

Published in final edited form as:

Nat Cell Biol. 2013 April ; 15(4): 363–372. doi:10.1038/ncb2709.

Characterisation of transcriptional networks in blood stem and progenitor cells using high-throughput single cell gene expression analysis

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Abstract

Cellular decision-making is mediated by a complex interplay of external stimuli with the intracellular environment, in particular transcription factor regulatory networks. Here we have determined the expression of a network of 18 key haematopoietic transcription factors (TFs) in 597 single primary blood stem and progenitor cells isolated from mouse bone marrow. We demonstrate that different stem/progenitor populations are characterised by distinctive TF expression states, and through comprehensive bioinformatic analysis reveal positively and negatively correlated TF pairings, including previously unrecognised relationships between *Gata2*, *Gfi1* and *Gfi1b*. Validation using transcriptional and transgenic assays confirmed direct regulatory interactions consistent with a regulatory triad in immature blood stem cells, where *Gata2* may function to modulate cross-inhibition between *Gfi1* and *Gfi1b*. Single cell expression profiling therefore identifies network states and allows reconstruction of network hierarchies involved in controlling stem cell fate choices, and provides a blueprint for studying both normal development and human disease.

Haematopoiesis has long served as a model system for studying cell fate decisions during stem cell differentiation^{1, 2}. At the molecular level, transcription factors (TFs) are major

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Author Contributions: V.M. designed and performed single cell experiments, performed analysis, and wrote the paper. B.G. conceived the study, designed experiments and wrote the paper. I.C.M. and S.E.J. designed and performed FACS and wrote the paper. F.B. and F.T. analysed GPLVM data and wrote the paper. S.K. and J.S. performed transgenic and luciferase analysis. A.J. and R.H. performed bioinformatic analysis. F.J.C.-N. performed ChIP-Seq experiments. G.S. and M.d.B. designed experiments and performed preliminary studies.

Competing financial interests: The authors declare no competing financial interests.

Accession Numbers

ChIP-seq data for mast cells have been deposited into the NCBI Gene Expression Omnibus portal under the accession number GSE42518.

drivers of cellular identity and cell fate transitions as exemplified by their key role in reprogramming³ and lineage switching experiments⁴⁻⁶. TFs function within wider regulatory networks, the connectivity of which can be revealed using classical transcriptional assays or inferred from global expression and TF binding profiling studies⁷⁻¹².

However, these experiments report population averages, while cell fate choices are made by individual cells. The importance of studying single cells is emphasised by the known functional heterogeneity in haematopoietic stem cells (HSCs) as well as other cell types, which manifests as relatively stable subpopulations with either balanced production of myeloid and lymphoid cells or a deficiency in lymphoid potential¹³⁻¹⁵. HSCs have also been reported to be heterogeneous in gene expression¹⁶⁻¹⁸, although previous studies have been limited in terms of numbers of genes, cells or populations analysed. Recent advances in microfluidics technologies have facilitated high-throughput Q-RT-PCR analysis of tens of genes in hundreds of single cells simultaneously¹⁹. This technology has recently been used to resolve cell populations in 64 cell mouse embryos²⁰, to dissect cellular heterogeneity in human colon cancer²¹, and to reveal significant variation in early erythroid gene expression, which resolved upon commitment²².

Here we analysed expression of a network of 18 densely interconnected TFs in 597 single cells from five primary haematopoietic stem and progenitor cell populations, which not only revealed characteristic expression states for the different cell populations, but also identified previously unrecognized regulatory relationships. This included a putative regulatory triad consisting of *Gata2*, *Gfi1* and *Gfi1b*, which was validated using cell line and transgenic mouse assays. Our 4 findings suggest that GATA2 may function in a regulatory loop to modulate *Gfi1/Gfi1b* cross-antagonism during entry into the myeloid/lymphoid lineages, thus demonstrating that high-throughput single cell TF expression analysis provides a powerful approach towards the identification of regulatory network links.

RESULTS

Single-cell expression analysis reveals heterogeneity in transcription factor expression in haematopoietic stem and progenitor cells

To study core regulatory circuits during early haematopoietic differentiation stages, we performed gene expression analysis for transcription factors in single primary haematopoietic stem/progenitor cells prospectively isolated from mouse bone marrow by fluorescence activated cell sorting (FACS). We analysed long-term haematopoietic stem cells (LSK CD150⁺CD48⁻ HSC²³), lymphoid-primed multipotent progenitors (LSK Flt3^{hi} LMPP²⁴), bipotential megakaryocyte/erythroid progenitors (CD16/32^{lo}CD41⁻CD150⁺CD105^{lo} PreMegE²⁵), granulocyte-monocyte progenitors (CD41^{lo}CD16/32^{hi} GMP^{25, 26}), and common lymphoid progenitor (Lin⁻ IL7R⁺ Kit^{lo} Sca-1^{lo} CLP²⁷) (Figure 1A and Supplementary Fig. 1). A total of 597 single cells (123 CLPs, 124 GMPs, 121 HSCs, 116 LMPPs, 113 PreMegEs) passed quality control measures (see Methods).

Single cell gene expression analysis was performed for 24 genes in all 597 cells (see Supplementary Table 3 for raw Ct data). Our gene set included 18 transcription factors (Figure 1B) with known key roles in haematopoiesis, as well as five housekeeping genes and the Stem Cell Factor receptor *c-Kit*, which is expressed on the surface of all analysed haematopoietic stem/progenitor subsets²³⁻²⁸. We have previously reported the potential for abundant regulatory linkages between many of the 18 TFs^{9, 11, 12, 29-33}, and a similarly densely interconnected network was obtained using data curated from the literature and protein interaction databases (Figure 1B). Importantly, while previous studies have examined individual HSCs and commitment to the erythroid lineage, they have been limited

in cell numbers¹⁶, were focussed on expression heterogeneity^{16, 17} or have examined a lineage-specific set of genes²². Moreover, the potential for identifying regulatory connections from single cell gene expression profiling had not been demonstrated, nor had the potential for dynamic changes of regulatory network states been studied during the differentiation of HSCs into the various multipotential blood progenitors.

Single cell gene expression analysis recovered expected expression patterns for the 18 TFs as well as housekeepers and *c-Kit* (Figure 2). For example, *c-Kit* expression was highest in HSCs and gradually reduced in the progenitor populations, consistent with the reported downregulation in progenitors²⁸. *Gata1* is known to be expressed at high levels in erythroid and megakaryocyte lineages, but not in HSCs³⁴, and here was expressed in around two thirds of PreMegE cells, yet absent in almost all cells of the other populations. Likewise, *Gata2* is known to be expressed in HSCs and during megakaryopoiesis^{35, 36}, and in our data was expressed in most HSCs and PreMegEs but at lower levels or not at all in LMPPs, GMPs and CLPs. GFI1B is important for the development of erythroid progenitors, while GFI1 is important for myeloid and T cell development, and the two factors are known to be mutually inhibitory^{37, 38}. Outside of the HSC population; *Gfi1* was expressed in the majority of LMPPs, CLPs and GMPs, but rarely in PreMegEs, while *Gfi1b* was expressed in most PreMegEs, with lower or absent expression in LMPPs, CLPs and GMPs.

