Chromosomal rearrangements are initiating events in acute lymphoblastic leukaemia (ALL). Here using RNA sequencing of 560 ALL cases, we identify rearrangements between MEF2D (myocyte enhancer factor 2D) and five genes (BCL9, CSFIR, DAZAP1, HNRNPUL1 and SS18) in 22 B progenitor ALL (B-ALL) cases with a distinct gene expression profile, the most common of which is MEF2D-BCL9. Examination of an extended cohort of 1,164 B-ALL cases identified 30 cases with MEF2D rearrangements, which include an additional fusion partner, FOXJ2; thus, MEF2D-rearranged cases comprise 5.3% of cases lacking recurring alterations. MEF2D-rearranged ALL is characterized by a distinct immunophenotype, DNA copy number alterations at the rearrangement sites, older diagnosis age and poor outcome. The rearrangements result in enhanced MEF2D transcriptional activity, lymphoid transformation, activation of HDAC9 expression and sensitive to histone deacetylase inhibitor treatment. Thus, MEF2D-rearranged ALL represents a distinct form of high-risk leukaemia, for which new therapeutic approaches should be considered.
**Results**

**RNAseq identifies MEF2D fusions in a novel B-ALL subgroup.** The 560 cases examined by RNAseq enrolled on the Children's Oncology Group (COG) trials included AALL0232 (N = 216), AALL0331 (N = 65), AALL0932 (N = 61), AALL1131 (N = 92), P9906 (N = 1) and adults (age of diagnosis ≥18, N = 122) or children (age of diagnosis <18, N = 3) with B-ALL enroled on various non-COG protocols (Supplementary Data 1).

This analysis identified multiple new recurring targets of rearrangement, including MEF2D, NUTM1 and ZNF384, and new partners of rearrangement with known genes such as PAX5 and IGH (Supplementary Data 1). We identified 22 cases (3.9%) with rearrangements of MEF2D at chromosome 1q21–22, encoding myocyte enhancer factor 2D (Table 1, Fig. 1, Supplementary Data 2 and Supplementary Table 1), representing 15 of 367 (4.1%) childhood cases (up to age 15 years), 4 of 62 (6.5%) adolescent cases (age 16–20 years), 2 of 73 (2.7%) young adults and a single case in 56 older adults. Five MEF2D fusion partners were identified: BCL9 (16 cases), HNRNPUL1 (3 cases), DAZAP1, CSF1R and SS18 (1 case each). Each fusion was confirmed by reverse transcription and PCR (RT–PCR) followed by Sanger sequencing (Fig. 1, Supplementary Fig. 1 and Supplementary Table 2). In each case, the amino terminus of MEF2D was fused in frame with the carboxy-terminal portion encoded by the partner gene.

The most common MEF2D rearrangement was MEF2D-BCL9, which was present in 16 cases. Four isoforms of this fusion were identified, involving exons 5 or 6 of MEF2D (NM_001271629) fused in-frame to exons 9 or 10 of BCL9 (NM_004326; Supplementary Data 2). MEF2D-BCL9 rearrangement was associated with deregulated expression of the C terminus of BCL9 distal to the rearrangement breakpoint (Supplementary Fig. 2). No fusions deregulating expression of full-length BCL9, such as IGH-BCL9 (ref. 5), were identified. Only one isoform of the MEF2D-HNRNPUL1 fusion was identified, involving exon 8 of MEF2D and exon 12 of HNRNPUL1. Expression of the fusion proteins in patient leukaemic cells was confirmed by immunoblotting (Supplementary Fig. 3) and the disruption of MEF2D and MEF2D-BCL9 rearrangements were shown to be present in the majority of leukaemic cells by fluorescence in situ hybridization (FISH), consistent with the fusions being leukaemia-initiating events acquired early in leukaemogenesis (Supplementary Fig. 4). Several patterns of probe hybridization were identified, consistent with diverse mechanisms of rearrangement, including insertion of the fusion into a different chromosome (for example, SJBALL020667), interstitial insertion at chromosome 1q (SJBALL020703) and reciprocal rearrangement (SJBALL020987).

**Genomic alterations of MEF2D-rearranged cases.** MEF2D and BCL9 are both located at 1q21.2–22 and the common pattern of interstitial insertion resulted in these rearrangements being cryptic on cytogenetic analysis. We observed that many rearrangements were accompanied by copy number alterations at the MEF2D and partner gene loci. Ten cases harbouring MEF2D rearrangements had single-nucleotide polymorphism (SNP) microarray data available (Table 1), with eight of ten exhibiting focal gains or loss of DNA at the partner genes (MEF2D, BCL9, DAZAP1 and HNRNPUL1; Supplementary Fig. 5). The genomic structure of the MEF2D-BCL9 rearrangements was confirmed in two cases by whole genome sequencing (Supplementary Fig. 6). Additional genetic alterations identified in MEF2D-rearranged cases included IKZF1 alterations in three cases, including exons 4–8 deletions in two cases and exons 4–7 deletion in one case (resulting in the expression of the dominant negative IK6 isoform). Mutational analysis of RNAseq data showed that 6 of 16 MEF2D-BCL9 cases harboured activating NRAS mutations, although these were frequently subclonal and thus secondary events in leukaemogenesis (Supplementary Data 1 and Supplementary Table 3). Analysis of Ras mutations (NRAS, KRAS, NFI and PTPN11) across the RNAseq cohort revealed a similar prevalence of Ras mutations as other subtypes of ALL previously reported to be enriched for Ras mutations (for example, hyperdiploid and MLL-rearranged ALL (Supplementary Fig. 7)).

