $\langle c \rangle$ is increasing with $\beta$ and $J$ (see inset in figure 6). Thus $\beta_c$ separates the ‘low-dose tolerance’ PM phase, where immune system is insensitive to Ag, from the ‘vigorous immune response’ FM phase where it is very sensitive to Ag.

So far we have been able to discuss the behaviour of T- and B-clones by making only weak assumptions about network topologies and interactions. This analysis is qualitative, and only covers the cases where we can map our model onto ferromagnetic Ising spin systems, which are quite well understood [34, 35]. Unfortunately, one cannot construct such arguments for the case where $\sigma_l \in \{0, 1\}$ with $\xi_l^m \in \{−1, 1\}$, which is again equivalent to an Ising model but is no longer ferromagnetic; the map $\sigma : \{−1, 1\} \rightarrow \{0, 1\}$, where $\sigma = \frac{1 + \epsilon}{2}$, here gives rise to site-dependent external fields, which could be positive or negative. Moreover, to make quantitative predictions about the observables such as fractions of T cells, $m_±$, and concentrations of B cells, $\langle c \rangle$, and more informative observables such as the B-clone distribution (29), we have to be more specific about the graph $G$ and the interaction parameters $\{\xi_l^m, J, \}$.

In the absence of microscopic knowledge about interactivity in real immune systems, we will follow the ‘Bayesian’ route and choose topological and interaction parameters randomly, but subject to biological constraints such as the average number of regulator T cells $p = \frac{1}{N} \sum_{l=1}^{N} \xi_l \xi_{l-1}$ or the number of T cells that a single B cell can interact with (for example in the graphs of figure 5 these numbers are 4 and 8) which can be deduced from [38]. In real immune systems the interactions of lymphocytes with Ag occur in a 3-dimensional volume, so it seems reasonable to choose a regular 3d lattice for our graph topology. However, even if we choose a simpler 2d lattice, there are still many possibilities for how to construct the graph $G$ (two of which are shown in figure 5). Systems interacting on $d$-dimensional lattices, with $d > 1$, are hard to study analytically. One therefore often uses locally tree-like [39] random topologies, an approximation which is expected to be good away from phase transitions [40].

3.1. Bethe–Peierls approximation

For systems interacting on trees, relevant local observables such as the distribution of fields (27) can be computed recursively. Let us first consider the distribution (28), which governs the (replicated) T-clone variables $\{\sigma_n^\mu\}$ in both the fast B-clone ($n = 1$) and fast T-clone ($n \in \mathbb{Z}^+$) equilibration regimes, defined on a factor-tree $T_\mu(r)$ of radius $r$ which is centred at the factor-node $\mu$ (see figure B1 in appendix B). The field $F_\mu(\{h_l\}) = f_\mu(\sum_{l=1}^{n} \xi_l h_l + \theta_\mu)$, where $h_l = \sum_{n=1}^{n} \sigma_n^\mu$, on this factor node is governed by the distribution.

8 Derivation details of the first three equations in this section are provided in appendix B.
where the cavity distribution $P_{\mu}[h_i]$ can be computed recursively, starting from the variables located in the boundary $\partial T_\mu$ of the tree $T_\mu$, via the equation

$$P_{\mu}[h_i] = \sum_{\{h_i\}} \prod_{v \in \partial T_\mu} \prod_{j \in \partial v} P_{\mu}[h_j] \sum_{\sigma_{i_1}, \ldots, \sigma_{i_{\mu}}} \delta_{\sigma_{i_1, \ldots, \sigma_{i_{\mu}}}^{\sum_{i_1}^\infty \sigma_{i_{\mu}}}} \frac{e^{\beta \sum_{i_1}^\infty \sigma_{i_{\mu}}} f(\sum_{j_{i_1}^\infty \sigma_{i_{\mu}}}^\infty \sigma_{i_{\mu}})}{\sum_{\sigma_{i_1, \ldots, \sigma_{i_{\mu}}}^{\sum_{i_1}^\infty \sigma_{i_{\mu}}}} e^{\beta \sum_{i_1}^\infty \sigma_{i_{\mu}}} f(\sum_{j_{i_1}^\infty \sigma_{i_{\mu}}}^\infty \sigma_{i_{\mu}})} - 1 \right).$$

Similarly the magnetization $\langle \sigma_i \rangle = \frac{1}{n} \sum h P(h) h$, where $P(h) = \sum_{\sigma} P(\sigma) \delta_{\sigma_{i_1, \ldots, \sigma_{i_{\mu}}}^{\sum_{i_1}^\infty \sigma_{i_{\mu}}}}$, of the tree $T_i$ can be computed via the equation

$$\langle \sigma_i \rangle = \sum_{h} \frac{1}{n} \sum_{\{h_i\}} \prod_{v \in \partial T_\mu} \prod_{j \in \partial v} P_{\mu}[h_j] \sum_{\sigma_{i_1}, \ldots, \sigma_{i_{\mu}}} \delta_{\sigma_{i_1, \ldots, \sigma_{i_{\mu}}}^\infty} \frac{e^{\beta \sum_{i_1}^\infty \sigma_{i_{\mu}}} f(\sum_{j_{i_1}^\infty \sigma_{i_{\mu}}}^\infty \sigma_{i_{\mu}})}{\sum_{\sigma_{i_1, \ldots, \sigma_{i_{\mu}}}^\infty} e^{\beta \sum_{i_1}^\infty \sigma_{i_{\mu}}} f(\sum_{j_{i_1}^\infty \sigma_{i_{\mu}}}^\infty \sigma_{i_{\mu}})} - 1 \right).$$

We note that the equations derived in this section can be used to compute local observables, such as the distribution $P_{\mu}(F)$ or the local magnetization $\langle \sigma_i \rangle$, on locally-tree like graphs. In these graphs a ‘ball’ of radius $r$ centred on any variable node $i$ (or factor node $\mu$) converges to the tree of radius $r$ centred at this node, $T_i(r)$, when $N \to \infty$ (or when $M \to \infty$ with $M/N < \infty$), for any $r \geq 39$. An observable associated with node $i$ in such graphs is usually approximated by the same observable computed on $T_i(\infty)^9$. Furthermore, local observables computed on random trees can be used to compute densities: for example, upon assuming that the density $P(F) = \lim_{M \to \infty} \frac{1}{M} \sum_{\mu=1}^M P_{\mu}(F)$ is self-averaging [39] we can replace this average with the average over the trees $T_\mu(\infty)$ for which $n \in \mathbb{Z}^+$ gives us the equation

$$P(F) = \sum_{K \geq 1} P(K) \left\{ \prod_{i=1}^K \int \{dP\} W(\{P\}) \right\} \int dJ \int d\theta \ P(\theta) \left( \frac{\delta_{\sum_{i=1}^K \xi_j h_i + \theta, f}}{\sum_{\{h_i\}} \prod_{i=1}^K P_{\mu}[h_i] \xi_1} \right).$$

The above equation was derived for the ensemble of random graphs with the prescribed distributions $P(L)$ and $P(K)$ of the variable-node and factor-node connectivities, respectively. We have also assumed that all parameters $\{\xi_j\}$, $\{J_j\}$ and $\{\theta_j\}$ are drawn independently at random from the distributions $P(\xi), P(J)$ and $P(\theta)$ respectively. The distribution $W(\{P\})$ is

---

9 This procedure is exact for ferromagnetic Ising models with uniform interactions [41].
drawn from $J. \text{Phys. A: Math. Theor.}$ variable nodes (T-clones). We also assume that this regularity imposes the constraint $P_i$ is drawn randomly and uniformly from the population which is obtained via a population dynamics algorithm \cite{42}, which at each step replaces a member $i$ of the population $\mathcal{P} = \{P_i|\xi\}$ with the new value

$$P_i[\xi] = \sum_{\{h_i\}} \left\{ \prod_{a=1}^{L-1} \prod_{j=1}^{K-1} P_{i_a}[h_j]^{[\xi_a]} \right\} \sum_{\sigma'_{\ldots \sigma_n}} \delta_{\xi \sum_{a=1}^{L-1} \rho_a \sum_{a=1}^{K-1} I_a \sum_{a=1}^{K-1} J_a + \theta_i + \theta}$$

which is computed using the $(L-1) \times (K-1)$ distributions $P_{i_a}[h_j]^{[\xi_a]}$. Here $i_{a,j} \in \{1, \ldots, N\}$ is drawn randomly and uniformly from the population $\mathcal{P}$, and $L$ and $K$ are random integers drawn from $Q(L) = LP(L) \sum_{L \geq 1} LP(L)$ and $Q(K) = KP(K) \sum_{K \geq 1} KP(K)$, respectively. The parameters $\{\xi_a\}$, $\{J_a\}$ and $\{\theta_a\}$ are also random numbers, drawn from the distributions $P(\xi)$, $P(J)$ and $P(\theta)$, respectively.

Finally, we note that Bethe–Peierls approximation also can be used to study the $n \in \mathbb{R}^+$ case (see appendix C).

### 3.2. Analysis of homogeneous systems on random regular factor-graphs

The simplest non-trivial interaction topologies that allows us to obtain more quantitative results are models defined on random regular factor-graphs. The ensemble of these graphs contains all graphs with fixed connectivities of variable-nodes, $|\partial i| = L$, and factor-nodes, $|\partial \mu| = K$. This regularity imposes the constraint $L/K = M/N$ on the ratio of factor-nodes (B-clones) to variable nodes (T-clones). We also assume that $J_a = J$ and $\theta_a = \theta$, and consider the case of $\sigma_i \in \{-1, 1\}$ with $n \in \mathbb{Z}^+$. For $N \to \infty$, random regular factor-graphs are locally tree-like \cite{39}, and we therefore expect that the ‘tree approximation’, described in the previous section will be exact on such graphs. Furthermore, the (local) topology of the T-clones system \cite{28} in this connectivity regime resembles a variant of the Bethe lattice \cite{34}, where each node is connected to exactly $L$ fully connected graphs (cliques) of size $K$ (see figure 7). From now on we will call such a graph a random $(K, L)$-regular clique graph.
the computation of \( P(\mu) \), see values The same is true for the average magnetization where \( \mu \) is the parameter of B-clone equilibration. This regime can be studied using \( \mu \) and \( \mu \) T-clone noise \( \mu \), with \( \mu = 0.0929 \), for the values \( \beta = 0.0639 \), with \( \mu = 0.1219 \) with \( \mu = 0.1 \) (centre) and \( \beta = 0.1219 \) with \( \mu = 0.9 \) (right).

