

## MYELOID NEOPLASIA

**CUX1 is a haploinsufficient tumor suppressor gene on chromosome 7 frequently inactivated in acute myeloid leukemia**

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**Key Points**

- CUX1 is a transcription factor encoded on a region of chromosome 7 that is frequently deleted in high-risk acute myeloid leukemia.
- Haploinsufficiency of CUX1/cut promotes hematopoietic overgrowth in both *Drosophila melanogaster* and human xenograft mouse models in vivo.

Loss of chromosome 7 and del(7q) [–7/del(7q)] are recurring cytogenetic abnormalities in hematologic malignancies, including acute myeloid leukemia and therapy-related myeloid neoplasms, and associated with an adverse prognosis. Despite intensive effort by many laboratories, the putative myeloid tumor suppressor(s) on chromosome 7 has not yet been identified. We performed transcriptome sequencing and SNP array analysis on de novo and therapy-related myeloid neoplasms, half with –7/del(7q). We identified a 2.17-Mb commonly deleted segment on chromosome band 7q22.1 containing CUX1, a gene encoding a homeodomain-containing transcription factor. In 1 case, CUX1 was disrupted by a translocation, resulting in a loss-of-function RNA fusion transcript. CUX1 was the most significantly differentially expressed gene within the commonly deleted segment and was expressed at haploinsufficient levels in –7/del(7q) leukemias. Haploinsufficiency of the highly conserved ortholog, cut, led to hemocyte overgrowth and tumor formation in *Drosophila melanogaster*. Similarly, haploinsufficiency of CUX1

gave human hematopoietic cells a significant engraftment advantage on transplantation into immunodeficient mice. Within the RNA-sequencing data, we identified a CUX1-associated cell cycle transcriptional gene signature, suggesting that CUX1 exerts tumor suppressor activity by regulating proliferative genes. These data identify CUX1 as a conserved, haploinsufficient tumor suppressor frequently deleted in myeloid neoplasms. (*Blood*. 2013;121(6):975-983)

**Introduction**

Loss of chromosome 7 and del(7q) [–7/del(7q)] was first recognized as a frequent event in acute myeloid leukemia (AML) nearly 40 years ago.<sup>1</sup> –7/del(7q) occurs in 8% of de novo AML<sup>2</sup> and 50% of therapy-related myeloid neoplasms (t-MNs).<sup>3</sup> –7/del(7q) is also found in myelodysplastic syndromes, AMLs arising from myeloproliferative neoplasms, the blast phase of chronic myelogenous leukemia, Ph<sup>+</sup> acute lymphoblastic leukemia, and AMLs associated with inherited syndromes.<sup>4-10</sup> –7/del(7q) is an adverse-risk prognostic indicator in myeloid disorders, and the long-term outcome for patients is typically poor. The median overall survival for patients with de novo AML or t-MNs with –7/del(7q) is ~ 6 months.<sup>2,3</sup>

Loss of 1 or more tumor suppressor gene(s) (TSGs) is thought to contribute to leukemic growth in myeloid malignancies with –7/del(7q). Several groups have mapped a commonly deleted segment (CDS) of chromosome band 7q22 using polymorphic markers, conventional cytogenetic analysis, and FISH analysis.<sup>11-13</sup> In one study of 81 patients with malignant myeloid disorders

characterized by chromosome 7 abnormalities, the CDS was mapped to a 2.52-Mb region of 7q22 by FISH using YAC clones.<sup>11</sup> However, deletion of a 2-Mb syntenic region in mice did not result in overt myeloid disease.<sup>14</sup> Other studies have mapped rearrangements involving 7q22 and identified similar<sup>15</sup> or slightly more centromeric intervals (Figure 1).<sup>12,13,16</sup>

In this study, we used single nucleotide polymorphism (SNP) arrays to map the 7q22 CDS with resolution of ~ 1 kb. Overlaying transcriptome-sequencing with copy number aberrations, we identified the gene encoding the transcription factor, CUX1, to be within the CDS and expressed at haploinsufficient levels. Haploinsufficient levels of the CUX1 ortholog, cut (ct), in *Drosophila melanogaster* hemocytes led to hemocyte overproliferation and melanotic tumor formation in vivo. Similarly, decreased expression of CUX1 led to an engraftment advantage for human hematopoietic progenitors transplanted into immunodeficient mice. Thus, from invertebrates to humans, CUX1/cut is a conserved, haploinsufficient hematopoietic TSG.

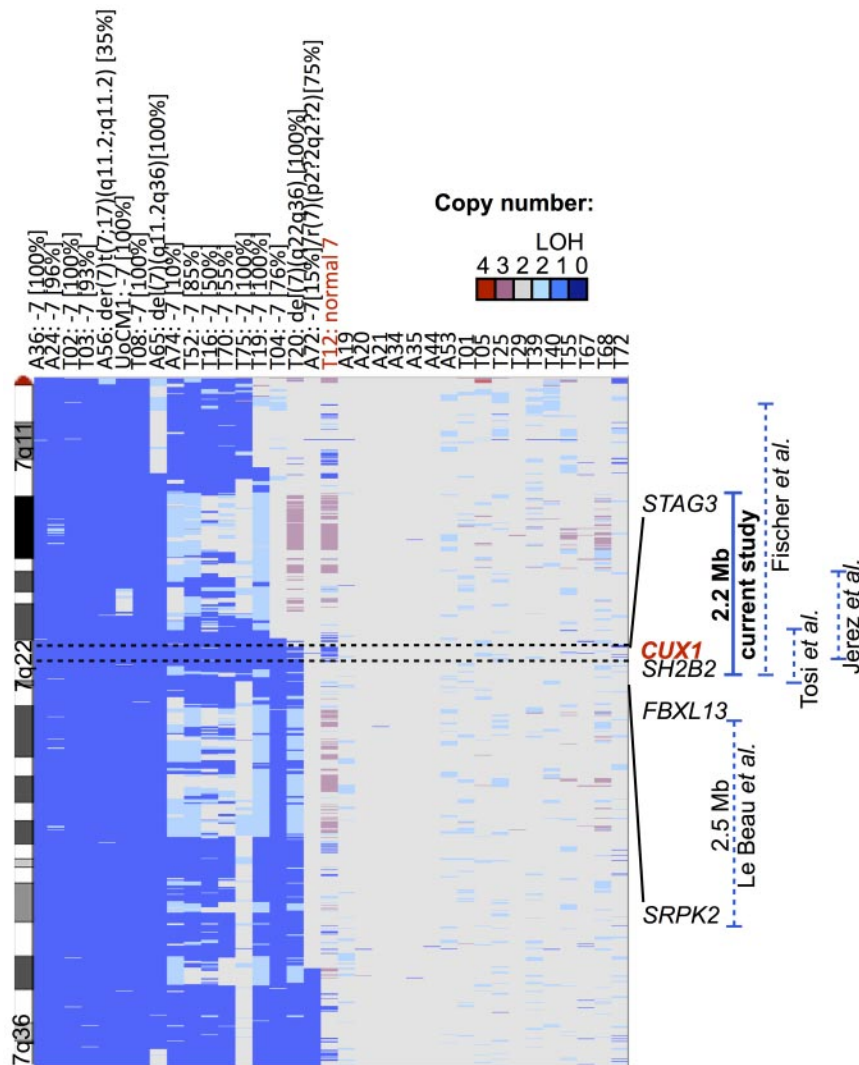
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**Figure 1. *CUX1* is within the 2.17-Mb CDS of 7q22.1.** Copy number analysis of 7q derived from SNP arrays of 35 samples of de novo AML or t-MNs. Thirty-four samples are from primary leukemia samples, and UoCM1 is a cell line derived from the leukemia cells of a patient with de novo AML. Seventeen samples have  $-7/\text{del}(7q)$  detected by cytogenetic analysis as depicted above the sample identifiers. The percentage of cells exhibiting the abnormality is indicated in brackets. T12 has a cryptic deletion in 7q22.1 not visible by conventional cytogenetic analysis. The CDS spans 49 genes; however, only the genes bordering the CDS are shown for clarity. Data are visualized with Integrated Genome Viewer Version 2.0.<sup>50</sup> Previously identified commonly deleted segments are shown on the right panel, and genomic coordinates are indicated in build hg19. LOH indicates loss of heterozygosity.

## Methods

### Patient samples

This research was approved by the University of Chicago Institutional Review Board. Leukemia samples were obtained from the University of Chicago Hematopathology and Cancer Cytogenetics Laboratories (supplemental Table 1, available on the *Blood* Web site; see the Supplemental Materials link at the top of the online article) with informed consent per the Declaration of Helsinki.

