Repression of chimeric transcripts emanating from endogenous retrotransposons by a sequence-specific transcription factor

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Abstract

Background: Retroviral elements are pervasively transcribed and dynamically regulated during development. While multiple histone- and DNA-modifying enzymes have broadly been associated with their global silencing, little is known about how the many diverse retroviral families are each selectively recognized.

Results: Here we show that the zinc finger protein Krüppel-like Factor 3 (KLF3) specifically silences transcription from the ORR1A0 long terminal repeat in murine fetal and adult erythroid cells. In the absence of KLF3, we detect widespread transcription from ORR1A0 elements driven by the master erythroid regulator KLF1. In several instances these aberrant transcripts are spliced to downstream genic exons. One such chimeric transcript produces a novel, dominant negative isoform of PU.1 that can induce erythroid differentiation.

Conclusions: We propose that KLF3 ensures the integrity of the murine erythroid transcriptome through the selective repression of a particular retroelement and is likely one of multiple sequence-specific factors that cooperate to achieve global silencing.

Keywords: Retrotransposons, Chimeric transcripts, KLF1, KLF3, KLFs, Gene regulation, Erythroid, Transcription factors
A notable example of this is the murine Agouti viable yellow (A\(^v\)) allele, in which an upstream intra-cisternal A particle (IAP) retrotransposon functions as a constitutively active promoter that drives ectopic expression of Agouti, resulting in yellow fur, obesity, and increased susceptibility to tumorigenesis [15,16]. The prevalence of this phenomenon, whereby retrotransposons serve as alternative promoters, has recently been revealed following the advent of high-throughput RNA sequencing and shown to occur primarily in embryonic cells but also to some extent in adult tissues [17]. During early embryonic development in particular, up to 20% of the transcriptome has been shown to initiate from within retrotransposons [17,18]. These retroelements frequently function as alternative promoters and show a propensity for tissue-specific activity, more so in fact than non-retrotransposon promoters [17]. In many instances, these retrotransposons have been co-opted by the host by exonization and they are transcribed and spliced to downstream genomic exons [17,19,20]. The resulting chimeric transcripts thus potentially encode isoform variants with spatio- or temporally-restricted expression profiles [21]. Indeed, a recent study of the *Drosophila melanogaster* transcriptome has revealed that several hundred LTR retrotransposons serve as promoters of annotated genes throughout development, exhibiting specific expression profiles depending on the different regulatory modules they carry [22].

However, while there are many reported instances of TEs being co-opted by the host for various biological functions, genomic integration of TEs can also be deleterious [23-25]. For instance, Hodgkin's lymphoma has been shown to arise from aberrant transcription of the *colony-stimulating factor 1 receptor* (*CSF1R*) gene driven by an internal LTR element known as *THEI1* [26].

Accordingly, higher eukaryotes have developed numerous defence mechanisms to silence TEs, typically involving DNA methylation and/or histone modification [27-29]. This silencing largely occurs early in embryonic development and is dependent on epigenetic modifiers including: DNA methyltransferases (DNMTs) [27,30]; histone modifying enzymes such as the demethylase LSD1/KDM1A, the deacetylase HDAC1, and the methyltransferases SETDB1 and G9A [31-35]; and Polycomb Group proteins [36]. Ablation of these factors in embryonic stem cells results in widespread de-repression of retrotransposon-derived transcripts.

However, while silencing of retroelements is broadly carried out by these epigenetic modifiers, little is known about the underlying mechanisms by which the diverse classes of retroelements are each specifically recognized [28]. Indeed the lack of sequence similarity between unrelated retroviral families suggests the existence of multiple recognition factors that participate in the silencing of retroelements. An accumulating body of evidence has pointed towards the possible role of DNA binding, tandem zinc finger proteins in providing this specificity. Thomas and Schneider have proposed a model of co-evolution between retroelements and C\(_2\)H\(_2\) zinc finger proteins based on striking correlations of their expansion throughout vertebrate genomes [37]. This model followed from the discovery that the Krüppel-associated box (KRAB)-zinc finger protein ZFP809 binds and represses a large number of retroelements in mouse embryonic stem cells [38]. ZFP809 achieves this through the recruitment of the corepressor TRIM28 (also known as KRAB-associated protein 1, KAP1). TRIM28 in turn silences ERVs through SETDB1 mediated trimethylation of H3K9 [31,39,40].

The Krüppel-like factors (KLFs) are a family of DNA binding, zinc finger transcription factors [41]. They lack a KRAB domain and are characterized by a set of three tandem C\(_2\)H\(_2\) zinc fingers at their C-termini that confer specificity towards CACCC-like and GC-rich sequences in regulatory elements [42]. While the DNA binding domain is highly conserved within the family, the N-terminal regulatory domains vary considerably such that the different KLFs recruit an assortment of corepressors to activate or repress genes [43].

The founding member of the family, KLF1, is an erythroid-specific transcriptional activator that drives the expression of genes required for red blood cell maturation [44]. One such gene is that encoding the related family member KLF3 [45]. KLF3 and KLF1 recognize similar sequences of DNA that adhere to the consensus 5'-NCN CNC CCN-3' [42,46]. However, unlike KLF1, KLF3 is a transcriptional repressor that recruits the co-repressor C-terminal binding protein (CTBP) [47]. CTBP forms part of a large repressor complex that includes the histone deacetylases HDAC1 and HDAC2, the histone methyltransferases EHMT1 and G9A/EHMT2, and the lysine-specific demethylase LSD1/KDM1A [48]. KLF1 and KLF3 exhibit opposing activities at a number of genes in erythroid cells and serve to fine-tune their expression during erythropoiesis [49,50]. Accordingly, loss of either factor disrupts this balance. *Klf1* null mice die of severe anemia *in utero* while mice lacking KLF3, though viable, exhibit erythroid defects in both fetal and adult tissues [49,51].

Here, we have further explored the interplay between KLF1 and KLF3 in regulating the erythroid transcriptome. We find that KLF1 activates, while KLF3 represses, transcription from a specific family of LTR elements known as *ORRIA*. Ablation of KLF3 results in widespread, de-repressed transcription from these LTRs in erythroid cells. Because the *ORRIA* element contains an intact splice donor site, these transcripts are spliced to exons of the genes in which they reside. We show...
that for the spleen focus forming virus proviral integration 1 (Sfpi1) gene, an ORR1A0-driven transcript is translated into a truncated variant of PU.1 which exhibits dominant negative activity and can functionally promote erythroid differentiation. These results suggest that KLF3 ensures normal murine erythropoiesis by preventing aberrant, chimeric transcription driven from ORR1A0 LTRs by KLF1.

Results
Increased expression of downstream Pu.1 exons in erythroid cells in the absence of KLF3

We recently identified a number of KLF3 target genes via microarray analysis of Klf3−/− TER119+ (erythroid) fetal liver cells at embryonic day E14.5 [49]. These genes were predominantly de-repressed in Klf3 null tissue, consistent with KLF3 being a repressor of transcription. One of the most highly de-repressed genes was that encoding the key hematopoietic regulator PU.1/SFPI1, hereafter referred to as PU.1.

We first sought to validate the upregulation of Pu.1 expression in Klf3−/− cells by quantitative real-time RT-PCR. Initial experiments, using primers that span the exon 2/3 junction of Pu.1, did not recapitulate the microarray results (Figure 1A). Unexpectedly, Pu.1 mRNA was detected at similar levels in Klf3+/+, Klf3+/−, and Klf3−/− Ter119+ E14.5 fetal liver cells. To resolve this discrepancy, we analyzed the individual probe intensities across the Pu.1 locus from the microarray data. The murine Pu.1 gene comprises five exons and of these, exons 2 to 5 are represented by probes on the arrays. Expression of only exons 3 to 5 of Pu.1 was found to be higher in Klf3−/− compared to Klf3+/+ tissue; however, expression of exon 2 was unchanged (Figure 1B). Real-time RT-PCR using primers specific for the exon 3/4 and exon 4/5 boundaries of Pu.1 (Figure 1C, D) confirmed that indeed, exons 3 to 5 exhibit upregulated expression in Klf3−/− cells while exon 2 does not (Figure 1A).