Let us consider the system (28) on one of such graphs. Since the variable nodes and factor nodes are now all equivalent, the cavity distribution \( P[h] \) in equation (47) is the same for all \( i \) and \( \mu \), i.e. \( P[h] = P[h] \), which gives us the recursive equation

\[
P[h] = \frac{\left[ \sum_{(b_i)} \left[ \prod_{j=1}^{K-1} P[h_j] \right] e^{\beta \left( \sum_{j=1}^{K} \rho_j b_j + \mu \right)} \right]^{L-1} \left( \frac{n}{n+h} \right)}{\sum_{(b_i)} \left[ \prod_{j=1}^{K-1} P[h_j] \right] e^{\beta \left( \sum_{j=1}^{K} \rho_j b_j + \mu \right)} \right]^{L-1} \left( \frac{n}{n+h} \right)},
\]

(51)

where \( h \in \{-n, -n+1, \ldots, n-1, n\}, \) and where the binomial coefficient \( \left( \frac{n}{n+h} \right) \) results from the computation of \( \sum_{\sigma_1, \ldots, \sigma_L} \delta_{h_k} \sum_{\sigma_1, \ldots, \sigma_L} \). The solution of this equation can be used to compute the field distribution (46), via the equation

\[
P(F) = \frac{\sum_{(b_i)} \left[ \prod_{j=1}^{K-1} P[h_j] \right] e^{\frac{\beta}{n} F} \delta \left( F - J \sum_{j=1}^{K} \rho_j b_j \right)}{\sum_{(b_i)} \left[ \prod_{j=1}^{K-1} P[h_j] \right] e^{\frac{\beta}{n} \sum_{j=1}^{K} \rho_j b_j}}.
\]

(52)

The same is true for the average magnetization \( m = \frac{1}{n} \sum_{h} P[h] h \), where \( P(h) \) is the RHS of the equation (51) with \( L = L + 1 \). The latter result is obtained by comparing the equations (B.10) and (47).

Let us first consider the case of fast B-clone equilibration. This regime can be studied using equations (51), (52) and (29), with \( n = 1 \). For \( n = 1 \) the T-clone system is in the paramagnetic (PM) phase when \( \beta < \beta_c \) (see appendix D). Here the distribution of fields \( P(F) \) is symmetric, see figure 8, which gives rise to the quadratic behaviour in the distribution of B-clones \( P(c) \) seen in figure 9. As we increase \( \beta \), the system will enter the ferromagnetic (FM) \( m = 0 \) phase when \( \beta > \beta_c \). Here the distribution \( P(F) \) is no longer symmetric, see figure 8, and the distribution of B-clones is seen to exhibit ‘power law’ behaviour when \( \beta \to \infty \). We note that in this regime the distribution (29) is dominated by the probability \( P(F) \) evaluated at the field values \( F = \pm JK \), corresponding to the magnetization \( m > 0 \) or \( m < 0 \). In particular when \( \beta \to \infty \), we have

\[
\]
We note that power law behaviour is also present in the empirical B-clone distributions \[30\]. Moreover, in the FM phase the average concentration of B cells \[31\] can be controlled by the fraction \(m-\) of regulating T-clones; see figure 9. Comparing with the results of fast T-clone equilibration, summarised in figure 6, we note that here \(\beta_c\) is larger and hence a larger amount of Ag (i.e. a larger value of \(J\)) is needed by the immune system to mount a vigorous immune response (see figure 9), i.e. the system is less sensitive\[10\]. Also the behaviour of B-clones in the 'low-dose tolerance' PM regime is different: the average number of B-cells, \(\langle c \rangle\), is increasing with \(\beta\) and \(J\) in figure 9, but in figure 6 it is a constant.

In the regime of fast T-clone equilibration with \(Z \in \mathbb{N}\) \((n > 2)\) we solve the recurrence equation \[51\] using the initial condition

\[P(c) \propto c^{-1+\beta} e^{-\frac{1}{2} \rho \beta \log(3)c},\]  

(53)

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(53)

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(53)

This aspect of the model can be used to distinguish between the equilibration regimes when comparing model with experimental data.
dependence on less pronounced, see figure 11.\[51\]

Finally, we note that equation (51), which thus far we were able to analyse only numerically, can be studied further analytically when the connectivity $L$ or $K$ is large. Also, this
analysis allows us to probe the extreme regime of large B-clone numbers $M \gg N$ or large T-clone numbers $(N \gg M)$.

### 3.3. Large $L$ and finite $K$ (or $M/N \to \infty$) regime

To start we note that equation (51) can also be written in the form

$$
P(h) = \sum_{(N)} \frac{(L-1)! \left\{ \Pi_{h_0}(\Pi_{j=1}^{h_0-1} P(h_j))^{M(h_0)} \right\} \frac{\beta^2}{\gamma^{\sum_{j=1}^K M(h_j)}} \left( \frac{n}{n+h} \right)^{\frac{n}{2}}}{\Pi_{h_0} M(h_0)!} \times \sum_{h} (L-1)! \left\{ \Pi_{h_0}(\Pi_{j=1}^{h_0-1} P(h_j))^{M(h_0)} \right\} \frac{\beta^2}{\gamma^{\sum_{j=1}^K M(h_j)}} \left( \frac{n}{n+h} \right)^{\frac{n}{2}},
$$

(55)
where $\sum_{\{h_j\}} N[\{h_j\}] = L - 1$ and $N[\{h_j\}] \in \{0, \ldots , L - 1\}$. Using $N! = \rho_N N^N e^{-N}$ with $\rho_0 = 1$ and $\rho_N = \sqrt{2\pi Ne^{1/2}}$, where $|\theta_N| < 1$ (see [43]), gives us

$$
\frac{(L - 1)!}{\prod_{\{h_j\}} N[\{h_j\}]!} = \frac{\rho_{L-1}(L - 1)^{L-1} e^{-(L-1)}}{\Pi_{\{h_j\}} \rho_N N[\{h_j\}] e^{-N[\{h_j\}]}} = e^{-(L-1) \sum_{\{h_j\}} \frac{N[\{h_j\}]}{L-1} \log \frac{N[\{h_j\}]}{\rho_N}} N[\{h_j\}] + R_{L-1} N[\{h_j\}],
$$

(56)

where $R_{L-1}(N[\{h_j\}]) = \log \rho_{L-1} - \sum_{\{h_j\}} \log \rho_N[N[\{h_j\}]]$ (it is easy to show that $R_{L-1}(N[\{h_j\}]) = \Omega([N/L - 1]) + O(\log(L - 1))$). Let us define the functional

$$
D \left( \frac{N}{L - 1} \parallel p^{K-1} \right) = \sum_{\{h_j\}} N[\{h_j\}] \log \frac{N[\{h_j\}]}{(L - 1) \prod_{\{h_j\}} \rho_N[N[\{h_j\}] + R_{L-1} N[\{h_j\}])},
$$

(57)

which is the Kullback–Leibler (KL) ‘distance’ [44] between the distributions $N[\{h_j\}]/(L - 1)$ and $\prod_{\{h_j\}} \rho_N[N[\{h_j\}]$, and use the identity (56) to compute the numerator in the equation (55). This computation is greatly simplified if we assume that $J^2 = 1/(L - 1)$ and take the limit $L \to \infty$, which gives us

$$
\sum_{\{h_j\}} e^{-(L-1) D \left( \frac{N}{L - 1} \parallel p^{K-1} \right) + R_{L-1} N[\{h_j\}]} \sum_{\{h_j\}} N[\{h_j\}] \frac{\rho_N[N[\{h_j\}]]}{L-1} \sum_{\{h_j\}} e^{\frac{\beta}{2\pi}(h + \sum_{j=1}^{K-1} h_j)}
$$

$$
= \sum_{\{h_j\}} e^{-(L-1) D \left( \frac{N}{L - 1} \parallel p^{K-1} \right) + R_{L-1} N[\{h_j\}]} \sum_{\{h_j\}} N[\{h_j\}] \frac{\rho_N[N[\{h_j\}]]}{L-1} \sum_{\{h_j\}} e^{\frac{\beta}{2\pi}(h + \sum_{j=1}^{K-1} h_j)}
$$

$$
= \sum_{\{h_j\}} \left[ \prod_{\{h_j\}} \rho_N[N[\{h_j\}] \frac{\rho_N[N[\{h_j\}]]}{L-1} \sum_{\{h_j\}} e^{\frac{\beta}{2\pi}(h + \sum_{j=1}^{K-1} h_j)} \right]^{-1}
$$

$$
= e^{\sum_{\{h_j\}} \frac{\beta}{2\pi}(\sum_{j=1}^{K-1} h_j) \sum_{\{h_j\}} e^{\frac{\beta}{2\pi}(h + \sum_{j=1}^{K-1} h_j)}} = e^{\frac{\beta}{2\pi}(h^2 + 2h(K-1)\delta)},
$$

(58)

where $\langle \tilde{h} \rangle = \sum_{\{h_j\}} \hat{h} \rho[\{h_j\}]$. Using the above result to compute the distribution (55) gives

$$
P[\hat{h}] = \frac{e^{\frac{\beta}{2\pi}(h^2 + 2h(K-1)\delta)}}{\sum_{\{h_j\} e^{\frac{\beta}{2\pi}(h^2 + 2h(K-1)\delta)}}} \left( \begin{array}{c} \frac{\beta}{2\pi} \\ \frac{\beta}{2\pi} \end{array} \right) \left( \begin{array}{c} n + \frac{\beta}{2\pi} \\ \frac{\beta}{2\pi} \end{array} \right)
$$

(59)

$$
\langle h \rangle = \frac{\sum_{\{h_j\} e^{\frac{\beta}{2\pi}(h^2 + 2h(K-1)\delta)}} \left( \begin{array}{c} \frac{\beta}{2\pi} \\ \frac{\beta}{2\pi} \end{array} \right) \left( \begin{array}{c} n + \frac{\beta}{2\pi} \\ \frac{\beta}{2\pi} \end{array} \right)}{\sum_{\{h_j\} e^{\frac{\beta}{2\pi}(h^2 + 2h(K-1)\delta)}}}
$$

(60)

We note that in this limit the cavity distribution $P[\hat{h}]$ and the physical distribution $P(h)$ are the same, and the average $\langle h \rangle / n$ equals the average magnetization $m$.

