

ORIGINAL ARTICLE

High levels of the adhesion molecule CD44 on leukemic cells generate acute myeloid leukemia relapse after withdrawal of the initial transforming event

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Multiple genetic hits are detected in patients with acute myeloid leukemia (AML). To investigate this further, we developed a tetracycline-inducible mouse model of AML, in which the initial transforming event, overexpression of *HOXA10*, can be eliminated. Continuous overexpression of *HOXA10* is required to generate AML in primary recipient mice, but is not essential for maintenance of the leukemia. Transplantation of AML to secondary recipients showed that in established leukemias, ~80% of the leukemia-initiating cells (LICs) in bone marrow stopped proliferating upon withdrawal of *HOXA10* overexpression. However, the population of LICs in primary recipients is heterogeneous, as ~20% of the LICs induce leukemia in secondary recipients despite elimination of *HOXA10*-induced overexpression. Intrinsic genetic activation of several proto-oncogenes was observed in leukemic cells resistant to inactivation of the initial transformation event. Interestingly, high levels of the adhesion molecule CD44 on leukemic cells are essential to generate leukemia after removal of the primary event. This suggests that extrinsic niche-dependent factors are also involved in the host-dependent outgrowth of leukemias after withdrawal of *HOXA10* overexpression event that initiates the leukemia.

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Introduction

Leukemia is a heterogeneous disease in which hematopoietic stem and progenitor cells acquire aberrant regulation of prominent transcription factors or genetic lesions that lead to a block in differentiation, increased self-renewal and dysregulated proliferation. The *HOX* family of Homeobox genes encodes transcription factors that regulate hematopoiesis and are expressed in specific compartments of the hematopoietic hierarchy.^{1–3} Several *HOXA* genes have been directly implicated in hematological malignancies as supported by the frequently observed elevation of *HOXA* gene expression in acute myeloid leukemia (AML) patients.^{4–6} A more direct involvement for *HOXA* genes in leukemia is supported by their frequent fusion to the nucleoporin gene *NUP98* in leukemia patients.^{7,8} Transgenic and retroviral transplantation mouse models have established that deregulation of *HOXA* genes^{9–11} or *NUP98-HOXA* fusion

genes^{12,13} initiates AML. *HOX* proteins contain a highly conserved DNA-binding homeodomain flanked by variable sequences that influence the DNA-binding specificity, by coordinating interactions to cofactors such as *PBX* and *MEIS*.^{14,15} The cofactor *MEIS1* has been described to synergize with multiple native *HOXA* and *NUP98-HOXA* fusion genes in promoting a differentiation block of *HOXA*-transformed myeloid progenitors accelerating the initiation of the leukemia in mice.^{9,13,16}

Several AML mouse models have recently identified a role for oncogenes in survival/maintenance of leukemia in addition to the initiation of leukemogenesis. This suggests that the growth of leukemic cells *in vivo* depends on continued activity of the oncogene and that efficient targeting of this activity may be a promising avenue to cure AML. *HOXA9* has been described to be required for survival in human mixed lineage leukemia (MLL)-rearranged acute leukemias.¹⁷ Similarly, a mouse model showed that AML induced by conditional expression of *MLL-ENL* can be cured by oncogene ablation, despite additional acquisition of complex genetic abnormalities.¹⁸

HOXA10 is expressed in the most primitive hematopoietic cell compartment in normal hematopoiesis, and its overexpression increases proliferation of human hematopoietic progenitor cells.¹⁹ By using a tet-operator mouse model where the expression level of *HOXA10* can be tightly regulated, we recently characterized the role of *HOXA10* as a critical regulator for normal hematopoietic stem cells and erythroid/megakaryocyte development *in vivo*.²⁰ In addition, it is known that deregulation of *HOXA10* initiates AML, as a significant proportion of recipient mice transplanted with retroviral vector-transduced *HOXA10*-overexpressing grafts developed AML with a latency of 19–50 weeks.¹¹ Therefore, we asked whether our tet-operator system-driven *HOXA10* mice would develop leukemia after long-term treatment with doxycycline to force expression of *HOXA10*. The development of leukemia was not observed even after doxycycline treatment for 1 year, suggesting that the *HOXA10* expression level generated in these mice might not be optimal for leukemic transformation *in vivo*.²⁰ To develop a mouse model of AML using the tet-operator system-driven *HOXA10* mice, we transduced the *HOXA10*-inducible bone marrow (BM) cells with the cofactor *MEIS1* and showed that *MEIS1* can potentiate leukemogenesis induced by moderate expression of *HOXA10*. However, the long latency period required to induce leukemia suggests that the cooperation with *MEIS1* is not sufficient to transform cells, but that additional hits might be required for full-blown AML to develop.

Recently, a mouse model using inducible expression of the *MLL-ENL* oncoprotein was used to show that withdrawal of induction of this powerful oncoprotein leads to elimination of the malignant clone because it depends entirely on this protein.¹⁸ Similarly, a recent report shows that another powerful

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oncoprotein, *HOXA9*, is required for survival in human MLL-rearranged acute leukemias, suggesting that targeting *HOXA9* may be a therapeutic option.¹⁷ Even if oncoproteins have the potential to drive hematopoietic cells into full malignant progression, the latency period to induce AML can be several months, indicating that additional events are required for leukemogenesis. Combination/cooperation of specific genetic events is known to be involved in the multi-step evolution of premalignant cells to full-blown AML. This concept is supported by clinical evidence from a variety of leukemias, in which multiple genetic hits are often diagnosed in AML patients.^{21–24} Moreover, mouse transplantation models have shown that several genetic changes collaborate to induce AML.^{25–28}

Here, we have developed a mouse model of AML and addressed whether elimination of the initial oncogenic event (*HOXA10* overexpression) caused by an oncoprotein that has a relatively weak activity will lead to elimination of established leukemia *in vivo* or whether acquisition of secondary events by leukemic cells renders them resistant to the removal of the oncoprotein overproduction that initiated the disease. The findings show that in 20% of cases, secondary mutations in additional proto-oncogenes and overproduction of the adhesion molecule CD44 contribute toward rendering the leukemic cells resistant to inactivation of the initial transformation event.

Materials and methods

Generation and screening of transgenic mice

The generation of the *Rosa26-rtTA-nls/tetO-HOXA10* mouse model and its characterization has been described previously.²⁰ Genomic DNA was isolated from tail biopsies and analyzed by polymerase chain reaction (PCR) for the detection of *tetO-HOXA10*²⁹ and *Rosa26-rtTA-nls*.²⁰ Lund University's Ethical Committee approved all animal experiments.

