Germline ETV6 Mutations Confer Susceptibility to Acute Lymphoblastic Leukemia and Thrombocytopenia

Sabine Topka1,2*, Joseph Vijai1,2*, Michael F. Walsh3, Lauren Jacobs1, Ann Maria1, Danylo Villano1, Pragna Gaddam1, Gang Wu3, Rose B. McGee3, Emily Quinn3, Hiroto Inaba3, Christine Hartford3, Ching-hon Pui3, Alberto Pappo3, Michael Edmonson3, Michael Y. Zhang4, Polina Stepensky5, Peter Steinherz1, Kasmintan Schrader6, Anne Lincoln1, James Bussel7, Steve M. Lipkin7, Yehuda Goldgur8, Mira Harit5, Zsofia K. Stadler1, Charles Mullighan3, Michael Weintraub5, Akiko Shimamura4,9, Jinghui Zhang3, James R. Downing3, Kim E. Nichols3*, Kenneth Offit1,2,7*

1 Department of Medicine, Memorial Sloan Kettering Cancer Center (MSKCC), New York, New York, United States of America, 2 Cancer Biology and Genetics Program, Sloan Kettering Institute, New York, New York, United States of America, 3 St Jude Children’s Research Hospital (SJCRH), Memphis, Tennessee, United States of America, 4 Fred Hutchinson Cancer Research Center and University of Washington, Seattle, Washington, United States of America, 5 Pediatric Hematology/Oncology Department, Hadassah-Hebrew University Medical Center, Jerusalem, Israel, 6 University of British Columbia, Vancouver, British Columbia, Canada, 7 Weill Cornell Medical College, New York, New York, United States of America, 8 Structural Biology Program, Sloan Kettering Institute, New York, New York, United States of America, 9 Seattle Children’s Hospital, Seattle, Washington, United States of America

☯ These authors contributed equally to this work.
‡ These authors are joint senior authors on this work.
* kim.nichols@stjude.org (KEN); offitk@mskcc.org (KO)

Abstract

Somatic mutations affecting ETV6 often occur in acute lymphoblastic leukemia (ALL), the most common childhood malignancy. The genetic factors that predispose to ALL remain poorly understood. Here we identify a novel germline ETV6 p. L349P mutation in a kindred affected by thrombocytopenia and ALL. A second ETV6 p. N385fs mutation was identified in an unrelated kindred characterized by thrombocytopenia, ALL and secondary myelodysplasia/acute myeloid leukemia. Leukemic cells from the proband in the second kindred showed deletion of wild type ETV6 with retention of the ETV6 p. N385fs. Enforced expression of the ETV6 mutants revealed normal transcript and protein levels, but impaired nuclear localization. Accordingly, these mutants exhibited significantly reduced ability to regulate the transcription of ETV6 target genes. Our findings highlight a novel role for ETV6 in leukemia predisposition.

Author Summary

Inherited mutations of transcription factors have recently been associated with susceptibility to acute leukemia. Here we report two unrelated kindreds with inherited mutations in ETV6, the gene encoding the transcription factor ETS variant 6. These families were
characterized by a low platelet count (thrombocytopenia) and acute lymphoblastic leukemia (ALL). Sequencing a panel of genes identified germline ETV6 mutations associated with leukemia and thrombocytopenia in multiple individuals tested. In one family, there was a substitution within the DNA binding domain of ETV6, termed L349P, and in the second there were five base pairs missing in ETV6 (N385fs), causing an abnormally truncated protein. We overexpressed the ETV6 mutants in the HeLa cell line and measured protein levels and localization within the cells. Instead of localizing to the nucleus, as expected for a transcription factor, the mutant proteins were found in the cytoplasm. The mutant proteins also showed decreased ability to regulate the expression of other genes typically suppressed by ETV6. These findings suggest that germline ETV6 mutations cause a new type of heritable leukemia. This discovery makes possible the pre-symptomatic diagnosis of leukemia susceptibility in families with germline ETV6 mutations, and also provides new information on the causes of leukemia.

Introduction

Acute leukemias comprise the most common form of pediatric cancer, among which acute lymphoblastic leukemia (ALL) makes up 80–85% of the cases[1,2]. It is well recognized that a proportion of affected children develop the disease due to an underlying predisposition. The currently recognized genes responsible for autosomal dominant transmission of childhood leukemia include TP53, CEBPA, PAX5 and GATA-2[3–8]. Occasionally, acute leukemia presents in the context of thrombocytopenia. Consistent with this feature, several heritable thrombocytopenia syndromes are known to exist, some of which are associated with an increased incidence of leukemia. Genes associated with these syndromes include RUNX1, ANKR26, GATA1, MPL, HOXA11 and RMB8A[9–16]. Despite the identification of these genes, there remain many cases for which the underlying mechanism remains unexplained. In this study, we analyzed one large kindred and one parent-child trio, both affected by ALL and thrombocytopenia. By exome sequencing and also sequencing plausible candidate genes such as those involved in B-lymphocyte development and differentiation, we identified germline mutations in the transcription factor ETV6 that co-segregated with disease in each kindred. Functional studies support a pathogenic role for the observed mutations, both of which affect the DNA binding domain. These findings are consistent with independent observations describing additional kindreds characterized by thrombocytopenia and predisposition to hematopoietic malignancy[17,18] and provide insights into the mechanisms of leukemia susceptibility and clinical phenotypes associated with germline ETV6 mutations[19].