Let us next define the average
\[
\langle g(h) \rangle_h = \frac{\sum_{n} e^{\frac{\beta n h}{n+\frac{1}{2}}}}{\sum_{n} e^{\frac{\beta (n+\frac{1}{2}) h}{2(n+\frac{1}{2})}}} g(h).
\] (61)

It allows us to write the RHS of equation (60) as \( \langle h \rangle_x \), where \( x = \langle \tilde{h} \rangle \). The function \( \langle h \rangle_x \) has the following properties for all \( x \geq 0 \):

(i) \( \langle h \rangle_{-x} = -\langle h \rangle_x \)

(ii) \( \frac{\partial}{\partial n} \langle h \rangle_x = \frac{\beta}{n \rho}(K-1)\langle (h-\langle h \rangle)\tilde{h} \rangle_x \geq 0 \)

(iii) \( \frac{\partial^2}{\partial n^2} \langle h \rangle_x |_{n=0} = \frac{(K-1)}{n \rho} \langle h^2 \rangle_0 + \frac{\beta(K-1)}{n \rho} \langle (h^2-\langle h \rangle^2) \rangle_0 > 0. \) 

From (i) it follows that \( \langle \tilde{h} \rangle = 0 \) is a fixed point of the recursion (60). According to (ii) this point becomes unstable when \( \left. \frac{\partial}{\partial n} \langle h \rangle_x \right|_{n=0} = \frac{\beta}{n \rho}(K-1)(h^2)_0 > 1 \). The derivative \( \left. \frac{\partial}{\partial n} \langle h \rangle_x \right|_{n=0} \) is a monotonic nondecreasing function of \( \beta \), by the inequality

\[
\frac{\partial^2}{\partial \beta^2} \langle h \rangle_x |_{n=0} = \frac{(K-1)}{n \rho} \langle h^2 \rangle_0 + \frac{\beta(K-1)}{n \rho} \left[ \langle (h^2-\langle h \rangle^2) \rangle_0 \right] \geq 0.
\] (62)

Furthermore, for \( \beta \) large

\[
\langle h^2 \rangle_0 = \frac{\sum_{n} e^{\frac{\beta n h}{n+\frac{1}{2}}} h^2}{\sum_{n} e^{\frac{\beta (n+\frac{1}{2}) h}{2(n+\frac{1}{2})}}} = \frac{\sum_{n} e^{\frac{\beta n h}{n+\frac{1}{2}}}}{\sum_{n} e^{\frac{\beta (n+\frac{1}{2}) h}{2(n+\frac{1}{2})}}} \langle h^2 \rangle_0 = \frac{\sum_{n=0} e^{\frac{\beta n h}{n+\frac{1}{2}}}}{\sum_{n=0} e^{\frac{\beta (n+\frac{1}{2}) h}{2(n+\frac{1}{2})}}} \langle h^2 \rangle_0 = n^2
\] (63)

and hence \( \left. \frac{\partial}{\partial n} \langle h \rangle_x \right|_{n=0} \in [0, (\beta n \rho)(K-1)] \) for \( \beta \in [0, \infty) \). Thus there exists a finite value \( \beta_c \) where the trivial solution \( \langle \tilde{h} \rangle = 0 \) of the recursion (60) becomes unstable. According to (i) and (iii) the function \( \langle h \rangle_x \) is concave (respectively convex) on the interval \( x \in (0, \infty) \) (respectively \( x \in (-\infty, 0) \), and hence this function crosses the diagonal \( x \) only once on this interval when \( \left. \frac{\partial}{\partial n} \langle h \rangle_x \right|_{n=0} > 1 \). This intersection corresponds to the stable solution \( |\langle \tilde{h} \rangle| \neq 0 \) of equation (60).

To prove (iii) we use the identity \( \langle n \rangle = n \beta \sigma \), where \( \langle n \rangle = n \beta \sigma \) in equation (61), which gives

\[
\langle g(h) \rangle_h = \frac{\sum_{\sigma} e^{\frac{\beta \sigma \sigma^*}{2}} \sum_{\sigma^*} e^{\frac{\beta \sigma^* \sigma^*}{2}} g(\sum_{\sigma^*} \sigma^* \sigma^* + \sum_{\sigma^*} \sigma^* \sigma^* + \sum_{\sigma^*} \sigma^* \sigma^*)}{\sum_{\beta^2} e^{\frac{\beta \sigma \sigma^*}{2}} \sum_{\sigma^*} e^{\frac{\beta \sigma^* \sigma^*}{2}} g(\sum_{\sigma^*} \sigma^* \sigma^* + \sum_{\sigma^*} \sigma^* \sigma^* + \sum_{\sigma^*} \sigma^* \sigma^*)}.
\] (64)

From this we infer that the average \( \langle h \rangle_x \) equals the average magnetization in ferromagnetic Ising system with interactions and external field given by \( 1/n \rho \) and \( (K-1)x \), respectively.

By the the Griffiths–Hurst–Sherman (GHS) theorem [45] the average magnetization in such systems is a concave function of a positive external field, and hence inequality (iii) is true.
Furthermore, upon using (64) to compute the average \( \langle h^2 \rangle_0 \) we obtain the identity
\[
(3\beta n \rho)(K - 1)\langle h^2 \rangle_0 = (3\beta \rho)(K - 1) + (2\beta n \rho)(K - 1) \sum_\alpha \langle \sigma^\alpha \sigma^\alpha \rangle_0.
\]
The correlation function \( \langle \sigma^\alpha \sigma^\alpha \rangle_0 \geq 0 \), by the Griffiths–Kelly–Sherman (GKS) theorem [35], and \( \langle \sigma^\alpha \sigma^\alpha \rangle_0 \leq 1 \), since \( \sigma^\alpha \in \{-1, 1\} \) for all \( \alpha \). From this follow the inequalities
\[
(3\beta \rho)(K - 1) \leq (3\beta n \rho)(K - 1) \leq \langle h^2 \rangle_0 < (n\beta \rho)(K - 1),
\]
which gives us the following lower and upper bounds on \( \beta L \); (\( \rho/n \) (\( K - 1 \)) < \( \beta \leq \rho/(K - 1) \)).

Next we define the function \( m(h) = h/n \) and the average \( m = \sum_h P[h]m(h) \). Using the recursive equation (60) we obtain
\[
m = \frac{\sum_h e^{n\beta/m^2(h)}m(h)\left(\frac{n}{n+\ln(h)}\right)}{\sum_h e^{n\beta/m^2(h)}m(h)\left(\frac{n}{n+\ln(h)}\right)}
\]
for this average. If in addition we define the function \( \phi_n(m(h)) = (\beta/2\rho)[m(h)^2 + 2m(h) (K - 1)m] + \frac{1}{n} \log \left(\frac{n}{n+\ln(h)}\right) \) then for \( n \to \infty \) we obtain the equation
\[
m = \frac{\sum_h e^{n\beta/\phi_n(m(h))}m(h)}{\sum_h e^{n\beta/\phi_n(m(h))}}
\]
\[
= \frac{\sum_h e^{-n(\sup_{M \in \{-1,1\}} \phi_n(M) - \phi_n(m(h)))}m(h)}{\sum_h e^{-n(\sup_{M \in \{-1,1\}} \phi_n(M) - \phi_n(m(h)))}}
\]
\[
= \text{arg sup}_{M \in \{-1,1\}} \left\{ \frac{\beta}{2\rho} [M^2 + 2M(K - 1)m] + S\left(\frac{1}{2}(1 + M)\right) \right\},
\]
where \( S(u) \) is the Shannon entropy of a binary variable \( s \in \{0, 1\} \) with average \( \langle s \rangle = u \) [44]. From the above it follows that \( m \) satisfies the equation
\[
m = \tanh(\beta Kn \rho)
\]
which recovers the \( \beta \to \infty \) equation (44), in the limit \( L \to \infty \) with \( J^2 = 1/(L - 1) \).

Finally, we note that the field distribution (52) converges to \( \delta(F) \) in the limit \( L \to \infty \) with \( J^2 = 1/(L - 1) \), and although the T-clones are responding to Ag in this regime (one can set \( J^2 = \tilde{J}^2/(L - 1) \)), they are unable to control the B-clones.