Isolation of BM cells

Femurs and tibias were harvested from 7- to 12-week-old *Rosa26-rtTA-nls/tetO-HOXA10*-positive mice and littermate control mice (Ly5.2). Bones were crushed, the cell suspension was filtered through a 100- μ m pore size Cell Strainer filter (Falcon; Becton Dickinson, Franklin Lakes, NJ, USA) and the lineage-negative (Lin⁻) population was depleted from BM cells, with a cocktail of antibodies: rat anti-mouse-B220, -Gr1, -Mac1, -CD4, -CD5, -CD8 and -Ter119 (Becton Dickinson). Anti-rat immunoglobulin G-conjugated immunomagnetic beads (DynaL Biotech, Oslo, Norway) were used for enrichment of the Lin⁻ population. For prestimulation before the retroviral transduction, cells were cultured in X-Vivo medium for 48 h, supplemented with 100 U/ml PenStrep (Sigma, St Louis, MO, USA), 1% glutamine, 1% bovine serum albumin, 50 ng/ml murine stem cell factor, 10 ng/ml murine interleukin-3 and 50 ng/ml human interleukin-6.

Retroviral transduction and BM transplantation

The *MEIS1* retroviral vector expressing both the *MEIS1* cofactor together with green fluorescent protein (GFP) was provided by Dr Corey Largman (Department of Medicine, University of California, San Francisco, CA, USA) and Dr Jeffrey Lawrence (UCSF, Roche Molecular Systems, San Francisco, CA, USA). Vectors were transfected into Phoenix Ampho cells (Nolan Lab, Stanford University, Palo Alto, CA, USA), and supernatants were harvested for transduction of *Rosa26-rtTA-nls/tetO-HOXA10*

Lin⁻ cells. A total of 2×10^5 transduced Lin⁻ cells (Ly5.2) were mixed with 2×10^5 support BM cells (Ly5.1) and transplanted into the tail vein of lethally irradiated (900 cGy) C57BL/6 \times B6SJL recipient mice (Ly5.1/Ly5.2). Recipient mice were continuously fed doxycycline (0.2 g/ml) in the drinking water to induce the expression of *HOXA10* or ciproxine as a negative control.²⁰ For the limiting dilution assay, leukemic cells (Ly5.2) were transplanted in escalating doses: 1, 2 or 8×10^3 cells with 2×10^5 support BM cells (Ly5.1) into lethally irradiated secondary recipient mice. To study the role of CD44, 2×10^4 leukemic cells were transplanted with 2×10^5 support BM cells into lethally irradiated secondary recipient mice. Different populations of the CD44 surface marker of leukemic cells from primary recipient mice were sorted (DiVa; Becton Dickinson) and sorted cells were injected in the tail vein of lethally irradiated secondary recipient mice fed doxycycline or ciproxine. For inhibition of CD44, leukemic cells were incubated during 20 min with high dose of the anti-CD44 phycoerythrin (PE) (20 μ g/ml) or control anti-immunoglobulin G2a PE (20 μ g/ml) before transplantation.

Flow cytometry

Engraftment and expansion of the transduced population (Ly5.2) was monitored by gating Ly5.1-negative cells using fluorescence-activated cell sorting (FACS) with PE-CD45.1 antibody and measuring the proportion of GFP⁺ cells. The phenotype of the leukemia was analyzed in peripheral blood or BM cells by staining with Mac1 adenomatous polyposis coli and c-Kit PE antibodies (Becton Dickinson). 7-Amino-actinomycin D was used for the detection of non-viable cells (Sigma). Stained cells were run throughout a FACS Calibur (Becton Dickinson Biosciences, Sparks, MD, USA) and the subsequent data were analyzed with the FlowJo software (Tree Star, Ashland, OR, USA). For the lineage staining analysis, the Lin⁻ population from BM cells was stained with a cocktail of antibodies: rat anti-mouse -B220, -Gr1, -Mac1, -CD4, -CD5, -CD8 and -Ter119 (Becton Dickinson) and goat anti-rat Tricolor antibody (Caltag Lab, Burlingame, CA, USA). Sca-1 PE-Cy5.5 and c-Kit adenomatous polyposis coli antibodies were used to gate LK and LSK compartments of Lin⁻ cells. CD34 PE-Cy7 and Fc γ R PE antibodies were used to gate granulocyte-monocyte progenitors, common myeloid progenitors and megakaryocyte-erythroid progenitors within the LK cell population. For cell cycle analysis, bromodeoxyuridine adenomatous polyposis coli antibody and 7-AAD were used using Bromodeoxyuridine Flow Kit (Becton Dickinson Pharmingen, San Diego, CA, USA). For CD44 surface glycoprotein expression, anti-CD44 PE antibody (Becton Dickinson) was used.

Hematology and histology

For *in vivo* studies, peripheral blood from the tail vein was collected in heparin (LEO Pharma, Thornhill, ON, USA) and put in a 96-well U-bottom plate. Following centrifugation, the supernatant was poured off, erythrocytes were lysed with ammonium chloride (NH₄Cl; Stem Cell Technologies, Vancouver, BC, Canada) and the cells were stained with conjugated antibodies for FACS analysis. Hematological parameters of peripheral blood were monitored to control the development of leukemia (KX-21N; Sysmex, Mundelein, IL, USA). For histology, smears of peripheral blood were generated and cells from BM, spleen and liver were subjected to cytopspin onto glass slides, and subsequently stained with May-Grünwald Giemsa for microscopic examination.

Western blot analysis

One million cells were lysed in 50 μ l Laemmli buffer (BioRad, Hercules, CA, USA). Proteins were separated by electrophoresis (10% SDS-polyacrylamide gel electrophoresis) and transferred onto nitrocellulose membrane. Specific proteins were detected by an anti-*HOXA10* antibody (Sc-17158; Santa Cruz Biotechnology, Santa Cruz, CA, USA) and an anti-matrix metalloproteinase 9 (MMP9) antibody (ab38898; Abcam, Cambridge, UK). For CD44 staining, an anti-CD44 (14-0441; eBioscience, San Diego, CA, USA) was used on PNGase F1 deglycosylated and heat-denatured proteins. Anti-actin (Ab-5; Becton Dickinson Biosciences) was used as a loading control. After blotting with a horseradish peroxidase-conjugated secondary antibody, chemiluminescence activity was detected using the enhanced chemiluminescence Advance Western blot detection kit (GE Healthcare, Piscataway, NJ, USA). Gel images were analyzed using Image J (Wright Cell Imaging Facility, www.uhnresearch.ca/wcif) for quantification.