**The transcriptional signatures of MEF2D and ZNF384 ALL.** Although RNAseq data were obtained from diverse library preparation (total versus messenger RNA, unstranded versus stranded) and sequencing (75 bp versus 100 bp paired end) methods, reproducible clustering of MEF2D-rearranged cases was observed in principal component analysis and hierarchical clustering (Fig. 2a and Supplementary Figs 8–10), with the exception of one sample harbouring an MEF2D-CSF1R fusion, which clustered with Ph-like ALL cases. Rearrangements of CSF1R, encoding the macrophage colony-stimulating factor receptor, are observed in Ph-like ALL and result in activation of tyrosine kinase signalling pathways inhibited by kinase inhibitors such as imatinib and dasatinib5,6,7. Thus, the clustering of this sample with other Ph-like ALL cases distinct from other MEF2D-rearranged cases was not surprising. Supervised analysis of MEF2D-rearranged cases identified a distinct gene expression
cases were discovered in this study that exhibit a distinct gene (Supplementary Data 6). Taken together, 42 exon 7 of the forkhead transcription factor gene, FOXJ2 of previously identified eight ALL subgroups, several of which stranded and unstranded sequencing methods (Fig. 2b and signature with substantial overlap between cases sequenced using stranded and unstranded sequencing methods (Fig. 2b and Supplementary Data 3–5).

Analysis of outlier gene expression in high-risk childhood ALL previously identified eight ALL subgroups, several of which harboured recurring genetic alterations including rearrangement of KMT2A (MLL) (ROSE group 1, R1), TCF3-PRX1 (R2), deregulation of ERG (R6) and Ph+/-Ph-like ALL (R8), and so on. Analysis of 1,164 childhood, adolescent and young adult ALL cases with available microarray gene expression data identified 31 cases in the R3 subgroup that lacked known chromosomal rearrangements (Supplementary Fig. 11), of which were in the discovery RNAseq cohort and had MEF2D rearrangements (Supplementary Data 1). RT–PCR identified known MEF2D rearrangements in 19 of the 20 remaining cases (Supplementary Data 6 and Supplementary Fig. 12). One of the two remaining cases had sufficient material for RNAseq, which identified a novel chimeric in-frame fusion of MEF2D exon 8 to exon 7 of the forkhead transcription factor gene, FOXJ2 (Supplementary Data 6). Taken together, 42 MEF2D-rearranged cases were discovered in this study that exhibit a distinct gene expression profile. Moreover, RNAseq identified 20 cases with rearrangements of ZNF384 to 6 different partners (ARID1B, CREBBP, EP300, SMA2CA1, TAF15 and TCF3), which were also associated with a distinct gene expression profile and exclusively observed in the R5 group (Supplementary Data 7 and Supplementary Fig. 13). Together, these findings suggest that MEF2D and ZNF384 rearrangements are founding alterations of the R3 and R5 subgroups, respectively, and represent distinct subtypes of B-ALL.

To explore the deregulated gene expression characteristics of MEF2D-rearranged ALL, we performed the gene set enrichment analysis (GSEA) and pathway analysis and demonstrated that MEF2D-rearranged cases exhibited a gene expression profile of a later maturation stage than other subtypes of ALL (Supplementary Data 8 and 9). We did observe negative enrichment of WNT signalling, consistent with the observation that the MEF2D-BCL9 fusion removes the β-catenin-interacting domains of BCL9. The clustering of gene expression profiles MEF2D-rearranged cases irrespective of C-terminal fusion partner suggests that deregulation of MEF2D function may in part mediate leukaemogenesis. To examine this, we compared the