### 3.4. Finite L and large K (or M/N → 0) regime

The starting point of our analysis in this parameter regime is to note that the equation (51) can be written in the following form:
\[
P[\tilde{h}] = \frac{\int Dz \ e^{\beta \sqrt{\tilde{\rho}} \left( \sum_h P[h] e^{\beta \sqrt{\tilde{\rho}} h} \right)^{K-1} \left(\frac{n}{n+\tilde{h}}\right)}}{\sum_h \int Dz \ e^{\beta \sqrt{\tilde{\rho}} \left( \sum_h P[h] e^{\beta \sqrt{\tilde{\rho}} h} \right)^{K-1} \left(\frac{n}{n+\tilde{h}}\right)}}
\]
(68)
where \( Dz = e^{-z^2/2}dz/\sqrt{2\pi} \). Using the multinomial theorem we can write
\[
\left( \sum_{h} P[h] e^{z_j \sqrt{\frac{z}{2\pi}}} \right)^{K-1} = \sum_{\{\lambda[h]\}} \frac{(K-1)!}{\prod_h \lambda[h]} \left\{ \prod_h P^{\lambda[h]}[h] e^{z_j \sqrt{\frac{z}{2\pi}}} \right\}^{\lambda[h]}
\]
\[
= \sum_{\{\lambda[h]\}} \frac{(K-1)!}{\prod_h \lambda[h]} \left\{ \prod_h P^{\lambda[h]}[h] \right\} e^{z_j \sqrt{\frac{z}{2\pi}}} \sum_h \lambda[h],
\]
where \( \sum_h \lambda[h] = K - 1 \) and \( \lambda[h] \in \{0, 1, \ldots, K - 1\} \). Using the above in equation (68) gives us
\[
P[h] = \left[ \sum_{\{\lambda[h]\}} \frac{(K-1)!}{\prod_h \lambda[h]} \left\{ \prod_h P^{\lambda[h]}[h] \right\} e^{z_j \sqrt{\frac{z}{2\pi}}} \sum_h \lambda[h] \right]^{L-1} \left( \frac{n}{n+\frac{h}{2}} \right)
\]
\[
\times \left[ \sum_{\{\lambda[h]\}} \frac{(K-1)!}{\prod_h \lambda[h]} \left\{ \prod_h P^{\lambda[h]}[h] \right\} e^{z_j \sqrt{\frac{z}{2\pi}}} \sum_h \lambda[h] \right]^{L-1} \left( \frac{n}{n+\frac{h}{2}} \right).
\]
Let us consider the multinomial distribution in equation (70). Using the formula \( K! = \rho_{K} K^{K} e^{-\beta} \), with \( \rho_{0} = 1 \) and \( \rho_{K > 0} = \sqrt{2\pi K} e^{\frac{\beta_{K}}{2}} \), where \( |\beta_{K}| < 1 \), we obtain
\[
\frac{(K-1)!}{\prod_h \lambda[h]} \left\{ \prod_h P^{\lambda[h]}[h] \right\} e^{z_j \sqrt{\frac{z}{2\pi}}} \sum_h \lambda[h] \log P[h]
\]
\[
= e^{-\beta(K-1)} \sum_{\{\lambda[h]\}} \lambda[h] \log \left( \frac{\lambda[h]}{(K-1)P[h]} \right) - \sum_{\{\lambda[h]\}} \lambda[h] \log P[h] - e^{-\beta(K-1)} \log \rho_{K-1}
\]
\[
= e^{-\beta(K-1)} \left\{ \frac{\sum_{\{\lambda[h]\}} \lambda[h] \log \left( \frac{\lambda[h]}{(K-1)P[h]} \right) - \sum_{\{\lambda[h]\}} \lambda[h] \log P[h]}{\rho_{K-1}} \right\} + \text{const}.
\]
The functional \( D\left( \frac{\mathcal{N}}{P} \right) = \sum_{\{\lambda[h]\}} \lambda[h] \log \left( \frac{\lambda[h]}{(K-1)P[h]} \right) \) is the KL distance between the distributions \( \mathcal{N}[h]/(K-1) \) and \( P[h] \). We have also defined the remainder functional
\[
R\left( \frac{\mathcal{N}}{K-1} \right) = \frac{1}{2} \sum_{\{\lambda[h]\}} \log \left( \frac{\lambda[h]}{(K-1)P[h]} \right) + \sum_{\{\lambda[h]\}} \frac{\lambda[h]}{(K-1)P[h]} - e^{-\beta(K-1)} \log \rho_{K-1}
\]
Let us next consider the average
\[
\frac{\sum_{\{\lambda[h]\}} \frac{(K-1)!}{\prod_h \lambda[h]} \left\{ \prod_h P^{\lambda[h]}[h] \right\} e^{z_j \sqrt{\frac{z}{2\pi}}} \sum_h \lambda[h] \}}{\sum_{\{\lambda[h]\}} \frac{(K-1)!}{\prod_h \lambda[h]} \left\{ \prod_h P^{\lambda[h]}[h] \right\} e^{z_j \sqrt{\frac{z}{2\pi}}} \sum_h \lambda[h]}
\]
\[
= \frac{\sum_{\{\lambda[h]\}} \frac{(K-1)!}{\prod_h \lambda[h]} \left\{ \prod_h P^{\lambda[h]}[h] \right\} e^{z_j \sqrt{\frac{z}{2\pi}}} \left( \frac{\sum_{\{\lambda[h]\}} \lambda[h] \log \left( \frac{\lambda[h]}{(K-1)P[h]} \right) - \sum_{\{\lambda[h]\}} \lambda[h] \log P[h]}{\rho_{K-1}} \right) + \text{const} \}}{\sum_{\{\lambda[h]\}} \frac{(K-1)!}{\prod_h \lambda[h]} \left\{ \prod_h P^{\lambda[h]}[h] \right\} e^{z_j \sqrt{\frac{z}{2\pi}}} \sum_h \lambda[h]}
\]
For \( K \) large, with \( J = 1/(K-1) \), the above average is dominated by the summands \( \mathcal{N}[h]/(K-1) = P[h] \) and is equal to \( \exp\left[ \frac{\beta_{K}}{2\rho_{K}} \left( \frac{h}{K-1} + \sum_h P[h]\right)^2 \right] \). This allows us to compute the distribution (70) for large \( K - 1 \), which satisfies the equation.
In this paper we study lymphocyte network models of the adaptive immune system. We derive dynamic equation for the B cell clones (B-clones), which depend on the T cell clones (T-clones) and antigen, and assume (following [9, 10]) that the Hamiltonian of this dynamic equation also governs the dynamics of the T-clones. Furthermore, we propose that the dynamics of B- and T-clones is subject to different thermal noise environments, and that they may evolve on different characteristic timescales. We compute the stationary distribution of the process in the limit of infinite (adiabatic) separation of timescales, which corresponds to an equilibrium scenario in which the fast (or slow) B-clone variables are interacting with the slow (or fast) T-clone variables. From the stationary distribution we obtain the average density of B-clone sizes, and the thermal averages of other macroscopic observables.

To simplify our analysis we consider the scenario where T-clones can be modelled by either Ising spin variables $\{-1,1\}$ and binary variables $\{0,1\}$. The former definition describes activated helper T-cells, modelled by $+1$, and activated regulator T-cells, modelled by $-1$. In the latter case we only have helper T-clones, which are either inactive, modelled by 0, or active, modelled by $+1$. We show that in the fast B-clone equilibration regime and in the fast T-clone equilibration regime, with the ratio of B clone noise to T-clone noise parameter $\beta$, the behaviour of T-clones is governed by an equilibrium distribution of $n \times N$ interacting Ising spins, in an effectively ferromagnetic model with inverse temperature $\beta$ and interactions $J_{fi}/\mu$.

As a consequence, there are network topologies for which there exists the critical noise $\beta_c$ such that for $\beta > \beta_c$ the fraction of helper (regulator) T cells is a monotonic function of the Ag concentration. Furthermore, we show that the average B-clone size (or concentration), $\langle c \rangle$, is a monotonic increasing function of the fraction of helper T-clones $m_+$ (i.e. a decreasing function of the fraction $m_-$ of regulators). This result is consistent with experimental data [27, 33]. Unfortunately, at present we are unable to carry out such a topology-independent analysis for the fast T-clone equilibration regime with non-integer ratios $n \in \mathbb{R}^+$. Obtaining distributions $P(c)$ of B-clone sizes requires a more detailed knowledge of the topology of the lymphocyte networks. Assuming this topology to be locally tree-like, we use the Bethe–Peierls (BP) approximation to derive equations for $P(c)$ and other observables in models on random networks. We solve these equations for the case when the model is homogeneous ($J_\mu = J$ for all $\mu$) and defined on a random regular graph, where each B-clone is connected to exactly $K$ T-clones and each T-clone is connected to exactly $L$ B-clones; here the ratio of B- to T-clones $\alpha = M/N$ equals $L/K$. We study this model in the fast B-clone equilibration regime and the fast T-clone equilibration regime, with $n \in \mathbb{Z}^+$ and $\beta \to \infty$. We find that for fast B-clone equilibration the distribution $P(c)$ in an ‘overregulated’ ($m_- = 0.9$) immune response regime is different from the distribution in a ‘normal’ ($m_- = 0.1$) regime: it behaves

$$P[h] = \frac{e^{\mu(L-1)\left(\frac{n}{(L-1)} + 2\frac{\beta}{(L-1)} \sum_h P(h)\right)}}{\sum_h e^{\mu(L-1)\left(\frac{n}{(L-1)} + 2\frac{\beta}{(L-1)} \sum_h P(h)\right)}}.$$  

Clearly $\lim_{K \to \infty} P[h] = \left(\frac{\sum_h n \rho(h)}{\sum_h n}^m\right)$, so this limit is equivalent to the infinite temperature regime $\beta = 0$, and the T-clones are unable to control the B-clones.

4. Discussion

In this paper we study lymphocyte network models of the adaptive immune system. We derive dynamic equation for the B cell clones (B-clones), which depend on the T cell clones (T-clones) and antigen, and assume (following [9, 10]) that the Hamiltonian of this dynamic equation also governs the dynamics of the T-clones. Furthermore, we propose that the dynamics of B- and T-clones is subject to different thermal noise environments, and that they may evolve on different characteristic timescales. We compute the stationary distribution of the process in the limit of infinite (adiabatic) separation of timescales, which corresponds to an equilibrium scenario in which the fast (or slow) B-clone variables are interacting with the slow (or fast) T-clone variables. From the stationary distribution we obtain the average density of B-clone sizes, and the thermal averages of other macroscopic observables.