Results/Discussion

Phenotypic features of the kindreds

As part of a collaborative study focusing on pedigree analysis and gene discovery in childhood leukemia, we identified a Polish/Moroccan kindred in which 10 individuals developed thrombocytopenia and 4 individuals developed thrombocytopenia and ALL (Kindred 1 in Fig 1A). In the 3 ALL cases in Kindred 1 for whom flow-cytometric data were available, all were of the pre-B-ALL subtype. In 3 cases with thrombocytopenia and no evidence of ALL, the mean corpuscular volume (MCV) was decreased in 1 case and normal in 2 others (Table 1). In 2 individuals with no evidence of hematologic abnormalities, there was a history of renal cell cancer and duodenal adenocarcinoma. A second unrelated Western European/Native American family
was identified in which a child developed ALL followed by myelodysplastic syndrome and acute myeloid leukemia (AML). This child’s mother, maternal aunt and maternal grandfather exhibited thrombocytopenia (Kindred 2 in Fig 1B). This patient was evaluated by a geneticist due to subtle dysmorphic features; however, clinical assessment did not suggest a known genetic syndrome, and microarray and karyotype did not reveal any large deletions, rearrangements or other structural chromosomal abnormalities.

Identification of germline \( ETV6 \) mutations

DNA from 16 individuals in Kindred 1 (9 individuals with thrombocytopenia and/or ALL and 7 unaffected individuals) was subjected to Sanger sequencing for all exons of a targeted panel of leukemia-associated genes (Methods). Co-segregation of identified variants was tested using an autosomal dominant mode of inheritance. Published demographic data and medical literature were manually reviewed for all variants observed. Only one variant chr12:12,037,415 T>C satisfied the criteria of segregation as well as rarity, as evidenced by its absence in public genomic databases such as dbSNP[20], 1000 genomes[21], Exome Sequencing Project[22] and...
Exome Aggregation Consortium (http://exac.broadinstitute.org). This variant, identified in 9 out of 9 (100%) affected family members tested, represents a heterozygous missense c. T1046C mutation in ETV6 (NM_001987). One individual (generation 3, individual 13) with thrombocytopenia and leukemia was not tested. This nucleotide change is predicted to result in the substitution of proline for leucine at codon 349 (L349P; Fig 1A and Table 1). Seven out of 7 (100%) unaffected family members tested exhibited a wild type (WT) ETV6 sequence.

Fibroblast and lymphocyte DNA from the proband with ALL and parents in Kindred 2 were analyzed by clinical whole exome sequencing (Ambry Genetics, Aliso Viejo, CA, USA). The proband and his mother harbored a heterozygous deletion of 5 nucleotides (c.1153-5_1153_1delAACAG) within ETV6. This deletion is predicted to lead to a frameshift at codon 385 and truncation of the ETV6 protein at codon 389 (N385fs; Fig 1B and Table 1). Genome-wide DNA copy alteration analysis using single nucleotide polymorphism microarrays of the diagnostic ALL sample from the proband in Kindred 2 revealed deletion of the wild type and

Table 1. Clinical features of individuals in the study kindreds.

| Kindred | ETV6 alteration status | Individual | Cytopenias | Platelet count (per L) | MCV (fL) | Flow cytometry | Malignancies | Additional features |
|---------|------------------------|------------|------------|------------------------|----------|----------------|--------------|-------------------|
| 1       | Not sequenced          | I.1        | Unknown    | not available          | n/a      |                |              |                   |
| 1       | Not sequenced          | II.1       | Unknown    | not available          | n/a      |                | RCC          |                   |
| 1       | Not sequenced          | II.2       | Thrombocytopenia | 11 x 10^9          | n/a      |                | ALL          |                   |
| 1       | L349P                  | II.3       | Thrombocytopenia | not available        | CD19+,CD20+,CD22+,CD34+,TdT+ |                | ALL          |                   |
| 1       | L349P                  | II.6       | Thrombocytopenia, anemia | 8 x 10^9          | n/a      |                | Arthritis     |                   |
| 1       | L349P                  | II.9       | Thrombocytopenia | not available        | CD10+,CD20+,CD22+,TdT+ |                | ALL          |                   |
| 1       | L349P                  | II.10      | Thrombocytopenia | 53 x 10^9          | 85       | n/a            | Ankylosing spondylitis, uveitis |                   |
| 1       | L349P                  | II.13      | Thrombocytopenia | not available        | n/a      |                |              |                   |
| 1       | L349P                  | II.14      | Thrombocytopenia | not available        | n/a      |                |              |                   |
| 1       | L349P                  | III.1      | Thrombocytopenia | 75 x 10^9          | 85.8     | n/a            | Secondary amenorrhea |                   |
| 1       | L349P                  | III.2      | Thrombocytopenia, anemia | 13 x 10^9          | 79       | n/a            |              |                   |
| 1       | L349P                  | III.12     | Thrombocytopenia | not available        | n/a      |                | Cleft lip/palate |                   |
| 1       | Not sequenced          | III.13     | Thrombocytopenia | not available        | CD10+,CD20+,CD79a+,TdT+ |                | Pancytopenia, MDS, ALL | Cleft lip/palate |
| 2       | Not sequenced          | I.2        | Thrombocytopenia | not available        | n/a      |                |              |                   |
| 2       | Not sequenced          | II.1       | Thrombocytopenia | not available        | n/a      |                |              |                   |
| 2       | N385fs                 | II.2       | Thrombocytopenia | not available        | n/a      |                |              |                   |
| 2       | N385fs                 | III.1      | Thrombocytopenia | not available        | CD10+,CD20+,CD79a+,TdT+ |                | ALL,MDS,AML | Craniofacial/skeletal dysmorphisms |

RCC = renal cell carcinoma; ALL = acute lymphoblastic leukemia; MDS = myelodysplastic syndrome; AML = acute myeloid leukemia; MCV = mean corpuscular volume. ETV6 status, hematologic phenotypes, and additional clinical features of patients in two kindreds with segregating germline ETV6 mutations.

doi:10.1371/journal.pgen.1005262.t001
retention of the mutant \(ETV6\) allele, as well as deletions of \(IKZF1\), \(PAX5\), \(BTG1\), and \(RB1\). Other than the 2 mutations in \(ETV6\), there were no pathologic genetic mutations associated with ALL or thrombocytopenia that co-segregated with disease in either kindred.