### Table 1 | Clinical and genetic features of patients with MEF2D-rearranged ALL.

| Sample ID               | Fusion              | Age  | WBC   | Sex    | Karyotype                                      | SNP results                        |
|------------------------|---------------------|------|-------|--------|------------------------------------------------|-----------------------------------|
| SJBALL013388           | MEF2D-BCL9<sup>a</sup> | 10.0 | 45.4  | Female | NA                                              | NA                                |
| SJBALL016252           | MEF2D-BCL9<sup>a</sup> | 14.0 | 42    | Female | NA                                              | NA                                |
| SJBALL016291           | MEF2D-BCL9<sup>a</sup> | 10.0 | 23    | Male   | NA                                              | NA                                |
| SJBALL020667           | MEF2D-BCL9<sup>a</sup> | 14.8 | 39    | Female | NA                                              | No MEF2D change                    |
| SJBALL020703           | MEF2D-BCL9<sup>a</sup> | 14.2 | 38    | Female | 46,XX,del(9)(p22)x2/45,idem, – del(9), + der(9)(q10;q10), – 17(16)/46, idem, – del(9), + mar(14) | No MEF2D change                    |
| SJBALL020711           | MEF2D-BCL9<sup>b</sup> | 13.9 | 96.4  | Male   | NA                                              | Focal gains at MEF2D and BCL9 Deletion between BCL9 and MEF2D | NA |
| SJBALL020906           | MEF2D-BCL9<sup>b</sup> | 15.1 | 9.1   | Male   | NA                                              | NA                                |
| SJCOGALLO10235          | MEF2D-BCL9<sup>a</sup> | 13.5 | 82.7  | Female | 46,XX,del(11)(q24q31).i(7)(q10),del(9)(p21p24), add(18)(q21)[8]/46,XX,del(1)(q24q31), del(9)(p22),del(9)(p21p24), del(10)(p13p14)[7]/46,XX[4] | NA                                |
| SJBALL014169           | MEF2D-BCL9<sup>b</sup> | 21.3 | 20.9  | Female | 46,XX,del(12)(p11.2)[2]/45,XX, – 9, del(12)(p11.2)[15]/45,XX, – 11[1] | NA                                |
| SJBALL020917           | MEF2D-HNRNPUL1       | 11.9 | 3.5   | Female | 46,XX,del(13)(q12q32)[24]/46,XX[6]              | NA                                |
| SJBALL020967           | MEF2D-BCL9<sup>b</sup> | 16.3 | 27    | Female | 46,XX,del(17)(q11q21)[12]/46,XX[6]              | NA                                |
| SJBALL020964           | MEF2D-HNRNPUL1       | 17.6 | 10.8  | Male   | 46,XY,del(2)(q13q33),del(12)(p11.2)[22]         | Focal gain at HNRNPUL1             |
| SJBALL016275           | MEF2D-BCL9<sup>f</sup> | 15.0 | 61.9  | Female | NA                                              | NA                                |
| SJBALL016306           | MEF2D-BCL9<sup>f</sup> | 12.0 | 61.9  | Female | NA                                              | NA                                |
| SJCOGALLO103195         | MEF2D-BCL9<sup>f</sup> | 14.0 | 7.4   | Male   | 48,XY,add(3)(q11.2), + 8,del(8)(p11.1), add(12)(p11.2), + mar[11]/94, idem[2] – 8, – 21, + mar[6]/46,XX[2] | NA                                |
| SJCOGALLO10884          | MEF2D-BCL9<sup>d</sup> | 13.8 | 107   | Female | 46,XX,del(9)(p13)[1]/46,XX, add(17)(q24q24)[5]/46,XX[14] | NA                                |
| SJBALL020987           | MEF2D-BCL9<sup>a</sup> | 18.5 | 22.5  | Female | 46,XX,del(9)(p13),t(10;14)(q23;q11.2)[17]/46,XX[11] | No MEF2D change                    |
| SJBALL021005           | MEF2D-BCL9<sup>a</sup> | 17.6 | 9.4   | Male   | NA                                              | Deletion at MEF2D, and adjacent gain to BCL9 | NA |
| SJCOGALLO10859          | MEF2D-HNRNPUL1       | 7.6  | 21.8  | Female | 46,XX,add(7)(q11.2),add(9)(p13)[8]/46,XX[5]     | NA                                |
| SJBALL012686           | MEF2D-SJ8           | 9.0  | 28.1  | Female | NA                                              | No MEF2D change                    |
| SJBALL020141           | MEF2D-DAZAP1<sup>*</sup> | 35.0 | 5.4   | Male   | NA                                              | No MEF2D change                    |
| SJBALL015117           | MEF2D-CSF1R<sup>a</sup> | 48.0 | Female | NA                                              | NA                                |

NA, not available; SNP, single-nucleotide polymorphism.

Age is shown in years. WBC, peripheral blood leukocyte count at diagnosis, shown as 10⁹/l.

*Previously reported<sup>26</sup>.

<sup>2</sup>Four different isoforms of MEF2D-BCL9 fusions were detected in 16 samples, as denoted by the superscript letters a–d.

<sup>3</sup>Previously reported<sup>3</sup>.
gene expression profile of MEF2D-rearranged ALL with existing chromat in immunoprecipitation and sequencing and RNAseq data sets for MEF2D studies in muscle, retinal and lymphoid cells, and observed partial overlap with known MEF2D-bound genes. Notably, HDAC9, one of the top upregulated genes was consistently represented in the data sets (Supplementary Data 10). HDAC9 is a direct transcriptional target of MEF2 and acts to inhibit the transcriptional activity of MEF2C9, consistent with the striking downregulation of this gene in MEF2D-rearranged ALL (Supplementary Data 3–5 and Supplementary Figs 14 and 15).

Clinical and pathological features of MEF2D ALL. The median age of MEF2D-rearranged cases at diagnosis was 14.0 years (range 4.8–48.0) for the 42 MEF2D-rearranged cases and 14.0 years (range 4.8–21.3) for 29 MEF2D-BCL9 cases with approximately equal occurrence in male and female patients (Supplementary Data 3–5 and Supplementary Figs 14 and 15).