### SNP array

DNA was analyzed on Infinium HumanOmni2.5-Quad Version 1.0 DNA Analysis BeadChips (Illumina). Log R ratio and B allele frequencies generated using GenomeStudio (Illumina) were used to identify copy number aberrations by GenoCN.<sup>17</sup> GEO accession number is GSE42482.

### RNA sequencing library preparation

Paired-end libraries were prepared following the standard protocol recommended by Illumina. In brief, purified mRNA (MicroPoly(A) Purist kit, Ambion) was fragmented for 7 minutes at 85°C, and first-strand cDNA generated (Superscript II Reverse Transcriptase, Invitrogen) with random hexamers (Invitrogen), followed by second-strand synthesis with RNaseH and DNA Polymerase I (New England Biolabs). cDNA was repaired and

polished with T4 DNA polymerase, Klenow, and T4 PolyNucleotide Kinase (New England Biolabs), followed by adenosine addition with Klenow Fragment 3' → 5' exo<sup>-</sup> (New England Biolabs). Paired-end adaptors (Illumina) were ligated with DNA T4 ligase, and libraries were amplified with p5 and p7 primers (Illumina) and Platinum Pfx taq polymerase (Invitrogen) for 18 PCR cycles; 450-bp fragments were gel-extracted using QIAquick Gel Extraction kit (QIAGEN).

### Sequence analysis

Paired-end reads of lengths 36-100 bp were generated on the Illumina Genome Analyzer II. Reads were individually trimmed from the 3' end, such that the trimmed 3' bases had an average Phred-scaled quality score < 15. Reads were aligned to the human reference genome hg18 using TopHat (Version 1.3.1)<sup>18</sup> and output in the BAM format.<sup>19</sup> Further alignment manipulation was performed with SAMTools (Version 0.1.18),<sup>19</sup> Picard (Version 1.52, <http://picard.sourceforge.net>), and custom perl scripts. Read alignments with mapping qualities < 10 were removed. Alignments were refined with the Genome Analysis Toolkit (Version 1.0.5) Realigner-TargetCreator and IndelRealigner tools.<sup>20</sup> Cufflinks (Version 1.3.0) was used to estimate transcript abundance using RefSeq gene models and output as Fragments Per Kilobase per Million mapped reads (FPKM) with upper-quartile normalization (supplemental Table 2).<sup>21</sup> Statistical tests were performed on log<sub>2</sub>-transformed data after addition of 1 pseudocount. RNA-sequencing data are in the Short Read Archive SRA061655.

## RT-PCR

cDNA was generated using the High Capacity RNA-to-cDNA Master Mix (Applied Biosystems). *CUX1* was amplified with Power SYBR Green PCR Master Mix (Invitrogen) on the ABI Step One Plus (Applied Biosystems). *cut* was amplified from cDNA from hemocytes pooled from 6 larvae per group, using EXPRESS qPCR Supermix (Invitrogen) and detected with a dual-labeled probe (IDT). Primer sequences are provided (supplemental Table 3). Samples were run in triplicate, and expression levels relative to a housekeeping control gene were determined using the comparative  $C_T$  method according to the manufacturer's software.

## Fusion identification

deFuse<sup>22</sup> (Version 0.5.0) was used to identify chimeric transcripts requiring > 7 split reads, > 7 spanning reads, split\_span  $P$  value > 0.2, split position  $P$  value > 0.2, breakpoint homology < 11, and breakpoints in coding regions. To identify reads mapping to the fusion breakpoint junction, exon-exon junctions between the 2 genes were constructed and all reads were mapped to this junction library with BWA (Version 0.5.9).<sup>23</sup> Reads with < 3 mismatches and > 14 bp anchor region around the junction were identified.

## Cell culture

K562, Kasumi-1, KG-1, NB4, Mono7, and U-937 human AML cell lines, and HeLa adenocarcinoma cells were maintained in RPMI 1640-GlutaMAX (Invitrogen) supplemented with 10% fetal bovine serum (Invitrogen), 100 U/mL penicillin, 100  $\mu$ g/mL streptomycin, and 0.25  $\mu$ g/mL amphotericin B (Invitrogen). Cells were grown at 37°C in 5% CO<sub>2</sub>.

## Western blot

A total of 2 million cells were lysed in RIPA with sodium orthovanadate, protease inhibitor cocktail, and phenylmethylsulfonyl fluoride (Santa Cruz Biotechnology). A total of 40  $\mu$ g of supernatant, quantified by Micro BCA kit (Thermo Scientific), was electrophoresed on a 4%-15% Mini-PROTEAN TGX gradient gel (Bio-Rad) and transferred to nitrocellulose. Blots were blocked with Odyssey Blocking buffer (LI-COR) and probed with anti-CUX1 at 1:200 (SC-6327, Santa Cruz Biotechnology), followed by donkey anti-goat AlexaFluor-680 (Invitrogen). Mouse anti- $\beta$ -actin (SC-4778, Santa Cruz Biotechnology) followed by goat anti-mouse IRDye 800CW (LI-COR) was used for normalization. Fluorescence was quantified by Odyssey imaging (LI-COR).

## Fly stocks

Fly stocks were obtained from the Bloomington Drosophila Stock Center. To express the RNAi in larval hemocytes, *hml $\Delta$ -Gal4<sup>24</sup>* or *cg-Gal4<sup>25</sup>* was used as the driver line to cross with the UAS-RNAi and UAS-2xEGFP lines. The following stocks were used: *cr<sup>RNAi A</sup>*, *cr<sup>RNAi B</sup>*, and luciferase RNAi (nonspecific control). RNAi-negative control larvae were from the crosses of *hml $\Delta$ -Gal4* or *cg-Gal4*, UAS-2xEGFP and *w<sup>1118</sup>*. All the crosses were set up at 25°C on standard cornmeal media.

## Larval hemocyte counts

Circulating hemocytes from wandering third instar larvae were collected by opening the larval cuticles in 20  $\mu$ L PBS. A total of 1  $\mu$ L of hemocytes was fixed in 4% paraformaldehyde (Electron Microscopy Sciences), stained with 10 ng/mL Hoechst 33342 (Invitrogen), and counted using EBImage<sup>26</sup> under a DM5000 B microscope (Leica).

## Sanger sequencing

Sanger sequencing of *CUX1* exons was performed on the 35 samples studied by SNP array plus an additional 109 leukemia samples consisting of 57 de novo AMLs, 47 t-MNs, and 5 AML cell lines (K562, Kasumi-1, KG-1, NB4, and U-937). Whole genome amplified DNA (REPLI-g, QIAGEN) was amplified using Taq 5X Master Mix (New England Biolabs) with primers for *CUX1*.<sup>27</sup> PCR product was treated with 3.3 units Exonuclease I and 0.3 units Shrimp Alkaline Phosphatase (Affymetrix) for 30 minutes

at 37°C followed by 15 minutes at 80°C. Sequencing traces were analyzed using MutationSurveyor Version 4.00 software (SoftGenetics).

## Xenografts

Lentiviral GFP-IRES-shRNA pGIPZ vectors purchased from Thermo Scientific are as follows: *CUX1* shRNA A (clone V2LHS\_151077), *CUX1* shRNA B (clone V2LHS\_151078), and nonspecific shRNA control. Lentivirus was produced by transient transfection of pGIPZ, pCMV-dR8.74, and pMD2 VSVG in HEK293T cells and ultracentrifugation of supernatant. Human cord blood was collected at the University of Chicago. Hematopoietic stem cells were purified from Ficoll gradient-separated white blood cells by Lineage Cell Depletion Kit (Miltenyi Biotec) and cultured for 24 hours in growth media: Stemline II media (Sigma-Aldrich) supplemented with 50 ng/mL each of IL-3, IL-6, thrombopoietin, SCF, and Fms-like tyrosine kinase 3 ligand (R&D Systems). Transduction was performed by spinning cells on Retronectin-coated plates (Takara) with viral stock in Stemline II media at 2500 rpm for 90 minutes at 30°C on day 1 and repeated on day 2. After the spin, supernatant was removed and replaced with growth media. On day 3, cells were assayed for GFP expression by flow and  $1 \times 10^5$  cells were injected via facial vein in 1- to 2-day-old NOD.Cg-Prkdc<sup>scid</sup>IL2rg<sup>tm1Wjl</sup>/SzJ (NOD scid gamma, NSG, The Jackson Laboratory) pups after irradiation with 115 cGy. Retro-orbital blood was collected and stained with the following antibodies (eBioscience): anti-mouse CD45-allophycocyanin-Cy7, anti-human CD45-peridinin chlorophyll protein, anti-human CD13-PE, anti-human CD33-allophycocyanin, and anti-human CD3-PE-Cy7. Flow cytometry was performed on an LSR II (BD Biosciences) and analyzed by FlowJo (TreeStar). Sorting was performed on a FACSAria II (BD Biosciences).