An ORR1A0 LTR element serves as an alternative promoter in the Pu.1 locus in the absence of KLF3

The upregulated expression of exons towards the 3′ end of Pu.1 raised the possibility that an alternative, internal promoter was driving transcription from the locus and that this promoter is repressed by KLF3. To investigate this, we conducted 5′ RACE on mRNA from Klf3+/+ and Klf3−/− Ter119+ fetal liver cells using a reverse primer specific for exon 3 of Pu.1. While electrophoretic...
separation of RACE products revealed a common transcript in both samples (an approximately 420 bp band), a smaller transcript (226 bp) was found in the Klf3−/− sample (Figure 2A). Sequencing of the two RACE products revealed that the larger band corresponds to exons 1 to 3 of a typical Pu.1 transcript (GenBank:NM_011355). The shorter transcript, however, was found to contain exon 3 of Pu.1 preceded by a novel sequence (shown in bold in Figure 2B). This sequence maps to intron 2 of Pu.1 and represents an alternative leader exon, hereby termed exon 2b, which is spliced to exon 3 and which has not been documented previously, to our knowledge (Figure 2C). Hereafter, we refer to this novel transcript as Pu.2.

Figure 2 A novel, internal Pu.1 promoter resides within an ORR1A0 LTR element and is repressed by KLF3. (A) RNA from Klf3+/+ (WT) and Klf3−/− (KO) TER119+ fetal liver cells was subjected to 5′ RACE using a reverse primer specific for exon 3 of Pu.1 and analyzed by agarose gel electrophoresis. The smaller band in the Klf3 KO lane was sequenced and found to contain a novel exon (exon 2b). (B) The sequence of the ORR1A0 LTR, in which Pu.1 exon 2b is shown in bold. Sequences which fit the KLF binding consensus 5′-NCN CNC CCN-3′ are boxed, and the TATA box at −30 is underlined. (C) Schematic of the murine Pu.1 locus showing the position of exon 2b. Exons are represented by blue boxes, transcription start sites by arrowheads and splicing events by broken lines. Start points of translation (ATGs) for the two alternative transcripts are also shown. (D) Real-time RT-PCR quantification revealing that transcripts containing exon 2b spliced to exon 3 of Pu.1 (that is, Pu.2 transcripts) are upregulated in Klf3−/− TER119+ E14.5 fetal liver cells compared to Klf3+/− (HET) and Klf3+/+. Values have been normalized to 18S RNA and the Klf3+/+ sample has been set to 1.0. n = 3 for each genotype. **, P <0.005 compared to both Klf3+/− and Klf3+/+ (Student’s two-tailed t-test). (E) ChIPs were performed on Klf3+/+ and Klf3−/− E14.5 fetal livers (n = 2 or 3 of each genotype per IP). Data are represented as the fold-change enrichment in Klf3−/− cells compared to Klf3+/+. The Fam132a and Klf8 promoters have been included as positive controls while Serpina9, Gapdh, and MyoD are negative control regions. *, P <0.05 compared to Gapdh (Student’s one-tailed t-test). In (D and E), error bars represent standard error of the mean.
Searches using the RepeatMasker program showed that exon 2b lies within a 343 bp long terminal repeat (LTR) element, named ORR1A0, belonging to the MakLR (mammalian apparent LTR retrotransposon) family (Figure 2B) [52]. The ORR1A0 element in the murine Pu.1 locus contains several hallmarks of an euchromatic core promoter including a TATA box at −30, an initiator sequence (5′-TCAGTY-3′) at the TSS and a downstream promoter element around +30 [53]. In addition, it contains several motifs fitting the KLF DNA-binding consensus 5′-NCN CNC CCN-3′ (Figure 2B).

In order to verify that this novel Pu.2 transcript is de-repressed in erythroid cells lacking KLF3, we performed real-time RT-PCR on Klf3+/+, Klf3+/- and Klf3−/− Ter119+ E14.5 fetal liver RNA using a forward primer specific for exon 2b and a reverse primer targeting exon 3 of Pu.1. Indeed, significant up-regulation of the Pu.2 transcript (>140-fold) was observed in Klf3−/− compared to Klf3+/+ and Klf3+/- samples (Figure 2D). While this transcript was not amplified from wild-type tissue by 5′ RACE (Figure 2A), we detected low amounts of it in wild-type and Klf3−/− mice, we observed marked upregulation of this chimeric transcript in erythroid organs (spleen and bone marrow) (Additional file 1: Figure S1A). In contrast, canonical Pu.1 mRNA was unaltered in these and other tissues examined (Additional file 1: Figure S1B).

KLF3 can repress transcription by recruiting CTBP, a co-repressor that silences genes through a number of different histone-modifying enzymes. We therefore analyzed a series of histone marks around the Pu.2 ORR1A0 promoter in Klf3−/− compared to Klf3+/+ E14.5 fetal liver cells (Figure 2E). In particular, we observed a marked increase specifically of histone 3 lysine 4 tri-methylation (H3K4me3) in Klf3−/− cells at the ORR1A0 promoter (approximately 12-fold). We found that this mark was also increased at the promoters of previously validated KLF3 target genes such as Klf8 [50] and Fam132a/adipolin [49,54] (Figure 2E). H3K4me3 is a mark typically found at actively transcribed promoters [55]. Moreover, loss of this mark, rather than the acquisition of repressive modifications, has been reported during the developmental silencing of retrotelemers [18]. In addition, the Pu.2 promoter displayed only a moderate level of H3K9me3 in wild-type cells (Additional file 2: Figure S2). This was not appreciably altered in Klf3−/− cells, suggesting that H3K9 tri-methylation is not the primary mechanism through which KLF3 silences transcription at this locus. Together, these results indicate that the ORR1A0 element is a functional, alternative promoter for the Pu.1 gene in erythroid cells and is highly de-repressed in the absence of KLF3.

KLF1 and KLF3 can bind to the CACCC-like boxes in the ORR1A0 LTR and activate and repress transcription, respectively

We next examined by electrophoretic mobility shift assay (EMSA) whether one or more of the four 5′-NCN CNC CCN-3′ sites in the ORR1A0 promoter are recognized by KLF3. Indeed, KLF3 was found to bind strongly to the two sites most distal to the TSS and weakly to the third CACCC-box (Figure 3A, B). We next assessed whether the related family member KLF1 also binds to these sites. KLF1 is highly expressed in erythroid cells and has a similar DNA-binding specificity to KLF3, such that the two proteins co-regulate overlapping genes in vivo [46,49]. We found that like KLF3, KLF1 binds to the two 5′ most sites (Figure 3C). No detectable binding was observed for the CACCC boxes closest to the TSS (Figure 3D).

Having established that both KLF1 and KLF3 can bind to motifs present in the ORR1A0 element, we next assessed whether they can functionally regulate this promoter in cellular assays. To do this, we cloned the ORR1A0 promoter upstream of a Firefly luciferase reporter gene in the pGL4.10[luc2] vector. This was then co-transfected together with increasing amounts of KLF1 in SL-2 cells, a cell line that is often used to examine KLF function due to minimal background CACCC-binding activity [45]. We found that KLF1 strongly activates expression from the ORR1A0 promoter but has little effect on empty pGL4.10[luc2] vector (Figure 4A). By titrating increasing dosage of KLF3 we found that it counters the activity of KLF1 at the ORR1A0 promoter and represses expression (Figure 4B).