Both \(ETV6\) variants were absent in the National Heart Lung Blood Institute (NHLBI) Exome Sequencing Project (ESP) (http://evs.gs.washington.edu/EVS/), Exome Aggregation Consortium (ExAC) (http://exac.broadinstitute.org/), or St. Jude Children’s Research Hospital–Washington University Pediatric Cancer Genome Project (PCGP) databases[23]. SIFT[24] and Polyphen prediction tools suggest the mutations to be deleterious and probably damaging to protein function. To understand how these two mutations might influence protein function, we modeled their effect on the \(ETV6\) protein structure. Both the L349P and the N385fs mutation are located in the ETS domain of \(ETV6\) (Fig 2A). The L349P mutation is predicted to cause significant conformational changes in areas adjacent to the ETS domain by introducing a kink in the H2 \(\alpha\)-helix, resulting in possible \(ETV6\) protein misfolding. The N385fs mutation affects the ETS domain and is predicted to truncate \(ETV6\) at a region involved in DNA interaction (Fig 2B).

### Functional assessment of the \(ETV6\) mutations

To evaluate the functional consequences of these mutations, we first assessed whether L349P and N385fs might impair transcriptional repression by \(ETV6\). HeLa cells were transiently co-transfected with constructs encoding the WT or mutant \(ETV6\), as well as constructs containing the \(PF4\) or \(MMP3\) promoters, which harbor multiple ETS binding sites and are natural \(ETV6\) targets. We compared the results to those obtained using other recently described germline \(ETV6\) variants, P214L, R369Q, R399C [17]. As expected, WT \(ETV6\) repressed expression of both reporters (Fig 3A), while each of the \(ETV6\) mutants exhibited significantly decreased repression. To further explore the effects of the \(ETV6\) mutations, we analyzed the expression of \(EGR1\) and \(TRAF1\), genes that are normally upregulated by WT \(ETV6\) [17]. Consistent with published reports, \(EGR1\) and \(TRAF1\) were upregulated 3-fold in cells transfected with WT \(ETV6\). In contrast, the mutants induced minimal to no upregulation for both of these target genes (Fig 3C). Indeed, the levels were significantly reduced compared to WT \(ETV6\). In each of these assays, we observed comparable levels of WT and \(ETV6\) mutant mRNA transcripts (Supporting Information 1). Thus, transcript stability appears to be unaffected by the \(ETV6\) mutations.

To examine whether the L349P and N385fs mutations negatively impact translation or alter subcellular localization of the \(ETV6\) protein, we performed cell fractionation assays and western blotting of HeLa cells transiently transfected to express WT or mutant \(ETV6\). Both proteins were detectable by Western blotting, with a smaller product observed for the N385fs mutation. Both mutants were undetectable in the nucleus (Fig 4A), but detected within the cytoplasmic fraction (Fig 4B). This is in contrast to the described mutants P214L, R369Q and R399C, which were detected in cytoplasmic as well as nuclear fractions. These patterns were quantitated and confirmed by measuring the nuclear to cytoplasmic ratio (Fig 4C).

### Evaluation of the incidence of germline \(ETV6\) mutations

Fusions involving \(ETV6\) in leukemia have long been recognized [25–27]. Other mutation types, including single nucleotide variations, insertions, deletions, frame-shifts and non-sense alterations are also becoming increasingly evident in hematologic malignancies[17,18,28]. We performed additional sequence analysis on exons 5–8 of \(ETV6\) in unrelated probands from 27 unrelated kindreds with a family history of ALL, but identified no mutations in this region of \(ETV6\). To further characterize the spectrum of germline and somatic \(ETV6\) mutations that
contribute to childhood leukemia, we screened a cohort of 588 leukemia patients evaluated through the PCGP, a genomic sequencing effort involving pediatric cancers [4,28-42](accession# EGAS00001000348, EGAS00001000654, EGAS00001000380, EGAS00001000253, EGAS00001000246, EGAS00001000447). Seventeen distinct somatic ETV6 variants and two rare germline variants were identified (V37M, R181H; Fig 2A). Both rare variants occurred in patients with B-ALL, but with no evidence for loss or mutation of the WT ETV6 allele within the leukemia samples. In one of these cases, there was a secondary vulvar squamous cell

---

Fig 2. Location of somatic and germline ETV6 mutations and structural modeling. (a) Schematic depicting the germline ETV6 mutations detected in the MSKCC and SJCRH kindred, reported separately or somatic mutations detected as part of the Pediatric Cancer Genome Project. Somatic and germline mutations are indicated by separate green and purple lines, respectively. (b) Structural modeling of ETV6 with the germline ETV6 L349P and N385fs mutations. The ETV6 L349P amino acid substitution is located on an α-helix within the DNA binding domain and causes extensive kinking of the protein structure. The ETV6 N385fs mutation results in truncation of the DNA binding domain.

doi:10.1371/journal.pgen.1005262.g002
carcinoma. There was no family history of leukemia or thrombocytopenia in either of these cases. Luciferase assays performed on these variants showed no significant changes in transcriptional repression activity when compared to WT ETV6 (Fig 3B). We queried several public variant databases for the presence of these two variants. The 1000 genomes project has a total of 2,819 samples from the world’s major populations. The current version of the Exome sequencing project (EVS/ESP) has a set of 2,203 African-American and 4,300 European-American unrelated individuals, totaling 6,503 samples. The Exome aggregation consortium (ExAC) has 60,706 unrelated individuals sequenced as part of various disease-specific and population genetic studies. In total, we have queried over 140,000 chromosomes. The V37M variant (chr12:11905459G>A) was seen only in the ExAC data at an allele frequency of 1.649x10^-05 and the R181H variant (chr12:12022436 G>A) was found at an allele frequency of 1.071x10^-04.