Figure 1 | MEF2D rearrangements in ALL. (a) Protein structure of wild-type MEF2D and fusion partners, (b) protein structure of each fusion protein and (c) electropherograms showing the results of RT–PCR and Sanger sequence (see also Supplementary Fig. 1). Four MEF2D-BCL9 isoforms were identified in 16 cases, with breakpoints involving exons 5 or 6 of MEF2D fused in frame to exons 9 or 10 of BCL9. Rearrangements involving MEF2D and other four fusion partners were identified in six samples. MEF2D-DAZAP1 and MEF2D-CSF1R fusions have previously been reported2,26. Protein region annotation: MADS box, immunoglobulin domain; Ig4, SCFR like, fourth Ig-like domain of stem cell factor receptor; PTK catalytic domain, tyrosine kinase catalytic domain.
RNAseq data, showing clustering of MEF2D-rearranged cases. A full list of cases and rearrangements identified by RNAseq is provided in Supplementary Data 1. (b) The gene expression signature for MEF2D-rearranged ALL was determined by comparing 19 MEF2D fusion samples (excluding 2 cases that failed quality-control threshold for gene expression analysis and one MEF2D-CSF1R rearranged case) and 199 control samples through Wald’s test and Benjamini–Hochberg adjustment. Genes with adjusted P-value < 0.01 are provided in Supplementary Data 3. The top 50 upregulated and top 50 downregulated genes (based on the adjusted P-value) were selected to perform supervised clustering and shown in the order from most significant upregulated genes (top) to the most downregulated ones (bottom). Stranded-PE100, total stranded paired-end 100 bp RNAseq; unstranded-PE75, unstranded mRNA paired-end 75 bp RNAseq. MEF2D, fusions involving MEF2D; JAK2/TYK2, fusions involving JAK2 or TYK2; PAX5, fusions with PAX5 as the N-terminal fusion partner with non-kinase genes; ABL1/2, fusions with ABL1 or ABL2 observed in Ph-like ALL; CRLF2, IGH-CRLF2 or P2RY8-CRLF2 fusions; ZNF384, fusions involving ZNF384 as the C-terminal fusion partner; EPOR, fusions deregulating EPOR observed in Ph-like ALL; PDGFRB, fusions involving PDGFRB; IGH, IGH fusions except IGH-CRLF2 and IGH-EPOR; MLL, fusions involving KMT2A(MLL) gene.

**Figure 2 | MEF2D-rearranged ALL cases have a distinct gene expression profile.** (a) Principal component analysis of samples subjected to unstranded RNAseq data, showing clustering of MEF2D-rearranged cases. A full list of cases and rearrangements identified by RNAseq is provided in Supplementary Data 1. (b) The gene expression signature for MEF2D-rearranged ALL was determined by comparing 19 MEF2D fusion samples (excluding 2 cases that failed quality-control threshold for gene expression analysis and one MEF2D-CSF1R rearranged case) and 199 control samples through Wald’s test and Benjamini–Hochberg adjustment. Genes with adjusted P-value < 0.01 are provided in Supplementary Data 3. The top 50 upregulated and top 50 downregulated genes (based on the adjusted P-value) were selected to perform supervised clustering and shown in the order from most significant upregulated genes (top) to the most downregulated ones (bottom). Stranded-PE100, total stranded paired-end 100 bp RNAseq; unstranded-PE100, unstranded mRNA paired-end 100 bp RNAseq; unstranded-PE75, unstranded mRNA paired-end 75 bp RNAseq. MEF2D, fusions involving MEF2D; JAK2/TYK2, fusions involving JAK2 or TYK2; PAX5, fusions with PAX5 as the N-terminal fusion partner with non-kinase genes; ABL1/2, fusions with ABL1 or ABL2 observed in Ph-like ALL; CRLF2, IGH-CRLF2 or P2RY8-CRLF2 fusions; ZNF384, fusions involving ZNF384 as the C-terminal fusion partner; EPOR, fusions deregulating EPOR observed in Ph-like ALL; PDGFRB, fusions involving PDGFRB; IGH, IGH fusions except IGH-CRLF2 and IGH-EPOR; MLL, fusions involving KMT2A(MLL) gene.
Data 1 and 6). The immunophenotype of **MEF2D**-rearranged ALL was distinct from other subtypes of B-ALL and was characterized by weak or absent expression of CD10 and high expression of CD38. Low or absent expression of CD10 is a feature of **MLL**-rearranged ALL and high CD38 expression is typically seen in normal, regenerating lymphoid cells (haematogones), but these features are otherwise uncommon in B-cell precursor ALL (Supplementary Table 4).