To investigate whether KLF1 indeed drives transcription of these chimeric Pu.2 transcripts from the ORR1A0 element in vivo, we employed a KLF1-inducible erythroid cell line known as B1.6 [56]. These cells were derived from Klf1−/− fetal liver and have been rescued with a transgene encoding a tamoxifen-inducible KLF1-ER (estrogen receptor) fusion protein. Upon addition of tamoxifen, KLF1-ER is activated and drives expression of KLF1 target genes to induce hemoglobinization and erythroid differentiation. KLF3 protein has not been detected in these cells, although Klf3 mRNA is induced after KLF1-ER activation [50].

Using real-time RT-PCR we observed a low level of Pu.2 mRNA in untreated B1.6 cells; however, tamoxifen induction resulted in a dramatic increase of these transcripts (Figure 4C). Moreover, despite decreased expression of canonical Pu.1 transcripts upon KLF1-ER induction (represented by exon 2/exon 3, Figure 4D), the total level of Pu.1 plus Pu.2 transcripts increased (represented by the exon 3/exon 4 junction, Figure 4E), albeit not significantly. This suggests that the chimeric transcripts contribute substantially to the total Pu.1 plus
Pu.2 mRNA pool in induced B1.6 cells. In addition, the induction of Pu.2 expression was rapid following tamoxifen addition (within 2 hours) and occurred in the presence of the translation inhibitor cycloheximide (Additional file 3: Figure S3A, B). Taken together, these data suggest that KLF1 directly activates Pu.2 transcription from the endogenous ORR1A0 promoter in erythroid cells in the absence of KLF3.

Widespread de-repression of chimeric transcripts from ORR1A0 elements in the absence of KLF3

A RepeatMasker survey revealed that there are approximately 2,130 ORR1A0 integrants in the mouse genome. The consensus sequence of ORR1A0 found in Repbase contains all of the core promoter sequences shown in Figure 2B as well as the four 5’-NCN CNC CCN-3’ motifs. In addition, there is little divergence between individual ORR1A0 elements with elements generally sharing greater than 97% sequence identity to the consensus [57]. We therefore hypothesized that KLF3 might play a broader role in silencing aberrant transcription from ORR1A0 LTRs.

To investigate this, we performed RNA-Seq on triplicate samples of Klf3+/+ and Klf3−/− E14.5 TER119+ fetal liver cells. In total, 1,025 genes were found to be significantly deregulated (FDR <0.05) in the absence of KLF3.
The majority of these (76.7%) were upregulated in Klf3−/− cells, concordant with the view from previous studies that KLF3 is predominantly a transcriptional repressor [49]. Importantly, previously validated KLF3 targets also displayed significant upregulation by RNA-Seq including Klf8 (108-fold), Lgals3 (33-fold), Fam132a/adipolin (7.5-fold), Hba-x (2.3-fold), and Hbb-y (1.8-fold) [49,50,54,58].

We next assessed whether the ORR1A0 LTR, and related ORR1A0-int, elements were enriched among the list of KLF3 target genes. We found that of the 786 significantly upregulated genes, 166 of these (21.1%) contained one or more ORR1A0 LTRs. In contrast, of the 239 downregulated genes, there was only one instance of an overlap with an ORR1A0 element (0.004%). Similarly, ORR1A0-int elements, which are typically flanked by ORR1A0 LTRs, were found in 96 upregulated genes (12.2%) and only in a single case of a downregulated gene. These results illustrate a clear enrichment of the ORR1A0 and ORR1A0-int retroelements specifically within genes that are normally repressed by KLF3 in erythroid cells.

Because of sequence conservation between ORR1A0 elements and difficulties associated with unambiguously assigning them to specific genomic loci, we instead looked for evidence of splicing between ORR1A0 transcripts and downstream genic exons. To do this, we confined our analysis to annotated genes which displayed differential isoform expression in Klf3−/− cells (greater than 10-fold upregulated compared to Klf3+/+). We identified 70 such genes (Additional file 5: Table S2). Of these, 34.3% contained transcribed ORR1A0 elements, and almost half of these (41.7%) were spliced to genic exons and a further 16.7% showed splicing to unannotated exons.

By real-time RT-PCR, we validated these results for a selection of candidate target genes. Using forward primers specific for the ORR1A0 exon and reverse primers specific for downstream genic exons, we observed striking upregulation for all three genes tested.
(Znrf2, Brca2, and Pqlc3) in E14.5 TER119+ cells lacking KLF3 (Figure 5A, C, E), mirroring our previous result for Pu.2 transcripts (Figure 2D). In addition, expression of all of these chimeric mRNAs increased considerably upon tamoxifen induction of B1.6 cells (Figure 5B, D, F). In these cells, their upregulation was rapid (Additional file 3: Figure S3C, E) and occurred in the presence of cycloheximide (Additional file 3: Figure S3D, F), suggesting that like Pu.2, their transcription is also directly driven by KLF1. Lastly, it should be noted that definitively mapping repetitive reads to their correct genomic loci is difficult and thus the RNA-Seq analysis is not anticipated to give an exhaustive list of genes for which ORR1A0 exons are spliced to downstream exons. Indeed, by targeted real-time RT-PCR we assessed a further five candidate genes for which the RNA-Seq analysis had not

![Image](http://genomebiology.com/2014/15/4/R58)

Figure 5 KLF1 activates while KLF3 represses chimeric transcripts from ORR1A0 LTRs in erythroid cells. RNA from Klf3+/+ (WT), Klf3+/− (HET), and Klf3−/− (KO) TER119+ E14.5 fetal liver cells (A, C, E) and from untreated and tamoxifen-treated KLF1-ER inducible B1.6 cells (B, D, F) was analyzed by quantitative real-time RT-PCR using forward primers which recognize the ORR1A0 exon and reverse primers specific for downstream exons of the Znrf2 (A, B), Brca2 (C, D) and Pqlc3 (E, F) genes. All values have been normalized to 18S rRNA levels and WT (A, C, E) and untreated (B, D, F) samples have been set to 1.0. Error bars represent standard error of the mean and n = 3 for each genotype or condition. *, P < 0.05 (Student’s two-tailed t-test) compared to both Klf3+/+ and Klf3+/− (A, C, E) and compared to untreated cells (B, D, F).
Figure 6 (See legend on next page.)
called splicing events (Cd59b, Tmx4, Bsw2, Cpe, and Tcf5). In each case, we found that in Klf3−/− cells, the ORR1A0 exon is spliced and the resulting chimeric transcripts are markedly upregulated compared to Klf3+/+ and Klf3+/− cells (Additional file 7: Figure S5A-C).

De-repressed transcription from ORR1A0 elements was found to affect local gene expression in a number of ways, shown in Figure 6 and Additional file 7: Figure S5. In many instances, ORR1A0 LTRs reside within the body of the gene and the new transcripts are spliced to downstream genic exons. This is the case for Pu.1, Thsd7b, Znrf2, and Brc2 (Figure 6A, B, Additional file 7: Figure S5A, B). In addition, ORR1A0 LTRs upstream of genes also act as novel transcriptional start sites for such chimeric transcripts, as is the case for Pqlc3 (Figure 6C).

We also observed spliced transcripts emanating from ORR1A0 elements in un-annotated regions (Figure 6D) and also detected novel transcripts antisense to known genes (Additional file 7: Figure S5C). Lastly, in several cases we observed significantly de-repressed transcription from and across ORR1A0 elements that did not appear to influence the expression of the surrounding gene, as for Drosha (Additional file 7: Figure S5D). This typically occurred either where two ORR1A0 LTRs exist as a cassette, flanking an ORR1A0-int element (Additional file 7: Figure S5D), or where they are in an antisense direction to a transcribed gene (Additional file 7: Figure S5C).