To simplify our analysis we consider the scenario where T-clones can be modelled by either Ising spin variables $\{-1,1\}$ and binary variables $\{0,1\}$. The former definition describes activated helper T-cells, modelled by $+1$, and activated regulator T-cells, modelled by $-1$. In the latter case we only have helper T-clones, which are either inactive, modelled by 0, or active, modelled by $+1$. We show that in the fast B-clone equilibration regime and in the fast T-clone equilibration regime, with the ratio of B clone noise to T-clone noise parameter $n = \beta/\beta \in \mathbb{Z}^+$, the behaviour of T-clones is governed by an equilibrium distribution of $n \times N$ interacting Ising spins, in a effectively ferromagnetic model with inverse temperature $\beta$ and interactions $J_{fi}/\mu$. As a consequence, there are network topologies for which there exists the critical noise $\beta_c$ such that for $\beta > \beta_c$, the fraction of helper (regulator) T cells is a monotonic function of the Ag concentration. Furthermore, we show that the average B-clone size (or concentration), $\langle c \rangle$, is a monotonic increasing function of the fraction of helper T-clones $m_+$ (i.e. a decreasing function of the fraction $m_-$ of regulators). This result is consistent with experimental data [27, 33]. Unfortunately, at present we are unable to carry out such a topology-independent analysis for the fast T-clone equilibration regime with non-integer ratios $n \in \mathbb{R}^+$. Obtaining distributions $P(c)$ of B-clone sizes requires a more detailed knowledge of the topology of the lymphocyte networks. Assuming this topology to be locally tree-like, we use the Bethe–Peierls (BP) approximation to derive equations for $P(c)$ and other observables in models on random networks. We solve these equations for the case when the model is homogeneous ($J_\mu = J$ for all $\mu$) and defined on a random regular graph, where each B-clone is connected to exactly $K$ T-clones and each T-clone is connected to exactly $L$ B-clones; here the ratio of B- to T-clones $\alpha = M/N$ equals $L/K$. We study this model in the fast B-clone equilibration regime and the fast T-clone equilibration regime, with $n \in \mathbb{Z}^+$ and $\beta \to \infty$. We find that for fast B-clone equilibration the distribution $P(c)$ in an ‘overregulated’ ($m_- = 0.9$) immune response regime is different from the distribution in a ‘normal’ ($m_- = 0.1$) regime: it behaves
as a ‘power law’ when $m_\infty = 0.9$. The overregulated regime corresponds to the ‘branch’ of the phase diagram (see figure 4) where the ‘signal’ from the T cells to the B cell clones is predominantly suppressive (see figure 8). We envisage that in the real immune system such a situation can occur when the B cell clones are self-reactive, which is possible in the immune response to tumours [29].

We also study the regimes when $\alpha \to \infty$ (‘large’ B-clone number limit) and $\alpha \to 0$ (‘large’ T-clone number limit), with rescaled interactions $\tilde{J}^2 = \frac{1}{L-1}$ and $\tilde{J} = \frac{1}{K-1}$ respectively (for technical reasons). We find that in both of these regimes the B-clones are operating independently of the T-clones. In the regime $\alpha \to \infty$ we find that the response of T-clones to Ag is following the same phase transition pattern as in the figure 4. Also, in this regime the ‘cavity’ distribution equation (B.7) simplifies significantly, and much can be learned about this equation analytically (although for finite $K$, $L$ and $n > 1$ we were unable to find an explicit solution, even in the ‘simple’ high temperature phase). The regime $\alpha \to 0$ is equivalent to the infinite temperature regime, where T cells are insensitive to the Ag stimulation.

In future studies we plan to compute phase diagrams for the model in the fast T-clone equilibration regime with non-integer $n \in \mathbb{R}^+$. A good starting point here would be to consider the regular case (see appendix E). Also, it would be interesting to consider the case where the fraction of T-regulator clones is fixed and all T-clones could be active or inactive, i.e. the case of $\sigma_i \in [0, 1]$ and $\xi_j \in [-1, 1]$. Here one could assume that that the networks and interactions are random, and compute the phase diagrams by solving equation (50) by population dynamics. In order to model the process of affinity maturation, a further important ingredient of the adaptive immune system which has not been included yet, one can also assume that, in addition to B clones and T clones, also the interactions $J_i$ evolve in time, and use the slow (or fast) variable assumptions to compute stationary distributions of the more complicated process. Furthermore, the assumption of separated time-scales can be relaxed, but to make progress in this purely dynamical scenario would require application of sophisticated analytical tools such as the (dynamical) path integral method [46] or the dynamical replica theory [47]. However, the most important next step would be to connect the theoretical framework developed in this article with concrete experimental data.

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Appendix A. Analysis of the function $(\mathcal{e}^{F/n})_\beta$

In this section we study the behaviour as a function of $\beta$ of the following average:

$$
(\mathcal{e}^{F/n})_\beta = \sum_{\{\sigma^\prime, \ldots, \sigma^n\} } \frac{1}{M} \sum_{\nu} \frac{1}{\nu} \mathcal{O}^\nu \mathcal{E}(\sigma^n)
$$

$$
= \frac{1}{M} \sum_{\nu} \frac{1}{\nu} \mathcal{O}^\nu \mathcal{E}(\sigma^n)
$$

$$
\mathcal{E}(\sigma^n) = \frac{\sum_{\nu} \sum_{\nu} \mathcal{O}^\nu \mathcal{E}(\sigma^n)}{\sum_{\nu} \mathcal{O}^\nu \mathcal{E}(\sigma^n)}.
$$

(A.1)
In particular we are interested in the derivative
\[ \frac{\partial}{\partial \beta} \langle e^{F_m} \rangle_\beta = \frac{1}{M} \sum_{\nu, \mu} \left\{ \left( \sum_{n=1}^{\infty} F_n(\sigma^\nu) \right)^2 e^{\frac{1}{\beta} \sum_{n=1}^{\infty} F_n(\sigma^\nu)} \right\} - \left\{ \left( \sum_{n=1}^{\infty} F_n(\sigma^\nu) \right)^2 \right\} \left( e^{\frac{1}{\beta} \sum_{n=1}^{\infty} F_n(\sigma^\nu)} \right). \]  

Using Taylor’s expansion of the exponential,
\[ e^{\frac{1}{\beta} \sum_{n=1}^{\infty} F_n(\sigma^\nu)} = \sum_{\ell \geq 0} \frac{1}{(\ell \rho)^\ell \ell!} \left( \sum_{n=1}^{\infty} F_n(\sigma^\nu) \right)^\ell, \]
this derivative can be written as the infinite sum
\[ 2n \rho \frac{\partial}{\partial \beta} \langle e^{F_m} \rangle_\beta = \sum_{\ell \geq 0} \frac{1}{(\ell \rho)^\ell \ell!} \frac{1}{M} \sum_{\nu, \mu} \left\{ \left( \sum_{n=1}^{\infty} F_n(\sigma^\nu) \right)^2 \left( \sum_{n=1}^{\infty} F_n(\sigma^\nu) \right) \right\} \]
\[ \times \left\{ \left( \sum_{n=1}^{\infty} F_n(\sigma^\nu) \right)^2 \right\} \left( e^{\frac{1}{\beta} \sum_{n=1}^{\infty} F_n(\sigma^\nu)} \right). \]  

Finally, we rewrite the products \( \left( \sum_{\alpha=1}^{\infty} F_n(\sigma^\nu) \right)^\ell = \sum_{\nu, \mu} \prod_{j=1}^{\ell} F_n(\sigma^\nu) \) as sums, using the identity
\[ \prod_{j=1}^{\ell} (x_j + \theta) = \sum_{S_\ell \subseteq [\ell]} \theta^{\ell - |S_\ell|} \prod_{j \in S_\ell} x_j, \]
where \( S_\ell \) is a subset of the set \( [\ell] = \{1, \ldots, \ell\} \). This leads us to the equation
\[ 2n \rho \frac{\partial}{\partial \beta} \langle e^{F_m} \rangle_\beta = \sum_{\ell \geq 0} \frac{1}{(\ell \rho)^\ell \ell!} \sum_{S_\ell \subseteq [\ell]} \sum_{\nu, \mu} \left( \sum_{n=1}^{\infty} F_n(\sigma^\nu) \right)^2 \left( \sum_{n=1}^{\infty} F_n(\sigma^\nu) \right) \]
\[ \times \left( \sum_{n=1}^{\infty} F_n(\sigma^\nu) \right)^2 \left( e^{\frac{1}{\beta} \sum_{n=1}^{\infty} F_n(\sigma^\nu)} \right) \] 
\[ \times \left( \sum_{n=1}^{\infty} F_n(\sigma^\nu) \right)^2 \left( e^{\frac{1}{\beta} \sum_{n=1}^{\infty} F_n(\sigma^\nu)} \right). \]  

Now, by the GKS theorem [35], the correlation terms in the above sum are positive, from which it follows that \( \frac{\partial}{\partial \beta} \langle e^{F_m} \rangle_\beta \geq 0 \) and hence the average \( \langle e^{F_m} \rangle_\beta \) is a monotonic non-decreasing function of \( \beta \). Furthermore, for \( J_\nu = J \sum_{\alpha=1}^{\infty} \sigma_\nu^\alpha \), the average \( \langle e^{F_m} \rangle_\beta \) is also a monotonic non-decreasing function of \( J \). To show this we consider the derivative
\[ J \frac{\partial}{\partial J} \langle e^{F_m} \rangle_\beta = 2 \beta \frac{\partial}{\partial \beta} \langle e^{F_m} \rangle_\beta + \frac{1}{M} \sum_{\mu=1}^{M} \left( e^{\frac{1}{\beta} \sum_{n=1}^{\infty} F_n(\sigma^\nu)} \right) \frac{1}{n \rho} \sum_{\alpha=1}^{n} F_n(\sigma^\nu) \].
Appendix B.