Microarray assay

Total RNA from BM cells of ciproxine- and doxycycline-receiving recipient mice suffering from leukemia was extracted from BM cells (Qiagen, Valencia, CA, USA). RNA concentration and integrity were, respectively, determined by NanoDrop spectrophotometry (NanoDrop Technologies, Wilmington, DE, USA) and a Bioanalyser (Agilent Technology, Waldronn, Germany). Reverse transcription was carried out using Illumina TotalPrep RNA Amplification Kit (Applied Biosystems/Ambion, Austin, TX, USA) and hybridization using Mouse WG-6 Expression v1. v1.2 BeadChips (Illumina, San Diego, CA, USA) interrogating more than 45 000 transcripts were performed by the Swegene Center for Integrative Biology at Lund University (SCIBLU). Signal intensity data was acquired with a BeadArray Scanner (Illumina) and experimental quality was performed using the BeadStudio Software (Illumina). The raw expression values mRNA abundance were submitted to the European Bioinformatics Institute (EMBL-EBI) ArrayExpress repository (accession no. E-MEXP-2491). Validation of proto-oncogene transcriptional modulation was performed by real-time PCR (LightCycler; Roche Diagnostic, Mannheim, Germany) with primers described in Supplementary Table 1.

Mass spectrometry

The experimental procedure has been described in detail.³⁰ Mass spectrometer (ThermoFisher Scientific, Bremen, Germany) was used to collect proteomics data from tryptic digests of leukemic whole-cell lysates. Identification of proteins by mass spectrometry was performed by matching experimental mass spectra against calculated spectra of all possible peptides in a protein database.

Statistical analyses

All data are expressed as the means \pm s.e.m. We assessed differences between the groups by Student's unpaired *t*-test. Statistical analysis of survival curves was performed using the Mantel–Haenszel log-rank test.

Results

Generation of leukemia in mice following *MEIS1* and *HOXA10* expression

HOXA10 transgenic mice were crossed with *Rosa26-rtTA-nls* mice to generate the *Rosa26-rtTA-nls/tetO-HOXA10* mouse

model.²⁰ With this tet-operator system, the expression level of *HOXA10* can be tightly regulated in hematopoietic cells using doxycycline.²⁰ These cells were then transplanted together with support cells to lethally irradiated recipients that were given doxycycline (0.2 g/ml) in the drinking water continuously for induced expression of *HOXA10* or ciproxine as a negative control for *HOXA10* induction (Figure 1a). All mice were then monitored for the development of leukemia. Induced expression of *HOXA10* mediated by a tet-operator system does not transform BM cells *in vivo*, as doxycycline-treated recipients of inducible *HOXA10* BM cells that had been transduced with the empty control vector remained healthy throughout a whole year (Figure 1b). This is in agreement with previous observations²⁰ that failed to detect leukemia in these same mice after long-term induction of *HOXA10*. The failure to develop leukemia could be due to the level of expression in the *in vivo* setting that might not be optimal for leukemic transformation. To create a leukemia model, we harvested BM cells from the *Rosa26-rtTA-nls/tetO-HOXA10* mice and transduced them with an *MEIS1* expressing vector. Seventy percent of recipient mice that received *MEIS1* and doxycycline developed leukemia, from around 12 weeks following transplantation. The average survival for mice in this group was 26 weeks (Figure 1b). We have therefore confirmed previous findings showing that *MEIS1* interacts with the *HOXA10* protein in leukemogenesis and we show here that *MEIS1* is required to initiate leukemia following tet-promoter-inducible expression of *HOXA10*. The disease has characteristics of acute leukemia, with a greater than eight-fold increase of total white blood cells in peripheral blood of moribund animals compared with healthy animals that have been fed ciproxine and killed 50 weeks after the transplantation (Figure 1c). In addition, the leukemic mice show lower red blood cell, hemoglobin and platelets levels, probably due to the outgrowth of leukemic cells and suppression of normal hematopoiesis. Symptomatic mice had severe splenomegaly (Figure 1d) and infiltrations of myeloid progenitors in different hematopoietic tissues, peripheral blood, BM, spleen and non-hematopoietic tissues like the liver (Figure 1e). Recipient mice that are given doxycycline continuously (*HOXA10*^{ON}) and have leukemia exhibit a clear upregulation of *HOXA10* protein levels in BM cells (Figure 1f), whereas only endogenous levels of *HOXA10* protein are detected in mice transplanted with *Rosa26-rtTA-nls/tetO-HOXA10* cells that are given ciproxine (*HOXA10*^{OFF}). Furthermore, no leakiness of the tet-promoter was observed in the *HOXA10*^{OFF} situation, as the *HOXA10* protein levels detected was similar to the endogenous *HOXA10* level found in unmanipulated wild-type mice (Figure 1f). FACS analysis of BM cells from moribund mice developing leukemia revealed a large accumulation of myeloblastic cells (Mac1⁺ c-Kit⁺), which correspond to a substantial expansion (>80%) of donor Ly5.1⁻ *tetO-HOXA10* cells (Figure 1g), and furthermore confirm transformation of BM with a complete disappearance of LSK stem cells (Lin⁻ c-Kit⁺ Sca-1⁺) and a total reconstitution with LK progenitors (Lin⁻ c-Kit⁺ Sca-1⁻), which in turn largely consist of granulocyte–monocyte progenitors (CD34⁺ Fc γ R⁺) (Figure 1h).

Twenty percent of the LICs induce leukemias in secondary transplant mice upon withdrawal of *HOXA10* overexpression

Leukemia-initiating cells (LICs) are capable of limitless self-renewal and are responsible for the maintenance of leukemia. Therefore, we asked whether all clones among the population of LICs are dependent on continuous expression of the *HOXA10*