## K562 transfection

K562 cells were transfected with pGIPZ-shRNA by Nucleofector (Lonza) per the manufacturer's instructions and stably selected with 3  $\mu$ g/mL puromycin (Invitrogen).

## Statistical analysis

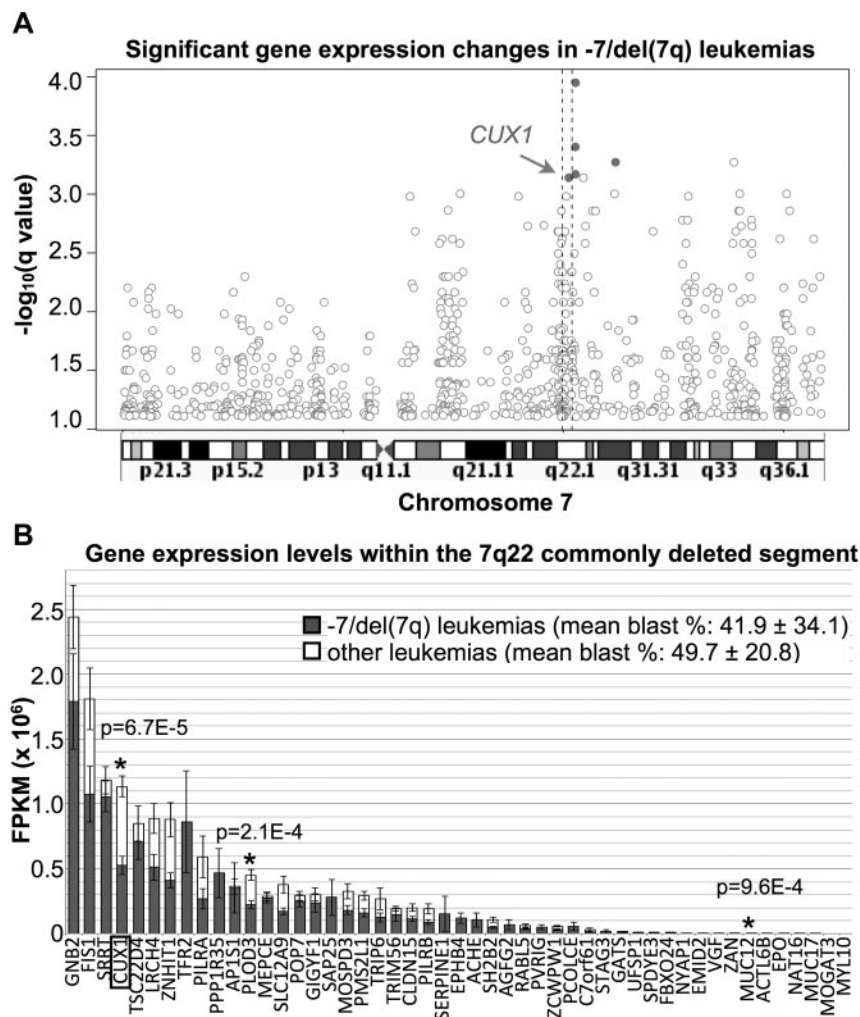
*CUX1*-associated genes were identified by regressing the expression level of each gene against the expression level of *CUX1* using the function lm in the statistical package R. The significance of regression coefficients was quantified by  $t$  test. False discovery rates (FDR) were estimated by the Storey  $q$ -value method.<sup>28</sup> Unless otherwise noted,  $\pm$  indicates SD, and  $P$  values were calculated by  $t$  test.

## Results

### *CUX1* resides within a commonly deleted segment of chromosome band 7q22

We performed SNP-array analysis of 35 de novo and t-MN leukemia samples and defined a 2.17-Mb CDS on chromosome band 7q22 (Figure 1). Cytogenetic analysis revealed clonal  $-7\text{del}(7q)$  in 17 of these. None of the samples showed biallelic deletions of chromosome 7. As seen in previous reports,<sup>11-13</sup> most samples had large deletions of chromosome 7 and were uninformative for defining the CDS of 7q22. One sample, T12, was found to have an interstitial deletion encompassing 7q22 not visible by cytogenetic analysis (Figure 1). Samples T12 and T20 (with a  $\text{del}(7)(q22q36)$ ) had similar proximal breakpoints and defined the proximal boundary, and sample T12 defined the distal boundary of the 2.17-Mb CDS (Figure 1). Although this CDS is defined by a small number of patients, this region overlaps with prior studies.<sup>12,13,16</sup> The CDS within the current study is only 500 kb centromeric to the CDS identified in Le Beau et al<sup>11</sup> (Figure 1). The relatively lower resolution of the 500- to 1.5-Mb YAC probes used in this earlier study may have contributed to imprecision in defining the borders of the deleted segment.





**Figure 2. The most significantly decreased transcripts, genome-wide, in -7/del(7q) leukemias reside on chromosome band 7q22.1, and *CUX1* is the most significantly decreased within the CDS.** (A) Genes on chromosome band 7q22.1 are the most significantly altered in leukemias with -7/(del)7q. Gene expression levels quantified by FPKM from RNA-sequencing were compared between patient samples with loss of -7/del(7q) (n = 11) and all other leukemias (n = 12). Only genes from chromosome 7 are shown. Each circle represents 1 gene plotted on the x-axis by the 5' genomic coordinate. The y-axis indicates the  $-\log_{10}(q \text{ value})$  of the comparison in gene expression levels between -7/del(7q) and other samples (Wilcoxon rank test). Solid circles indicate the top 5 most significant genes (genome-wide) with at least a 1-fold change in expression level. Hyphenated lines represent the CDS as defined by SNP-array analysis in Figure 1. (B) *CUX1* transcripts are the most significantly decreased of the products of genes residing within the CDS. The expression levels of 49 RefSeq genes within chromosome 7:99,807,993-101,974,292 (hg19) are shown. Patient samples are divided into those with loss of 7q22 (gray bars) and others (white bars). Genes are ranked on the x-axis by the mean expression level of the gene in all samples. The graph is shown on a linear scale to illustrate relative expression differences between -7/del(7q) versus other leukemias. Error bars represent SEM. \*Significant P value after Bonferroni correction (Wilcoxon rank test). The nominal P value is shown.

### *CUX1* is expressed at haploinsufficient levels in -7/del(7q) leukemias

To identify genes that are differentially expressed in samples with -7/del(7q), RNA-sequencing was performed on 23 of the leukemia samples analyzed by SNP array. Eleven of the sequenced samples have loss of 7q22. Although sample A72 was reported to have -7[15%]t(7)(p22q22)[75%] by cytogenetic analysis, we categorized it as having diploid 7q22 based on the SNP array (Figure 1). The percentage of blasts was similar in the 2 groups with  $41.9\% \pm 30.1\%$  and  $49.7\% \pm 20.8\%$  blasts in -7/del(7q) and other leukemias, respectively. Alignment statistics are provided in supplemental Table 4.

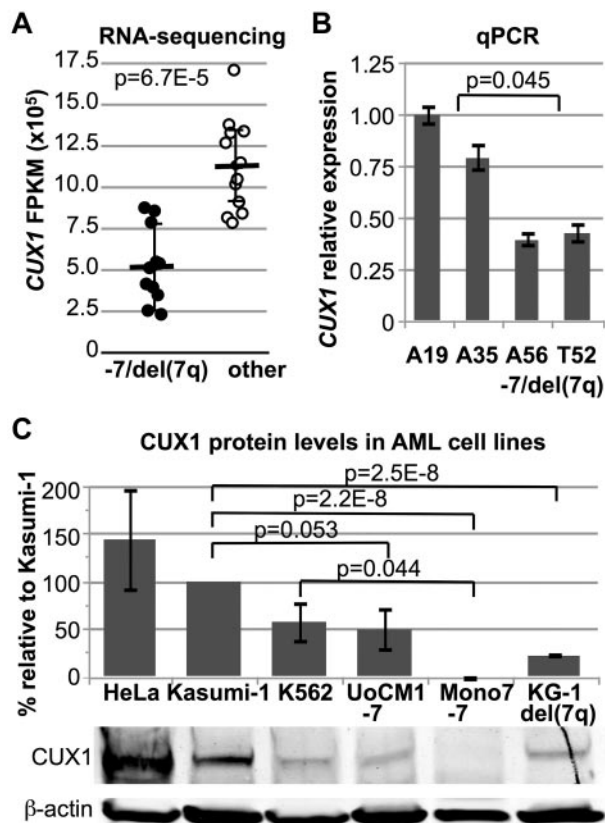
We performed an unbiased, genome-wide comparison of transcript levels in samples with -7/del(7q) versus other samples to define regions of haploinsufficient gene expression. Across all 20 139 expressed transcripts, 4 of the 5 most significantly, differentially expressed transcripts with > 1 fold change are located on chromosome band 7q22.1 and include *CUX1* (Figure 2A). The RNA-sequencing dataset thereby provides an independent validation of the CDS defined by SNP-array (Figure 1). The 2.17-Mb CDS contains 49 RefSeq genes, 3 of which exhibited significantly decreased expression after Bonferroni correction (Figure 2B). *CUX1* is the most significantly differentially expressed transcript within the CDS.