Computation on a factor tree for $n \in \mathbb{Z}^+$

Let us assume that the system (28) is defined on a factor tree $\mathcal{T}_n(r)$ of radius $r$ rooted at factor-node $\mu$ (see figure B1). Then the distribution of (replicated) fields on $\mu$

$$P_n(F) = \sum_{\sigma^1, \sigma^n} P(\sigma^1, \ldots, \sigma^n) \delta \left( F - \sum_{\alpha=1}^{n} F_\sigma(\sigma^\alpha) \right)$$

(B.1)

can be computed recursively as follows. Firstly, we compute the ‘partition’ function

$$Z_n(F) = \sum_{\{\sigma, \sigma^1, \ldots, \sigma^n\} \in \mathcal{T}_n} e^{\sum_{\alpha=1}^{n} F_\sigma(\sigma^\alpha)} \delta \left( F - \sum_{\alpha=1}^{n} F_\sigma(\sigma^\alpha) \right)$$

$$= \sum_{\{\sigma, \sigma^1, \ldots, \sigma^n\} \in \mathcal{T}_n} e^{\sum_{\alpha=1}^{n} F_\sigma(\sigma^\alpha)} \delta \left( F - \sum_{\alpha=1}^{n} F_\sigma(\sigma^\alpha) \right)$$

(B.2)

In the above we defined the factor tree $\mathcal{T}_n$ (of radius $r-1$), rooted at variable node $i$. The partition function $Z_n(F)$ can be used to construct the distribution of fields.
Figure B1. The interaction topology of a replicated spin system on a factor-tree $T_{\mu}$, rooted at factor-node $\mu$. All spins are represented by circular ‘variable’ nodes, and each term in the ‘Boltzmann factor’ $\exp\left[\frac{1}{k_B T} \sum_{\nu \in \partial T} (\sum_{\alpha = 1}^{n} F_{\nu}(\sigma^{\alpha}))^{2}\right] = \prod_{\nu \in \partial T} E_{\nu}((\sigma : j \in \partial \nu))$ corresponds to a square ‘factor’ node. A link between variable node $\ell$ and factor node $\nu$ implies that $\sigma_{\nu}$ acts as an argument of $E_{\nu}$.

$$P_{\mu}(F) = \frac{Z_{\mu}(F)}{\int Z_{\mu}(F) \, dF} = \frac{\exp^{\sum_{\nu \in \partial \mu} \left\{ \prod_{i \in \mu} Z_{\mu}[\pi_{i}] \right\} \delta(F - \sum_{\alpha = 1}^{n} F_{\nu}(\sigma^{\alpha}))} {\sum_{\nu \in \partial \mu} \left\{ \prod_{i \in \mu} Z_{\mu}[\pi_{i}] \right\} \exp^{\sum_{\nu \in \partial \mu} (F_{\nu}(\sigma^{\alpha}))^{2}}}.$$ \hspace{1cm} (B.3)

In the same way we can define the ‘cavity’ distribution, $P_{\mu}[\sigma]$, corresponding to the topology that would be found if the edge $(i, \mu)$ were removed (see figure B1),

$$P_{\mu}[\sigma] = \frac{Z_{\mu}[\sigma]}{\sum_{\sigma} Z_{\mu}[\sigma]},$$ \hspace{1cm} (B.4)

and thereby obtain the equation

$$P_{\mu}(F) = \left\{ \delta(F - \sum_{\alpha = 1}^{n} F_{\nu}(\sigma^{\alpha})) \right\}^{n} = \frac{\sum_{\sigma} \prod_{i \in \mu} P_{\mu}[\sigma_{i}] \exp^{\sum_{\nu \in \partial \mu} \left\{ \prod_{i \in \mu} Z_{\mu}[\pi_{i}] \right\} \delta(F - \sum_{\alpha = 1}^{n} F_{\nu}(\sigma^{\alpha}))} {\sum_{\sigma} \prod_{i \in \mu} P_{\mu}[\sigma_{i}] \exp^{\sum_{\nu \in \partial \mu} \left\{ \prod_{i \in \mu} Z_{\mu}[\pi_{i}] \right\} \delta(F - \sum_{\alpha = 1}^{n} F_{\nu}(\sigma^{\alpha}))^{2}}}.$$ \hspace{1cm} (B.5)

We note that in order to distinguish ‘cavity’ distributions, such as $P_{\mu}[\sigma]$, from the corresponding distributions on graphs with all links intact, such as $P(\sigma)$, we will use square brackets $[\ldots]$ in referring to the former, throughout this paper.

We next we compute the cavity partition function
\[ Z_{\mu}[\sigma] = \sum_{\{\sigma_j\} \in \mathcal{T}_I} e^{\beta J_{\mu \nu} \sum_{j=1}^{n_{\nu}} E(\sigma_{\nu j})} \delta_{\sigma_{\mu j}, \sigma_{\nu j}} \]
\[ = \sum_{\{\sigma_j\} \in \mathcal{T}_I} e^{\beta J_{\mu \nu} \sum_{j=\partial \mu, j \in \partial \nu} E(\sigma_{\nu j})} \times \prod_{j=\partial \mu} \prod_{j \in \partial \nu} e^{\beta e_{T}(\sum_{j=1}^{n_{\nu}} E(\sigma_{\nu j}))} \delta_{\sigma_{\mu j}, \sigma_{\nu j}} \]
\[ = \sum_{\{\sigma_j\}} e^{\beta \sum_{j=1}^{n_{\nu}} E(\sigma_{\nu j}) j} \left\{ \prod_{j=\partial \mu} \prod_{j \in \partial \nu} Z_{\mu j}[\sigma_j] \right\} \delta_{\sigma_{\mu j}, \sigma_{\nu j}} \]  \hspace{1cm} (B.6)

Using this expression in definition (B.4) gives us equation

\[ P_{\mu j}[\sigma_j] = \sum_{\{\sigma_j\}} \left\{ \prod_{j=\partial \mu} \prod_{j \in \partial \nu} Z_{\mu j}[\sigma_j] \right\} \frac{e^{\beta \sum_{j=1}^{n_{\nu}} E(\sigma_{\nu j})}}{Z_{\mu}[\sigma]} \times \prod_{j=\partial \mu} \prod_{j \in \partial \nu} P_{\mu j}[\sigma_j] \left\{ \frac{e^{\beta \sum_{j=1}^{n_{\nu}} E(\sigma_{\nu j})}}{Z_{\mu}[\sigma]} \right\}^{-1} \]  \hspace{1cm} (B.7)

Furthermore, we note that the partition function

\[ Z_\nu(\sigma) = \sum_{\{\sigma_j\} \in \mathcal{T}_I} e^{\beta \sum_{j=1}^{n_{\nu}} E(\sigma_{\nu j})} \delta_{\sigma_{\mu j}, \sigma_{\nu j}} \]
\[ = \sum_{\{\sigma_j\}} \left\{ \prod_{j=\partial \mu} \prod_{j \in \partial \nu} Z_{\mu j}[\sigma_j] \right\} \frac{e^{\beta \sum_{j=1}^{n_{\nu}} E(\sigma_{\nu j})}}{Z_{\mu}[\sigma]} \times \prod_{j=\partial \mu} \prod_{j \in \partial \nu} e^{\beta \sum_{j=1}^{n_{\nu}} E(\sigma_{\nu j})} \delta_{\sigma_{\mu j}, \sigma_{\nu j}} \]
\[ = \sum_{\{\sigma_j\}} e^{\beta \sum_{j=1}^{n_{\nu}} E(\sigma_{\nu j})} \left\{ \prod_{j=\partial \mu} \prod_{j \in \partial \nu} Z_{\mu j}[\sigma_j] \right\} \frac{e^{\beta \sum_{j=1}^{n_{\nu}} E(\sigma_{\nu j})}}{Z_{\mu}[\sigma]} \times \prod_{j=\partial \mu} \prod_{j \in \partial \nu} e^{\beta \sum_{j=1}^{n_{\nu}} E(\sigma_{\nu j})} \delta_{\sigma_{\mu j}, \sigma_{\nu j}} \]  \hspace{1cm} (B.8)

which is defined on a factor-tree \( T_\mu(R) \) rooted at the variable node \( \nu \), can be used to construct the distribution

\[ P(\sigma) = \frac{Z_\nu(\sigma)}{\sum_\sigma \sigma Z_\nu(\sigma)} \]  \hspace{1cm} (B.9)

which gives the (local) magnetization equation

\[ \langle \sigma \rangle = \sum_\sigma \frac{P(\sigma)}{\sum_\sigma \sigma Z_\nu(\sigma)} \sum_{a=1}^{n} \sigma^a \]
\[ = \sum_{\{\sigma_j\}} \left\{ \prod_{j=\partial \mu} \prod_{j \in \partial \nu} P_{\mu j}[\sigma_j] \right\} \sum_{a=1}^{n} \sigma^a \]
\[ \times e^{\beta \sum_{j=\partial \mu} \sum_{j \in \partial \nu} J_{\mu \nu} \sum_{j=1}^{n_{\nu}} E(\sigma_{\nu j})} \delta_{\sigma_{\mu j}, \sigma_{\nu j}} \]
\[ \times \left\{ \prod_{j=\partial \mu} \prod_{j \in \partial \nu} P_{\mu j}[\sigma_j] \right\} \sum_\sigma e^{\beta \sum_{j=\partial \mu} \sum_{j \in \partial \nu} J_{\mu \nu} \sum_{j=1}^{n_{\nu}} E(\sigma_{\nu j})} \delta_{\sigma_{\mu j}, \sigma_{\nu j}} \]  \hspace{1cm} (B.10)
The structure of the equations derived in this section is not affected by the choice for the \( \sigma^0 \) variables, but in the Ising case \( \sigma^0 \in \{-1, 1\} \) with \( \theta^0 = 0 \) the recursion (B.7) preserves the spin-reversal symmetry, i.e. the equality \( P_\beta[\sigma] = P_\beta[-\sigma] \) implies that \( P_\beta[\sigma] = P_\beta[-\sigma] \). For \( n = 1 \) this implies that \( P_\beta[\sigma] = \frac{1}{2} \) is a solution of this map for any \( \beta \), but for \( n > 1 \) the uniform distribution \( P_\beta[\sigma] = \frac{1}{2} \) is a solution of (B.7) only when \( \beta = 0 \). The consequence of the symmetry \( P_\beta[\sigma] = P_\beta[-\sigma] \) is that the local magnetization becomes \( \langle \sigma \rangle = 0 \), and that the distribution (B.5) is symmetric, i.e. \( P_\beta(F) = P_\beta(-F) \).

Finally, we note that the equations (B.5), (B.7) and (B.10) can be simplified if we define the replica ‘magnetization’ distribution

\[
P[h] = \sum_\sigma P[\sigma] \delta_h \sum_{i=1}^n \sigma_i.
\]

Using above definition in these equations gives the equations (46)–(48).