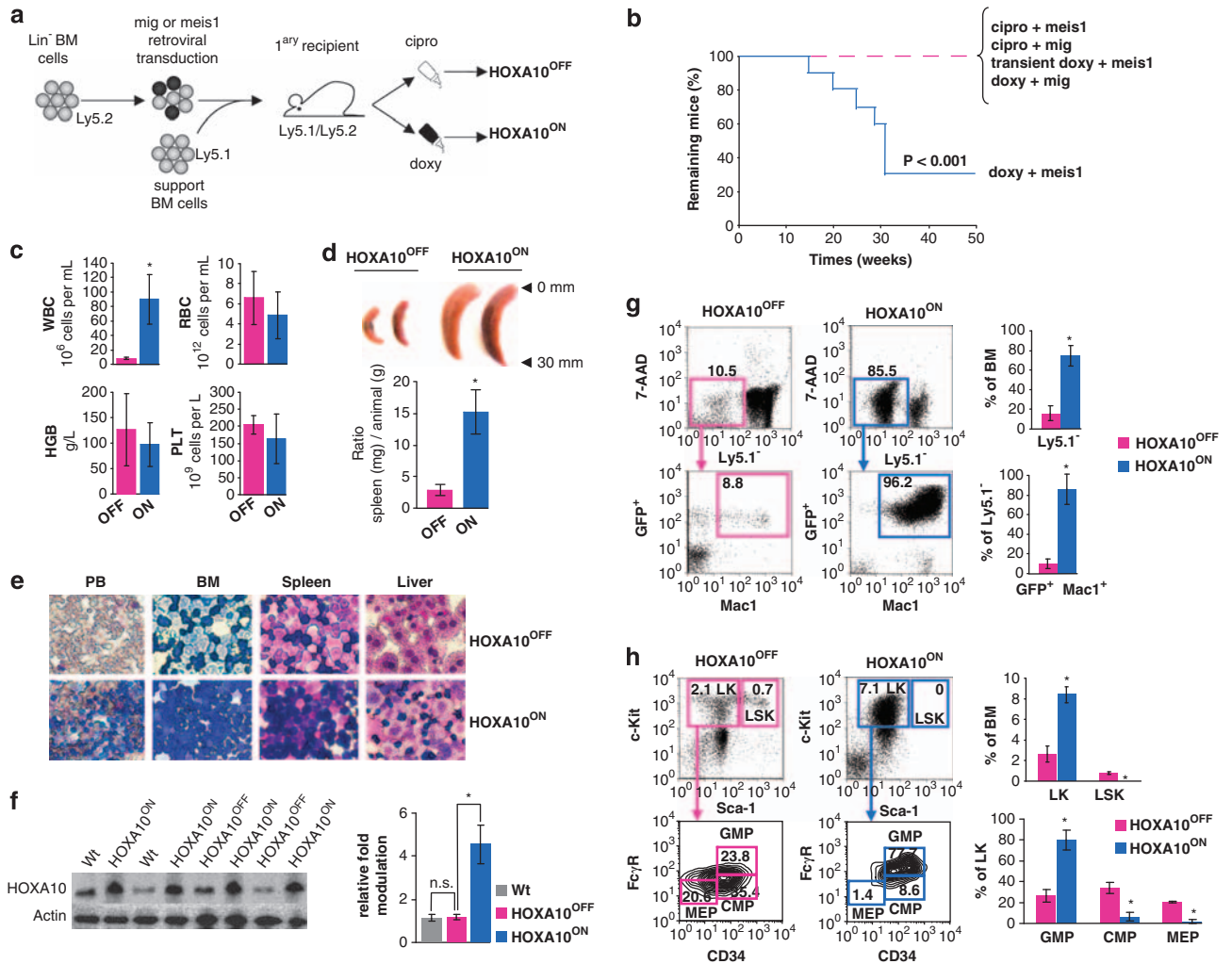


Figure 1 Generation of AML in primary recipients following *MEIS1* transduction requires continuous induction of *HOXA10* overexpression. (a) Illustration of the method used to generate AML in primary recipient mice. Mice were transplanted with BM cells from the *Rosa26-rtTA-*nls*/tetO-HOXA10* mice, transduced either with an *MEIS1*-expressing vector (*MEIS1*) or an empty control vector (*mig*). Following transplantation, mice were continuously fed doxycycline (*doxy*) in the drinking water (*HOXA10*^{ON}), ciproxine (*cipro*) as a negative control for *HOXA10* induction (*HOXA10*^{OFF}) or doxycycline transiently during 5 weeks (transient *doxy*). (b) Kaplan–Meier plot of recipient survival over time; $n = 10$ mice for *doxy* + *MEIS1* and $n = 6$ mice for all the other groups. P -value is measured by Mantel–Haenszel log-rank test. (c) Analysis of peripheral blood parameters, white blood cell (WBC), red blood cell (RBC), hemoglobin (HGB) and platelets (PLT) observed in moribund mice from the doxycycline group (*HOXA10*^{ON}, $n = 6$) and healthy mice killed 50 weeks after the transplantation from the control group (*HOXA10*^{OFF}, $n = 6$). (d) Severe splenomegaly in mice developing AML, representative pictures of two spleens of each group and weight ($n = 5$). (e) Giemsa stainings showing infiltration of myeloid progenitors in hematopoietic organs, in the peripheral blood (PB), bone marrow (BM), spleen and liver in mice with AML. Data are representative of five mice. (f) Western blot of BM cells showing induced expression of the *HOXA10* protein in moribund mice receiving doxycycline (*HOXA10*^{ON}), in contrast to healthy mice killed 50 weeks after the transplantation from the ciproxine group (*HOXA10*^{OFF}) or wild-type control mice (Wt). Representative western blot (left panel) and presentation of the relative increase (right panel, $n = 5$ for each group). (g) FACS analysis showing robust reconstitution of BM with the donor *tetO-HOXA10* cells (Ly5.1⁻) expressing *MEIS1* (GFP⁺) for mice receiving continuous doxycycline (*HOXA10*^{ON}), whereas no accumulation of donor cells is observed in mice given ciproxine in the drinking water (*HOXA10*^{OFF}). (h) FACS analysis showing the transformation of Lin⁻ cells for mice suffering from AML. Representative FACS (left panel) and pooled data from five mice (right panel). In this figure, data show means \pm s.e.m. * $P < 0.01$, as measured by unpaired t -test. n.s., nonsignificant variation.

transgene or whether the LICs' population is heterogeneous with respect to the dependence on the primary event that has generated the leukemia.

We previously described that the *HOXA10* transgene transcription and protein production is inducible and reversible in this model and can be terminated upon withdrawal of doxycycline.²⁰ GFP⁺ leukemic cells from two independent AML donor mice (nos. 1 and 2) were injected into lethally irradiated secondary recipients that were fed either doxycycline for continuous induced expression of *HOXA10* or ciproxine to eliminate the expression (Figure 2a). We asked how many LICs

per recipient were required to cause disease by transplanting limiting dilutions of the leukemic cells derived from mice, which continuously received doxycycline together with support cells. We tested the repopulative capacity of LICs from two independent donors. Time to onset of leukemia following BM transplantation and the number of mice that developed AML were quite dependent on the quantity of leukemic cells injected (Figure 2b). After enumerating the LICs per donor and testing the proportion of mice that developed leukemia in secondary transplant recipients, we could determine that the percentage of leukemic cells that represent LICs capable of generating