*CUX1* transcript levels in samples with a single *CUX1* allele are present at half the level ( $46.3\% \pm 20.2\%$ ) of samples with 2 copies

of the gene (Figure 3A) consistent with a recent report.<sup>16</sup> Transcript expression was confirmed by quantitative RT-PCR. Leukemia samples with loss of 7q22 express  $45.8\% \pm 4.7\%$  of the level of *CUX1* expressed by other samples (Figure 3B). Loss of 1 copy of *CUX1* is also associated with haploinsufficient protein levels in human AML cell lines (Figure 3C).

### The *CUX1* gene is disrupted by a translocation resulting in a chimeric transcript

As an orthogonal dataset to identify the putative TSG on chromosome 7, RNA-sequencing data were mined for chimeric fusion reads involving genes within the CDS. Nine high-confidence fusion events were identified: 4 validated by Sanger sequencing, 2 were false positives, and there was insufficient patient material to confirm the remaining (Table 1; supplemental Table 5). *MLL-MLL1* and *USP42-RUNX1* fusions have been previously described in AML.<sup>29</sup> Three expressed fusion events were identified in sample T20, 1 of which contains the first exon of *CUX1* (7q22) upstream of exons 2-4 of *CLDN7* (chromosome band 17p13.1), in the same orientation (Figure 4). The resulting transcript is predicted to contain the codons for the first 21 amino acids of *CUX1*, followed by 8 amino acids in the wrong reading frame from *CLDN7*, followed by a premature stop codon, thus truncating 98.6% of the *CUX1* transcript. The second allele of *CUX1* in T20 probably remains intact based on RNA-sequencing (data not shown) and RT-PCR



**Figure 3. CUX1 RNA and protein levels are haploinsufficient in AML cells with loss of 7q22.** (A) CUX1 expression levels in RNA-sequenced leukemia samples. Samples are divided into those with  $-7/del(7)(q22)$  (●) and other samples (○). Each circle represents an individual sample. P value calculated by Wilcoxon rank test. (B) Quantitative RT-PCR of CUX1 in RNA from primary patient samples. A19 and A35 have a normal karyotype, and A56 and T52 have  $-7/del(7q)$ . One representative experiment of 3 independent experiments is shown. Samples with  $-7/del(7q)$  have  $45.8\% \pm 2.4\%$  of the level of CUX1 expressed by other samples ( $P = .045$ , comparing the mean within 1 experiment, A19 and A35 vs A56 and T52). (C) Protein lysates from human AML cell lines were probed for CUX1 protein by Western blot. UoCM1, Mono7, and KG-1 have  $-7/del(7q)$  karyotypes. HeLa, K562, and Kasumi-1 cell lines were used as controls. The full-length p180 protein isoform of CUX1 is shown. The same blot was probed for and normalized to  $\beta$ -actin. Error bars represent  $\pm$  SEM from 3 independent experiments.

(Figure 4B). The level of CUX1 in T20 is 47.6% of the level in samples without loss of 7q22 as estimated by FPKM.

deFuse<sup>22</sup> did not identify CUX1 in fusion events in any of the other RNA-sequenced samples. Sanger sequencing of CUX1 exons in 144 AML/t-MNs samples, 39 of which have  $-7/del(7q)$ , did not reveal any somatic mutations (data not shown). Similarly, there are no mutations in CUX1 reported in 200 AML samples by the Cancer Genome Atlas (tcga-data.nci.nih.gov, Level 2 DNA sequencing somatic mutations, Version 2.2.0). There is 1 reported case of CUX1 mutation acquired during leukemic transformation of a myeloproliferative neoplasm.<sup>30</sup> Thus, CUX1 function is disrupted

primarily by chromosome loss or deletion (Figure 1) and infrequently by a translocation event (Figure 4) or mutation.<sup>30</sup>

**CUX1 is associated with a promitotic transcriptional gene signature in leukemia**

CUX1 is a compelling candidate TSG: it is a homeodomain-containing transcription factor that regulates cell cycle progression and apoptosis and has been implicated in tumorigenesis of solid tumors.<sup>31</sup> Consistent with a TSG, we uncovered a CUX1-associated cell cycle program within the RNA-sequencing data. Of the 662 genes significantly associated with CUX1 expression ( $FDR < 0.20, P < .0083$ ), there is a substantial enrichment for the mitotic cell cycle pathway (DAVID, GO:0000278, Benjamini  $P = 2.57 \times 10^{-5}$ ).<sup>32</sup>

CUX1-associated genes replicate in an independent microarray dataset of 38 subtypes of normal human hematopoietic cells.<sup>33</sup> There is a significant overlap ( $P = .0063$ , permutation test) between the top 1000 CUX1-associated genes ( $FDR < 1.58 \times 10^{-10}$ ) in normal hematopoietic cells and CUX1-associated genes in leukemia cells (2300 genes at  $P < .05$ ). 73.8% of the 126 overlapping genes have a concordant direction (Pearson  $\chi^2$  test  $P = 6.24 \times 10^{-7}$ ). CUX1-associated genes in normal cells are also enriched for cell cycle (GO:0007049, Benjamini  $P = .017$ ). Thus, the CUX1-associated cell growth transcriptional profile is present in both leukemia and normal cells.

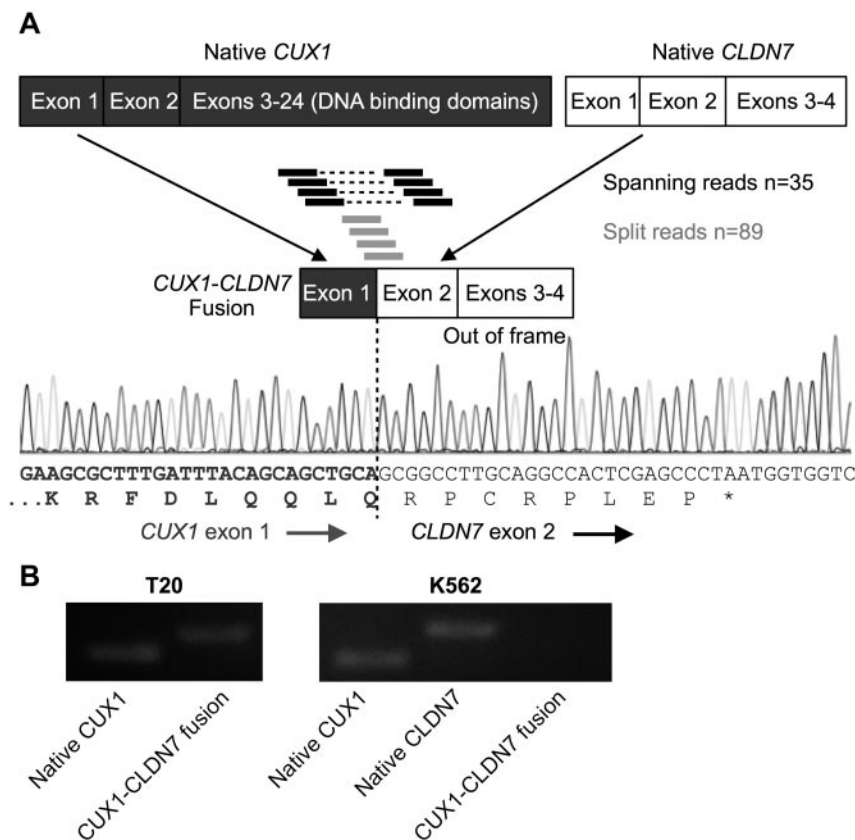
This transcriptional program is probably a result of direct regulation by CUX1, as many of these genes are bound by CUX1. We overlaid the CUX1-correlated leukemia gene set with previously reported CUX1 genome-wide promoter occupancy data from 8 human cancer cell lines, 5 of which are hematopoietic.<sup>34</sup> There is a significant enrichment for direct targets of CUX1 (defined as binding with a  $P < .0005$  in any cell type) within the leukemia CUX1-associated genes ( $P = .0016$ , permutation test). The 125 overlapping genes are also enriched for cell cycle genes (GO:0022403, Benjamini  $P = .05$ , supplemental Table 6). Nine of the 10 cell cycle genes in the overlapping gene set are inversely correlated with CUX1 expression, further implicating CUX1 as a negative regulator of hematopoietic proliferation.