Many genes exhibited heterogeneous expression within cell populations, with some cells expressing the gene at high levels and undetectable expression in others, in line with previous reports of expression heterogeneity in blood stem and progenitor populations^{16-18, 22}. Several TFs, including *Runx1* and *Fli1*, had a very similar gene expression distribution in all cell types, and these genes were expressed in almost all analysed cells. Conversely, genes including *Erg*, *Lmo2* and *Meis1*, differed in expression level between cell types. Genes including *Gfi1*, *Gfi1b* and *Scf* (also known as *Tal1*) showed bimodal expression amongst the cells that expressed the gene, with the potential therefore to generate three distinct expression states (high, medium, not-expressed) within a single population that is pure based on FACS analysis. Importantly, such detailed insights into the dynamical nature of TF gene expression in primary blood stem and progenitor cells could not have been obtained from population studies.

Cell populations can be resolved by differential network activity states

To establish cell type-specific patterns of gene expression that may aid our understanding of network activity and cell state transitions, we next performed hierarchical clustering and principal component analysis using the expression data for our TFs in all 597 haematopoietic stem/progenitor cells. The relatedness of cells is determined using only the gene expression values, without prior knowledge of which population a cell originates from. Hierarchical clustering demonstrated that mRNA levels for these 18 key TFs allow the partitioning of cells largely by sorted population (Figure 3A). This was particularly clear for the GMPs, which formed a distinct cluster. HSCs and PreMegEs formed a cluster separate from the myeloid and lymphoid lineages in which the two populations were also largely separated from each other, while LMPPs and CLPs showed significant overlap. There was some mixing of HSCs with LMPPs and PreMegEs, in line with the evidence that LMPPs and the megakaryocyte/erythroid lineage may be generated as early and alternative fates of HSCs²⁴ and so are both closely related to HSCs but distinct from one another.

Principal component analysis confirmed the above results, where each data point represents a single cell, colour-coded according to its flow cytometric phenotype (Figure 3B, upper panel). Principal component 1, which captures the largest proportion of the variation in the data, separates the HSCs and PreMegEs from the lymphoid and myeloid populations, and partially separates the GMPs from the LMPPs and CLPs. Principal component 2 further

separates the HSCs and PreMegEs. However, although individual populations can be distinguished there is also significant overlap, particularly between the LMPPs and CLPs, both of which contain lympho-myeloid-restricted progenitors³⁹, indicating that cells at the edges of adjacent populations have similar network activity states. *Gata2*, *Gfi1b*, *Scl* and *Gfi1* contribute to separation of the HSCs and PreMegEs from the myelolymphoid populations along component 1, while *Erg*, *Hhex* and *Gata1* are important across component 2 (Figure 3B, lower panel), consistent with known expression patterns in these populations. *Runx1* and *Fli1* contributed little to the separation of cell types, consistent with their similar expression distributions between cell types (Figure 2).

Gaussian Process Latent Variable Models (GPLVMs) are a non-linear generalisation of PCA⁴⁰ and were recently shown to be a powerful alternative, in particular for resolving non-linear differences in single-cell gene expression patterns⁴¹. GPLVM resulted in a better separation of the different cell types than PCA (Figure 3C), with the greatest improvement found for GMPs. However, LMPPs and CLPs cells could not be resolved completely. Our ability to separate populations was further confirmed by calculating the spatial median, a robust multivariate measure of the ‘centre’ of the distribution for each cell type, and then analysing distances between cell types. These GPLVM map distances reflected the differentiation hierarchy shown in Figure 3A with cell types close in the hierarchy located close together in the map (Supplementary Fig. 2).

While for standard PCA the relevance of different genes to the separation of the data (quantified by component loadings in Figure 3B) can only be found for the entire PCA map, the relevance of each gene can change across the GPLVM map, providing a greater resolution of the changes separating cell types. The GPLVM relevance map (Figure 3D) shows the most important gene at each point of the map, and illustrates for example that *Gata2*, *Gfi1* and *Gfi1b* feature frequently throughout the map and particularly in the region bordering the HSC and LMPP populations. Indeed, expression maps for individual genes (Figure 3E) demonstrate that high *Gata2* expression occurs mostly in the HSC and PreMegE populations, while high *Gfi1* is restricted mostly to LMPPs, GMPs and a subset of CLPs. Taken together therefore, bioinformatic analysis of single cell gene expression allowed us to correlate distinct expression states of a core set of 18 key haematopoietic TFs with some of the earliest blood stem and progenitor populations, including the earliest known lineage restriction stage from HSCs resulting in distinct MegE and lympho-myeloid restricted pathways.

Single cell analysis reveals dynamic regulatory relationships

We next hypothesised that single cell expression data could be used to identify regulatory linkages by identifying pairs of factors with correlated expression, where a positive correlation suggests that one factor may activate another and a negative correlation indicates an antagonistic relationship. Correlation analysis and hierarchical clustering of the eighteen transcription factors across all 597 haematopoietic stem/progenitor cells revealed both positive and negative correlations (Figure 4A, top left panel). Among the positive correlations is a group of seven genes (*Scl*, *Gata2*, *Nfe2*, *Eto2*, *Gfi1b*, *Gata1* and *Ldb1*) known to be important in the erythroid/megakaryocytic lineages^{35, 38, 42-49}, while there was a negative correlation between *PU.1* and *Gata1*, which are thought to function as a switch controlling erythroid and myelomonocytic fates^{50, 51}. To establish whether such regulatory relationships were stable or dynamic during differentiation, we repeated the correlation analysis for each of the five stem/progenitor populations individually (Figure 4A). While many of the strong positive correlations identified in the whole data set remained stable between cell types, there were some clear differences, particularly in negative correlations, which could suggest that repression, or relief of repression, of some TFs by others is a vital step in cell fate transitions. For example, the strong negative correlation between *Gfi1* and

Gata2 present in the whole dataset is seen only in HSCs. *Gfi1b* and *Gata1* are negatively correlated with *PU.1*, *Mitf*, *Gfi1*, *Ly11* and *Lmo2* in GMPs, and to some extent in CLPs, but are either not correlated or positively correlated in the earlier progenitors and in the erythroid/megakaryocyte lineage. Together these results indicate that expression of the core haematopoietic transcriptional regulatory network is dynamic (Figures 2 and 3), which is presumably intimately connected to the dynamics of regulatory interactions between the components of the network (Figure 4).