Several of these chimeric transcripts have previously been reported as ESTs that have typically been detected in genic regions (Additional file 7: Figure S5B). However, some show upregulated expression in KO samples. In (A) and (B), internal ORR1A0 elements are transcribed and spliced to downstream genic exons which show a marked increase in expression in KO samples. In (C), an ORR1A0 element serves as an upstream promoter and transcripts are spliced to genic exons. In (D), an ORR1A0 is transcribed solely in KO cells and is spliced to unannotated exons.

We thus sought to determine whether these chimeric transcripts are in fact translated in vivo using the Pu.1 gene as an example, given the role of this transcription factor as a master regulator of hematopoietic differentiation [59].

The chimeric Pu.2 transcript contains a potential ATG start codon within exon 3 (Figure 2C) and is predicted to encode a truncated isoform (Pu.2) that lacks 88 amino acids at its N-terminus. Since the ETS DNA-binding domain lies at the C-terminus of Pu.1, we anticipated that Pu.2 would retain DNA-binding ability. We cloned and expressed Pu.1 and Pu.2 in COS cells and tested their ability to bind to a radiolabelled probe containing the Pu.1 DNA-binding consensus sequence (5′-GAGGAA-3′) by EMSA. Indeed, Pu.2 is able to bind to DNA and migrates more rapidly than Pu.1 (Figure 7A). Moreover, while Pu.1 is recognized and supershifted by antibodies raised against both the N- and C-terminus of Pu.1, Pu.2 is only supershifted by the antibody specific for the C-terminus (Figure 7A).

To determine whether Pu.2 protein is expressed in vivo, we analyzed nuclear extracts from E14.5 Klf3−/− fetal livers. Extracts from these cells formed bands that co-migrated with both Pu.1 and Pu.2 (Figure 7A). The upper band is supershifted by both antisera while the lower band is only supershifted by the C-terminal antisera, confirming their identities as Pu.1 and Pu.2, respectively (Figure 7B, C). Pu.2 protein was also detected in nuclear extracts from Klf3+/+ fetal liver cells (Figure 7B, C) albeit at a lower level than in Klf3−/− samples. Lastly, we also observed marked induction of Pu.2 protein upon tamoxifen activation of KLF1-ER in B1.6 cells (Figure 7D) consistent with the upregulation of Pu.2 transcripts (Figure 4C). Taken together, these results indicate that Pu.2 is indeed translated in erythroid cells in vivo.

PU.2 can act as a dominant negative protein in erythroid cells

The PU.2 protein lacks the N-terminal activation domain of PU.1, a region that interacts with the general
Figure 7 (See legend on next page.)
transcription factor TFII D [60]. We therefore postulated that PU.2 might not function as a transcriptional activator and might antagonize the activity of PU.1 at its target genes. To investigate this, we first conducted reporter assays using the promoter of a previously characterized PU.1 target gene, CLECSA [61]. We found that while PU.1 robustly activated expression, PU.2 repressed this promoter in a dose-dependent manner (Additional file 8: Figure S6A, B).

We next sought to examine the possible dominant negative activity of PU.2 in a hematopoietic system. To do this, we ectopically expressed PU.2, with or without PU.1, in human K562 cells and derived stable clones (Figure 8B, C). Forced expression of PU.1 has previously been shown to promote monocytic differentiation of these cells while inhibiting erythroid maturation [62]. Strikingly, we found that expression of PU.2, both by itself and when co-expressed with PU.1, caused spontaneous erythroid differentiation of these cells in the absence of any chemical-inducing agents (Figure 8A).

Microarray analysis and real-time RT-PCR validation of these cell lines confirmed the upregulation of multiple erythroid genes including the globins, ALAS2, and erythroblast membrane-associated protein (ERMAP) (Additional file 9: Table S3, Figure 8D-G). These results suggest that the LTR-driven PU.2 protein that is upregulated in the absence of KLF3 can oppose the normal function of PU.1 and promote erythroid differentiation.

Discussion
It has recently been shown that TEs frequently act as promoters of genic transcription and are dynamically transcribed during ontogeny [17,18,20,22]. Typically, the majority of retroelement silencing occurs early in gestation [28] and numerous studies have demonstrated the role of histone- and DNA-modifying enzymes in this process [27,30-34,36]. KRA B domain zinc finger proteins, which interact with the co-repressor TRIM28, have been proposed to play a role in the specific recognition and repression of distinct retroelement families [31,35,37,39,40]. This has indeed shown to be the case for ZFP809 in embryonic stem cells [38]. Other DNA-binding zinc finger proteins, including REX1/ZFP42, have also been implicated, but their mechanism of action remains unclear and they appear to affect multiple, unrelated retroviral families [63].

Here we show that the zinc finger protein KLF3, which lacks a KRA B domain, is required for the silencing of transcription from ORR1A0 elements of the MalR family. In the absence of KLF3, there is a pronounced increase in chimeric transcripts generated from these LTRs. The phenomenon of deregulated chimeric transcription has previously been observed upon ablation of epigenetic modifiers such as SETDB1 and LSD1 in embryonic stem cells [27,34]. For example, in cells lacking SETDB1, 15% of de-repressed genes arise due to failed silencing of promoter-proximal ERVs and half of these genes exhibit chimeric transcripts [27]. However, in both of these cases the effects described were more global than we observe for KLF3 and covered diverse retroelement families.

Silencing of the ORR1A0 LTR by KLF3 appears to occur largely independently of the TRIM28/SETDB1 pathway and DNA methylation. Analysis of over 23,000 TRIM28 binding sites in ES cells compiled from two studies [40,64] revealed that only eight lie within 100 bp of an ORR1A0 element. This may partially reflect the difficulties of detecting ChIP peaks that extend beyond the boundaries of repeat elements. However, a separate examination of regions of TRIM28-dependent H3K9 tri-methylation, which typically spread beyond repeat borders, revealed that only 62 of the 2,140 ORR1A0 LTRs (that is, 2.9%) lie within 100 bp [40]. Similarly, analysis of the DNA methylation status of hematopoietic stem cells and erythroblasts revealed that only 1.0% (22) and 0.3% (7) of ORR1A0s, respectively, lie proximal to regions of DNA-methylation as determined by MBD-Seq [65].

From a number of Klf3−/− tissues examined, the upregulation of ORR1A0 transcription appeared to be restricted to erythroid cells. This is of particular interest given that most retroelement silencing has been demonstrated in embryonic stem and germ cells [27,30,31,33]. This suggests that KLF3 plays an active role in somatic repression of retroviral transcription, both in fetal and adult erythroid tissues. Consistent with its erythroid restricted profile, we found that the master erythroid
Forced expression of PU.2 in K562 cells induces spontaneous erythroid differentiation. (A) Cell pellets of K562 lines infected with pMSCVpuro-Pu.1, pMSCVhyg-Pu.2, pMSCVpuro-Pu.1 plus pMSCVhyg-Pu.2, and empty vector (pMSCVpuro plus pMSCVhyg). Hemoglobinization (signified by red hue) is apparent in the lines expressing PU.2 and PU.1 plus PU.2. (B-G) Total RNA was extracted from duplicate lines and was subjected to quantitative real-time RT-PCR analysis using primers specific for: (B) the 5′ end (exon 2/3) of the murine Pu.1 gene (recognizes Pu.1 only and not Pu.2); (C) the 3′ end (exon 3/4) of Pu.1/Pu.2 (recognizes both Pu.1 and Pu.2); (D) ALAS2; (E) ζ-globin (HBZ); (F) ERMAP; and (G) δ-globin (HBD). In (B and C) the duplicate lines have been shown separately while in (D-G) the average of the two lines has been determined. Levels have been normalized to 18S rRNA and set to 1.0 for the lowest samples. Error bars represent standard error of the mean. *, P < 0.05 (Student’s two-tailed t-test) compared to empty vector.
regulator KLF1 drives expression of ORR1A0-originating transcripts in the absence of KLF3.