**The CUX1 ortholog, cut, is a conserved hematopoietic TSG in D melanogaster**

The hematopoietic system in Drosophila is composed of hemocytes with functions similar to human myeloid cells.<sup>35</sup> The development of hemocytes involves signaling molecules and transcription factors that are highly conserved in Drosophila and humans, such as GATA2, RUNX1, ZFPM1 (FOG-1), and NOTCH- and WNT-family orthologs.<sup>35</sup> Several genetic lesions in human leukemia result in hemocyte hyperproliferation in Drosophila, including gain-of-function mutations of RAS- and JAK-family genes, and expression of the human RUNX1-RUNX1T1 fusion protein.<sup>35</sup> Most human TSGs have Drosophila orthologs with

**Table 1. Fusions identified in RNA-sequenced samples**

Sample	Gene 1	Gene 2	Gene 1 chromosome	Gene 2 chromosome	PCR validated	Open reading frame
A44	MLL	MLLT1	11q23	19p13	Yes	Yes
T19	KEAP1	ACER1	19p13	19p13	Yes	Yes
T20	CLDN7	CUX1	17p13	7q22	Yes	No
T20	GNAS	PCDH9	20q13	13q21	Insufficient patient material	No
T20	ATP7B	TRPC4	13q14	13q13	Yes	Yes
T55	USP42	RUNX1	7p22	21q22	Insufficient patient material	Yes
T55	PTMA	CXCR4	2q37	2q21	Insufficient patient material	No



**Figure 4.** The *CUX1* gene on 7q22 is disrupted by a fusion event involving *CLDN7* on 17p13.1 resulting in a chimeric transcript in 1 sample. (A) Model of the chimeric fusion transcript. The fusion contains the first exon of *CUX1* upstream of exons 2-4 of *CLDN7*. The fusion was identified by 35 RNA-sequencing reads wherein one end of the paired-end reads maps to one gene whereas the mate-pair maps to the other gene (indicated by black bars connected by hyphenated lines). In addition, 89 junction spanning reads (indicated as single bars) were identified that map across the exon-exon boundaries of the fused genes. Sanger sequencing of the PCR product confirmed the fusion. The *CLDN7* exons are out of frame with respect to the *CUX1* start site, resulting in a premature stop codon. (B) RT-PCR of RNA from t-MN sample T20 confirms the presence of both the *CUX1-CLDN7* fusion transcript and native *CUX1*. K562 cells, which express both endogenous *CUX1* and *CLDN7*, but not the fusion transcript, were used as a control.

evolutionarily conserved networks that regulate proliferation and differentiation; indeed, much of our understanding of TSGs comes from genetic screens in *Drosophila*. Ct is highly conserved between humans and *Drosophila*,<sup>36</sup> and ectopic expression of human *CUX1* can rescue *ct*-deficient phenotypes in *Drosophila*.<sup>37</sup>

To test the hypothesis that *ct* is a TSG in vivo, *ct*, which is expressed in hemocyte progenitors,<sup>38</sup> was targeted by RNAi specifically in developing *Drosophila* hemocytes. Expression of *ct*<sup>RNAi A</sup> in developing hemocytes led to the development of melanotic pseudotumors<sup>35</sup> in both third instar larvae (Figure 5A) and pupae (data not shown). In a small number of cases, tumors were seen as early as the second instar (data not shown). A total of 23.8% (10 of 42) of larvae developed tumors, whereas none (0 of 30) of the control larvae did. To confirm that tumor formation was not the result of an off-target effect of the *ct* shRNA, we tested a second RNAi construct. Expression of *ct*<sup>RNAi B</sup> in hemocytes also led to tumor formation in 16.1% (10 of 62) of larvae.

Melanotic tumors are a phenotype of hemocyte overgrowth<sup>35</sup>; however, we confirmed that there are increased numbers of circulating hemocytes. *ct*<sup>RNAi A</sup> and *ct*<sup>RNAi B</sup> knockdown larvae have significantly increased numbers of hemocytes compared with nonspecific shRNA-expressing larvae (Figure 5B). In addition, we tested a second hemocyte-specific GAL4-driver.<sup>39</sup> Nine of 16 *cg-GAL4/UAS-ct*<sup>RNAi A</sup> larvae developed tumors and had significantly increased numbers of circulating hemocytes (33 737 ± 5719 SE in *cg-GAL4/UAS-ct*<sup>RNAi A</sup>, n = 7, vs 18 826 ± 1984 *cg-GAL4* wild-type, n = 6, *P* = .042, data not shown). Residual expression of *ct* transcripts in hemocytes was 38.3% ± 3.8% (*ct*<sup>RNAi A</sup>, Figure 5C) and 28.9% ± 6.8% (*ct*<sup>RNAi B</sup>) of control, indicating that hemocyte overgrowth was a result of haploinsufficient levels of *ct*.

### *CUX1* is a haploinsufficient TSG in human hematopoietic stem cells

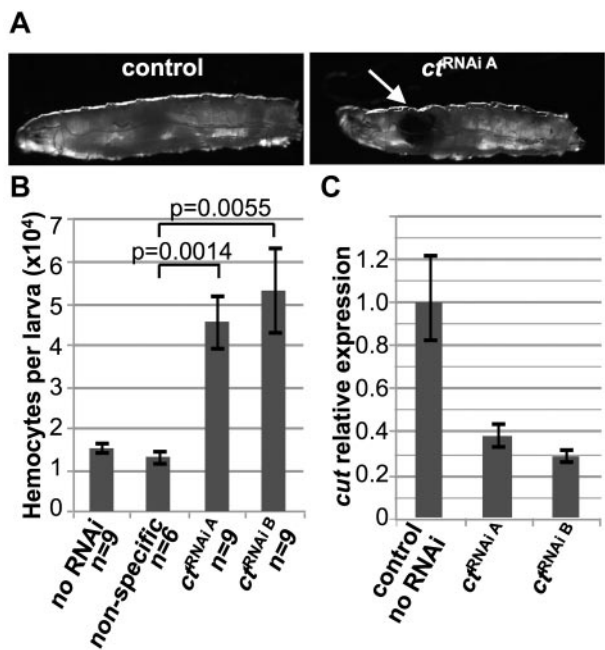
To determine whether *CUX1* is a conserved TSG in humans, we targeted *CUX1* for knockdown in human hematopoietic progenitors. *CUX1* is normally expressed highly in multipotent hematopoietic stem cells (Figure 6A).<sup>33</sup> *CUX1* expression decreases in common myeloid progenitors and varies significantly in terminally differentiated myeloid cells (Figure 6A).

We used 2 different *CUX1*-targeting shRNAs, shRNA A and shRNA B. K562 cells expressing shRNA A or shRNA B retain 24.1% ± 3.7% or 55.3% ± 10.9% residual *CUX1* protein, respectively (Figure 6B). These shRNAs were transduced in a GFP-expressing lentiviral vector into lineage-depleted human cord blood progenitors and transplanted into NOD scid γ (NSG) mice. *CUX1* shRNA-transduced progenitors retained 47.0%-66.9% residual *CUX1* transcripts at the time of transplantation (Figure 6C). Transduction efficiency was similar in all groups (Figure 6D). Total engraftment of all human cells was not significantly different in *CUX1*-targeted recipient mice (Figure 6E). However, there was a significant expansion of the GFP<sup>+</sup> population in the *CUX1*-knockdown xenografts (Figure 6E). Compared with nonspecific shRNA-transduced cells, there is a 40% increase in the percentage of *CUX1*-shRNA-transduced cells. All lineages appeared to be equally expanded in the peripheral blood (Figure 6F), suggesting that *CUX1* TSG activity is not restricted to myeloid cells.