**Appendix C. Computation on a factor tree for** \( n \in \mathbb{R}^+ \)

Let us assume that the distribution (24), as given by

\[
P(b) = \frac{1}{Z} e^{-\frac{1}{\beta} \sum_{\mu} \sum_{ij} b_{ij} + n \sum_{i=1}^n \log 2 \cosh(\beta \sum_{\mu \in \mathcal{I}} b_{\mu i})}
\]

when adopted to the Ising case \( \sigma_i \in \{-1, 1\} \), is defined on a factor-tree \( \mathcal{T}_\mu(r) \) of radius \( r \) rooted at variable-node \( \mu \) (see figure C1). We note that all equations in this section are derived for
σᵣ ∈ {−1, 1}, but they can be used trivially also for σᵣ ∈ {0, 1}, by making the transformations 2 cosh(ξ) → 1 + eξ and 2 sinh(ξ) → eξ. To compute the marginal distribution

\[ P_n(b) = \int db \, P(b) \, \delta(b - b_\mu) \]  

(C.2)

we consider the partition function

\[ Z_n(b) = \left\{ \prod_{\nu \in \mathcal{T}_\mu} \int db_\nu \right\} \delta(b - b_\nu) e^{-\frac{1}{2} \mu \beta \sum_{\nu \in \mathcal{T}_\mu} \left( b_\nu - \frac{J_\nu}{\beta} \right)^2} \left\{ \prod_{\nu \in \partial \mu} 2 \cosh \left( \beta \sum_{\nu \in \partial \mu} J_\nu \xi_\nu^\mu b_\nu \right) \right\}^n \]

\[ = e^{-\frac{1}{2} \beta \mu \sum_{\nu \in \mathcal{T}_\mu} \left( b_\nu - \frac{J_\nu}{\beta} \right)^2} \left\{ \prod_{\nu \in \partial \mu} \int db_\nu \right\} \delta(b - b_\nu) \left\{ \prod_{\nu \in \partial \mu} 2 \cosh \left( \beta \sum_{\nu \in \partial \mu} J_\nu \xi_\nu^\mu b_\nu \right) \right\}^n \]

\[ \times \left\{ \prod_{i \in \partial \mu} \prod_{\nu \in \partial \mu} \left[ e^{-\frac{1}{2} \mu \beta \sum_{\nu \in \mathcal{T}_\mu} \left( b_\nu - \frac{J_\nu}{\beta} \right)} \left\{ \prod_{\nu \in \partial \mu} 2 \cosh \left( \beta \sum_{\nu \in \partial \mu} J_\nu \xi_\nu^\mu b_\nu \right) \right\}^n \right] \right\} \]

\[ = e^{-\frac{1}{2} \beta \mu \sum_{\nu \in \mathcal{T}_\mu} \left( b_\nu - \frac{J_\nu}{\beta} \right)^2} \left\{ \prod_{i \in \partial \mu} \prod_{\nu \in \partial \mu} \int db_\nu \, Z_n[b_\nu] \right\} \]

\[ \times \int db_\nu \left\{ \prod_{\nu \in \partial \mu} 2 \cosh \left( \beta \sum_{\nu \in \partial \mu} J_\nu \xi_\nu^\mu b_\nu \right) \right\}^n \delta(b - b_\nu) \]  

(C.3)

which subsequently gives us the marginal distribution via the equation

\[ P_n(b) = \frac{Z_n(b)}{\int_{-\infty}^{\infty} db \, Z_n(b)} \]

\[ = e^{-\frac{1}{2} \beta \mu \sum_{\nu \in \mathcal{T}_\mu} \left( b_\nu - \frac{J_\nu}{\beta} \right)^2} \left\{ \prod_{i \in \partial \mu} \prod_{\nu \in \partial \mu} \int db_\nu \, Z_n[b_\nu] \right\}  

\times \int db_\nu \left\{ \prod_{\nu \in \partial \mu} 2 \cosh \left( \beta \sum_{\nu \in \partial \mu} J_\nu \xi_\nu^\mu b_\nu \right) \right\}^n \delta(b - b_\nu) \]

\[ \times \left\{ \prod_{i \in \partial \mu} \prod_{\nu \in \partial \mu} \int db_\nu \, Z_n[b_\nu] \right\} \int db_\nu \, e^{-\frac{1}{2} \beta \mu \sum_{\nu \in \mathcal{T}_\mu} \left( b_\nu - \frac{J_\nu}{\beta} \right)^2} \left\{ \prod_{\nu \in \partial \mu} 2 \cosh \left( \beta \sum_{\nu \in \partial \mu} J_\nu \xi_\nu^\mu b_\nu \right) \right\}^n \]  

(C.4)

This immediately gives us the equation
\[ P_n(b) = \left\{ \prod_{i, \beta} \prod_{\nu} \int_{-\infty}^{\infty} P_{\nu}[b_{\nu}] \, db_{\nu} \right\} e^{-\frac{1}{2} m(b - \beta \frac{\nu}{\mu})^2} \]
\[ \times \left\{ \prod_{i, \beta} 2 \cosh \left( \beta \sum_{\nu} J_i \xi^\nu b_{\nu} + \beta J_i \xi^\nu \right) \right\}^\nu \]
\[ \times \left\{ \prod_{i, \beta} \int_{-\infty}^{\infty} P_{\nu}[b_{\nu}] \, db_{\nu} \int_{-\infty}^{\infty} \, db \, e^{-\frac{1}{2} m(b - \beta \frac{\nu}{\mu})^2} \right\}^{\nu n} \]
\[ \times \left\{ \prod_{i, \beta} 2 \cosh \left( \beta \sum_{\nu} J_i \xi^\nu b_{\nu} + \beta J_i \xi^\nu \right) \right\}^{n - 1}. \]

Equation (C.5)

If we insert the definition
\[ P_n[b] = \frac{Z_{\nu}[b]}{\int_{-\infty}^{\infty} db Z_{\nu}[b]} \]  

Equation (C.6)

In order to derive an equation for the distribution \( P_n[b] \) we compute the partition function, associated with the factor-tree \( T_r \) of radius \( r - 1 \) (see figure C1), as follows

\[ Z_{\nu}[b] = \left\{ \prod_{\beta \in T_r} \int_{-\infty}^{\infty} db_{\beta} \right\} \delta(b - b_{\beta}) \times e^{-\frac{1}{2} m(b - \beta \frac{\nu}{\mu})^2 + n \sum_{\nu} 2 \cosh \left( \beta \sum_{\nu} J_i \xi^\nu b_{\nu} \right)} \]
\[ \times \left\{ \prod_{j, \beta} \prod_{\nu} \int_{-\infty}^{\infty} P_{\nu}[b_{\nu}] \, db_{\nu} \right\} e^{-\frac{1}{2} m(b - \beta \frac{\nu}{\mu})^2} \times \left\{ \prod_{\nu} \log 2 \cosh \left( \beta \sum_{i, \beta} J_i \xi^\nu b_{\nu} \right) \right\}^{\nu} \]
\[ = \left\{ \prod_{j, \beta} \prod_{\nu} \int db_{\nu} Z_{\nu}[b_{\nu}] \right\} e^{-\frac{1}{2} m(b - \beta \frac{\nu}{\mu})^2} \]
\[ \times \left\{ \prod_{i, \beta} 2 \cosh \left( \beta \sum_{\nu} J_i \xi^\nu b_{\nu} + \beta J_i \xi^\nu \right) \right\}^{n} \]. \]  

Equation (C.7)

Using this expression in definition (C.6) gives us the recursive equation

\[ P_n[b] = \left\{ \prod_{i, \beta} \prod_{\nu} \int_{-\infty}^{\infty} P_{\nu}[b_{\nu}] \, db_{\nu} \right\} e^{-\frac{1}{2} m(b - \beta \frac{\nu}{\mu})^2} \]
\[ \times \left\{ \prod_{i, \beta} 2 \cosh \left( \beta \sum_{\nu} J_i \xi^\nu b_{\nu} + \beta J_i \xi^\nu \right) \right\}^\nu \]
\[ \times \left\{ \prod_{i, \beta} \int_{-\infty}^{\infty} P_{\nu}[b_{\nu}] \, db_{\nu} \int_{-\infty}^{\infty} \, db \, e^{-\frac{1}{2} m(b - \beta \frac{\nu}{\mu})^2} \right\}^{\nu n} \]
\[ \times \left\{ \prod_{i, \beta} 2 \cosh \left( \beta \sum_{\nu} J_i \xi^\nu b_{\nu} + \beta J_i \xi^\nu \right) \right\}^{n - 1}. \]  

Equation (C.8)
For $n \in \mathbb{Z}^+$, we can use the above equation, together with the identity $2^n \cosh^n(x) = \sum_{\sigma_1, \ldots, \sigma_n} e^{\sum_{i=1}^n \sigma_i}$, and the definition

$$P_n[\sigma] = \frac{\left\{ \prod_{i \in \partial_i} \mu_i \int_{-\infty}^{\infty} P_n[b_i] \, db_i \right\} e^{\beta \sum_{i \in \partial_i} J_i \xi_i b_i \sum_{i=1}^n \sigma_i}}{\left\{ \prod_{i \in \partial_i} \mu_i \int_{-\infty}^{\infty} P_n[b_i] \, db_i \right\} \left[ 2 \cosh(\beta \sum_{i \in \partial_i} J_i \xi_i b_i) \right]^n}$$

(C.9)

to recover equation (B.7) and all other equations of $n \in \mathbb{Z}^+$ which involve the cavity distribution $P_n[\sigma]$. Furthermore, the above approach can be used to compute any marginal of (C.1). In particular, the joint distribution of the variables in the set \{b_\mu : \mu \in \partial i\} is derived by considering the distribution (C.1) defined on a factor-tree $T_i$ rooted at factor-node $i$:

$$P_i(\{b_\mu : \mu \in \partial i\}) = \frac{\left\{ \prod_{\mu \in \partial i} P_n[b_\mu] \right\} \cosh^\beta(\sum_{\mu \in \partial i} J_\mu \xi_\mu b_\mu)}{\left\{ \prod_{\mu \in \partial i} \int_{-\infty}^{\infty} P_n[b_\mu] \, db_\mu \right\} \cosh^\beta(\sum_{\mu \in \partial i} J_\mu \xi_\mu b_\mu)}.$$

(C.10)