The specific recognition of the ORR1A0 element by KLF3 and KLF1 appears to involve two 5’-CACNCCC-3’ boxes upstream of the TSS (Figures 2B and 3). The similar DNA-binding specificities of KLF1 and KLF3 have previously been noted and indeed, approximately 50% of KLF3 target genes in erythroid cells are also regulated by KLF1 [49]. Interestingly, the ORR1A0-related LTR ORR1A1 lacks one of the 5’-CACNCCC-3’ boxes and additionally lacks the TATA box. Despite being 97% homologous with ORR1A0 and occurring at twice the frequency in the mouse genome, we did not observe enrichment of ORR1A1 in KLF3 repressed transcripts, alluding to the functional importance of these two promoter motifs and the extraordinary specificity of KLF3 for the ORR1A0 LTR.

Moreover, although the ORR1 retroelement family is abundantly represented throughout diverse rodent species, the ORR1A0 LTR is specifically only found in the mouse. It is interesting to note that the DNA-binding domain of KLF3 shows complete sequence conservation between Mus musculus, Rattus norvegicus, and Homo sapiens. This suggests that in the mouse, the DNA-binding specificity of KLF3 has not altered in response to the emergence of the ORR1A0 retroelement. Rather, it appears that KLF3-mediated repression of ORR1A0 occurred intrinsically from the initial appearance of the retroelement, and in this context, the spread of the ORR1A0 LTR may have been tolerated without deleterious impacts.

While KLF3 appears to efficiently silence ORR1A0 transcription in spleen and bone marrow cells, it is possible that at particular stages of development or cellular maturation, KLF1 predominates and these chimeric transcripts are expressed at higher levels, as observed in the B1.6 erythroblast line. These chimeric transcripts potentially encode functional protein isoforms and indeed, in the case of the Pu.1 locus, we have detected a truncated isoform expressed in fetal liver. PU.2 counters the normal activity of PU.1 and instead promotes erythroid differentiation when ectopically expressed in myeloid leukemic K562 cells (Figure 8). This is consistent with the role of its transcriptional activator, KLF1, in driving terminal erythroid differentiation [44]. From this study, we cannot discount the possibility that the chimeric transcripts driven by KLF1 may be biologically functional in some circumstances. Indeed, host expatation of TEs by exonization or through the remodelling of expression programs is a phenomenon of which multiple instances have been described [5,13,14,19]. Incidentally, although the consensus sequence of the ORR1A0 exon contains a number of short open reading frames (see Figure 2B), each ATG is ultimately succeeded by an in-frame stop codon. Thus ORR1A0 promoters are predicted to drive expression of either full-length endogenous proteins (where a canonical translation start site lies in a downstream, spliced exon), or N-terminal truncated protein isoforms, in cases where internal ATGs are suitable start sites of translation, as for PU.2.

The importance of appropriate retrotransposon control is apparent from numerous examples in which dysregulation results in disease [25,26]. It is unclear as to the extent to which the dysregulation of ORR1A0 transcription affects the physiology of the Klf3 null mice. However, these animals do display an erythroid phenotype, with impaired maturation, reticulocytosis, increased Howell-Jolly bodies and decreased lifespan of erythrocytes [49]. These defects have not been attributed to any single gene and it is more likely that they arise as a complicated result of multiple defects including the widespread expression of aberrant, chimeric transcripts that we have presented here.

Conclusions

In summary, these findings emphasize the non-redundant role that KLF3 plays in preventing widespread, promiscuous transcription specifically from the ORR1A0 LTR. We suggest that KLF3 is likely one of a patchwork of zinc finger proteins including ZFP809 that together collaborate to silence the diverse collection of TEs that constitute such a large fraction of vertebrate genomes.

Materials and methods

Klf3<sup>−/−</sup> mice

Generation and maintenance of the Klf3<sup>−/−</sup> mouse line has been described previously [66]. Ethical approval for animal use was obtained from the appropriate Animal Care and Ethics Committees (University of Sydney, approval number L02/7-2009/3/5079; and University of New South Wales, approval number 09/128A). Genomic DNA was extracted from tail snips using DirectPCR Lysis Reagent (Viagen Biotech Inc, Los Angeles, CA, USA) as per the manufacturer’s instructions and genotyping was performed as described previously [66].

Sorting of TER119<sup>+</sup> fetal liver cells

TER119<sup>+</sup> cells were sorted from whole fetal livers (E13.5 or E14.5) using anti-TER119 Microbeads with MS or LS columns (Miltenyi Biotec Australia, North Ryde, NSW, Australia) as per the manufacturer’s protocol. After eluting, cells were centrifuged at 300 g for 10 min at 4°C and RNA was extracted from the cell pellet as described below.

RNA extraction and quantitative real-time RT-PCR

RNA was extracted, purified and subjected to DNase treatment as described previously [45,49]. Extracted RNA was then used as a template for cDNA synthesis.
using the SuperScript® VILo™ cDNA Synthesis Kit (Invitrogen, Carlsbad, CA, USA) as instructed by the supplier. Quantitative real-time RT-PCR reactions were set up as described previously [45,49] but using FastStart Universal SYBR Green Master (ROX) (Roche Diagnostics Australia Pty Ltd, Castle Hill, NSW, Australia) or Power SYBR® Green PCR Master Mix (Life Technologies, Gladesville, NSW, Australia). Reactions were run using the Applied Biosystems 7500 Fast Real-Time PCR System (Life Technologies) and data were analyzed using 7500 Software v2.0.4 (Life Technologies). Relative quantification was calculated using 18S rRNA levels and standard curves derived from serial dilutions of amplicon as described previously [45].

Real-time RT-PCR primers
Gene-specific primer pairs were designed using the Primer-BLAST tool [67] or PrimerExpress v3.0 (Applied Biosystems, Foster City, CA, USA) software to cross exon boundaries where possible. Primers were included in reactions at a concentration of 400 nM. The sequences of forward and reverse primers for each gene are: 18S, 5'-CACGGCCGGTTACGTGGAAC-3' and 5'-AGAGGGACGCGCAACAA-3'; Pu.1 exons 2/3, 5'-CCTCTCTGTGGCCAGCGATGA-3' and 5'-GAACGTGCTTACGCTGGGGA-3'; Pu.1 exons 3/4, 5'-GAGCTGGATGCTCAGTGCT-3' and 5'-GGGCACTTTTTCT-3'; Tcfl5 exons 2b/3, 5'-GCCATGCTGCCTGGATA-3' and 5'-GGGTACAAGGTTTGAT-3'; Klf8 promoter 1a and 5'-AGCAGTCGGAAACTGGGAAG-3'; Serpina9 exons 2/3, 5'-TGTGCTGGACCTGGTTTGTA-3'; and 5'-TGTGGGCTTTGCAGCTCTTACTCAGATTCCT-3'; 5'-GGAG-3' and 5'-CTCTCAA-3'.