### Discussion

Loss of part or all of chromosome 7 is a frequent event in myeloid malignancies and is associated with a poor prognosis, yet it is





**Figure 5. Haploinsufficiency of the *Drosophila* *CUX1* homologue, *cut*, leads to melanotic tumor formation and hemocyte hyperproliferation in vivo.** *cut* knockdown was performed by crossing flies expressing *ct* RNAi in a construct containing a UAS promoter and GFP tag with flies expressing the GAL4 driver under the hemocyte-specific *hml* promoter. Two different *ct* targeting constructs were tested: *ct*<sup>RNAi A</sup> and *ct*<sup>RNAi B</sup>. (A) *ct*<sup>RNAi A</sup> but not wild-type (*w*<sup>1118</sup>; *hml*-GAL4,UAS-GFP/+), *Drosophila* third instar larvae develop melanotic tumors (indicated by the arrow). (B) *cut* knockdown leads to increased numbers of hemocytes. Circulating hemocytes bled from wild-type, nonspecific RNAi, *ct*<sup>RNAi A</sup>, and *ct*<sup>RNAi B</sup> developing larvae were stained with Hoechst and quantified (mean  $\pm$  SEM). (C) Haploinsufficient levels of *cut* are confirmed by quantitative RT-PCR of *cut* from hemocyte RNA. One representative experiment of 3 is shown.

unknown how these deletions contribute to leukemogenesis. We used high-density SNP arrays and transcriptome sequencing to identify a transcription factor gene, *CUX1*, as a haploinsufficient TSG within a CDS of chromosome 7. We also characterized a t-MNs sample with an inactivated *CUX1* gene resulting from a translocation (Figure 4). *CUX1* is normally highly expressed in multipotent hematopoietic progenitors and exhibits dynamic dosage changes during the course of differentiation (Figure 6A). Deletion of a single allele of *CUX1* is associated with haploinsufficient transcript and protein levels (Figure 3). To test the TSG activity of *CUX1*, we first used *Drosophila* to demonstrate that haploinsufficient levels of the highly conserved *Drosophila* homologue, *cut*, leads to hemocyte overproliferation and melanotic tumor formation in developing larvae (Figure 5). In a second in vivo model, partial knockdown of *CUX1* in human hematopoietic progenitors leads to increased hematopoietic engraftment on transplantation into NSG mice (Figure 6). These findings indicate that *CUX1/cut* is a conserved, haploinsufficient TSG that is essential for the regulation of normal hematopoietic cell growth. A proliferative gene signature was identified within *CUX1*-associated genes, a finding that was replicated in an independent dataset of normal hematopoietic cells (data not shown). A significant number of these genes are direct targets of *CUX1* binding.<sup>34</sup> This suggests that *CUX1* may act as a TSG in myeloid cells by regulating cell cycle components at the transcriptional level.

Deletions of chromosome 7 are typically large with heterogeneity in the breakpoints in myeloid diseases, making it difficult to map the CDS. It is probable that multiple TSGs on chromosome 7 cooperate in leukemogenesis. There may be at least 3 different

CDS on chromosome 7, including an interval at 7q36 containing the *EZH2* gene.<sup>4,16</sup> A precedent for this is seen in 5q deletions, which are also heterogeneous.<sup>4</sup>

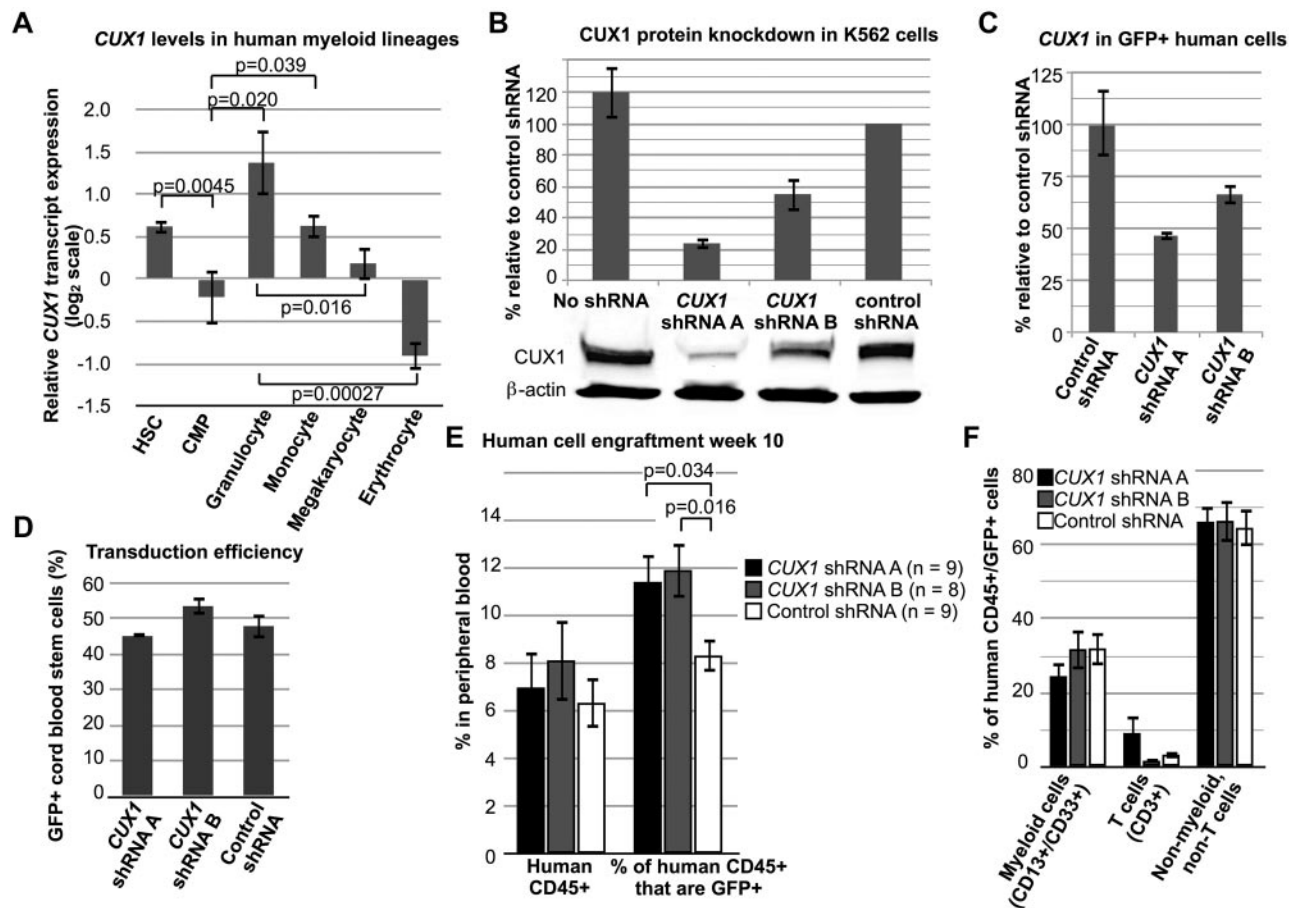
We have not excluded the possibility that other differentially expressed genes within the CDS of 7q22.1 may also play a role in disease pathogenesis. However, *CUX1*, which was implicated as early as 1996,<sup>40</sup> was identified in a narrower CDS in 2 recent studies of AML arising from myeloproliferative neoplasms.<sup>7,41</sup> Four cryptic deletions were identified: 3 samples had deletions that spanned just the *CUX1* and *SH2B2* genes,<sup>41</sup> and 1 sample had a deletion containing only *CUX1*.<sup>7</sup> The CDS defined in the current study is larger, most probably because of smaller sample size.

We note that biallelic inactivation of *CUX1* appears to be infrequent (Figure 1).<sup>7,41</sup> The remaining *CUX1* allele is unlikely to be silenced by epigenetic modifications, as samples with  $-7/\text{del}(7q)$  express *CUX1* at  $46.3\% \pm 20.2\%$  of the level of samples with both copies of *CUX1* (Figure 3), consistent with a gene dosage effect and expression from the remaining allele. However, rare instances of homozygous deletions of the *CUX1* locus and a *CUX1* homeodomain mutation in a region of acquired uniparental disomy have been reported in AML arising from myeloproliferative neoplasms.<sup>7,30</sup> Mutations in *CUX1* were not demonstrated by Sanger sequencing of *CUX1* exons in 144 AML/t-MNs samples, 39 with  $-7/\text{del}(7q)$  (data not shown). A previous study also failed to find *CUX1* mutations in pediatric myeloid neoplasias with  $-7$ .<sup>42</sup> *CUX1* is therefore most often inactivated by chromosome loss or deletion but can infrequently be inactivated by mutation<sup>30</sup> or translocation (Figure 4). As biallelic inactivation of *CUX1* occurs only rarely in myeloid diseases, *CUX1* appears to be a haploinsufficient TSG.