This is then used to compute the local magnetization $\langle \sigma_i \rangle = \sum_{\sigma} P(\sigma) \sigma$ from the distribution (26) which gives us the equation

$$\langle \sigma_i \rangle = \left\{ \prod_{\mu \in \partial i} \int_{-\infty}^{\infty} P_n[b_\mu] \, db_\mu \right\} \cosh^\beta(\sum_{\mu \in \partial i} J_\mu \xi_\mu b_\mu) \times \frac{\left\{ \prod_{\mu \in \partial i} \int_{-\infty}^{\infty} P_n[b_\mu] \, db_\mu \right\} \cosh^\beta(\sum_{\mu \in \partial i} J_\mu \xi_\mu b_\mu)}{\left\{ \prod_{\mu \in \partial i} \int_{-\infty}^{\infty} P_n[b_\mu] \, db_\mu \right\} \cosh^\beta(\sum_{\mu \in \partial i} J_\mu \xi_\mu b_\mu)}.$$  

(C.11)

Finally, to compute the distribution of fields (27) we need to know the joint distribution $P_p(b, \{b_\mu\})$ of the variable $b = b_\mu$ and of all its immediate neighbours $b_\nu$ (see figure C1). This distribution can be ‘read off’ from equation (C.5), using the identity $P_p(b) = \left\{ \int_{-\infty}^{\infty} \right\} P_p(b, \{b_\mu\})$, which gives

$$P_p(F) = \prod_{i \in \partial_p} \prod_{\mu \in \partial_i} \int_{-\infty}^{\infty} \int_{-\infty}^{\infty} \frac{1}{\mathcal{Z}_p} \exp \left\{ J_i b_i + \sum_{\mu \in \partial i} \xi_\mu b_\mu - \frac{1}{2} m i \left( b_i - \frac{b_\mu}{2} \right)^2 \right\} \prod_{\mu \in \partial i} \cosh^\beta(\sum_{\nu \in \partial i} J_\nu \xi_\nu b_\nu) \times \prod_{\mu \in \partial i} \cosh^\beta(\sum_{\nu \in \partial i} J_\nu \xi_\nu b_\nu) \times \left\{ \prod_{\mu \in \partial i} \int_{-\infty}^{\infty} P_n[b_\mu] \, db_\mu \right\} \left\{ \int_{-\infty}^{\infty} \int_{-\infty}^{\infty} \frac{1}{\mathcal{Z}_p} \exp \left\{ J_i b_i + \sum_{\mu \in \partial i} \xi_\mu b_\mu - \frac{1}{2} m i \left( b_i - \frac{b_\mu}{2} \right)^2 \right\} \prod_{\mu \in \partial i} \cosh^\beta(\sum_{\nu \in \partial i} J_\nu \xi_\nu b_\nu) \times \prod_{\mu \in \partial i} \cosh^\beta(\sum_{\nu \in \partial i} J_\nu \xi_\nu b_\nu) \right\}^{-1}.$$ 

(C.12)

Equation (C.8) preserves the symmetry $P_p[b] = P_p[-b]$ when $b_\mu = 0$. This implies that for such $P_p[b]$ the marginal distribution (C.5) and the distribution of fields (C.12) are both symmetric functions, and that the local magnetization (C.11) is zero.
Appendix D. Analysis of \( n = 1 \) case

For \( n = 1 \), equation (51) can be simplified further by noticing that here \( h \in \{-1, 1\} \) and the distribution \( P[h] \), with \( h = \sigma \), can be written in the form

\[
P[\sigma] = \frac{e^{-\phi}}{2 \cosh(\phi)}.
\]

where \( \phi \) is a ‘cavity field’ parameter. Using this in (51) gives us the equation

\[
P[\sigma] = \frac{\left[ \sum_{(\sigma)} e^{i J \sum_{j=1}^{K} \sigma_j \delta} (\phi \sum_{j=1}^{K-1} \sigma_j) \right]^{L-1}}{\sum_{\sigma} \left[ \sum_{(\sigma)} e^{i J \sum_{j=1}^{K} \sigma_j \delta} (\phi \sum_{j=1}^{K-1} \sigma_j) \right]^{L-1}}
\]

from which, via the identity \( \phi = \frac{1}{2} \log(P[+1]/P[-1]) \), we can derive

\[
\phi = \frac{L - 1}{2} \log \left( \frac{\sum_{\sigma} e^{i J \sum_{j=1}^{K} \sigma_j \delta} (\phi \sum_{j=1}^{K-1} \sigma_j)}{\sum_{\sigma} e^{i J \sum_{j=1}^{K} \sigma_j \delta} (\phi \sum_{j=1}^{K-1} \sigma_j)} \right).
\]

The solution \( \phi \) of the above equation can be used to compute the marginal distribution

\[
P(\sigma) = \frac{\left[ \sum_{(\sigma)} e^{i J \sum_{j=1}^{K} \sigma_j \delta} (\phi \sum_{j=1}^{K-1} \sigma_j) \right]^{L}}{\sum_{\sigma} \left[ \sum_{(\sigma)} e^{i J \sum_{j=1}^{K} \sigma_j \delta} (\phi \sum_{j=1}^{K-1} \sigma_j) \right]^{L}}.
\]

Comparing this with the cavity distribution (D.2) gives us the magnetization formula

\[
m = \sum_{\sigma} P(\sigma) \sigma = \tanh \left( \frac{L \phi}{L - 1} \right).
\]

Also, using definition (34) in the distribution of fields (52) gives us the equation

\[
P(F) = \frac{\sum_{(\sigma)} e^{i J \sum_{j=1}^{K} \sigma_j \delta} (F - J \sum_{j=1}^{K} \sigma_j)}{\sum_{(\sigma)} e^{i J \sum_{j=1}^{K} \sigma_j \delta} (\phi \sum_{j=1}^{K-1} \sigma_j)}.
\]

Upon defining the RHS of (D.3) as \( f(\phi) \), this equation takes the form \( \phi = f(\phi) \). The value \( \phi = 0 \), which corresponds to the paramagnetic (PM) \( m = 0 \) phase, is always a solution. However, it becomes unstable at the point where \( f'(0) = 1 \), in which

\[
f'(0) = (L - 1)(K - 1) \sum_{(\sigma)} e^{i J \sum_{j=1}^{K} \sigma_j \delta} \sinh \left( \frac{\beta J}{\rho} \sum_{j=1}^{K-1} \sigma_j \right) \sum_{(\sigma)} e^{i J \sum_{j=1}^{K} \sigma_j \delta} \cosh \left( \frac{\beta J}{\rho} \sum_{j=1}^{K-1} \sigma_j \right).
\]

For \( f'(0) > 1 \) equation (D.3) has two stable solutions \( \phi = 0 \), which correspond to the ferromagnetic (FM) \( m = 0 \) phase. Solving the equation \( f'(0) = 1 \) gives us the critical inverse temperature \( \beta_c \) (\( J = \rho = 1 \)) where the PM to FM transition occurs (see figure D1). We note
Appendix E. Homogeneous systems on random regular factor-graphs, with \( n \in \mathbb{R}^+ \)

The equations derived in appendix \( C \) can be easily adopted to study the system (C.1), which is homogeneous and defined on a random regular factor-graph (see section 3.2 for details). Here the marginal distribution \( P(b) = P_m(b) \) is given by

\[
P(b) = e^{-\frac{1}{2} \sum_{\nu=1}^{L-1} \int db_{\nu} P[b_{\nu}] \cosh \left( \beta \left( \sum_{\nu=1}^{L-1} b_{\nu} + b \right) \right)} \times \left\{ \int dB e^{-\frac{1}{2} \sum_{\nu=1}^{L-1} \int d\tilde{b}_{\nu} P[\tilde{b}_{\nu}] \cosh \left( \beta \left( \sum_{\nu=1}^{L-1} \tilde{b}_{\nu} + \tilde{b} \right) \right)} \right\}^{-1}
\]

and the cavity distribution \( P[b] = P_d[b] \) can be computed recursively via

\[
P[b] = \frac{Z[b]}{\int db Z[b]} \quad \text{(E.2)}
\]

\[
Z[b] = e^{-\frac{1}{2} \sum_{\nu=1}^{L-1} \int db_{\nu} P[b_{\nu}] \cosh \left( \beta \left( \sum_{\nu=1}^{L-1} b_{\nu} + b \right) \right)} \quad \text{. (E.3)}
\]

that for \( L > 2 \) and \( K = 2 \) the equation (D.7) recovers the result \( \beta_i = \tanh^{-1}(1/(L - 1)) \) of ferromagnetic Ising models on Bethe lattices [42].
We note that $P(b)$ and $P[\beta]$ are related by the transformation $K - 1 \rightarrow K$. Finally, once we know the distribution $P[\beta]$, then we can also compute the magnetization

$$\langle \sigma \rangle = \left\{ \prod_{i=1}^{L} \int db_{i} P(b_{i}) \right\} \cosh^{\alpha} \left( \beta J \sum_{i=1}^{L} b_{i} \right) \frac{\tanh \left( \beta J \sum_{i=1}^{L} b_{i} \right)}{\left\{ \prod_{i=1}^{L} \int db_{i} P(b_{i}) \right\} \cosh^{\alpha} \left( \beta J \sum_{i=1}^{L} b_{i} \right)} \tag{E.4}$$

and the distribution of fields

$$P(F) = \left\{ \prod_{i=1}^{K} \prod_{l=1}^{L-1} \int db_{i} P(b_{i}) \right\} \int db \ e^{-\frac{1}{2\mu\delta^{2}} \left( K \sum_{i=1}^{L-1} b_{i} + b \right)} \left( F - J \sum_{i=1}^{K} \left\{ \prod_{l=1}^{L-1} \prod_{j=1}^{K} \int db_{j} P(b_{j}) \right\} \right) \int db \ e^{-2\mu\delta^{2} \left( K \sum_{i=1}^{L-1} b_{i} + b \right)} \left( F - J \sum_{i=1}^{K} \left\{ \prod_{l=1}^{L-1} \prod_{j=1}^{K} \int db_{j} P(b_{j}) \right\} \right)^{K-1} \tag{E.5}$$

References

[27] Vanderleyden I, Linterman M A and Smith K G C 2014 Arthritis Res. Ther. 16 471