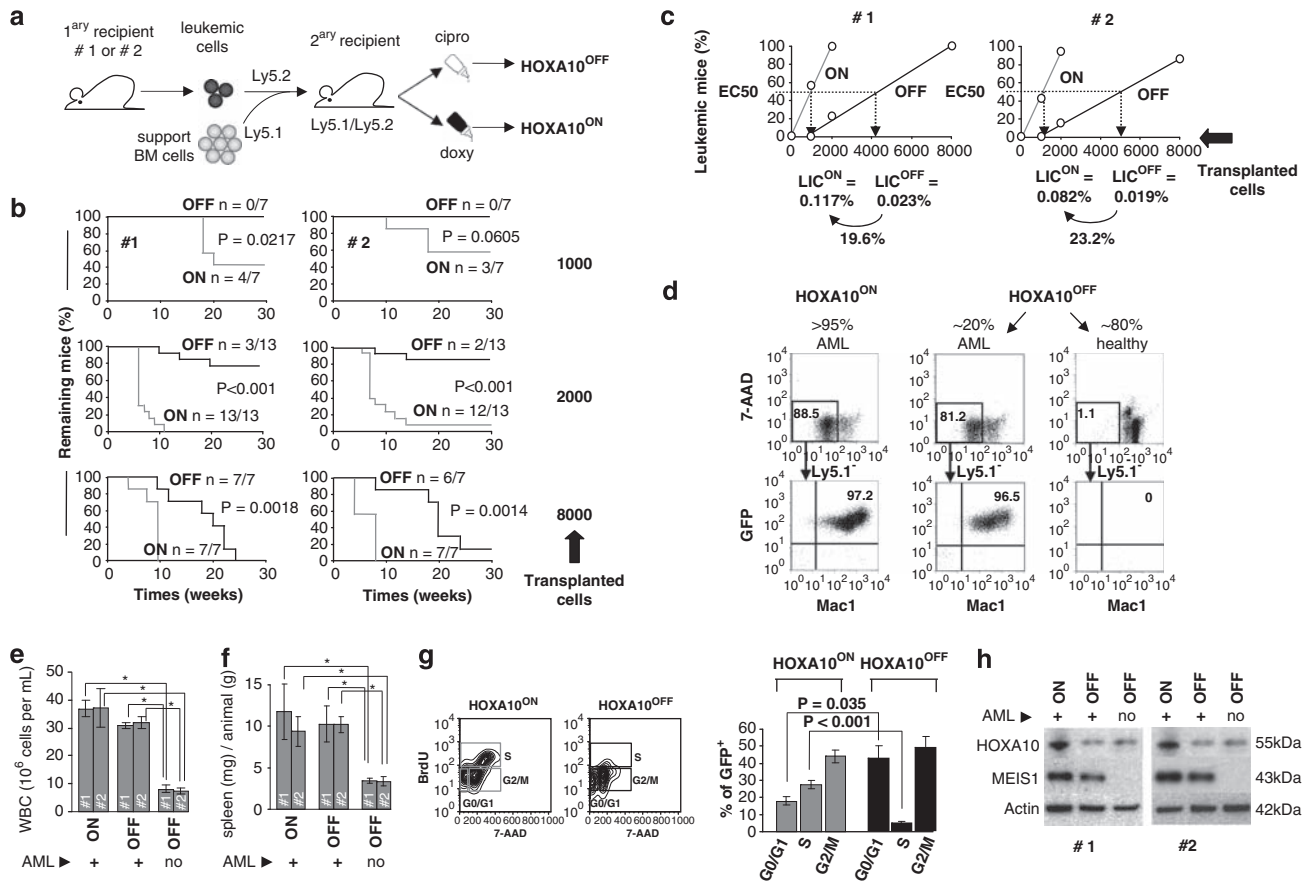


Figure 2 Twenty percent of the LICs generate a leukemia after withdrawal of *HOXA10*-induced expression in secondary transplant recipients. **(a)** Schematic overview of the transplantation strategy into secondary recipient mice. Leukemic cells were transplanted into lethally irradiated mice that were given either doxycycline for continuous expression of *HOXA10* (*HOXA10*^{ON}) or ciprofloxacin as a negative control (*HOXA10*^{OFF}). **(b)** Kaplan–Meier plot of survival mice over time when mice were transplanted with 1000, 2000 and 8000 leukemic cells. *P*-value is measured by Mantel–Haenszel log-rank test. **(c)** Limiting dilution assay for quantification of transplanted leukemic cells to induce leukemia in 50% of the mice in *HOXA10*^{ON} (EC50^{ON}) or *HOXA10*^{OFF} (EC50^{OFF}) recipients. Quantification of transplanted leukemic cells to induce 50% of leukemic mice, when *HOXA10*^{ON} (EC50^{ON}) or *HOXA10*^{OFF} (EC50^{OFF}) conditions as 1 LIC/total number of cells injected to induce EC50. The percentage of LICs resistant upon withdrawal of the initial *HOXA10*-transforming event is calculated as percent of LIC^{OFF} compared with LIC^{ON} (total LICs). **(d)** FACS analysis of secondary recipient mice BM transplanted with 2000 leukemic cells showing expansion of GFP⁺ cells only for ~20% of mice that developed AML. Increase in the WBC level **(e)** and enlarged spleen size **(f)** in leukemic mice. *HOXA10*^{ON} (ON; *n* = 10), *HOXA10*^{OFF} (OFF; *n* = 3, donor no. 1, *n* = 2, donor no. 2) and healthy mice from the ciprofloxine group killed at week 30 (*n* = 10). **(g)** Analysis of cell cycle stages showing a reduction in the proportion of cells in the S phase and an increased percentage of cells in the G0/G1 phase in AML from mice given ciprofloxine (*HOXA10*^{OFF}) compared with doxycycline (*HOXA10*^{ON}). Representative FACS analysis is shown (left) and pooled data from four mice per group are displayed (right). **(h)** Representative western blot showing expression of the *HOXA10* protein in BM of leukemic secondary recipient mice. High expression is restricted to the group that was given doxycycline (ON), whereas endogenous *HOXA10* expression is observed in leukemic mice that were given ciprofloxine (OFF). In this figure, the findings are from two independent donor leukemias (nos. 1 and 2). Data show means ± s.e.m.; **P* < 0.01 as measured by unpaired *t*-test.

leukemias was 0.117% for no. 1 and 0.082% for no. 2 in the *HOXA10*^{ON} condition. The percentage of LICs that can generate secondary leukemias was 0.023% for no. 1 and 0.019% for no. 2 in the *HOXA10*^{OFF} condition. In conclusion, approximately 20% of LICs (average of two donors; Figure 2c and Supplementary Table 2) were able to cause secondary leukemias after removal of the primary events.