Studies in murine models support the hypothesis that *CUX1* is a myeloid TSG in mammals. *Cux1*<sup>ΔHD</sup> mice, which have a *Cux1* truncation, have increased myeloid cell numbers and myeloid colony-forming units.<sup>43</sup> Transgenic expression of the p75 isoform of *Cux1*, containing 1 Cut repeat and the homeodomain, leads to a myeloproliferative disease in 33% of mice.<sup>44</sup> It is not clear whether the myeloid expansion is the result of increased *Cux1* activity or a dominant negative effect of the transgene.

*CUX1/cut* was first described in *Drosophila*, wherein *cut* is essential for lineage specification in multiple tissues.<sup>31</sup> Consistent with a developmental role in several cell types, altered *Cux1* levels led to abnormal growth of multiple cell types in murine models.<sup>31</sup> In humans, *CUX1* mutations have been reported in 6.7% of lung squamous cell carcinoma<sup>45</sup> and 2.8% of colorectal carcinoma<sup>46</sup>; however, the functional significance of these mutations remains to be tested. The TSG activity of *CUX1* may have several mechanisms, including regulation of differentiation, cell cycle progression, genomic stability, and apoptosis, and the mechanism may be cell type specific.<sup>31</sup> In one report, *cut* deletion in a *Drosophila* cancer model promoted invasive tumorigenesis because of both a block in differentiation and increased proliferation.<sup>47</sup>

In conclusion, this is the first biologic confirmation of a haploinsufficient myeloid TSG on chromosome band 7q22. The concentrations of hematopoietic transcription factors play a critical role in hematopoietic cell fate, regulating lineage decisions, proliferative signals, and leading to leukemia when concentrations are above or below crucial thresholds. Examples of the detrimental consequences of abnormal gene product dosage can be found in many master regulators of hematopoiesis, including *SPI1*, *RUNX1*, and *CBFB*.<sup>48</sup> Intriguingly, these genes are typically not completely inactivated when mutated in leukemia but instead retain some level of activity.<sup>48</sup> *CUX1* levels during normal human hematopoiesis



**Figure 6. Knockdown of *CUX1* leads to increased engraftment of human hematopoietic cells in immunodeficient mice.** (A) *CUX1* is highly expressed in normal human hematopoietic stem cells. The *CUX1* transcript expression profile was extracted from published microarray data.<sup>33</sup> Hematopoietic stem cells (HSC, lineage<sup>-</sup>, CD133<sup>+</sup>, CD34<sup>dim</sup>, n = 10), common myeloid progenitors (CMP, lin<sup>-</sup>, CD34<sup>+</sup>, CD38<sup>+</sup>, IL-3R<sup>lo</sup>, CD45RA<sup>-</sup>, n = 4), granulocytes (FSC<sup>hi</sup>, SSC<sup>hi</sup>, CD16<sup>+</sup> CD11b<sup>+</sup>, n = 4), monocytes (FSC<sup>hi</sup>, SSC<sup>lo</sup>, CD14<sup>+</sup>, CD45<sup>dim</sup>, n = 5), megakaryocytes (CD34<sup>-</sup>, CD41<sup>+</sup>, CD61<sup>+</sup>, CD45<sup>-</sup>, n = 6), and erythrocytes (CD34<sup>-</sup>, CD71<sup>-</sup>, GlyA<sup>+</sup>, n = 6). Data were normalized such that the mean of each gene, across all 38 hematopoietic cell types tested, is zero. (B) shRNA constructs targeting *CUX1* lead to haploinsufficient *CUX1* protein levels in K562 cells. Western blot was performed on lysates of K562 cells stably expressing *CUX1* shRNA A, shRNA B, or control shRNA. Data represent the mean  $\pm$  SD from 3 experiments. (C) Residual *CUX1* transcript levels in human hematopoietic stem cells transduced with lentivirus expressing shRNA targeting *CUX1*. Lineage-depleted human cord blood cells were transduced with 2 different shRNA constructs targeting *CUX1* (shRNA A or shRNA B), or a nonspecific control shRNA. RT-PCR was performed on sorted GFP<sup>+</sup> cells on the day of transplantation. Each sample was performed in triplicate, and 1 of 2 experiments is shown. (D) Transduction efficiency was similar across the 3 groups of transplanted cord blood stem cells. Transduction efficiency was determined by flow cytometry for GFP<sup>+</sup> cells before transplantation. (E) Knockdown of *CUX1* leads to increased engraftment of human hematopoietic cells in NSG mice. Peripheral blood from mice transplanted with human xenografts was collected on week 10 after transplantation and tested for human CD45 expression (left bars) and coexpression of human dCD45 and GFP (right bars). The numbers of mice per group are indicated. Data are mean  $\pm$  SEM. (F) Peripheral blood human CD45<sup>+</sup>/GFP<sup>+</sup> cells from *CUX1* shRNA A, *CUX1* shRNA B, or control shRNA transduced xenografts were stained for the lineage markers CD13/CD33 (myeloid cells) and CD3 (T cells).

appear to change significantly, depending on the developmental stage and lineage (Figure 6A). A gradient of Ct has been shown to be important in neural development in *Drosophila*: high, medium, and low levels of Ct, measured at the single-cell level, lead to distinct classes of neural cells.<sup>49</sup> Thus, changes in *CUX1* levels may have important phenotypes during hematopoiesis and deleterious consequences when altered.

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## Authorship

Contribution: M.E.M. designed research, performed experiments, analyzed and interpreted data, and wrote the manuscript; C.D.B. designed research, analyzed, and interpreted RNA-sequencing data, and edited the manuscript; X.W. performed *Drosophila* crosses and hemocyte counting; E.T.B. assisted in analyzing RNA-sequencing data; S.K. assisted in generating sequencing libraries; C.B. identified fusion genes; S.Y. assisted in analyzing SNP data and fusion validation; B.P.S., J.K., and J.M.C. designed and performed xenograft experiments; T.S. assisted in designing research; J.A. performed morphologic analysis and collected biospecimens; M.M.L. performed cytogenetic analysis of leukemia



samples, designed research, collected biospecimens, interpreted data, and edited the manuscript; R.L.G. provided computational infrastructure and bioinformatics support; and K.P.W. designed research, interpreted data, and edited the manuscript.