Significant positive and negative correlations between TFs were displayed as a putative interaction network (Figure 4B). Among these is the known *Scl-Gata2* relationship and the *PU.1-Gata1*⁵¹ and *Gfi1-Gfi1b*^{37, 38} inhibitory relationships, indicating that additional relationships identified may indeed signify previously unrecognised interactions. Two newly predicted regulatory links (*Gata2-Gfi1* and *Gata2-Gfi1b*) suggested possible involvement of GATA2 in modulating the cross-inhibitory relationship between GFI1 and GFI1B (Figure 4B), which was of particular interest to us because we had seen downregulation of *Gata2* and *Gfi1b* accompanied by reciprocal upregulation of *Gfi1* in a recent transcriptomic and epigenomic analysis of leukaemia development in an MLL-ENL-driven mouse model of acute myeloid leukaemia (AML)⁵² (see Supplementary Fig. 3).

Direct repression of *Gata2* distal enhancer elements by GFI1 provides a likely mechanism for negatively correlated expression

To investigate whether downregulation of *Gata2* might be mediated directly through GFI1 binding to *Gata2* regulatory elements, we interrogated existing CHIP-Sequencing data for GFI1 in MLL-ENL transduced cells^{52, 53}. Across the *Gata2* locus, a prominent peak was identified 83kb upstream of the *Gata2* transcriptional start site (Supplementary Fig. 3B). This -83 kb region had been shown previously to loop to the *Gata2* promoter⁵⁴, and was bound in the HPC7 haematopoietic progenitor cell line^{11, 55} by multiple TFs (Supplementary Fig. 3)¹¹. TF CHIP-Seq studies are currently not possible with the small numbers of cells that can be obtained for the highly purified blood stem cell populations used here for single cell expression analysis. We nevertheless wanted to confirm binding of GFI1 to the *Gata2* gene locus in primary blood cells, and therefore performed GFI1 CHIP-Seq in primary mast cells, which like HSCs express the stem cell factor receptor c-KIT and a number of TFs important for HSCs including GFI1 and GATA2. This GFI1 CHIP-Seq experiment confirmed GFI1 binding to the *Gata2* -83 kb region in primary mouse blood cells (Figure 5A).

As no *in vivo* activity has as yet been reported for the *Gata2* -83 kb region, we generated a LacZ reporter construct with the -83 kb region fused to a minimal SV40 promoter/LacZ reporter cassette. Analysis of LacZ expression in E11.5 transgenic mouse embryos demonstrated consistent staining in the midbrain, hindbrain and spinal cord (Figure 5B), all known domains of endogenous *Gata2* expression⁵⁶. However, no haematopoietic staining was seen in any of the transgenic embryos. We had shown previously that a *Gata2* -3 kb enhancer is active at E11.5 in the dorsal aorta endothelium including budding haematopoietic cells, but not the foetal liver⁹. Given the prominent TF binding to the -83 kb region in haematopoietic cells, we next asked whether a combination of the -83 and -3 kb enhancers was able to drive expression to foetal liver haematopoietic cells. Transgenic embryos carrying a combined enhancer construct (-3/SV/lacZ/-83) displayed the neural activities of both of the individual enhancers (Figure 5B). Moreover, staining was not only seen in the dorsal aorta but also in foetal liver haematopoietic cells. Transgenic analysis therefore confirmed the *Gata2* -83 kb region as a candidate enhancer element involved in haematopoietic expression of *Gata2*.

To investigate whether GFI1 could repress activity of this element, we next generated a luciferase reporter construct (–3/SV/luc/–83) and performed transfection assays in the HPC7 progenitor cell line which expresses high levels of *Gata2* but very low levels of *Gfi1*³³. Compared with control transfection assays, we observed that co-transfection of a *Gfi1* expression construct caused a 40% reduction in reporter activity (Figure 5C). Taken together therefore, ChIP-Seq, transgenic and transfection studies validated the previously unrecognised regulatory interaction between GFI1 and *Gata2*.

Direct activation of *Gfi1b* distal enhancer elements by GATA2 provides a likely mechanism for positively correlated expression

To investigate whether positively correlated expression of *Gfi1b* and *Gata2* might be mediated directly through GATA2 binding to the *Gfi1b* gene locus, we interrogated existing ChIP-Seq data for the HPC7 cell line¹¹ which demonstrated binding of GATA2 to the *Gfi1b* promoter as well as three 3 candidate enhancer regions 13kb, 16kb and 17kb downstream of the start of the *Gfi1b* gene (Supplementary Fig. 4). To confirm binding in primary mouse blood cells, we again turned to primary mast cells, and indeed, significant GATA2 binding was observed at all four regions bound by GATA2 in HPC7 as well as a region in the first intron not bound in HPC7 (Figure 6A). Taken together, TF-binding in both mast cells and HPC7 therefore identified 4 candidate regulatory regions that might be involved in mediating control of *Gfi1b* expression in stem/progenitor cells by GATA2.

To assess whether the four regions bound by GATA2 correspond to bona fide gene regulatory sequences, we performed transgenic assays. LacZ reporter constructs were generated for the *Gfi1b* promoter as well as the +13, +16 and +17 kb candidate enhancer regions and assayed in E11.5 mouse embryos. The promoter region alone did not mediate any haematopoietic expression, a phenomenon we have observed before for both *Runx1* and *Scl/Tal1* promoters^{57, 58}. By contrast, all three distal regions mediated haematopoietic expression: the +13 kb region showed weak expression in a subset of circulating (likely primitive) blood cells, the +16 kb region displayed strong staining in haematopoietic clusters in the dorsal aorta as well as a subset of foetal liver cells, and the +17 kb region showed staining in a small subset of foetal liver haematopoietic cells (Figure 6B). Transgenic analysis therefore provided *in vivo* validation of GATA2-bound regions, with the +16 and +17 kb regions being particularly relevant due to their activity in the anatomical sites of early definitive haematopoietic development.

Since transgenic analysis had focussed our attention on the +16 and +17 kb enhancer regions as possible mediators of *Gfi1b* activation by GATA2, we investigated the possible presence of conserved GATA motifs. Both the +16 and +17 kb regions showed extensive sequence conservation across a wide range of mammalian species consistent with their function as gene regulatory elements (Supplementary Fig. 5). Moreover, both enhancers contained two completely conserved GATA motifs. To investigate the role of the GATA sites in enhancer activity, we generated luciferase reporter constructs with both the wild type and GATA mutant +16 and +17 kb enhancers (Figure 6C). Following stable transfection assays in the myeloid progenitor cell line 416B, both regions showed substantial enhancer activity that was almost completely lost in the GATA mutant constructs. GATA2 binding in ChIP assays, activity in transgenic assays and presence of conserved GATA motifs essential for enhancer function are therefore all consistent with a model whereby GATA2 directly activates *Gfi1b* expression through the +16 and +17 kb enhancer regions.