Fig 3. Effect of germline ETV6 mutations on transcription. (a) The effects of germline mutations on ETV6 function were examined using a Dual Luciferase Reporter Assay. Each of the mutants tested exhibited significantly (** * P < 0.0001) impaired transcriptional repression from the PF4 and MMP3 promoter constructs when contrasted with the WT ETV6 in the co-transfection experiment. The experiment was performed with 6 replicates for each condition and repeated 3 times. Statistical analysis was done using an unpaired t-test, the error bars show the Standard Error of Mean (SEM). (b) The effects of V37M and R181H germline mutations on ETV6 function were examined using a Dual Luciferase Reporter Assay. The experiment was performed with 6 replicates for each condition and repeated twice. Statistical analysis was done using an unpaired t-test, the error bars show the Standard Deviation (SD). (c) Quantitative PCR of ETV6 transcriptional targets EGR1 and TRAF1 showed reduced transcriptional abundance in the mutants when contrasted with the WT. The effect was most pronounced in the frameshift mutant. The experiment was performed in triplicate for each condition and repeated three times. Statistical analysis was done using an unpaired t-test, the error bars show the Standard Error of Mean (SEM).

doi:10.1371/journal.pgen.1005262.g003
in ExAC. It was also found 2 times in NHLBI-ESP (AF = 1.162x10^{-04}) and assigned as rs150089916. While V37M was predicted in silico as benign by SIFT, R181H was classified as deleterious. Based on these preliminary findings the clinical significance of these two additional rare germline variants remains to be determined and at this time is classified as variants of unknown significance.

**Discussion**

*ETV6* encodes an ETS family transcription factor that is frequently rearranged or fused with other genes in human leukemias of myeloid or lymphoid origin[28]. Also known as the *TEL* oncogene, *ETV6* is a sequence specific transcriptional repressor, regulated by auto-inhibition and self-association[43,44]. Descriptions of *ETV6* largely focus on the *ETV6/RUNX1* fusion, which is a product of a t(12;21) chromosomal translocation, the most common genetic abnormality in pediatric ALL[25]. While somatic deletions or mutations in *ETV6* are increasingly
recognized in ALL, nothing is known regarding the impact of germline ETV6 mutations [17,28]. Here we extend the description of the clinical phenotype and functional effects associated with novel germline ETV6 L349P and ETV6 N385fs mutations, both of which reside in the highly conserved ETS DNA binding domain and co-segregate with disease in 2 unrelated kindreds affected by thrombocytopenia and ALL.

In both kindreds ETV6 mutations were inherited in an autosomal dominant manner with variable expression of thrombocytopenia and/or ALL. There was no evidence for parent of origin or sex-delimited expression, as males and females equally transmitted the putative predisposing alleles with associated phenotypes manifesting in daughters as well as sons. Interestingly, in addition to his leukemia, the proband in Kindred 2 exhibited craniofacial and musculoskeletal anomalies (anterior placement of the right ear, downward shaped mouth, joint hypermobility and CNS heterotopias seen on magnetic resonance imaging). No other obvious pathogenic variants were identified in this individual by whole exome sequencing. In addition to atypical physical features, the proband in Kindred 2 developed grade 3 myelosuppression following exposure to anti-metabolite therapy; this feature of chemotherapy hypersensitivity was shared by another patient with T-/myeloid mixed phenotype leukemia and a germline ETV6 mutation (P214L) [17]. In addition, two of the three individuals affected with ALL and harboring ETV6 mutations in the kindreds reported here required bone marrow transplantation, and 1 of the 3 expired from disease, in contrast to the 90% rate of cure with chemotherapy alone in more typical ALL. Whether germline ETV6 mutations might serve as markers for toxicity and outcome will require larger studies controlling for other prognostic variables.

In vitro studies revealed impaired function of the ETV6 mutants identified in both kindreds. While ETV6 L349P and N385fs exhibited normal mRNA levels, both mutations were associated with decreased transcriptional regulation (repression and activation). Structural modeling suggests that both ETV6 mutations would impair transcriptional activity by altering the conformation of the ETV6 protein or truncating it within the DNA binding domain. Interestingly, neither mutant localized to the nucleus. Although the precise mechanism for this behavior remains unclear, it seems likely that these two mutations may affect intracellular transport. Consistent with its putative role as a tumor suppressor, examination of the diagnostic leukemia sample in the proband from Kindred 2 revealed retention of the mutant and deletion of the WT ETV6 allele. Our findings are in agreement with 2 recent reports describing additional ETV6 mutations, including R399C, R369Q[17] and R418G[18] in the ETS DNA binding domain and P214L[17,18], located in a serine-proline phosphorylation motif present in the internal linker domain. In the 3 reports of germline ETV6 mutations to date (including the current series), a mixed phenotype of thrombocytopenia and ALL is observed. An association with elevated MCV was not observed in 3 cases included here, which is in contrast to one of the other recent reports[18]. The 2 additional germline variants reported here in patients with ALL (V37M and R181H) did not impair transcriptional repression of ETV6. While this was expected given that these mutations are not located in or close by the ETS DNA binding domain, we cannot exclude that these variants impair ETV6 function on another functional level.

The discovery of mutations in ANKRD26, RUNX1, and the ETS family transcription factors has led to an increased understanding of the genetic basis of hereditary syndromes involving thrombocytopenia, red cell macrocytosis and leukemia [9,10,17,18] and of the pathways regulated by these genes [17,45]. Constitutional alterations in RUNX1 predispose individuals to thrombocytopenia and hematological malignancies, mainly myelodysplastic syndrome and AML, but also T-ALL [3,9,10,46,47]. Mutations in RUNX1 have been shown to result in either haploinsufficiency or can act in a dominant-negative manner, the latter resulting in an increased risk of hematological malignancies [48,49]. Inherited mutations in ANKRD26...
Germline ETV6 Mutations Predispose to Hematological Disease

[10,45], which is transcriptionally regulated by RUNX1 lead to a similar clinical phenotype, in which thrombocytopenia is often associated with AML and in some cases, with chronic myelogenous leukemia, chronic lymphocytic leukemia and myelodysplastic syndrome [38]. However, there remain additional kindreds affected by thrombocytopenia and/or leukemia that do not demonstrate germline mutations of RUNX1 or ANKRD26. Our data suggest that at least a proportion of these cases result from ETV6 mutations. To date, it is not known whether the ETV6 pathway contributes to non-leukemic cancer phenotypes. We observed no pathogenic germline ETV6 mutations in children with cancers other than ALL in the PCGP. Therefore, the contribution of ETV6 mutations to solid tumor predisposition remains to be determined.