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## References

- Rowley JD. Deletions of chromosome 7 in haematological disorders [letter]. *Lancet*. 1973; 2(7842):1385-1386.
- Byrd JC, Mrozek K, Dodge RK, et al. Pretreatment cytogenetic abnormalities are predictive of induction success, cumulative incidence of relapse, and overall survival in adult patients with de novo acute myeloid leukemia: results from Cancer and Leukemia Group B (CALGB 8461). *Blood*. 2002;100(13):4325-4336.
- Smith SM, Le Beau MM, Huo D, et al. Clinical-cytogenetic associations in 306 patients with therapy-related myelodysplasia and myeloid leukemia: the University of Chicago series. *Blood*. 2003;102(1):43-52.
- Bejar R, Levine R, Ebert BL. Unraveling the molecular pathophysiology of myelodysplastic syndromes. *J Clin Oncol*. 2011;29(5):504-515.
- Preiss BS, Bergmann OJ, Friis LS, et al. Cytogenetic findings in adult secondary acute myeloid leukemia (AML): frequency of favorable and adverse chromosomal aberrations do not differ from adult de novo AML. *Cancer Genet Cytogenet*. 2010;202(2):108-122.
- Luna-Fineman S, Shannon KM, Lange BJ. Childhood monosomy 7: epidemiology, biology, and mechanistic implications. *Blood*. 1995;85(8):1985-1999.
- Klampfl T, Harutyunyan A, Berg T, et al. Genome integrity of myeloproliferative neoplasms in chronic phase and during disease progression. *Blood*. 2011;118(1):167-176.
- Mitelman F. The cytogenetic scenario of chronic myeloid leukemia. *Leuk Lymphoma*. 1993; 11(Suppl 1):11-15.
- Niemeyer CM, Arico M, Basso G, et al. Chronic myelomonocytic leukemia in childhood: a retrospective analysis of 110 cases. European Working Group on Myelodysplastic Syndromes in Childhood (EWO-GMDS). *Blood*. 1997;89(10):3534-3543.
- Wetzler M, Dodge RK, Mrozek K, et al. Additional cytogenetic abnormalities in adults with Philadelphia chromosome-positive acute lymphoblastic leukaemia: a study of the Cancer and Leukaemia Group B. *Br J Haematol*. 2004;124(3):275-288.
- Le Beau MM, Espinosa R 3rd, Davis EM, Eisenbart JD, Larson RA, Green ED. Cytogenetic and molecular delineation of a region of chromosome 7 commonly deleted in malignant myeloid diseases. *Blood*. 1996;88(6):1930-1935.
- Tosi S, Scherer SW, Giudici G, Czepulkowski B, Biondi A, Kearney L. Delineation of multiple deleted regions in 7q in myeloid disorders. *Genes Chromosomes Cancer*. 1999;25(4):384-392.
- Fischer K, Frohling S, Scherer SW, et al. Molecular cytogenetic delineation of deletions and translocations involving chromosome band 7q22 in myeloid leukemias. *Blood*. 1997;89(6):2036-2041.
- Wong JC, Zhang Y, Lieu KH, et al. Use of chromosome engineering to model a segmental deletion of chromosome band 7q22 found in myeloid malignancies. *Blood*. 2010;115(22):4524-4532.
- Liang H, Fairman J, Claxton DF, Nowell PC, Green ED, Nagarajan L. Molecular anatomy of chromosome 7q deletions in myeloid neoplasms: evidence for multiple critical loci. *Proc Natl Acad Sci U S A*. 1998;95(7):3781-3785.
- Jerez A, Sugimoto Y, Makishima H, et al. Loss of heterozygosity in 7q myeloid disorders: clinical associations and genomic pathogenesis. *Blood*. 2012;119(25):6109-6117.
- Sun W, Wright FA, Tang Z, et al. Integrated study of copy number states and genotype calls using high-density SNP arrays. *Nucleic Acids Res*. 2009;37(16):5365-5377.
- Trapnell C, Pachter L, Salzberg SL. TopHat: discovering splice junctions with RNA-Seq. *Bioinformatics*. 2009;25(9):1105-1111.
- Li H, Handsaker B, Wysoker A, et al. The Sequence Alignment/Map format and SAMtools. *Bioinformatics*. 2009;25(16):2078-2079.
- McKenna A, Hanna M, Banks E, et al. The Genome Analysis Toolkit: a MapReduce framework for analyzing next-generation DNA sequencing data. *Genome Res*. 2010;20(9):1297-1303.
- Trapnell C, Williams BA, Pertea G, et al. Transcript assembly and quantification by RNA-Seq reveals unannotated transcripts and isoform switching during cell differentiation. *Nat Biotechnol*. 2010;28(5):511-515.
- McPherson A, Hormozdiani F, Zayed A, et al. deFuse: an algorithm for gene fusion discovery in tumor RNA-Seq data. *PLoS Comput Biol*. 2011; 7(5):e1001138.
- Li H, Durbin R. Fast and accurate short read alignment with Burrows-Wheeler transform. *Bioinformatics*. 2009;25(14):1754-1760.
- Sinenko SA, Mathey-Prevot B. Increased expression of Drosophila tetraspanin, Tsp68C, suppresses the abnormal proliferation of ytr-deficient and Ras/Raf-activated hemocytes. *Oncogene*. 2004;23(56):9120-9128.
- Yasothornsrikul S, Davis WJ, Cramer G, Kimbrell DA, Dearolf CR. viking: identification and characterization of a second type IV collagen in Drosophila. *Gene*. 1997;198(1):17-25.
- Pau G, Fuchs F, Sklyar O, Boutros M, Huber W. EBIImage: an R package for image processing with applications to cellular phenotypes. *Bioinformatics*. 2010;26(7):979-981.
- Jones S, Zhang X, Parsons DW, et al. Core signaling pathways in human pancreatic cancers revealed by global genomic analyses. *Science*. 2008;321(5897):1801-1806.
- Storey JD. A direct approach to false discovery rates. *J R Stat Soc B*. 2002;64(3):479-498.
- Palanisamy N. Chromosomal translocations in AML: detection and prognostic significance. *Cancer Treat Res*. 2010;145:41-58.
- Thoennissen NH, Lasho T, Thoennissen GB, Ogawa S, Tefferi A, Koeffler HP. Novel CUX1 missense mutation in association with 7q- at leukemic transformation of MPN. *Am J Hematol*. 2011; 86(8):703-705.
- Hulea L, Nepveu A. CUX1 transcription factors: From biochemical activities and cell-based assays to mouse models and human diseases. *Gene*. 2012;497(1):18-26.
- Huang da W, Sherman BT, Lempicki RA. Systematic and integrative analysis of large gene lists using DAVID bioinformatics resources. *Nat Protoc*. 2009;4(1):44-57.
- Novershtern N, Subramanian A, Lawton LN, et al. Densely interconnected transcriptional circuits control cell states in human hematopoiesis. *Cell*. 2011;144(2):296-309.
- Harada R, Vadrnais C, Sansregret L, et al. Genome-wide location analysis and expression studies reveal a role for p110 CUX1 in the activation of DNA replication genes. *Nucleic Acids Res*. 2008; 36(1):189-202.
- Crozatier M, Vincent A. Drosophila: a model for studying genetic and molecular aspects of haematopoiesis and associated leukaemias. *Dis Model Mech*. 2011;4(4):439-445.
- Neufeld EJ, Skalnik DG, Lievens PM, Orkin SH. Human CCAAT displacement protein is homologous to the Drosophila homeoprotein, cut. *Nat Genet*. 1992;1(1):50-55.
- Ludlow C, Choy R, Blochlinger K. Functional analysis of Drosophila and mammalian cut proteins in flies. *Dev Biol*. 1996;178(1):149-159.
- Jung SH, Evans CJ, Uemura C, Banerjee U. The Drosophila lymph gland as a developmental model of hematopoiesis. *Development*. 2005; 132(11):2521-2533.
- Asha H, Nagy I, Kovacs G, Stetson D, Ando I, Dearolf CR. Analysis of Ras-induced overproliferation in Drosophila hemocytes. *Genetics*. 2003; 163(1):203-215.
- Lewis S, Abrahamson G, Boulwood J, Fidler C, Potter A, Wainscoat JS. Molecular characterization of the 7q deletion in myeloid disorders. *Br J Haematol*. 1996;93(1):75-80.
- Thoennissen NH, Krug UO, Lee DH, et al. Prevalence and prognostic impact of allelic imbalances associated with leukemic transformation of Philadelphia chromosome-negative myeloproliferative neoplasms. *Blood*. 2010;115(14):2882-2890.
- Hindersin S, Niemeyer CM, Germing U, Gobel U, Kratz CP. Mutation analysis of CUTL1 in childhood myeloid neoplasias with monosomy 7. *Leuk Res*. 2007;31(9):1323-1324.
- Sinclair AM, Lee JA, Goldstein A, et al. Lymphoid apoptosis and myeloid hyperplasia in CCAAT displacement protein mutant mice. *Blood*. 2001; 98(13):3658-3667.
- Cadioux C, Fournier S, Peterson AC, Bedard C, Bedell BJ, Nepveu A. Transgenic mice expressing the p75 CCAAT-displacement protein/Cut homeobox isoform develop a myeloproliferative disease-like myeloid leukemia. *Cancer Res*. 2006;66(19):9492-9501.
- Hammerman PS, Hayes DN, Wilkerson MD, et al. Comprehensive genomic characterization of squamous cell lung cancers. *Nature*. 2012; 489(7417):519-525.
- Cancer Genome Atlas Network. Comprehensive molecular characterization of human colon and rectal cancer. *Nature*. 2012;487(7407):330-337.
- Zhai Z, Ha N, Papagiannouli F, et al. Antagonistic regulation of apoptosis and differentiation by the cut transcription factor represents a tumor-suppressing mechanism in Drosophila. *PLoS Genet*. 2012;8(3):e1002582.
- Rosenbauer F, Koschmieder S, Steidl U, Tenen DG. Effect of transcription-factor concentrations on leukemic stem cells. *Blood*. 2005;106(5):1519-1524.
- Grueber WB, Jan LY, Jan YN. Different levels of the homeodomain protein cut regulate distinct dendrite branching patterns of Drosophila multidendritic neurons. *Cell*. 2003;112(6):805-818.
- Robinson JT, Thorvaldsdottir H, Winkler W, et al. Integrative genomics viewer. *Nat Biotechnol*. 2011;29(1):24-26.