The outcome of **MEF2D**-rearranged ALL was inferior to that of other ALL subtypes. Although the number of **MEF2D**-rearranged cases was small compared with several other subtypes, analysis of children with ALL enroled on the AALL0232 study of high-risk pre-B ALL showed that the 5-year event-free survival (EFS) of **MEF2D**-rearranged ALL was 71.6% (s.e. ± 10.2%, n = 22), compared with that of **BCR-ABL1** (59.9 ± 8.9%, n = 48), **MLL** (77.5 ± 10.6%, n = 25), **Ph-like** (60.3 ± 5.5%, n = 115) and other pre-B ALL cases (87.3 ± 1.6%, n = 612; log rank test P < 0.0001; Fig. 3). Notably, Ras mutations were not associated with inferior outcome in this cohort (5 year EFS 89.7 ± 4.2% for 62 cases with Ras mutations compared with 76.2 ± 4.4% for 112 cases without, log rank test, P = NS). Through multivariable analysis with the Cox proportional-hazards regression model10, **MEF2D** rearrangements were not significantly associated with poor outcome following correction for age (≥ 10 years unfavourable, P = 0.0034), MRD positivity (P < 0.0001), peripheral blood leukocyte count (≥ 100 mm⁻³) unfavourable, P = 0.0067) and sex (P = NS). Further analysis of larger cohorts of **MEF2D**-rearranged cases will be required to determine whether this subset is independently associated with poor outcome.

Deregulation of expression of **MEF2D** target genes. The clustering of gene expression profiling data of **MEF2D**-rearranged cases with this diverse range of partner genes strongly suggests the fusions exert similar oncogenic effects. All fusions preserve the **MEF2D** MADS-box domain that mediates DNA binding and potentially dimerization. Thus, aberrant function mediated by **MEF2D** transcriptional activation is likely to be central in leukaemogenesis. To examine this, we performed transcriptional activation assays in which the ability of **MEF2D** fusion proteins, wild-type **MEF2D** and **BCL9**, and the rearranged N terminus of **MEF2D** to activate transcriptional targets were examined. This showed that each **MEF2D** fusion protein was significantly more potent in activating expression than wild-type **MEF2D**, and that truncated **MEF2D** was inactive (Fig. 4a). Thus, each fusion partner stabilizes and augments **MEF2D** transcriptional activation. In colony-forming assays, **MEF2D**-**BCL9** sustained serial replating of lymphoid colonies, indicating that this fusion confers haematopoietic self-renewal (Fig. 4b).

**Sensitivity to HDAC inhibition.** The poor outcome of **MEF2D**-rearranged ALL suggests that new therapeutic approaches directed against deregulated cellular pathways observed in this form of leukaemia are warranted. To explore this, we established xenografts of leukaemic cells of five cases with **MEF2D**-rearranged ALL engrafted in immunocompromised mice. Following engraftment, leukaemic cells were harvested from the bone marrow and spleen, purified and exposed to chemotherapeutic agents ex vivo. This demonstrated sensitivity to the HDAC inhibitor panobinostat (but little sensitivity to **BCL9** inhibitors), suggesting HDAC inhibition as a therapeutic option in this high-risk form of leukaemia (Fig. 4c and Supplementary Fig. 16a–d). Although HDAC inhibitors such as panobinostat are broadly active in many haematopoietic tumours, including **ALL**11 (data for non-**MEF2D** ALL subtypes are shown in Supplementary Table 4).
Vincristine
Vorinostat
MIG
B-1 through M/B-3, three most common
more xenografted samples were also tested by the four drugs (Supplementary Fig. 16). Error bars represent mean
MIG, MSCV-IRES-GFP, represents empty vector; MEF2D-WT, vectors express wild-type MEF2D; MEF2D
in mean
factor genes in ALL, including
rearranged to a variety of cytokine receptors and transcription
IGH
leukaemia cell line CEMO-1 (refs 5,17). Although
of a t(1;14)(q21;q32)
located on chromosome 1q21 and was first identified as the target
IGH.
involved
IGH
in B-lymphoid maturation at the pre-B-cell stage15,16. MEF2D is expressed throughout B-cell
differentiation15 and inactivation of
IGH
rearrangement has previously been identified
MEF2D-DAZAP1
is located at 19p13.3 and encodes
DAZ and DAZL, and other RNA-binding proteins 25. The
heterogeneous nuclear ribonucleoprotein family that may exert a
role in nucleocytoplasmic RNA transport and DNA repair 28.
HNRNPUL1 encodes a nuclear RNA-binding protein that interacts with the infertility factors
DAZ and DAZL, and other RNA-binding proteins25. The
MEF2D-DAZAP1 rearrangement has previously been identified and shown to result in proliferation of fibroblasts26,27. HNRNPUL1 encodes a nuclear RNA-binding protein of the heterogeneous nuclear ribonucleoprotein family that may exert a role in nucleocytoplasmic RNA transport and DNA repair28. SS18 is fused with SSX1 or SSX2 in synovial sarcoma, and has putative roles in chromatin remodelling and WNT/β-catenin signalling29,30. FOXJ2 is a transcription factor that has been reported to affect migration and invasiveness of various tumours31,32 but, unlike other forkhead transcription factor genes, has not previously been identified as a target of rearrangement in cancer.