Maintenance of *HOXA10* overexpression is not essential for growth of resistant leukemic cells in vivo
When we analyzed further mice transplanted with 2000 leukemic cells, ~80% (*n* = 21/26) of the ciprofloxine-treated control mice stayed healthy over the 30 weeks they were monitored, showing that a large majority of LICs from the primary recipient mice require continuous overexpression of

HOXA10 to develop leukemia in secondary recipients. Using FACS analysis, we could not detect any significant expansion of the donor cells in healthy mice, and there were no detectable GFP⁺ cells in the BM after 30 weeks (Figure 2d). The total clearance of the LICs in BM was moreover confirmed since the re-administration of doxycycline in the drinking water of these healthy mice during 10 weeks did not re-activate expansion of GFP⁺ LICs cells (*n* = 10 mice tested, data not shown). This shows eradication of the LICs when *HOXA10*-induced overexpression is withdrawn in ~80% of the secondary transplanted mice, which consequently highlights the importance of the primary event in the maintenance of majority of the LICs. However, ~20% (*n* = 5/26) of secondary recipients developed AML despite withdrawal of *HOXA10* and those mice showed accumulation of Ly5.1⁻ GFP⁺ donor leukemic cells in the BM (Figure 2d), increased total white blood cells in peripheral blood

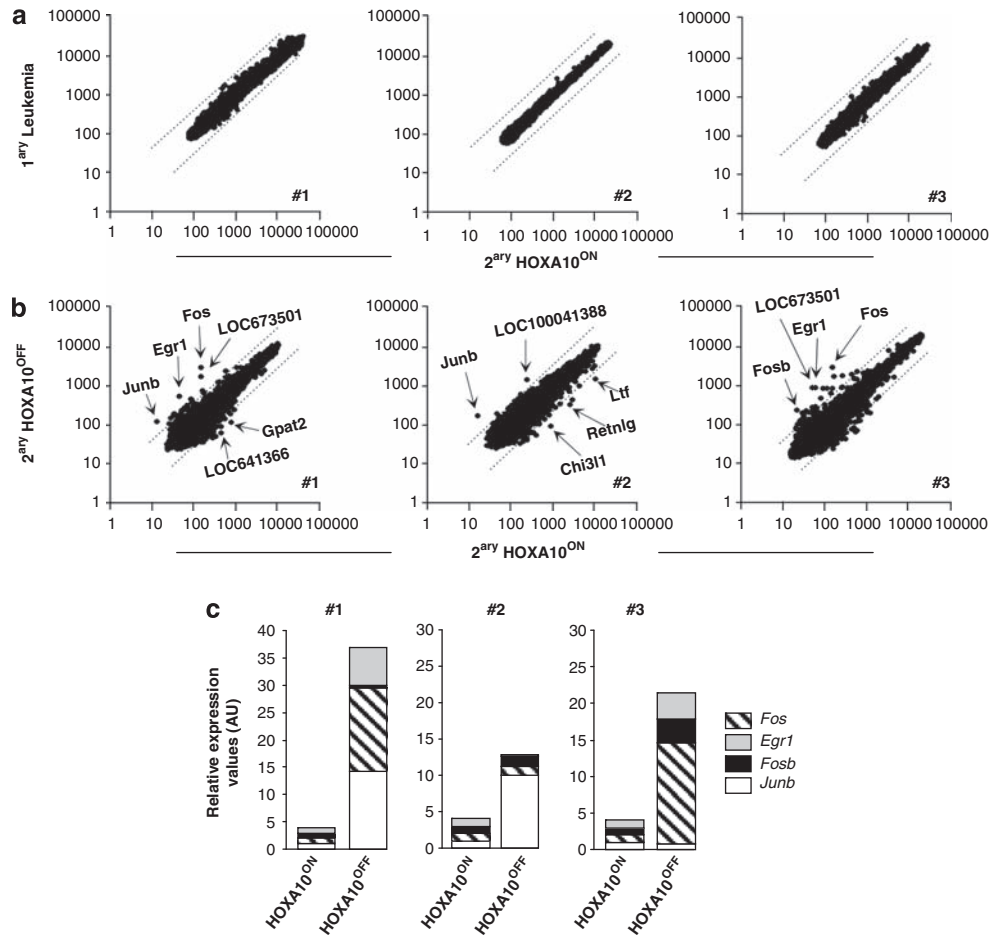


Figure 3 Leukemias generated upon withdrawal of *HOXA10* overexpression exhibit elevated expression of several proto-oncogenes. (a) Microarray scatter plots showing that the global RNA expression profile of leukemic cells from the *HOXA10*^{ON} secondary recipient mice are similar to leukemic cells generated from primary donors. (b) Microarray scatter plots showing upregulation of several proto-oncogenes in leukemias from *HOXA10*^{OFF} secondary recipient mice when compared with leukemic cells from *HOXA10*^{ON} secondary recipient mice. Scatter plots show normalized raw data. Line represents a fold change value of 5. (c) Validation of the transcriptional activation of several proto-oncogenes observed in resistant leukemic cells by real-time PCRs. Data are expressed in arbitrary units (AU): real-time PCRs from *HOXA10*^{OFF} are normalized to *HOXA10*^{ON}.

(Figure 2e) and splenomegaly (Figure 2f). Using FACS to detect cycling and non-cycling leukemic cells *ex vivo*, we observed that *HOXA10*^{OFF} leukemic cells had an increased proportion of cells in the G0/G1 phase and a decreased proportion of cells in the S phase (Figure 2g). This clearly shows that the cell cycling capacity is greatly reduced when overexpression of *HOXA10* is stopped, but this does not affect outgrowth capacity of resistant leukemic cells. High levels of the *HOXA10* protein were observed in leukemic mice under the administration of doxycycline, whereas only endogenous expression of the *HOXA10* protein was detected in BM from leukemic mice fed ciproxine (Figure 2h). This clearly provides evidence that maintenance of *HOXA10* overexpression is not essential for the growth of resistant leukemic cells.

Leukemias generated upon withdrawal of HOXA10 overexpression exhibit elevated expression of several proto-oncogenes

A clonality analysis of BM cells was carried out by linear amplification-mediated PCR to identify if resistance might be explained by the expansion of specific leukemic cells owing to insertional mutagenesis caused by retroviral integration in

genomic DNA. In secondary recipient mice, we observed that leukemias of the *HOXA10*^{ON} and the *HOXA10*^{OFF} groups were both induced by expansion of LICs generated in primary recipient mice (Supplementary Figure 1). Assignment of sequencing data to mouse chromosomes shows that the retroviral integration sites are at the 5' side of the *eukaryotic translation elongation factor 1 alpha 1 (EEF1A1)* gene for donor no. 1 and at the 5' side of *RAB1*, a member of *RAS* oncogene family, for donor no. 2. However, integration did not seem to affect expression of these target genes, as transcription of *EEF1A1* and *RAB1* genes was not affected in leukemias upon withdrawal of *HOXA10* expression. Very high levels of *MEIS1* were still detected in those leukemic cells (Figure 2h). High levels of *MEIS1* can theoretically cooperate with dysregulation in the expression of other *Hoxa* genes, but we never observed dysregulation of several tested *Hoxa* genes in those samples (*Hoxa1*, *Hoxa5*, *Hoxa7* and *Hoxa9* and endogenous murine *Hoxa10*; Supplementary Figure 1). Next, the cytogenetic status of leukemic cells resistant upon withdrawal of the *HOXA10* expression was characterized by molecular karyotyping using a murine comparative genomic hybridization array. Interestingly, we could not detect any difference in whole chromosome gain or loss, unbalanced subtelomeric rearrangements or