Taken together therefore, single cell gene expression analysis of primary blood stem and progenitor cells suggests the existence of a regulatory triad including *Gata2*, *Gfi1* and *Gfi1b* (Figure 6D), the connectivity of which has been validated using transgenic and transcriptional assays. In this triad, the reported mutual inhibition of GFI1 and GFI1B^{37, 38}

is retained, but is modulated by GATA2 through its activation of *Gfi1b* and repression by GFI1.

DISCUSSION

Cellular phenotypes are controlled by networks of interacting TFs. Development can therefore be described as a procession through multiple dynamic regulatory states, which in the case of multilineage differentiation may give rise to multiple distinct outcomes. However, classical networks derived from gene expression or ChIP-Seq data provide a population average, giving little insight into cellular heterogeneity and possible regulatory interactions likely to be critical for the lineage commitment/restriction steps of individual progenitor cells⁵⁹.

In this study, we investigated a core transcriptional network of 18 TFs in single cells of five related stem and progenitor populations. Bioinformatic analyses were able to broadly distinguish sorted cell populations based only on the expression of those TFs, indicating that early haematopoietic stem/progenitor cells are characterised by related but distinct network activity states. While gross gene expression patterns were consistent with published population studies, we also found significant heterogeneity in TF expression within populations. This confirms previous observations that FACS sorted populations are molecularly and functionally heterogeneous at the single cell level¹⁶⁻¹⁸, and suggests either that heterogeneity is an inherent characteristic of stem cells, or that previously unresolved subpopulations may be present that could represent intermediate differentiation steps.

Importantly, heterogeneity did not confound our ability to resolve populations based on their distinct gene expression patterns. Thus, further analysis of defining TF expression patterns and interactions at important lineage restriction stages should facilitate the unravelling of critical TF interactions decisive for lineage commitment steps. Index sorting for example permits the tracing back of each cell to its position in the sorting data, and may therefore allow us in future to link heterogeneous gene expression states to novel subpopulations. Strikingly, in our study several transcription factors had similar expression levels in all 597 cells analysed, regardless of cell type of origin, including *Runx1* and *Fli1*, while others were much more variable. This may suggest that the blood network requires, or is able to tolerate, variation in some factors but not others. Furthermore, correlation analysis revealed both stable and dynamic TF relationships across the cell types analysed, together suggesting that some TF interactions may be vital for the general stability of the network and so remain constant, while others are important for network dynamics and state transitions and are therefore more variable.

As transcriptional regulatory networks control the spatiotemporal regulation of lineage-specific genes, successful reconstruction of regulatory hierarchies represents a major step towards gaining a mechanistic understanding of cellular decision-making processes. For example, detailed experimental and computational analysis of a core circuit of *Gata2*, *Scf* and *Fli1* revealed that this circuit is able to function as a bistable switch, where the internal wiring enables the network to filter noise when responding to external cues^{9, 60}. However, the generation of large-scale experimentally validated network models is impeded by the relatively low-throughput of experiments that can provide detailed experimental information on the functionality of individual regulatory elements. Network inference, where transcriptional interactions are inferred from statistical dependencies in large expression profiling datasets, provides an alternative approach⁷. However, in addition to the very substantial costs of performing hundreds of expression profiling experiments, standard microarray or RNA-Seq profiling requires large cell numbers, which may not be feasible for rare stem and progenitor populations.

Here we have demonstrated that gene expression data from a large set of single cells presents an alternative approach to network reconstruction through the analysis of pairwise correlations between network components. While hundreds of samples are still required, the fact that these are single cells makes this approach applicable to rare stem cell populations. The robustness of inferred regulatory links was demonstrated by our ability to recover key known regulatory relationships such as the *Gata1-PU.1* and *Gfi1-Gfi1b* antagonisms. Furthermore, we validated two novel putative regulatory interactions predicted from bioinformatic analysis of our single cell expression profiling data. This demonstrated that GFI1 directly represses *Gata2* expression through the -83 kb regulatory region, while GATA2 activates *Gfi1b* through the +16 kb and +17 kb regulatory regions. The resultant connectivity resembles a type 2 coherent feed forward loop⁶¹, and suggests that GATA2 works within this regulatory triad to modulate the antagonism between GFI1 and GFI1B. Moreover, GATA2 inhibits lymphopoiesis⁶², and along with *Gfi1b* is down-regulated concurrent with *Gfi1* upregulation in the myelolymphoid lineages in our data. This suggests that direct downregulation of *Gata2* and *Gfi1b* by GFI1 may represent a key event during the specification of early lymphoid cells, similar to the role of Tif1 in modulating the PU.1-GATA1 antagonism in the erythroid versus myeloid fate choice in zebrafish⁶³.

Loss of function mutations in *Gata2* have recently been reported to predispose carriers to developing acute myeloid leukaemia⁶⁴, whereas inhibition of *Gfi1* prolonged survival in leukaemia mouse models⁶⁵. This suggests that identification of network states and reconstruction of network hierarchies from single cell expression profiling will not only enhance our knowledge of normal differentiation and development, but also provide a blueprint for understanding the subversion of cell fate control likely to underlie many degenerative and malignant pathologies.

METHODS

Purification of stem cells and progenitor cells

Bone marrow cells were isolated from the bones (femurs, tibiae and crista ileaca) of 9-12 week old C57BL/6 mice. For the isolation of LMPPs and HSCs, cells were enriched for CD117+ (c-Kit) cells by MACS bead separation with anti-CD117 immunomagnetic beads (Miltenyi Biotec); for the other cell populations, unenriched bone marrow was used. Cells were pre-incubated with Fc-block for the HSC, LMPP and CLP stains but not for the myeloid progenitor stain (GMP and Pre-MegE isolation). Cells were stained with antibodies to mouse antigens to allow separation of the individual populations (Supplementary Table 1). A FACSAria II (BD Biosciences) was used for all cell sorting. Fluorescence-minus-one controls and unstained populations were used as gate-setting controls. Single cells were seeded by an automated cell deposition unit directly into the Fluidigm assay mixture (see below). Test sorts before and after single cell sorts verified the purity of all populations at >98% based on expression of cell-surface markers (Supplementary Fig. 1). Fluorescent beads were sorted into 96-well plates before and after samples to verify that only single events were sorted into each well.