5' RACE (Rapid Amplification of cDNA Ends)
600 ng total RNA from TER119+ fetal liver cells obtained from Klf3+/+ and Klf3−/− embryos (E13.5) was used as a template for first strand cDNA synthesis using the SMARTer™ RACE cDNA Amplification Kit (Clontech, Mountain View, CA, USA). The RACE PCR was performed as directed by the supplier but using 0.05 unit/μL REDTaq DNA Polymerase (Sigma Aldrich, St Louis, MO, USA). Thermal cycling settings were 24 cycles of 94°C/30 s, 64°C/30 s, and 72°C/1 min. Amplified products were resolved by electrophoresis through a 1.2% agarose gel and were purified using the Wizard SV Gel and PCR Clean-up System (Promega Corporation, Madison, WI, USA) as per the manufacturer's manual. Nested RACE PCR was then performed as described above with slight alterations to the thermal cycler parameters: 24 cycles of 94°C/30 s, 60°C/30 s, and 72°C/1 min. First round and nested RACE PCR primers are listed respectively: reverse primer targeting exon 4 of Pu.1, 5'-GTGGGGCTGGGGGACAAAGGTGGTGAT-3'; reverse primer targeting exon 3 of Pu.1, 5'-GCTGTAGCTGCTGGGGGACAAAGGTGGTGAT-3'; reverse primer targeting exon 2 of Pu.1, 5'-TGAGTCCTGCTGGGGGACAAAGGTGGTGAT-3'; reverse primer targeting exon 1 of Pu.1, 5'-GGGACAAAGGTGGTGAT-3'; reverse primer targeting exon 2 of Klf3, 5'-GGGAACTGGTACAGGCGAA-3'; reverse primer targeting exon 3 of Klf3, 5'-GCCACCTCTCCCGTACG-3'; reverse primer targeting exon 4 of Klf3, 5'-ACCGTGGGGC-3'.

Chromatin immunoprecipitation (ChIP)
ChIP was performed with slight modifications based on Schmidt et al. [68]. Briefly, one cross-linked E14.5 liver was used per IP with antibodies as follows: IgG, 5 μg sc-2027 (Santa Cruz Biotechnology, Santa Cruz, CA, USA); anti-H3K4me3, 5 μg C42D8 (Cell Signaling Technology, Danvers, MA, USA), or 5 μg ab12209 (abcam, Cambridge, MA, USA); anti-H3K9me3, 5 μg ab8898 (abcam); anti-H3K27me3, 3 μg C36B11 (Cell Signaling Technology) or 5 μg ab6002 (abcam); anti-H3K27ac, 0.5 μg #4353 (Cell Signaling Technology) or 5 μg ab4729 (abcam). Real-time PCR quantification of chromatin pull-down was performed as described above and amounts were normalized to the level of input material prior to immunoprecipitation. ChIP primer sequences have been described previously for Klf8 promoter 1a and Fam132a [49,50]. Other primers used are as follows: Pu.2 ORR1A0 -250 bp, 5'-GAGCTCTTCTGCTCCCCATCT-3' and 5'-CTGCTCGCTCCTCACG-3'; Pu.2 ORR1A0 -120 bp, 5'-GGCGGACATGCTGGTGG-3' and 5'-GGGGACCAAGTGCCCTGC-3'; Tcfβ5, 5'-GACATGGCTCGCTGGTGG-3' and 5'-GGCGGACATGCTGGTGG-3'; H3K4me3, 5'-AGCTCTCTGCCTCTCTCTCT-3' and 5'-GGCGGACATGCTGGTGG-3'.

Bzw2, 5'-GCCATGCTGCCTGGAT-3'; 5'-CTGCCACACGATGATAATGG-3'; 5'-CTGACCGTACCGTACG-3'; 5'-CGCTCTCCTGAGCTTC-3'; 5'-GAGTCTGCTCTCCACG-3'; 5'-TGGACCCCACTACGCTATC-3'; and 5'-ACCCCGTGTTTATGGAAGCTGGGGGACAAAGGTGGTGAT-3'.

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Vectors and cloning of PU.1 and PU.2

The vectors pPac and pPac-Klf1 were provided by Menie Merika and Stuart Orkin (Harvard Medical School, Boston, MA, USA). The plasmid pPac-Klf3 and the mammalian expression vectors pMT3 and pMT3-Klf3 were gifts from Dr José Perdomo (St George Clinical School, Sydney, Australia). The mammalian expression vector pSG5-Klf1 was supplied by James Bieker (Mount Sinai School of Medicine, New York, NY, USA). Firefly and Renilla luciferase vectors used were pGL4.10[luc2], and pGL4.23[luc2/minP] and pGLA.74[hRluc/TK] as transfection controls (Promega Corporation). The vector pEF-IREs-puro5 (pEF1α) was kindly provided by Dr Daniel Peet (University of Adelaide, Adelaide, Australia). The Renilla luciferase reporter construct pLightSwitch-Clec5a, containing approximately 1.1 kb of the human CLEC5A promoter, was purchased from Switchgear Genomics (Menlo Park, CA, USA). The Pu.2 ORR1A0 promoter (−140 to +23) was synthesized by GeneArt® (Life Technologies) and subcloned into KpnI/Xhol pGLA.10[luc2] to create pGLA.10-ORR1A0.

Full length Pu.1 and Pu.2 were cloned from the cDNA generated during the 5′ RACE described above. PCRs were set up using Phusion High-Fidelity DNA Polymerase (Finzymes OY, Espoo, Finland) as directed by the supplier. Primers used in the first round of amplification included: 10X Universal Primer A Mix (UPM) supplied from SMARTer™ RACE cDNA Amplification Kit (Clontech) as a forward primer, and a reverse primer specific for exon 5 of Pu.1; 5′-TCCGGGCGCCGGCAGCGGT-TAATGCTAT-3′. Thermal cycler parameters were 98°C/30 s, followed by 25 cycles of 98°C/10 s, 59°C/30 s, and 72°C/1 min, and a final cycle of 72°C/5 min. Amplified Pu.1 and Pu.2 products were resolved by electrophoresis, purified as described above and subjected to nested PCR using forward primers covering the start points of transcription for PU.1 (5′-ATTACTCGAGGGTCGTGGATGTACAGGGCTGGAAA-3′) and PU.2 (5′-ATTACTCGAGGGCCACCATGGAGCTGGAACAGATGCAC-3′) together with the common reverse primer 5′-TAATGCTCAACACATTGGAGCGTGAACAGATGCAC-3′ (which includes the Pu.1 stop codon in exon 5). Parameter settings were 98°C/30 s, followed by 25 cycles of 98°C/10 s, 69°C/30 s, and 72°C/1 min, and a final cycle of 72°C/5 min. The fragments were subsequently cloned into XhoI/EcoRI pMT3 to form pMT3-Pu.1 and pMT3-Pu.2, respectively. Pu.1 and Pu.2 were then subcloned into XhoI/EcoRI pEF1α to generate pEF1α-Pu.1 and pEF1α-Pu.2. Similarly, Pu.1 and Pu.2 were cloned into BglII/HpaI pMSCVpuro and pMSCVhyg (Clontech), respectively, using the forward primers 5′-ATTAAAGATCTCAGCTCGATGTTATACAGCCGTGCAA-3′ and the reverse primer 5′-TAATGTTAACACGCTTGCGGTCTCTGCCGGCGATCA-3′.

Cell culture

COS cells were cultured as described previously [45], K562 and HL60 cells were maintained similarly but in RPMI 1640 culture medium (Gibco-BRL Life Technologies, Grand Island, NY, USA). Culture conditions for B1.6 erythroblast cells have been described elsewhere [56]. B1.6 cells were induced with tamoxifen as described previously [45] and, with the exception of the time-course and cycloheximide experiments, were harvested for RNA or nuclear extracts after 48 h. SL2 cells were cultured in Schneider’s Drosophila medium (Gibco-BRL Life Technologies, Grand Island, NY, USA) supplemented with 10% heat-inactivated FCS and 1% penicillin/streptomycin/glutamine solution at 24°C.