Discussion
The distinct gene expression profile, tumour cell immunopheno-
type, older age of onset and poor outcome of MEF2D-rearranged ALL cases together suggest that MEF2D-rearranged ALL represents a biologically distinct form of leukaemia. The fusion partners of MEF2D have a diverse range of biological functions. MEF2D is a member of a family of four myocyte enhancer factor transcription factors with an important role in neuronal differentiation13,14. MEF2D is expressed throughout B-cell differentiation15 and inactivation of Mef2c/d results in an arrest in B-lymphoid maturation at the pre-B-cell stage15,16. BCL9 is located on chromosome 1q21 and was first identified as the target of a t(1;14)(q21;q32) IGH-BCL9 rearrangement in the pre-B-cell leukemia cell line CEMO-1 (refs 5,17). Although IGH is rearranged to a variety of cytokine receptors and transcription factor genes in ALL, including CRLF2, EPOR and members of the CEBP family of transcription factors18–21, and multiple known and novel IGH rearrangements, were identified (Supplementary Data 1), none of the BCL9 rearrangements identified in this study involved IGH. BCL9 is a component of the WNT/β-catenin signalling cascade that has important roles in development, stem cell self-renewal and oncogenesis. BCL9 is part of the nuclear complex consisting of T-cell factor/lymphoid enhancer factor, β-catenin, BCL9/BCL9L and PYGO, which activates transcription of canonical WNT target genes such as FGF20, DKK1, WISP1, MYC, CCND1 and GCG (Glucagon)22,23. BCL9 has also been shown to have an oncogenic role in multiple myeloma24.

The MEF2D-BCL9 rearrangement results in deregulated expression of the C terminus BCL9 expression (Supplementary Fig. 2), raising the possibility that these fusions may perturb WNT/β-catenin signalling. However, the MEF2D-BCL9 rearrangements involve only the last one or two of ten exons of BCL9 exons resulting in loss of the BCL9 domains that mediate interaction with PYGO1 and β-catenin, consistent with the lack of enrichment of WNT/β-catenin signalling in the gene expression profile of MEF2D-rearranged ALL. Thus, IGH-BCL9 and MEF2D-BCL9 probably exert different roles in leukemic transformation. DAZAP1 is located at 19p13.3 and encodes an RNA-binding protein that interacts with the infertility factors DAZ and DAZL, and other RNA-binding proteins25. The MEF2D-DAZAP1 rearrangement has previously been identified and shown to result in proliferation of fibroblasts26,27.

Figure 4 | Luciferase and drug-sensitivity assays. (a) Luciferase assay to test MEF2 binding activities. The average levels of normalized luciferase activities ± s.d. from six replicates are indicated. MEF2D-rearranged isoforms are more potent transcriptional activators than wild-type and truncated MEF2D. (b). Colony-forming assay for retroviral vectors transduced lineage negative bone marrow cells. Colony number counted from three replicate dishes is shown in mean ± s.d. Western blotting was performed to verify expression of proteins by each construct (Supplementary Fig. 18). RLU, relative luminescence units; MIG, MSCV-IRES-GFP, represents empty vector; MEF2D-WT, vectors express wild-type MEF2D; MEF2DA, truncated MEF2D; BCL9-WT, wild-type BCL9; M/B-1 through M/B-3, three most common MEF2D-BCL9 fusion isoforms; M/H, MEF2D-HNRNPUL1. (c) Ex vivo drug sensitivity assays from two xenografted samples. Four chemical compounds including ABT-199 (selective BCL-2 inhibitor), vincristine and two HDAC inhibitors (panobinostat and vorinostat) were applied to measure growth inhibition curves in xenografted samples. The MEF2D rearrangements were confirmed in xenografted samples by RT-PCR and three more xenografted samples were also tested by the four drugs (Supplementary Fig. 16). Error bars represent mean ± s.d.

and are being explored as a therapeutic option for a range of relapsed and refractory tumours, we observed exquisite sensitivity in MEF2D-rearranged ALL (Fig. 4c). In contrast, and consistent with a lack of activation of β-catenin signalling from MEF2D-BCL9 fusions, we observed a lack of efficacy of selective, potent BCL9 inhibitors17 ex vivo (Supplementary Fig. 16e–h).
These data demonstrate the power of transcriptome sequencing to identify novel rearrangements that define new genetic subtypes of ALL. The finding of MEF2D rearrangements as a hallmark of a subtype of ALL with a distinct gene expression profile provides another example of ALL subtypes that are cryptic (in this case, due to the common involvement of rearrangement of two genes co-located on one chromosome) and represented by rearrangement to diverse partners rather than a single fusion observed in other subtypes of ALL. Despite the common involvement of the WNT co-factor BCL9, there was no evidence of dysregulated WNT/β-catenin signalling in leukaemogenesis. In contrast, deregulated MEF2D activity, with activation of HDAC9 and resulting inhibition of MEF2C, were characteristic of this form of leukaemia. Further studies are required to dissect the relative role of MEF2D and partner gene involvement in leukaemogenesis, but our existing data suggest the potential for HDAC inhibition in this form of leukaemia.

Methods

Patients and samples. Patients with ALL were enrolled on Alliance—Cancer and Leukemia Group B, COG, City of Hope, Eastern Cooperative Oncology Group, MD Anderson Cancer Center, Medical Research Council UK, Northern Italian Leukemia Group, Southwestern Oncology Group and University of Toronto Protocols (Supplementary Data 1). Patients and/or guardians provided informed consent/assent. The study was approved by the St. Jude Children’s Research Hospital Institutional Review Board. Leukaemic cells from bone marrow aspirates obtained at diagnosis were processed by density gradient centrifugation. Samples of <70% tumour cell content were purified by fluorescence-activated cell sorting before RNA extraction.