Single cell gene expression analysis

Single-cell gene expression analysis was performed using 48.48 Dynamic Array integrated fluidics chips (M48, Fluidigm Corporation) on the BioMark HD platform (Fluidigm Corporation), which facilitates the simultaneous analysis of 48 genes in each of 48 samples. cDNA synthesis and specific target amplification (preamplification) of genes of interest were performed using the CellsDirect One-Step qRT-PCR kit (Invitrogen). Single cells were sorted by FACS directly into individual wells of 96-well plates containing 5µL CellsDirect 2× reaction mix (Invitrogen), 0.1µL SUPERase RNase inhibitor (Ambion), 2.5µL 0.2× assay

mix, 1.2µL TE buffer (Invitrogen) and 1.2µL SuperscriptIII/Platinum Taq (Invitrogen). The 0.2× assay mix contained a pool of 24 TaqMan assays (Applied Biosystems; details available on request) at a 1:100 dilution of each assay in TE buffer. Reverse transcription and specific target amplification were performed in the same plates immediately after sorting as follows: 50°C for 15 minutes, 95°C for 2 minutes, 22 × (95°C for 15 seconds, 60°C for 4 minutes). cDNA was diluted 1:5 with TE prior to qPCR on the BioMark HD. cDNA was stored at -20°C before processing on the BioMark HD. 6 positive controls of 20 cells per well, 14 negative controls (no cell sorted) and 124 single cells were sorted for each population. This corresponds to 3 M48 Dynamic Arrays per population, each containing 2 positive controls, 4-6 negative controls and 40-42 single cells. For the qPCR, 3µL of each TaqMan assay was mixed with 3µL Gene Expression Assay Loading Reagent (Fluidigm). 2.7µL of diluted cDNA was mixed with 3µL 2× TaqMan Universal Mastermix (Applied Biosystems) and 0.3µL Gene Expression Sample Loading Reagent (Fluidigm). 5µL of each sample and assay were loaded into individual sample and assay inlets on the M48 Dynamic Array. Samples and assays were then loaded into the reaction chambers of the Dynamic Array using the IFC Controller MX (Fluidigm), and then transferred to the BioMark HD for qPCR (95°C for 10 minutes; 40 cycles of 95°C for 15 s and 60°C for 60 s).

Testing TaqMan Assays

TaqMan assays were tested for single cell PCR by performing standard curves on the BioMark of cDNA from a population of 100 cells of the HPC7 haematopoietic progenitor cell line⁵⁵, reverse transcribed and preamplified as described above. Assays were selected based on the amplification efficiency and lack of background expression in no template controls. The linear range of all assays used was within the sensitivity of the BioMark HD (Ct 7-27).

Bioinformatic analysis of single cell gene expression data

Single cell expression data were initially analysed with the Fluidigm Data Collection software. For quality control, amplification curves were quality filtered using a threshold of 0.75 and Ct thresholds were set for each assay, with the same thresholds used across all experiments and cell populations. Data were then exported to Excel as .csv files. A table with all of the Ct values is available in the supplemental material accompanying this paper as Supplementary Table 3. Samples not expressing any genes (likely as a result of a failure of the sorter to put a cell in the well) were excluded from the analysis (n=6), as were cells not expressing the housekeeping genes *Ubc* (n=0) or *Polr2a* (n=17). Expression values over the cut off of the machine or beyond the linear range of the TaqMan assays (Ct>27) were set to 28. Each assay was performed in duplicate, and the mean of the duplicates was used for subsequent analysis. Following these quality control measures, Ct values were calculated as previously described²⁰ by cell-wise normalisation to the mean expression level of *Ubc* and *Polr2a*, as the two most robustly expressed housekeeping genes. Briefly, Ct values were subtracted from the assumed no template background of the BioMark of 28²⁰, followed by subtraction of the mean Ct value of *Ubc* and *Polr2a* for each cell. The Ct value for genes that were not expressed was then set to 15, representative of the detection limit of the BioMark. Hierarchical clustering and principal component analysis were performed in R (www.r-project.org). Hierarchical clustering and principal component analysis were performed only on the data for the 18 transcription factors, excluding *c-Kit* and housekeeping genes. Hierarchical clustering was performed on cells using Spearman Rank correlation. Positive and negative correlations between pairs of genes were tested with Spearman Rank correlation, with P-values calculated based on 10,000 permutations. Positive correlations with a Z-score above 12 ($P < 3E-33$) and negative correlations with a Z-score below -4 ($P < 6.09E-05$) were considered significant, with known antagonistic relationships recovered beyond these values. PCA was performed using the `prcomp` function.

A non-linear generalisation of PCA, a Gaussian Process Latent Variable Model (GPLVM), was employed to generate a nonlinear embedding of the expression data using the FGPLVM toolbox (<http://staffwww.dcs.shef.ac.uk/people/N.Lawrence/fgplvm/>). A probabilistic mapping was constructed from a 2 dimensional latent space to the 18 dimensional data space using Gaussian processes⁴⁰. To allow for non-linear effects we used a radial basis function (rbf) kernel to construct the covariance matrix. The 2D coordinates of the latent points corresponding as well as the kernel parameters were determined by optimising the likelihood of the data-set using 1000 iterations of a scaled conjugate gradient optimiser⁴⁰. A gene relevance map – corresponding to a generalised loadings plot for standard PCA – was generated; for this the mapping from latent space to data space was used to calculate the gradient of the expected value for 20×20 regularly spaced points across the GPLVM-map and the most important gene (greatest norm of gradient) was plotted⁴¹. Single-gene expression maps were generated by calculating the expected value of a single gene for 50×50 equally spaced points across the GPLVM-map. For each cell type the spatial median *m* in 2D was calculated by minimising the expected distance between *m* and the 2D coordinates of all cells of the respective cell type using an extended Weiszfeld algorithm⁶⁸.

Mouse bone marrow derived mast cells (BMMC)

Bone marrow cells were collected from tibias and femurs of 3- to 5-month old adult mice. Cells were cultured in Iscove's modified Dulbecco's medium (IMDM) supplemented with 10% fetal bovine serum (Sigma), 1% penicillin/streptomycin (Sigma), 150 μM MTG (Sigma), 10% stem cell factor conditional media from BHK/MKL cells and 10ng/ml of recombinant mIL-3 (Peprotech). Cells were frequently transferred to new flasks to remove adherent cells and experiments were performed after 3 weeks, when cultures were homogenous. Homogeneity of culture was confirmed by presence of FcERI by FACS and toluidine blue staining of cytopins.

Chromatin Immunoprecipitation Sequencing

ChIP assays were performed as previously described¹¹ using polyclonal antibodies against GATA2 (Santa Cruz, sc9008x) and GFI1 (Abcam, ab21061) and control nonspecific rabbit IgG (Sigma, I5006). Each sample was amplified⁷ and sequenced using the Illumina 2G Genome Analyzer, following manufacturer's instructions. Sequencing reads were mapped to the mm9 mouse reference genome using Bowtie⁶⁹, converted to a density plot, and displayed as UCSC genome browser custom tracks.