Improved understanding of the heritable nature of childhood cancers has important clinical implications pertaining to genetic counseling and testing of other family members, therapeutic decisions, donor selection for hematopoietic transplantation, and long-term monitoring for therapy-associated or second primary neoplasms [17,50,51]. Evaluation for germline alterations of ETV6 is therefore warranted in families with acute lymphoblastic leukemia, particularly when there is preceding evidence of thrombocytopenia.

Materials & Methods

Patients and controls

All individuals analyzed for purposes of our research were formally consented to Memorial Sloan Kettering Cancer Center’s IRB approved research Protocol, Protocol #00–069, "Ascertainment of Families for Genetic Studies of Familial Lymphoproliferative Disorders", or St. Jude Children’s Research Hospital IRB approved research Protocols NR14-132, "Case report of child with novel ETV6 mutation associated with development of leukemia" and/or NR14-162, “ETV6 germline variants in children with acute lymphoblastic leukemia”, respectively. For Kindred 1, we included 9 affected and 7 unaffected individuals for sequencing. For Kindred 2, we included the proband with ALL, his mother with thrombocytopenia and his unaffected father. For both kindreds, the presence and subtype of leukemia were confirmed by review of pathology reports, while thrombocytopenia was confirmed by medical history.

**Sequencing.** DNA from the proband of Kindred 1 was collected by buccal swab and extracted using the buccal swab DNA isolation kit, (Isohelix, Cat-# DDK-50SK2). DNA from all other family members in Kindred 1 was extracted from saliva using the Oragene DNA extraction kit (DNA Genotek, Cat# OG-250). DNA sequencing of all exons of the leukemia associated genes PAX5, ETV6, HOXA11, CDKN2A, TAL1 and ERG was performed using Sanger sequencing. Clinical exome sequencing (Kindred 2) was performed by Ambry Genetics (Ambry Genetics, Aliso Viejo, CA, USA). To this end, DNA libraries were prepared using 2μg of blood derived DNA (Paired End DNA Sample preparation Kit; Illumina). DNA was fragmented and libraries prepared. Target enrichment was carried out utilizing the TruSeq Exome enrichment Kit, which targets 62 megabases of the human genome. Captured DNA libraries were PCR amplified using the supplied paired end PCR primers.

**Variant assessment.** For Kindred 2, sequence reads were aligned to the human genome reference GRCh37.1 using the Burrows-Wheeler Aligner (BWA) [52], The resulting binary alignment format (BAM files) were jointly called for single nucleotide variants and insertion/deletions (indels) using the Genome Analysis Toolkit (GATK) v.3.1 [53], while structural variations were detected using Clipping Reveals Structure (CREST) [37,54]. Variant level annotations were performed using in-silico tools, such as ANNOVAR[55]. These annotations were used to predict the effects of identified germline variants on gene function and the relevant medical literature was reviewed. Variants were manually reviewed against the medical literature and disease locus specific databases.
**Plasmids.** The pHAGE-CMV-MCS-IRES-ZsGreen lentiviral expression plasmids containing WT human ETV6 and the ETV6 P214L, R369Q and R399C mutants as well as the pGL3-MMP3, pGL3-PF4 and pCS2-Renilla luciferase plasmids were provided by A. Shimamura[17]. ETV6 L349P and N385fs mutants were generated from the WT ETV6 plasmid using QuickChange II XL Site-Directed Mutagenesis Kit (Agilent).

**Cell culture and transfections.** The HeLa cells used in this study (a gift from A.Ventura, MSKCC) were derived from a subculture of the HeLa cell line (ATCC Cat# CCL-2) and subsequently tested for mycoplasma before being used in experiments. These cells were cultured in Dulbecco’s Modified Eagle Medium supplemented with 10%FBS, 1mM L-Glutamine and 1% penicillin-streptomycin. Cell cultures were maintained in a humidified incubator at 37°C in 5% CO₂. Transfections were carried out with FuGENE 6 transfection reagent (Promega) or Lipofectamine 2000 (Life Technologies) according to the manufacturer’s instructions.

**Real-time PCR.** RNA was extracted 24 h after transfection using the RNeasy Mini Kit (Qiagen) and reverse transcribed with the ReadyScript cDNA Synthesis Mix (Sigma-Aldrich). Quantitative Real-time PCR analyses were performed on an ABI PRISM 7900HT Sequence Detection System using the Power SYBR Green PCR Master Mix (Life Technologies) according to the manufacturer’s instructions. Following initial incubation for 10 min at 95°C, amplification was performed for 40 cycles at 95°C for 15 s and 60°C for 1 min. The Rpl32 gene was used as the internal standard and normalization for transfection efficiency was carried out with ZsGreen. Analysis was performed as per the based on the comparative Ct method. The following primer sequences were used:

- **Rpl32 F**, 5'-CATCTCCTTCTCGGCATCA-3';
- **Rpl32 R**, 5'-AACCTGTGGCTCAATGCTC-3';
- **ZsGreen F**, 5'-CTACTTCAAGAAGACTCTGCCC-3';
- **ZsGreen R**, 5'-TCGTGGTACATGCAGTTCTC-3';
- **TRAF1 qRT F**, 5'-AAGATCACCAATGTCACCAGG-3';
- **TRAF1 qRT R**, 5'-GCCATCTCCATTCAGGTACAG-3';
- **EGR1 qRT F**, 5'-CAGCACCTTCAACCCTCAG-3';
- **EGR1 qRT R**, 5'-AGTCGAGTGGTTTGGCTG-3';

**Western blotting.** To isolate protein lysates from nuclear and cytoplasmic subcellular fractions, transfected HeLa cells were lysed 48 hours after transfection (transfection efficiency was measured by ZsGreen positive cells using Guava easyCyte Flow Cytometer) and fractionation was performed using the NE-PER Nuclear and Cytoplasmic Extraction Reagents (Pierce). Samples were run on 4–12% gradient Bis-Tris SDS-PAGE gels, transferred onto PVDF membranes (Bio-Rad) and probed with antibodies against ETV6 (AF7945; 1:200; R&D systems), GAPDH (V-18; 1:200; R&D systems), GAPDH (V-18; 1:200; R&D systems), GAPDH (V-18; 1:200; R&D systems), and Lamin B (C-20; 1:400) (Santa Cruz Biotechnology). Probes were detected using ECL Prime Western Blotting Detection Reagent (GE Healthcare).

**Luciferase assay.** HeLa cells were co-transfected with pHAGE expression constructs, pGL3 reporter constructs and pCS2-Renilla luciferase construct and were harvested 48h after transfection using passive lysis buffer (Promega). Measurement of Firefly and Renilla luciferase expression levels was performed using the Dual-Luciferase Reporter Assay System (Promega) on a GloMax-96 Microplate Luminometer (Promega).

**Supporting Information**

**S1 Fig. Quantitative PCR analysis.** Quantitative PCR analysis using cDNA derived from transiently transfected HeLa cells reveals comparable transcript levels for WT ETV6, the L349P and N385fs mutants identified in the MSKCC and SJCRH kindreds, as well as mutants...
described in a recent separate report (P214L, R369Q, R399C)\textsuperscript{17}.

**Acknowledgments**

We thank the families who contributed samples for this study; we are also grateful to Layla Shahmirzadi (Ambry Genetics) for assistance with whole exome sequencing; Megan Harlan and Rachel Kobos (MSKCC) for contribution to genetic evaluation of Kindred 1 and Christine Odom (SJCRH) for contribution to genetic evaluation of Kindred 2.

**Author Contributions**

Conceived and designed the experiments: ST JV MFW AM DV KEN KO. Performed the experiments: ST JV DV CM YG. Analyzed the data: ST JV KEN KO JZ GW ME MW YG. Contributed reagents/materials/analysis tools: AS. Wrote the paper: ST JV MFW LJ AM DV PG GW RBM EQ HI CH ChP AP ME MYZ PStep PStei KS AL JB SML YG MH ZKS CM MW AS JZ JRD KEN KO. Provided care to the patients and collected/provided phenotypic data: PStep PStei LJ JB MH MW AS.