Transcriptome sequencing and data analysis. Transcriptome sequencing was performed using TruSeq library preparation and HiSeq 2000 and 2500 sequencers. All sequencing was paired-end and was performed using (1) total RNA and stranded RNA sequencing (100 bp reads), and (2) poly-A-selected mRNA (100 or 75 bp reads). Sequencing reads were mapped to the GRCh37 human genome reference by STAR35 (version 2.5.1b) through the two pass mapping pipeline. Gene annotation downloaded from Ensembl website (http://www.ensembl.org/) was used for STAR mapping and the following reads count evaluation. Cicero36 and FusionCatcher34,35 were used to detect fusions. To assess gene expression profile, reads count for each annotated gene was extracted by HTSeq package38 and gene expression level normalization and differential expression analysis was carried out using DESeq2 R package35. To evaluate the relative gene expression level, regularized log-transformed (log) value was calculated by DESeq2.

Gene expression profile between samples with and without MEF2D fusions were examined in two ALL cohorts. These included a cohort of total stranded RNAseq (30-fold coverage on exon regions (D30C > 20%) and cases with unstranded mRNA sequencing (D30C > 30%). Two MEF2D-rearranged cases failed to meet the requirement of D30C > 20% were excluded, both of which were sequenced by unstranded 75 bp paired-end protocol. Two hundred and nineteen samples were available for gene expression profiling analysis, including 95 samples with total stranded RNAseq (this cohort was enriched for Ph-like ALL and included 8 cases with MEF2D fusions) and 124 samples with unstranded RNAseq (enriched for other B-ALL and included 12 cases with MEF2D fusions) data are available for the gene expression profiling and gene signature analysis.

GSEA and pathway analysis. Normalized gene expression levels were evaluated from RNAseq data and subjected to DESeq for differential gene expression analysis. As recommended from GSEA34, genes ranked according to the significance of differential expression between samples with and without target rearrangements and then input to GSEA to calculate potentially enriched gene sets from a homemade gene sets database. To assess the pathway enrichment, genes with P-value ≤ 0.01 and fold change ≥ 2 were selected to run DAVID pathway analysis37.

Mutation detection from RNAseq data. The mutations were called according the GATK33 forum recommended pipeline for calling variant in RNAseq data (http://www.broadinstitute.org/gatk/discussion/3891/calling-mutations-in-rna-seq). Specifically, STAR mapped bam files were processed by Picard (https://broadinstitute.github.io/picard) to mark duplicate reads, then a GATK module SplitNCigarReads was used to splits reads into exon segments and hard-clip any sequences overhanging into the intronic regions. Variant calling was performed by the HaplotypeCaller module in GATK and then the variants were quality controlled by the following steps: (1) at least three reads support the mutation and the mutant allele frequency is ≥ 0.05; (2) not observed in common SNP database (from UCSC version 142); and (3) not observed in ≥ 2 samples from our germline exome sequence cohort (775 samples). After filtering, all the mutations were annotated to genes according to their genomic positions. Non-silent mutations were compared with COSMIC somatic mutation database (GRCh37-V74) and the overlapped cancer relevant genes (http://cancer.sanger.ac.uk/census) were kept for further analysis.

Retroviral constructs and immunoblotting. MEF2D-rearranged isoforms, wild-type MEF2D and BCL9 were amplified by RT-PCR using Phusion High-Fidelity DNA Polymerase (New England Biolabs, Inc.) and primers (Supplementary Table 5) from either leukemic cell or MOLT4 cell line complementary DNA. Amplification products were purified by Wizard SV Gel and PCR Clean-up system (Promega) and verified by Sanger sequencing. Purified PCR products were cloned into pCR-Blunt II-TOPO (Life Technologies) and sub-cloned into pLenti6-MCFpuro (Promega) and terminated stem cell virus long terminal ribonucleotide entry site-green fluorescent protein (MSCV-ires-GFP) retroviral vector.

Transfected BCL9 isoform was amplified from MSCV-MEF2D-BCL9-ires-GFP vector (Mu-B1 vector, the one with the most common MEF2D-BCL9 fusion isoform) by the same PCR system and specially designed primers for Gateway cloning (Vector NTI, Thermoscientific). BCL9 isoform primers for truncated MEF2D were designed to amplify from the 5’ terminus of MEF2D to the last amino acid fused to BCL9 with a stop codon inserted at the fusion point. Purified PCR products were cloned into the Gateway pDONR221 vector (Thermo Fisher) by BP Clonase Enzyme (Thermo Fisher). Transfected MEF2D and pDONR221 vector was shuttled into a Gateway compatible MSCV-ires-GFP vector using the LR Clonase enzyme (Thermo Fisher). Constructs were verified by Sanger sequencing (Supplementary Table 5).