Transgenic Mouse Analysis and Luciferase assays

Luciferase and LacZ reporter constructs were generated using standard recombinant DNA techniques. Coordinates of chromosomal regions cloned are given in Supplementary Table 2. Luciferase⁷ assays were performed as described⁷⁰. E11.5 transgenic mouse embryos were generated and LacZ-stained by Cyagen Biosciences. Staining patterns were analysed as described⁷¹.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgments

We thank Sally Ann Clark for help with FACS sorting. Research in the authors' laboratory is supported by the Medical Research Council, Leukaemia and Lymphoma Research, The Leukaemia and Lymphoma Society, Cancer Research UK, and core support grants by the Wellcome Trust to the Cambridge Institute for Medical Research and Wellcome Trust - MRC Cambridge Stem Cell Institute.

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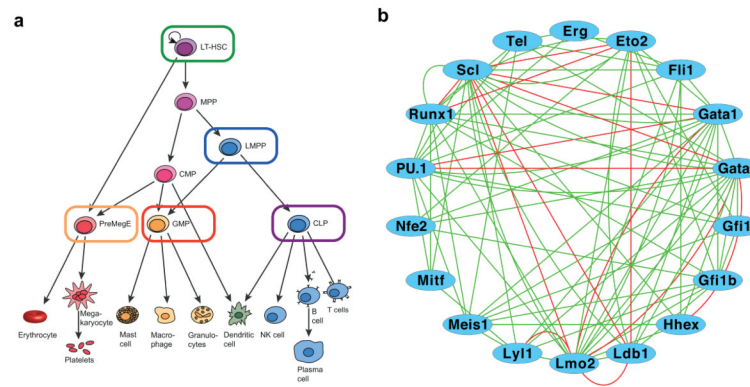


Figure 1. Single cell gene expression analysis of a core haematopoietic transcriptional regulatory network

(a) Schematic of the haematopoietic hierarchy, with the megakaryocyte-erythroid lineage in red, the myeloid lineages in orange and the lymphoid lineage in blue. Cell types investigated in this study are outlined in the colours used to represent these populations in subsequent figures, and encompass both early multipotent stem and progenitors and committed progenitors for each of the major haematopoietic lineages. Cell surface phenotypes were LSK CD150⁺CD48⁻ HSC (also gated as CD34^{lo}Flt3⁻), LSK Flt3^{hi} LMPP, Lin⁻IL7R⁺Kit^{lo}Sca-1^{lo} CLP, CD41^{lo}CD16/32^{hi} GMP (also gated Lin⁻c-Kit⁺CD150⁻), CD16/32^{lo}CD41⁻CD150⁺CD105^{lo} PreMegE (also gated Lin⁻c-Kit⁺). LT-HSC, long-term haematopoietic stem cell; MPP, multi-potent progenitor; LMPP, lymphoid-primed multipotent progenitor; CMP, common myeloid progenitor; CLP, common lymphoid progenitor; GMP, granulocyte-monocyte progenitor; PreMegE, pre megakaryocyte erythroid progenitor; NK cell, natural killer cell. (b) Network diagram of data curated from the literature and protein interaction databases (STRING⁶⁶ and FunctionalNet⁶⁷) illustrating the complex interactions between 18 core haematopoietic transcription factors. Green lines indicate functional relationships and red lines indicate direct protein-protein interactions. Activating and inhibitory connections are not distinguished.

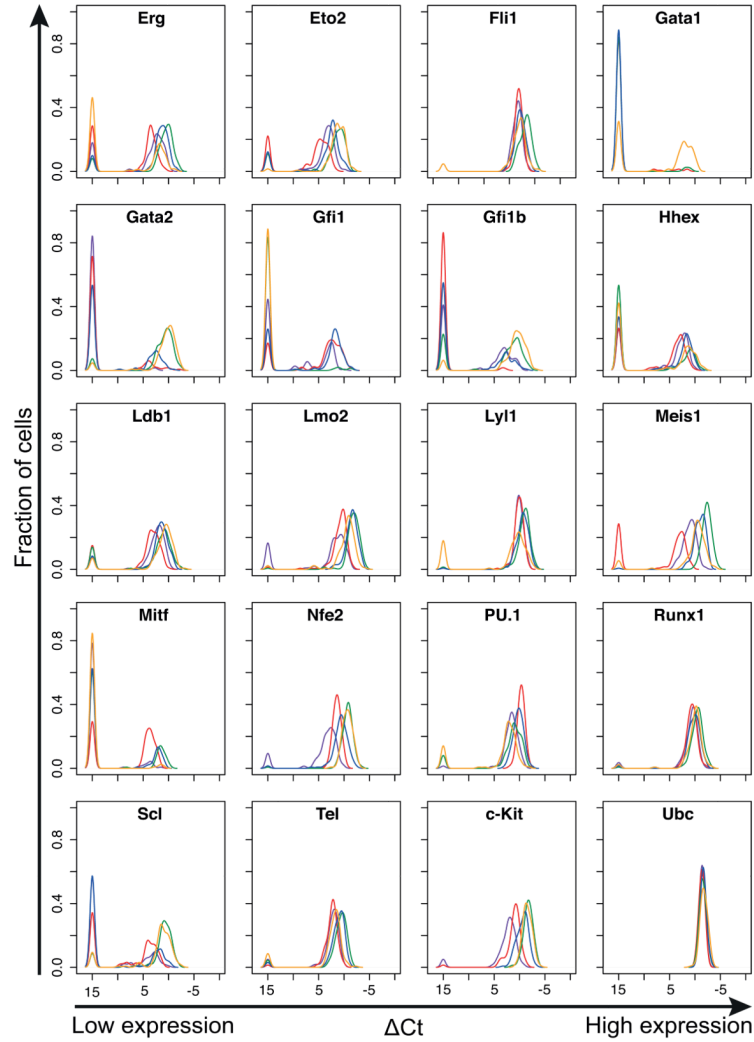


Figure 2. Haematopoietic transcription factors show heterogeneous expression in haematopoietic stem and progenitor cells

Density plots for 18 transcription factors, the stem cell factor receptor *c-Kit*, and the housekeeping gene *Ubc*, in five haematopoietic stem and progenitor populations. The density indicates the fraction of cells at each expression level, allowing direct comparison of the expression level of each gene in all five populations. Green, HSC; Blue, LMPP; Purple, CLP; Red, GMP; Orange, PreMegE.