Transfections and retroviral infections

COS cells were transfected with 5 μg pMT3-Pu.1, pMT3-Pu.2 or pMT3 empty using FuGENE6 (Roche Diagnostics) as instructed by the supplier. Cells were harvested after 48 h for nuclear extracts. For retroviral infection of K562 cells, Phoenix A packaging cells were transfected with 12 μg total vector DNA (12 μg pMSCVpuro-Pu.1, 12 μg pMSCVhyg-Pu.2, 6 μg pMSCVpuro plus 6 μg pMSCVhyg, or 6 μg pMSCVpuro-Pu.1 plus 6 μg pMSCVhyg-Pu.2) using Lipofectamine 2000 (Life Technologies) according to the manufacturer’s instructions. Target K562 cells were seeded at 1 × 10⁶ cells/mL in 6-well Plates 24 h prior to infection. Forty-eight hours following infection of packaging cells, virus-containing media (VCM) were collected and passed through a 0.45 μm low protein binding filter. Non-tissue culture treated 6-well dishes were then coated with retroNectin® (Clontech) as instructed by the manufacturer. Half the volume of VCM was then added to the 6-well dishes for 30 min at room temperature. The target K562 cells were resuspended in the remaining VCM with the addition of protamine to a final concentration of 8 μg/mL. After 30 min, the VCM was aspirated from the coated wells and was seeded with the K562 cells. The plates were subsequently centrifuged at 400 g for 1.5 h at 30°C and then incubated at 37°C with 5% CO₂ overnight. After 12 h, VCM was collected from the packaging cells for a second round of infection as described above. Eight hours after the second spinoculation, the K562 cells were replated with additional VCM and incubated for 48 h before replating in RPMI 1640 culture medium with 10% (v/v) heat-inactivated FCS (Gibco) and 1% (v/v) penicillin/streptomycin/glutamine solution (Gibco). After another 48 h, cells were subjected to antibiotic selection and maintained in 1 μg/mL puromycin dihydrochloride (Sigma) and 200 μg/mL hygromycin B (Life Technologies) as appropriate.

Separately, K562 cells were also transfected with pEF1α-Pu.2 or pEF1α empty using the Gene Pulser Xcell electroporation system (Bio-Rad, Hercules, CA, USA).
Briefly, 10⁶ cells and 20 μg plasmid were resuspended in Dulbecco’s modified eagle medium (DMEM) low glucose (LG) (Gibco-BRL Life Technologies) without serum and in a total volume of 400 μL. Cells were electroporated at 200 V, 950 μF and subsequently cultured and maintained in 2 μg/mL puromycin dihydrochloride to generate monoclonal lines.

Nuclear extracts and electrophoretic mobility shift assays (EMSAs)

Nuclear extracts were obtained and EMSAs were performed as described previously [46]. The radiolabelled probe containing the PU.1 consensus binding site comprises sense and antisense oligonucleotides for the sequence 5′-GCTCGAGGACTTTCTCTTTCAGTGCC-3′ as described elsewhere [69]. ORR1A0 double stranded CACCC probe sequences are as follows: CACCC#1, 5′-ATGGTGCCACTCTGCTTGC-3′; CACCC#2, 5′-CA GTGACACACCCATTCCA-3′; CACCC#3, 5′-TTATAC CCACACCCACAGTG-3′; CACCC#4, 5′-TAAGATCC TCACCTAGTTG-3′. The positive control KLF3 binding site in the Fam132a promoter is ‘Probe C’ from [54]. The antibodies used that were specific for the N-terminus and C-terminus of PU.1 were 9G7 (Cell Signaling Technology) and T-21 (Santa Cruz Biotechnology), respectively. KLF1- and KLF3-specific antisera have been described previously [46].

Western blotting

Western blots were performed by standard methods. Briefly, nuclear extracts were separated by SDS-PAGE and were electrotransferred to PVDF membrane, which was then blocked with 5% skim milk in 50 mM Tris-HCl (pH 7.4), 150 mM NaCl, and 0.05% Tween 20. PU.2 protein was probed by overnight incubation of membrane in 5% skim milk with 0.2 μg/mL PU.1 antibody (T-21) (Santa Cruz Biotechnology) at 4°C. Detection was achieved using Immobilon Western Chemiluminescent HRP Substrate (Millipore Corporation, Billerica, MA, USA) and subsequently, membranes were stripped in 0.2 M NaOH for 10 min and were probed with β-actin antibody (Sigma).

Reporter assays

SL2 cells were split into 6-well plates at a concentration of 5 × 10⁵/mL and 24 h later were transfected with pPac-Klf1 (0, 50, or 250 ng) and pPac-Klf3 (0, 12.5, 25, 50, or 100 ng) supplemented to equal loads with pPac empty vector, together with 100 ng pGL4.74 [luc2] or pGL4.10-ORR1A0 using FuGene6 (Roche Diagnostics) as instructed by the supplier. In competition assays, the pGL4.10-ORR1A0 vector (and pGL4.10 [luc2]) were driven by co-transfection of 200 ng pPac-Klf1. After 48 h, cells were harvested and lysates analyzed using the Dual-luciferase® Reporter Assay System (Promega Corporation) and a TD20/20 luminometer (Turner Biosystems, Sunnyvale, CA, USA). HEK293 cells were similarly transfected but with 0, 10, 100, and 1,000 ng pEFlα-Pu.1 (supplemented with pEFlα to a total of 1,000 ng vector) together with 1 μg pLightSwitch (empty vector) or pLightSwitch-Clec5a. As a control, 100 ng pGL4.23 [luc2/minP] Firefly luciferase vector was co-transfected. In competition assays, HEK293 cells were transfected and analyzed as above but with 1 μg pEFlα-Pu.1 together with 0, 10, 100, and 1,000 ng pEFlα-Pu.2.

Microarrays

Microarray data from TER119⁰ E14.5 fetal liver cells from Klf3+/+ and Klf3−/− embryos have previously been described [49]. Monoclonal K562 cell lines stably transfected with pEFlα or pEFlα-Pu.2 (n = 3 each) were harvested for total RNA which was subsequently hybridized to Affymetrix Human Gene 1.0 ST arrays according to the manufacturer’s instructions (Affymetrix, Santa Clara, CA, USA). Hybridization and processing were performed by the Ramaciotti Centre for Gene Function Analysis (University of New South Wales, Sydney, Australia). Data were analyzed using Affymetrix® Expression Console™ software (Affymetrix). Microarray data are available in the Gene Expression Omnibus database [70] under accession number GSE50083.

High throughput RNA-sequencing (RNA-Seq)

RNA was extracted from TER119⁰-sorted liver cells from three Klf3+/+ and three Klf3−/− litter-matched E14.5 embryos (two litters total). Libraries were prepared using 1 μg total RNA using the TruSeq RNA Sample Prep Kit v2 (Illumina, San Diego, CA, USA) according to the manufacturer’s instructions. The six libraries were multiplexed into two lanes using sample specific adapters such that there were three samples per lane. 100 bp paired end reads were sequenced using TruSeq v3 SBS reagents on the Hiseq 2000 (Illumina, San Diego, CA, USA). Library preparation and sequencing were performed by the Ramaciotti Centre, University of New South Wales, New South Wales, Australia. Quality control on the reads was performed using FastQC v0.10.1 available from [71].

Two separate strategies were used for alignment and RNA-seq analysis. The first of these sought to uniquely map reads from repeat elements by using high stringency alignment cut-offs and was primarily employed to visualize chimeric splicing events using Integrative Genomics Viewer [72]. Reads were aligned to the mm10 Mus musculus genome using tophat2 (v2.0.4) using the default settings except for the following -r -40 --segment-length 50 --coverage-search --segment-mismatches 0 -g 50 --genome-read-mismatches 0 --read-mismatches 0 -i
Bioinformatics
The sequence and genomic positions of ORR1A0 LTR elements were determined using the RepeatMasker program [75] in conjunction with the University of California Santa Cruz (UCSC) Genome Browser [76]. Sequence information of retroelement families was obtained using Repbase Update [57,77]. The overlap between differentially expressed loci as measured by RNA-Seq and ORR1A0 and ORR1A0-int elements was established using the intersect command in Bedtools v2.17.0 [78]. Similarly, overlaps (using 100 bp windows) were determined between ORR1A0 elements and: TRIM 28 ChIP peaks (from [40,64]); TRIM28-dependent H3K9me3 peaks (from [40]), and regions of DNA methylation in hematopoietic stem cells and erythroblasts (from [65]). Gene Expression Omnibus accession numbers for the data sets analyzed are GSM1032198, GSM773067, GSM1032190, and GSE38354, respectively.