**References**

1. Malkin D, Nichols KE, Zelley K, Schiffman JD (2014) Predisposition to pediatric and hematologic cancers: a moving target. Am Soc Clin Oncol Educ Book: e44-55.
2. Pui C-H (2012) Childhood leukemias. Cambridge, UK; New York: Cambridge University Press. xi, 880 p., 824 p. of col. plates p.
3. Arepally G, Rebbeck TR, Song W, Gilliland G, Maris JM, et al. (1998) Evidence for genetic homogeneity in a familial platelet disorder with predisposition to acute myelogenous leukemia (FPD/AML). Blood 92: 2600–2602. PMID: 9746808
4. Holmfeldt L, Wei L, Diaz-Flores E, Walsh M, Zhang J, et al. (2013) The genomic landscape of hypodiploid acute lymphoblastic leukemia. Nat Genet 45: 242–252. doi: 10.1038/ng.2532 PMID: 23334668
5. Shah S, Schrader KA, Waanders E, Timms AE, Vijai J, et al. (2013) A recurrent germline PAX5 mutation confers susceptibility to pre-B cell acute lymphoblastic leukemia. Nat Genet 45: 1226–1231. doi: 10.1038/ng.2754 PMID: 24013638
6. Smith ML, Cavenagh JD, Lister TA, Fitzgibbon J (2004) Mutation of CEBPA in familial acute myeloid leukemia. N Engl J Med 351: 2403–2407. PMID: 15575056
7. Stiegitz E, Loh ML (2013) Genetic predispositions to childhood leukemia. Ther Adv Hematol 4: 270–290. doi: 10.1177/2040620713498161 PMID: 23926459
8. Powell BC, Jiang L, Muzny DM, Trevino LR, Dreyer ZE, et al. (2013) Identification of TP53 as an acute lymphocytic leukemia susceptibility gene through exome sequencing. Pediatr Blood Cancer 60: E1–E3. doi: 10.1002/pbc.24417 PMID: 23255406
9. Song WJ, Sullivan MG, Legare RD, Hutchings S, Tan X, et al. (1999) Haploinsufficiency of CBFA2 causes familial thrombocytopenia with propensity to develop acute myelogenous leukaemia. Nat Genet 23: 166–175. PMID: 10508512
10. Noris P, Perrotta S, Seri M, Pecci A, Gnan C, et al. (2011) Mutations in ANKRD26 are responsible for a frequent form of inherited thrombocytopenia: analysis of 78 patients from 21 families. Blood 117: 6673–6680. doi: 10.1182/blood-2011-02-336537 PMID: 21467542
11. Ciocavaco WA, Raskind WH, Kacena MA (2008) Human phenotypes associated with GATA-1 mutations. Gene 427: 1–6. doi: 10.1016/j.gene.2008.09.018 PMID: 18930124
12. Ballmaier M, Schulze H, Strauss G, Cherkaoui K, Wittner N, et al. (1997) Thrombopoietin in patients with congenital thrombocytopenia and absent radii: elevated serum levels, normal receptor expression, but defective reactivity to thrombopoietin. Blood 90: 612–619. PMID: 9226161
13. Go RS, Johnston KL (2003) Acute myelogenous leukemia in an adult with thrombocytopenia with absent radii syndrome. Eur J Haematol 70: 246–248. PMID: 12656750
14. Fujino T, Suzuki A, Ito Y, Oyashiki K, Hatano Y, et al. (2002) Single-translocation and double-chimeric transcripts: detection of NUP98-HOXA9 in myeloid leukemias with HOXA11 or HOXA13 breaks of the chromosomal translocation t(7;11)(p15;p15). Blood 99: 1428–1433. PMID: 11830498
15. Albers CA, Newbury-Ecob R, Ouwehand WH, Ghevaert C (2013) New insights into the genetic basis of TAR (thrombocytopenia-absent radii) syndrome. Curr Opin Genet Dev 23: 316–323. doi: 10.1016/j.gde.2013.02.015 PMID: 23602329
16. Baldini CL, Savoia A (2012) Genetics of familial forms of thrombocytopenia. Hum Genet 131: 1821–1832. doi: 10.1007/s00439-012-1215-x PMID: 22886561
17. Zhang MY, Churpek JE, Keel SB, Walsh T, Lee MK, et al. (2015) Germline ETV6 mutations in familial thrombocytopenia and hematologic malignancy. Nat Genet 47: 180–185. doi: 10.1038/ng.3177 PMID: 25581430
18. Noetzli L, Lo RW, Lee-Sherick AB, Callaghan M, Noris P, et al. (2015) Germline mutations in ETV6 are associated with thrombocytopenia, red cell macrocytosis and predisposition to lymphoblastic leukemia. Nat Genet.
19. Walsh T, Casadei S, Lee MK, Pennil CC, Nord AS, et al. (2011) Mutations in 12 genes for inherited ovarian, fallopian tube, and peritoneal carcinoma identified by massively parallel sequencing. Proc Natl Acad Sci U S A 108: 18032–18037. doi:10.1073/pnas.1115052108 PMID: 22006311
20. Sherry ST, Ward MH, Kholodov M, Baker J, Phan L, et al. (2001) dbSNP: the NCBI database of genetic variation. Nucleic Acids Res 29: 308–311. PMID:11125122
21. Genomes Project C, Abecasis GR, Auton A, Brooks LD, DePristo MA, et al. (2012) An integrated map of genetic variation from 1,092 human genomes. Nature 491: 56–65. doi:10.1038/nature11632 PMID: 23128226
22. Exome Variant Server, NHLBI GO Exome Sequencing Project (ESP), Seattle, WA (URL:http://evs.gs.washington.edu/EVS/).
23. Downing JR, Wilson RK, Zhang J, Mardis ER, Pui CH, et al. (2012) The Pediatric Cancer Genome Project. Nat Genet 44: 619–622. doi:10.1038/ng.2287 PMID: 22641210
24. P. C. Ng SH (2003) SIFT: Predicting amino acid changes that affect protein function. Nucleic Acides Res 31: 3812–3814. PMID:12824425
25. Shurtleff SA, Buijs A, Behm FG, Rubnitz JE, Raimondi SC, et al. (1995) TEL/AML1 fusion resulting from a cryptic t(12;21) is the most common genetic lesion in pediatric ALL and defines a subgroup of patients with an excellent prognosis. Leukemia 9: 1985–1989. PMID:8609706
26. Barbany G, Andersen MK, Autio K, Borgstrom G, Franco LC, et al. (2012) Additional aberrations of the ETV6 and RUNX1 genes have no prognostic impact in 229 t(12;21)(p13;q22)-positive B-cell precursor acute lymphoblastic leukaemias treated according to the NOPHO-ALL-2000 protocol. Leuk Res 36: 936–938. doi:10.1016/j.leukres.2012.03.024 PMID: 22521551
27. De Braekeleer E, Douet-Guilbert N, Morel F, Le Bris MJ, Basinko A, et al. (2012) ETV6 fusion genes in hematological malignancies: a review. Leuk Res 36: 945–961. doi:10.1016/j.leukres.2012.04.010 PMID: 22578774
28. Wang Q, Dong S, Yao H, Wen L, Qiu H, et al. (2014) ETV6 mutation in a cohort of 970 patients with hematologic malignancies. Haematologica 99: e176–178. doi: 10.3324/haematol.2014.104406 PMID: 24997145
29. Parker M, Chen X, Bahrami A, Dalton J, Rusch M, et al. (2012) Assessing telomeric DNA content in pediatric cancers using whole-genome sequencing data. Genome Biol 13: R113. doi:10.1186/gb-2012-13-12-r113 PMID: 23232524
30. Wu G, Diaz AK, Paugh BS, Rankin SL, Ju B, et al. (2014) The genomic landscape of diffuse intrinsic pontine glioma and pediatric non-brainstem high-grade glioma. Nat Genet 46: 444–450. doi:10.1038/ng.2938 PMID: 24705251
31. Zhang J, Ding L, Holmfeldt L, Wu G, Heatley SL, et al. (2012) The genetic basis of early T-cell precursor acute lymphoblastic leukaemia. Nature 481: 157–163. doi:10.1038/nature10725 PMID: 22237106
32. Cancer Genome Atlas Research N (2013) Genomic and epigenomic landscapes of adult de novo acute myeloid leukemia. N Engl J Med 368: 2059–2074. doi:10.1056/NEJMoa1301689 PMID: 23634996
33. Lu C, Zhang J, Nagahawatte P, Easton J, Lee S, et al. (2014) The Genomic Landscape of Childhood and Adolescent Melanoma. J Invest Dermatol.
34. Chen X, Bahrami A, Pappo A, Easton J, Dalton J, et al. (2014) Recurrent somatic structural variations contribute to tumorigenesis in pediatric osteosarcoma. Cell Rep 7: 104–112. doi: 10.1016/j.celrep.2014.03.003 PMID: 24703847
35. Gruber TA, Larson Gedman A, Zhang J, Koss CS, Marada S, et al. (2012) An Inv(16)(p13.3q24.3)-encoded CSF1R/2T3-GLI32 fusion protein defines an aggressive subtype of pediatric acute megakaryoblastic leukemia. Cancer Cell 22: 683–697. doi:10.1016/j.ccr.2012.10.007 PMID: 23155540
36. Zhang J, Mullighan CG, Harvey RC, Wu G, Chen X, et al. (2011) Key pathways are frequently mutated in high-risk childhood acute lymphoblastic leukemia: a report from the Children’s Oncology Group. Blood 118: 3080–3087. doi: 10.1182/blood-2011-03-341412 PMID: 21680795
37. Huether R, Dong L, Chen X, Wu G, Parker M, et al. (2014) The landscape of somatic mutations in epigenetic regulators across 1,000 paediatric cancer genomes. Nat Commun 5: 3630. doi: 10.1038/ncomms4630 PMID: 24710217