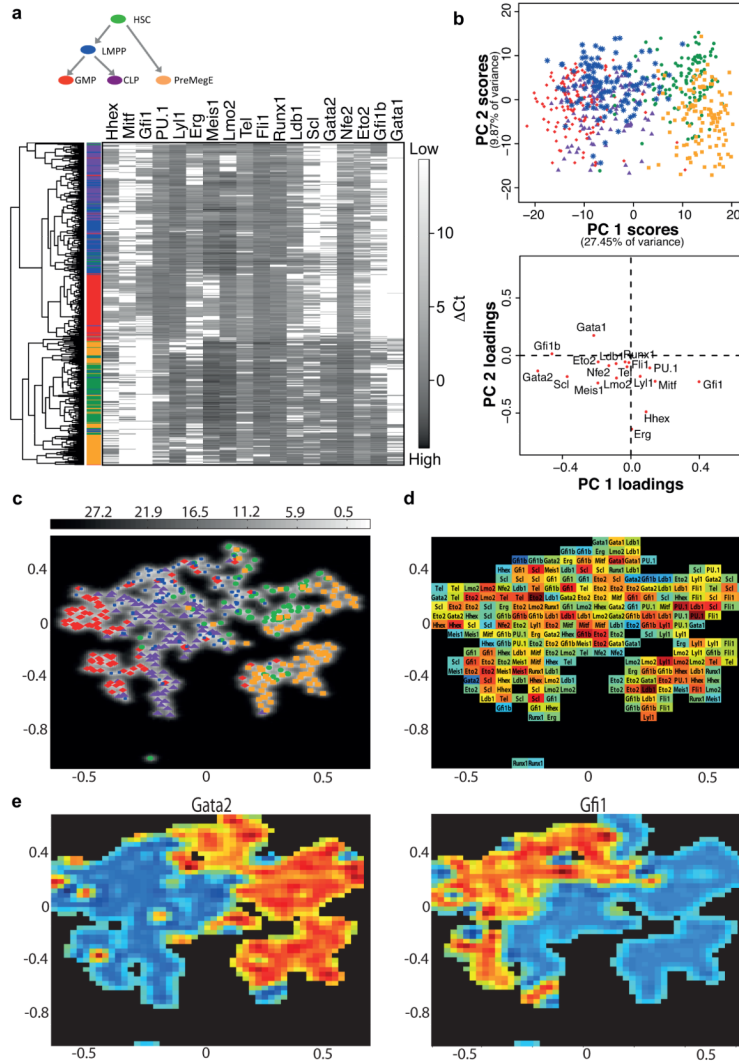


Figure 3. Single cell gene expression analysis reveals cell type-specific regulatory codes
 (a) Hierarchical clustering of 597 haematopoietic stem and progenitor cells according to the expression of the 18 TFs. Coloured bar indicates cell type of origin: Green, HSC; Blue, LMPP; Purple, CLP; Red, GMP; Orange, PreMegE. (b) Principal component projections of the 597 haematopoietic stem/progenitor cells, in the first and second components (top), from the expression of all 18 TFs. Principal component loadings (bottom) indicate the extent to which each gene contributes to the separation of cells along each component. (c) Gaussian process latent variable model (GPLVM) illustrating variations of 18 dimensional gene expression patterns between and within cell types in 2D. GPLVMs are a non-linear generalisation of PCA that allow for the analysis of more complex gene expression patterns than PCA and can thus potentially better represent variations between and within populations of different cell types. The uncertainty of the mapping from 2D to the 18 dimensional TF space is encoded in grey (white low uncertainty, grey high uncertainty). (d) Relevance map showing the most important genes across the GPLVM map. The colours correspond to the distance of the respective gene from the origin in a standard loadings plot (red far away/important, blue close to origin). (e) Expression maps for *Gata2* (left) and *Gfi1* (right), with high expression in red and low or absent expression in blue. *Gata2* is expressed primarily in the HSC and PreMegE clusters, and *Gfi1* in LMPPs, GMPs and some CLPs.

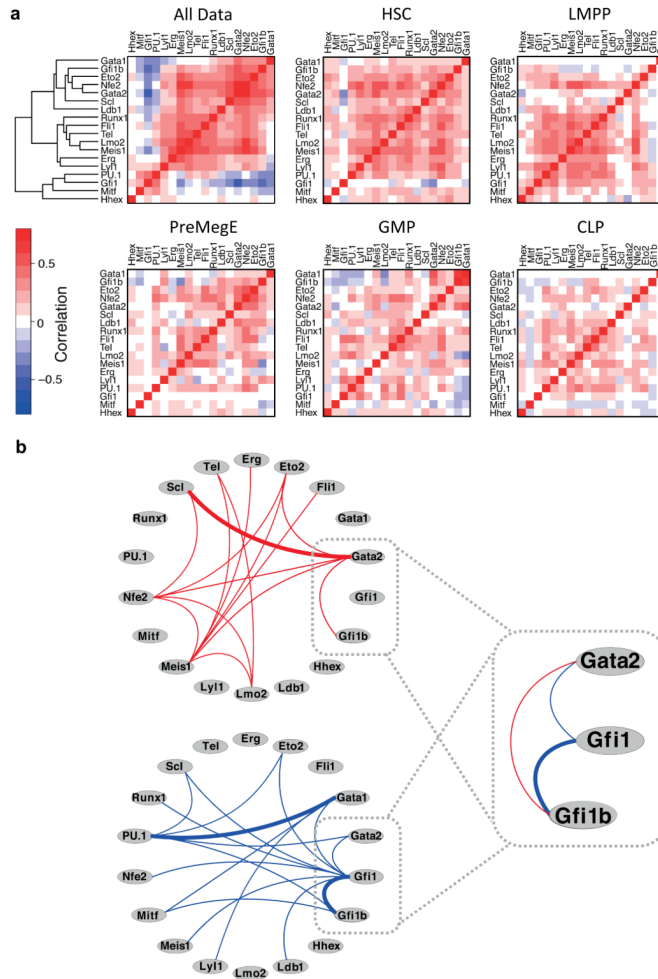


Figure 4. Single cell expression analysis of haematopoietic TFs identifies previously unrecognised putative regulatory interactions between key TFs

(a) Hierarchical clustering of Spearman Rank correlations between pairs of TFs for all 597 cells together and for the different cell types individually as indicated. Genes in all heatmaps are ordered according to the clustering performed for all data. Positive correlations (red) may result from the coordinate expression or lack of expression of pairs of factors in individual cells, while negative correlations (blue) can result either from the expression of one factor in the absence of the other, or from high expression of one factor and reciprocal low expression of the other in the same cell. (b) Network diagrams showing putative activating relationships between TFs suggested by significant positive correlations (top, red lines) and antagonistic relationships suggested by significant negative correlations (bottom, blue lines) in the whole data set. Known relationships are highlighted with bold lines. This highlights a putative transcription factor triad in which *Gfi1* is negatively correlated with *Gata2* and *Gfi1b*, but *Gata2* and *Gfi1b* are positively correlated.