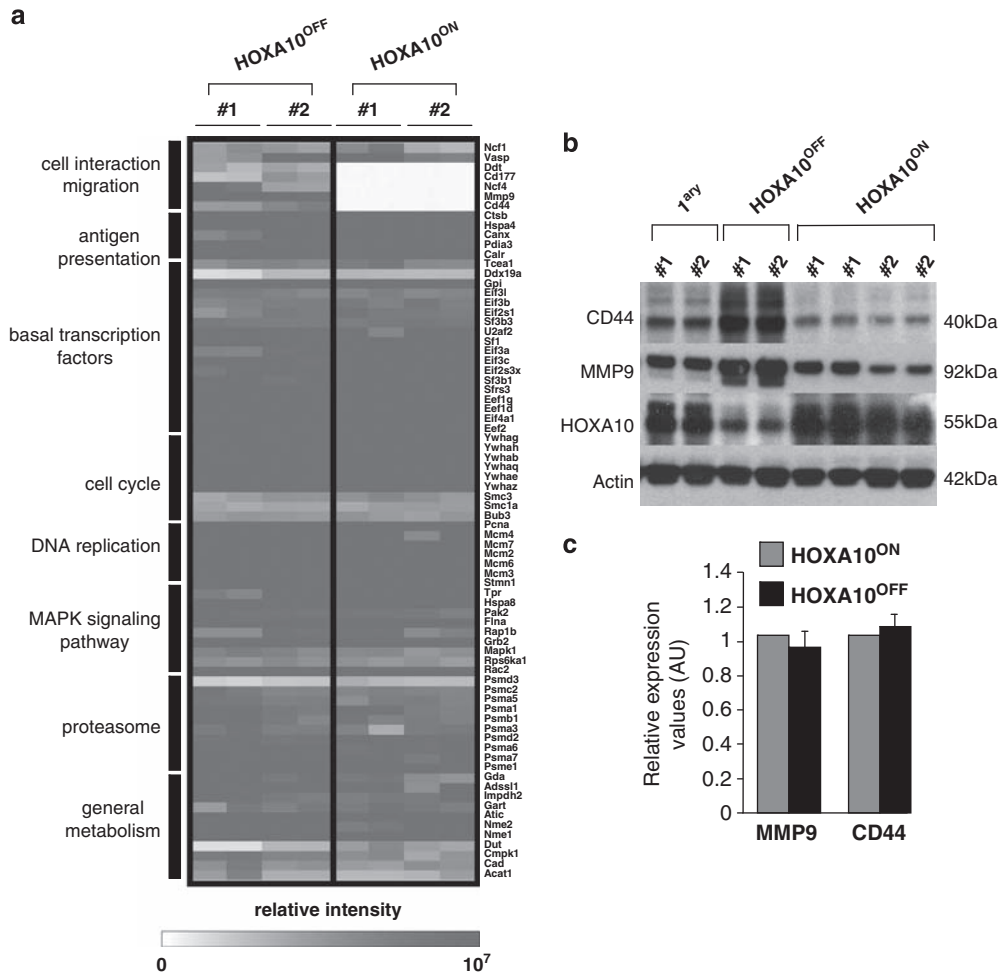


Figure 4 High levels of the CD44 transendothelial marker correlate with increased leukemia rate generated by LICs after withdrawal of the initial *HOXA10* transforming event. **(a)** Functional clustering of proteins detected by mass spectrometry comparing leukemic cells from *HOXA10*^{ON} secondary recipient mice to leukemic cells from the *HOXA10*^{OFF} group (two donors are tested: nos. 1 and 2). The secondary leukemias show high expression of subset of proteins involved in transendothelial migration, whereas proteins from other functional clusters do not exhibit a difference. **(b)** Western blot showing high levels of the CD44 protein in leukemic cells from the *HOXA10*^{OFF} mice. A representative western blot is displayed using recipients of cells from donors: nos. 1 and 2. **(c)** Transcriptional analysis of *CD44* and *MMP9* genes. Results are expressed in arbitrary units (AU); raw data from *HOXA10*^{ON} microarrays are normalized to *HOXA10*^{OFF}. Data show means ± s.e.m. of three donors.

microdeletions (Supplementary Figure 2). In conclusion, neither insertional mutagenesis nor subchromosomal changes can explain resistance of LICs.

For several years, investigators have used gene expression profiling to define genes that are dysregulated in AML cells. We tested the modulation of gene expression by microarray analysis and no difference was observed between the transcriptome of leukemic cells originating from donors in primary and secondary recipient mice when the *HOXA10* overexpression continues to be maintained (Figure 3a). When leukemic cells from secondary recipient mice from the *HOXA10*^{ON} group were compared with leukemic cells from the *HOXA10*^{OFF} group, the global gene expression profiles were found altered for each individual donor (Figure 3b). To define markers that characterized resistant *HOXA10*^{OFF} leukemic cells, we performed a Significance Analysis of Microarrays, but could not find any gene commonly dysregulated among the three tested donors (Supplementary Figure 3). Although variability seems to be more donor specific, we analyzed modulation in gene expression for each individual donor. To identify possible targets that were altered in resistant leukemias upon withdrawal of *HOXA10*

overexpression, we selected genes showing great degree of dysregulation (fold-change value >5). We found that *Jun*, *Fos* and *Egr1* genes were activated in different samples of the *HOXA10*^{OFF} group, with variability among the three tested donors (Figure 3a). Dysregulation of these proto-oncogenes was furthermore confirmed by real-time PCRs to be donor specific (Figure 3c).

High levels of CD44 on leukemic cells correlate with AML in secondary transplant mice upon withdrawal of the initial *HOXA10* transforming event

To further investigate if another possible mechanism is involved in leukemia, upon withdrawal of the primary oncogenic event, we performed proteomic analysis using mass spectrometry to define proteins differently expressed. Interestingly, we observed that leukemias generated upon removal of the primary event produced high levels of several proteins involved in cell–cell and cell–matrix interactions (Figure 4a). The CD177 and CD44 proteins expressed on the cell surface participate in cell transmigration. The MMP9 protein is involved in the breakdown of extracellular matrix and colocalized with CD44 on the