Transfected/transduced cells and human leukemic cells were lysed in RIPA buffer supplemented with phosphatase inhibitors (Sigma) and 30 μg protein of patient samples (15 μg of lysate from NIH-3T3 cells) electrophoresed through 4–12% NuPage Bis-Tris gels (Life Technologies) at 190 V for 80 min. Blots were probed with anti-MEF2D (H-57, sc-366368), anti-actin-I (1-9) (both, Santa Cruz Technologies) and anti-BCL9 (Abcam ab54833). For human leukemic leukaemia, 1,250, 1,500 and 1,2,000 dilutions were used for MEF2D, BCL9 and β-antibodies, respectively. For transfected/transduced cells, the 3 antibodies were 1,1,000 dilutions. Five percent BSA in Tris-buffered saline with 0.1% Tween 20 solution was used to dilute the antibodies.

Luciferase assays. In 96-well dishes, 2 × 104 293T cells were transfected with 0.04 pmols of MSCV-ires-GFP (MIG) vectors wild-type MEF2D, truncated MEF2D (MEF2DΔA), wild-type BCL9, three most common MEF2D-BCL9 fusion isoforms, MEF2D-HNRPL1 or empty vector along with 250 ng of a 3XMEF2 firefly luciferase reporter construct (Addgene plasmid #32967) and 50 ng of a pRL-TK Renilla luciferase reporter plasmid (Promega) using FuGene HD (Roche Diagnostics). Forty-eight hours after transfection, measurement of firefly and Renilla luciferase activity was performed using the Dual-Glo Luciferase Assay System (Promega E2920) according to the manufacturer’s instructions. All transfections were performed in 96-well plates in technical replicates and repeated three independent experiments. Firefly luciferase activity was normalized according to corresponding Renilla luciferase activity and reported as TB-S·T·s·d from one representative experiment.

Generation of xenografted mice. One million patient-derived leukemic cells were transplanted into three mice each by tail vein injection into 8–10-week-old subcutaneously irradiated (250 rads) female immunodeficient NOD.Cg-Pkd−/−Irbg-wt119/C2 (NSG) mice. Disease progression was monitored by flow cytometric analysis of peripheral blood cells stained for McD45, hCD45, hCD19 and hCD7, and the bone marrow and spleen were harvested when > 80% of hCD45 + hCD19 − cells were observed in peripheral blood. Mice were housed in an American Association of Laboratory Animal Care-accredited facility and were treated on Institutional Animal Care and Use Committee-approved protocols in accordance with NIH guidelines. The study was approved by institutional review board of St Jude Children’s Research Hospital.

Ex vivo drug sensitivity assays. Human xenograft cells were flushed from tibiae and femora of moribund mice, subjected to red cell lysis, washed and immediately plated 1 × 105 per well in 96-well plates containing media with the indicated concentrations of compounds in triplicate. Cells were treated for 48 h, incubated with resazurin for 4 h and read on a Synergy HT (Biotek).

Colony-forming assays. Bone marrow from 10-week-old C57BL/6 wild-type mice was extracted from the tibiae and femora. Red blood cells were lysed and the remaining bone marrow cells were incubated with a cocktail of biotinylated anti-mouse antibodies (Gr-1, B220, Ly-1, Ter119, Mac-1, diluted 1:100 in PBS with 5% rat serum and 5% FCS; BD Biosciences) followed by mixing with streptavidin-coated magnetic beads (Beckman Coulter) and then M280 Streptavidin; Life Technologies). Cells were separated on a magnet and unbound cells were collected and incubated at 37 °C, 8% CO2 for 2 days in the presence of interleukin (IL)-3, IL-6, stem cell factor
Statistical analysis. Associations between ALL subtype and treatment outcome (EFS and overall survival) were performed using the Kaplan–Meier estimator with Peto's estimator of s.d. and the log-rank test. Analyses were performed using Prism v6.0 (GraphPad), R (www.r-project.org) and SAS (v9.1.2, SAS Institute, Cary, NC).

Data availability. Genome and RNAseq data are deposited at the European Genome Phenome archive, accession EGAS000001000654 (ref. 3), EGAS00001001952 and the database of Genotypes and Phenotypes (dbGaP) under accession code phs000208. Data from the samples, including gene expression and SNP microarray data, may also be accessed through the Therapeutically Applicable Research to Generate Effective Treatments data matrix (https://oegcancer.gov/programs/target). The remaining data are available within the article and its Supplementary Files or available from the authors upon request.

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Author contributions

Z.G., R.C.H., Y. Li. and Y. Liu analysed genomic data. Z.G., I-M.C., M.C., K.M., I.I., D.P.-T., K.R., S.R. and E.S. performed experiments. M.V. performed fluorescence in situ hybridization. J.E., D.S.G., A.M., K. Mungall, M.M., R.M., Y.M. and Y.S. performed transcriptome sequencing. M.D. and Y.D. performed biostatistical analysis. M.B. and B.W. performed immunophenotyping. E.L., K. Maloney, L.A.M., A.A., W.L.S., M.J.R., J. G.-F., A.R., O.S., J.P.R., M.D.M., A.Y.M., B.P., A.R.F., J.M.R., S.L., R.B., I.A., J.K., K. Mrózek, G.M., C.D.B., W.S., S.K., H.M.K., M.K., W.P., C.L.W., M.L.L. and S.P.H. provided clinical samples and data. Z.G. and C.G.M. wrote the manuscript.

Additional information

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