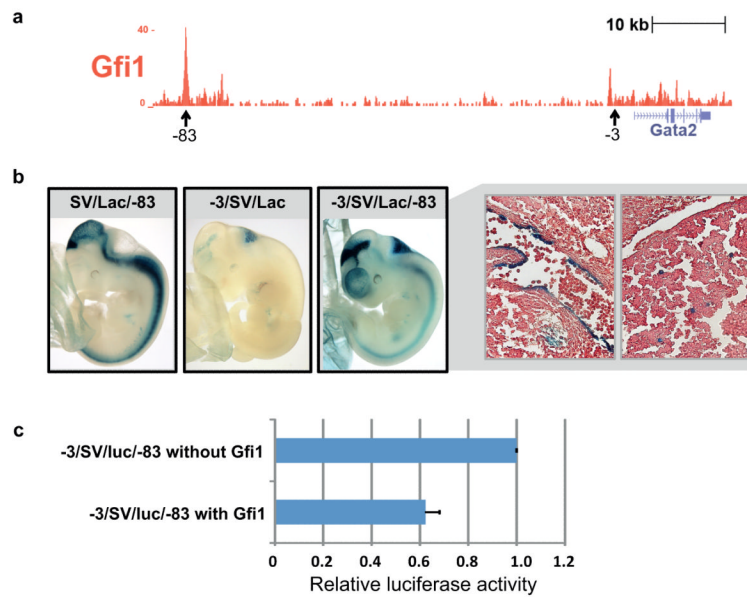


Figure 5. Direct repression of *Gata2* by GFI1 through a distal enhancer element provides a mechanism for negatively correlated expression

(a) ChIP-seq analysis of Gfi1 in primary mast cells indicates that GFI1 binds to the *Gata2* locus at the -83 kb regulatory element. (b) Representative embryos demonstrating LacZ staining for the *Gata2* -83kb, -3kb and combined -3/-83kb regulatory element reporter constructs. The -83kb region alone (SV/LacZ/-83) showed consistent staining of the midbrain, hindbrain and spinal cord, but no haematopoietic staining. The -3kb enhancer (-3/SV/LacZ) had only hindbrain staining. The -83/-3kb combined element (-3/SV/LacZ/-83) showed the neural activities of both individual enhancers, but also staining in the dorsal aorta (right-hand top panel) and foetal liver haematopoietic cells (right-hand bottom panel). Images of sections taken at 40× magnification. (c) A luciferase reporter construct carrying both regulatory elements (-3/SV/luc/-83) was transfected into the HPC7 haematopoietic progenitor cell line, which expresses high levels of *Gata2* but low levels of *Gfi1*. Co-transfection with a Gfi1 expression construct caused a 40% reduction in reporter activity. Luciferase activity is shown relative to -3/SV/luc/-83 and bars are the mean and standard deviation of three biological replicates.

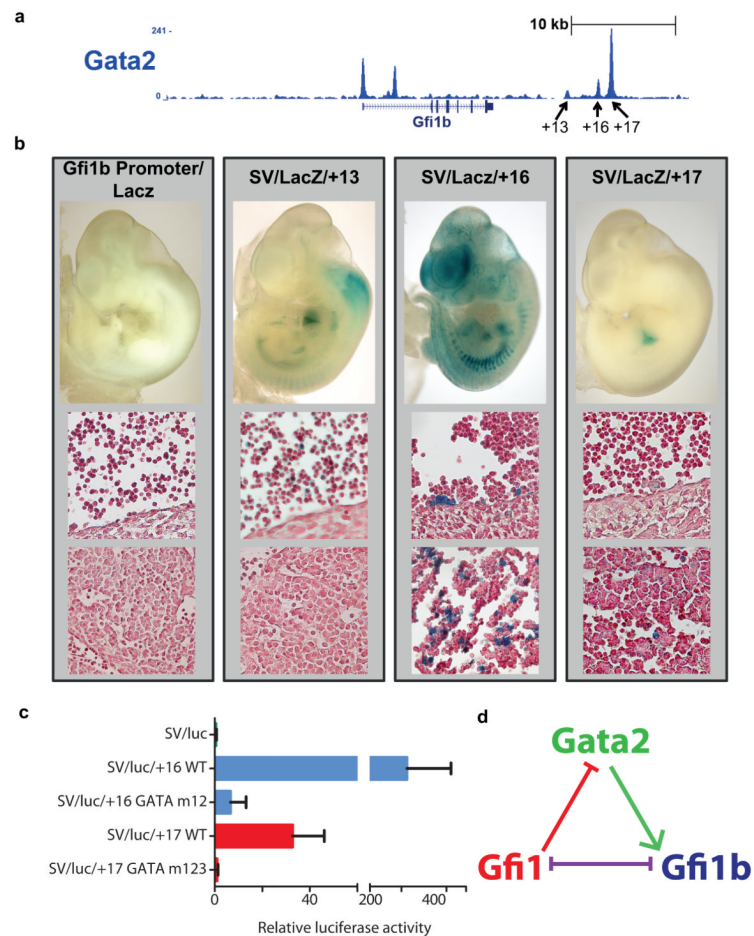


Figure 6. Direct activation of *Gfi1b* by GATA2 through distal enhancer elements

(a) ChIP-seq analysis of GATA2 in primary mast cells indicates that GATA2 binds to the *Gfi1b* locus at multiple locations, including the promoter, a region in the first intron and the +13, +16 and +17 kb regulatory elements. (b) Representative embryos demonstrating LacZ staining for the *Gfi1b* promoter as well as the +13, +16 and +17 kb regulatory elements. The promoter shows no staining. The +13 kb region shows weak expression in a subset of circulating blood cells. The +16 kb region has strong staining in haematopoietic clusters in the dorsal aorta, and in a subset of foetal liver cells. The +17 kb region shows staining in a small subset of foetal liver haematopoietic cells. Images of sections taken at 40× magnification. (c) Luciferase reporter constructs carrying the wild type +16kb (SV/luc/+16 WT) and +17 kb (SV/luc/+17 WT) kb regulatory regions transfected in 416B cells displayed high levels of luciferase activity, particularly for the +16 kb region. Mutation of the two conserved GATA sites in the +16 kb region (SV/luc/+16 GATA m12) reduced luciferase activity by >95%. Mutation of the two conserved and one partially conserved GATA sites in the +17 kb region (SV/luc/+17 GATA m123) also reduced luciferase activity by >95%. m12 and m123 indicate that GATA sites 1, 2 and 3 were mutated. Luciferase activity is shown relative to SV/luc. Experiments were performed in biological duplicate or triplicate on two separate occasions. Shown is one representative experiment displaying the mean and standard deviation for three biological replicate transfections. (d) A putative regulatory triad including GATA2, GFI1 and GFI1B suggested by the data. In this regulatory triad, GFI1 and GFI1B are mutually inhibitory, while GATA2 can activate expression of *Gfi1b* and GFI1 can repress expression of *Gata2*.