Additional files
Description of additional files
The following additional data are available with the online version of this paper. Additional file 1: Figure S1 shows that the chimeric Pu.1 transcript (Pu.2) is predominantly upregulated in erythroid tissues in the absence of KLF3. Additional file 2: Figure S2 shows levels of H3K9 tri-methylation at the Pu.2 promoter in Klf3−/− and wild-type E14.5 fetal liver cells. Additional file 3: Figure S3 shows that Pu.2 and other ORR1A0 chimeric transcripts are rapidly activated by KLF1 and in the presence of cycloheximide, suggesting that they are direct targets. Additional file 4: Figure S4 provides validation of genes to which the ORR1A0 exon is spliced in Klf3−/− E14.5 TER119+ fetal liver cells. Additional file 7: Figure S5 gives further examples of de-repressed ORR1A0 transcripts in the absence of KLF3. Additional file 8: Figure S6 contains reporter assay data demonstrating the opposing transcriptional activities of PU.1 and PU.2. Additional file 4: Table S1 shows the list of genes that are significantly, differentially expressed in Klf3−/− TER119+ fetal liver cells by RNA-Seq. Additional file 5: Table S2 shows the list of genes that have significantly de-repressed isoforms. Additional file 9: Table S3 shows microarray results from K562 cell stably expressing PU.2.

Additional file 1: Figure S1. The chimeric Pu.1 transcript (Pu.2) is predominantly upregulated in erythroid tissues in the absence of KLF3. Total RNA was extracted from adult tissue from three Klf3−/− (WT) and three Klf3−/− (KO) mice and analyzed by quantitative real-time RT-PCR using primers specific for the exon 2b/3 junction (A) or exon 2/3 junction (B) of Pu.1. Levels have been normalized to 18S rRNA and the lowest detectable reading in each chart has been set to 1.0. Error bars represent standard error of the mean. *, P<0.05 (Student’s two-tailed t-test) compared to wild-type.

Additional file 2: Figure S2. The relative enrichment of H3K9me3 at the Pu.2 promoter in Klf3−/− and Klf3−/− E14.5 fetal liver cells. ChIP data have been expressed as percentage input for each locus (n = 2 for each IP for Klf3−/− (WT) or Klf3−/− (KO)). Error bars represent standard error of the mean. The Pu.2 promoter shows a moderate level of H3K9 tri-methylation relative to positive (Sepin9) and negative (Gadph) control loci.

Additional file 3: Figure S3. Pu.2 and ORR1A0 chimeric transcripts are induced rapidly by KLF1-ER and in the presence of cycloheximide. RNA from KLF1-ER inducible B1.6 erythroid cells was analyzed by qRT-PCR using primers specific for Pu.2 (A, B), ORR1A0-2nr2 (C, D), and ORR1A0-Bxs/2 (E, F). In (A, C, and E), RNA was harvested at 0, 2, 4, and 24 h following tamoxifen treatment (n = 2). In (B, D, and F), cells were treated with cycloheximide for 30 min prior to tamoxifen addition (or ethanol for untreated), with samples being taken 8 h thereafter (n = 4 for each condition). All values have been normalized to 18S rRNA levels and t = 0 h time points (A, C, and E) and tamoxifen-untreated samples (B, D, and F) have been set to 1.0. Error bars represent standard error of the mean. *, P<0.05 (Student’s one-tailed t-test) compared to t = 0 h (A, C, and E) and tamoxifen-untreated (B, D, and F). NS, not significant.

Additional file 4: Table S1. Genes that are significantly, differentially expressed in Klf3−/− TER119+ fetal liver cells compared to Klf3+/+. RNA-Seq was performed on triplicate samples and differentially expressed genes were determined using a FDR cut-off of 0.05.

Additional file 5: Table S2. The list of annotated genes which have significantly de-repressed isoforms (>10-fold) in Klf3−/− TER119+ fetal liver cells compared to Klf3+/+. Eighty eight differentially expressed isoforms (FDR <0.05) were determined, covering 70 different genes.

Additional file 6: Figure S4. Confirmation of further ORR1A0 splicing events in Klf3−/− erythroid cells that were not detected by RNA-Seq analysis. RNA from Klf3−/− (WT), Klf3−/− (HE1), and Klf3−/− (K0) TER119+ E14.5 fetal liver cells were analyzed by real-time RT-PCR using forward primers specific for ORR1A0 and reverse primers recognizing downstream exons of the Cd55b (A), Tm8sf (B), and Bsw2, Cpe, and Td5 (C) genes. (A and B) Values have been normalized to 18S rRNA levels and WT samples have been set to 1.0. Error bars represent standard error of the mean (n = 2 WT, 3 HET, and 3 KO). *, P<0.05 (Student’s two-tailed t-test) compared to both Klf3−/− and Klf3+/+. (C) For these genes, the spliced transcripts were below the level of detection in Klf3−/− cells and thus could not be quantified. RT-PCR products were electrophoresed on a 3% agarose gel and stained with ethidium bromide.

Additional file 7: Figure S5. Further examples of de-repressed ORR1A0 transcripts in the absence of KLF3. As in Figure 6, tracks represent RNA-Seq reads and splicing events for Klf3−/− (WT) and Klf3−/− (KO) E14.5 TER119+ fetal liver cell samples. (A, B) ORR1A0 elements are transcribed and spliced to
downstream exons of Znvrf2 (A) and Brca2 (B), which in turn are expressed at a significantly higher level in KO cells. (C) Spliced transcripts initiating nearly to an ORR1A0 LTR are antisense to an annotated gene (Dxrosa). (D) De-repressed transcription of an ORR1A0/ORR1A0-int cassette does not alter the expression of the surrounding gene (Dxrosa).

Additional file 8: Figure S6. PU.2 opposes the transcriptional activity of PL1. HEK293 cells were transfected with pLightSwitch Rmelii luciferase reporter vector (promoter-less or containing the CLECA5 promoter). In (A), increasing amounts of pEF1α-Pu.1 have been co-transfected, while in (B), a steady amount of pEF1α-Pu.1 has been co-transfected together with increasing doses of pEF1α-Pu.2. In all experiments, pGL2(+)Luc(−) minP Firefly luciferase vector was included as a transfection control and used for normalization. The means of triplicate experiments are shown and error bars represent standard error of the mean. * P<0.05 (Student's two-tailed t-test) compared to pLightSwitch-Clec5a wells transfected with 0 ng pEF1α-Pu.1 (A) or 0 ng pEF1α-Pu.2 (B).

Additional file 9: Table S3. Erythroid genes are upregulated upon forced expression of PU.2 in K562 cells. Microarrays were performed on monoclonal K562 cell lines stably transfected with pEF1α-Pu.2 or pEF1α. A selection of erythroid genes that are upregulated >2-fold in cells expressing PU.2 is shown.

Competing interests
The authors declare that they have no competing interests.

Authors’ contributions
KSM, APWF, JB, and LJN performed experiments and analyzed data. Specifically, APWF, KSM, and LJN performed cell sorting experiments and qRT-PCR. KSM conducted 5’ RACE; histone ChIPs, reporter assays, western blots, and microarrays. APWF and KSM carried out EMSA experiments and generated the stable K562 lines. APWF performed B16 cell work. APWF and JB designed, performed and analyzed the RNA-Seq and conducted bioinformatics comparisons. APWF, MC, and RCM contributed to overall design of the study and directed research. APWF, KSM, JB, MC, and RCM wrote the paper. All authors read and approved the final manuscript.

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