38. Robinson G, Parker M, Kranenburg TA, Lu C, Chen X, et al. (2012) Novel mutations target distinct subgroups of medulloblastoma. Nature 488: 43–48. doi: 10.1038/nature11213 PMID: 22722829

39. Wu G, Broniscer A, MeEachron TA, Lu C, Paugh BS, et al. (2012) Somatic histone H3 alterations in pediatric diffuse intrinsic pontine gliomas and non-brainstem glioblastomas. Nat Genet 44: 251–253. doi: 10.1038/ng.1102 PMID: 22286216

40. Zhang J, Benavente CA, McEvoy J, Flores-Otero J, Ding L, et al. (2012) A novel retinoblastoma therapy from genomic and epigenetic analyses. Nature 481: 323–324. doi: 10.1038/nature09737 PMID: 22370222

41. Roberts KG, Li Y, Payne-Turner D, Harvey RC, Yang YL, et al. (2014) Targetable kinase-activating lesions in Ph-like acute lymphoblastic leukemia. N Engl J Med 371: 1005–1015. doi: 10.1056/NEJMoa1403088 PMID: 25207766

42. Chen X, Stewart E, Shelat AA, Qu C, Bahrami A, et al. (2013) Targeting oxidative stress in embryonal rhabdomyosarcoma. Cancer Cell 24: 710–724. doi: 10.1016/j.ccr.2013.11.002 PMID: 24332040

43. Green SM, Coyne HJ 3rd, McIntosh LP, Graves BJ (2010) DNA binding by the ETS protein TEL (ETV6) is regulated by autoinhibition and self-association. J Biol Chem 285: 18496–18504. doi: 10.1074/jbc.M109.096958 PMID: 20400516

44. Wang LC, Swat W, Fujiwara Y, Davidson L, Visvader J, et al. (1998) The TEL/ETV6 gene is required specifically for hematopoiesis in the bone marrow. Genes Dev 12: 2392–2402. PMID: 9694803

45. Bluteau D, Balduini A, Balayn N, Currao M, Nurden P, et al. (2014) Thrombocytopenia-associated mutations in the ANKRD26 regulatory region induce MAPK hyperactivation. J Clin Invest 124: 580–591. doi: 10.1172/JCI71861 PMID: 24430186

46. Mullighan CG, Zhang J, Kasper LH, Lerach S, Payne-Turner D, et al. (2011) CREBBP mutations in relapsed acute lymphoblastic leukaemia. Nature 471: 235–239. doi: 10.1038/nature09727 PMID: 21390130

47. Liew E, Owen C (2011) Familial myelodysplastic syndromes: a review of the literature. Haematologica 96: 1536–1542. doi: 10.3324/haematol.2011.043422 PMID: 21606161

48. Cohen MM Jr. (2009) Perspectives on RUNX genes: an update. Am J Med Genet A 149A: 2629–2646. doi: 10.1002/ajmg.a.33021 PMID: 19830829

49. Matheny CJ, Speck ME, Cushing PR, Zhou Y, Corpora T, et al. (2007) Disease mutations in RUNX1 and RUNX2 create nonfunctional, dominant-negative, or hypomorphic alleles. EMBO J 26: 1163–1175. PMID: 17290219

50. Offit K, Sagi M, Hurley K (2006) Preimplantation genetic diagnosis for cancer syndromes: a new challenge for preventive medicine. JAMA 296: 2727–2730. PMID: 17164459

51. Savage SA, Alter BP (2009) Dyskeratosis Congenital. Hematology-Oncology Clinics of North America 23: 215–+. doi: 10.1016/j.hoc.2009.01.003 PMID: 19327580

52. Li H, Durbin R (2010) Fast and accurate long-read alignment with Burrows-Wheeler transform. Bioinformatics 26: 589–595. doi: 10.1093/bioinformatics/btp698 PMID: 2080505

53. DePristo MA, Banks E, Poplin R, Garimella KV, Maguire JR, et al. (2011) A framework for variation discovery and genotyping using next-generation DNA sequencing data. Nat Genet 43: 491–498. doi: 10.1038/ng.806 PMID: 21478889

54. Wang J, Mullighan CG, Easton J, Roberts S, Heatley SL, et al. (2011) CREST maps somatic structural variation in cancer genomes with base-pair resolution. Nat Methods 8: 652–654. doi: 10.1038/nmeth.1628 PMID: 21666668

55. Wang K, Li M, Hakonarson H (2010) ANNOVAR: functional annotation of genetic variants from high-throughput sequencing data. Nucleic Acids Res 38: e164. doi: 10.1093/nar/gkq603 PMID: 20601685