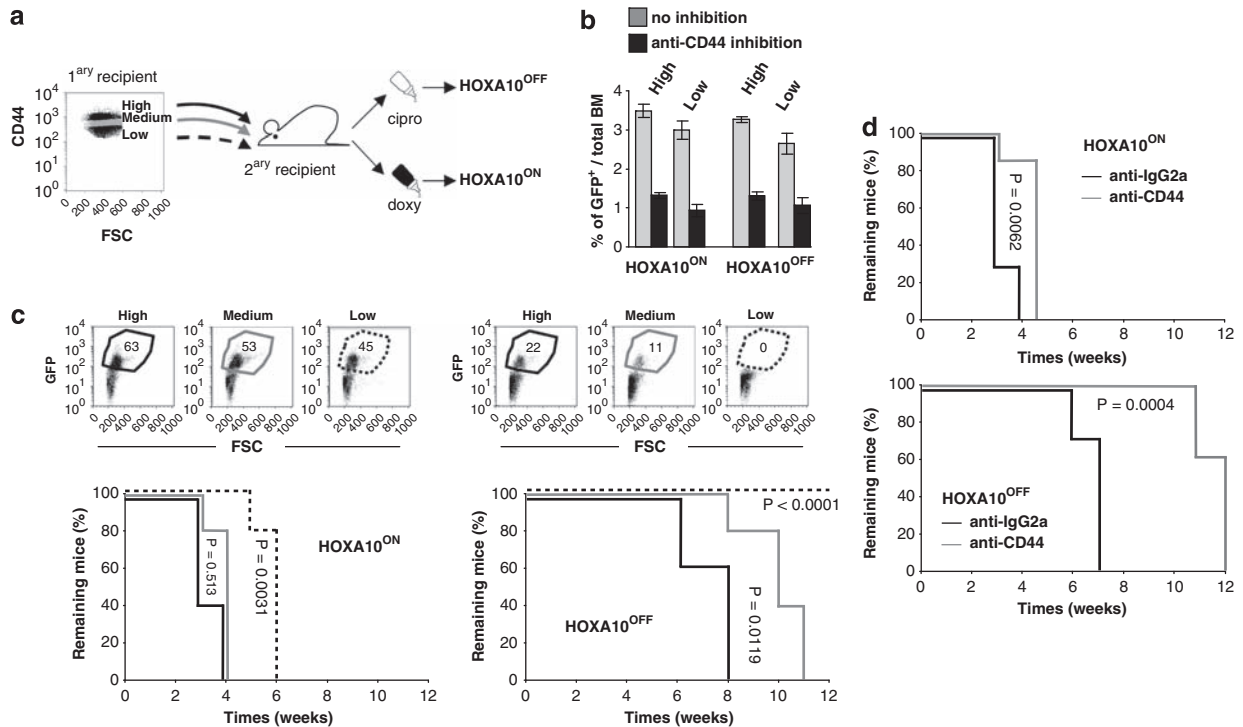


Figure 5 Withdrawal of *HOXA10* induces the outgrowth of leukemic cells characterized by high expression of the CD44 transendothelial marker. (a) Schematic overview of the transplantation of leukemic cells expressing different levels of the CD44 surface marker. Leukemic cells are sorted by FACS and transplanted into lethally irradiated mice that were given either doxycycline for continuous expression of *HOXA10* (HOXA10^{ON}) or ciprofloxacin as a negative control (HOXA10^{OFF}). (b) Homing of GFP⁺ leukemic cells in BM from HOXA10^{ON} and HOXA10^{OFF} mice 24 h after transplantation ($n=5$), with or without inhibition with anti-CD44 antibody. (c) Outgrowth of leukemic cells in BM 2 weeks after transplantation (top panel, representative FACS of one mouse for each group) and Kaplan–Meier plot of survival over time (bottom panel) displaying findings from mice transplanted with the different population and that were given doxycycline or ciprofloxacin. P -value is measured by Mantel–Haenszel log-rank test and compared with mice transplanted with CD44^{high} leukemic cells ($n=5$, each group). (d) Kaplan–Meier plot of survival over time displaying findings from mice transplanted with leukemic cells following CD44 inhibition. Leukemic cells were incubated during 20 min with high dose of the anti-CD44 PE (20 μ g/ml) or control anti-immunoglobulinG2a PE (20 μ g/ml) before tail vein injection ($n=7$, each group).

surface of HOXA10^{OFF} cells (Supplementary Figure 4). The neutrophil cytosol factor 4 protein is involved in leukocyte transendothelial migration.

We decided to focus on the CD44 surface marker, as this surface glycoprotein mediates cell adhesion, migration and homing of hematopoietic cancer cells.^{31,32} Using western blot analysis (Figure 4b), we confirmed that the outgrowth of cells from HOXA10^{OFF} leukemias is restricted to a population of cells expressing high levels of the CD44 adhesion protein on the surface, despite unaltered transcription levels of the *CD44* gene (Figure 4c). Even if different variants of CD44 are correlated to different risk in human AML,³³ resistance cannot be explained by the outgrowth of a population of leukemic cells expressing a variant of CD44. Using RT-PCR, we observed that HOXA10^{OFF}-resistant leukemias as well as leukemic cells generated in primary and secondary recipient mice fed with doxycycline were expressing the same CD44 transcript isoform (variant 3; NM_001039151) (Supplementary Figure 5).

Withdrawal of the initial *HOXA10* transforming event promotes outgrowth of leukemic cells expressing high levels of CD44 in secondary transplantation settings

To determine whether an increase in CD44 is a key mechanism by which LICs are resistant, we performed a functional test by FACS sorting leukemic cells generated in primary donors and transplanted 20 000 cells expressing different levels of the CD44 surface marker (CD44^{low}, CD44^{medium} and CD44^{high}) in the tail

vein of lethally irradiated secondary recipient mice fed doxycycline or ciprofloxacin (Figure 5a). When we analyzed engraftment 24 h after the transplantation, we observed that CD44^{low} leukemic cells have strong potential to home and engraft, as there is only a slight decrease in their capacity to engraft compared with CD44^{high} leukemic cells (Figure 5b). There was furthermore no difference in homing between mice fed with doxycycline or ciprofloxacin; therefore, when *HOXA10* overexpression was turned off within 24 h, homing of leukemic cells was not affected.

When we monitored mice for the occurrence of leukemia (Figure 5c), outgrowth of leukemic cells was not dependent on CD44 level of expression in the HOXA10^{ON} condition. Consistent with this, onset of leukemia was not delayed for mice transplanted with CD44^{low} leukemic cells. When mice were fed with ciprofloxacin to turn off *HOXA10* overexpression (Figure 5c), all mice injected with CD44^{high} leukemic cells developed leukemia, whereas all mice injected with CD44^{low} leukemic cells remained healthy ($n=5$ for each group, $P<0.005$) and no detectable GFP⁺ leukemic cells were observed in the BM of those healthy mice 2 weeks after the transplantation. In conclusion, we confirmed that withdrawal of the initial *HOXA10* oncogene promotes the outgrowth of leukemic cells expressing high levels of CD44. To rule out that the differences in the leukemogenic capacity of transplanted cells was not owing to secondary effects, we performed detailed immunophenotypic analysis of the three CD44 fractions by FACS and could not find a significant difference in the

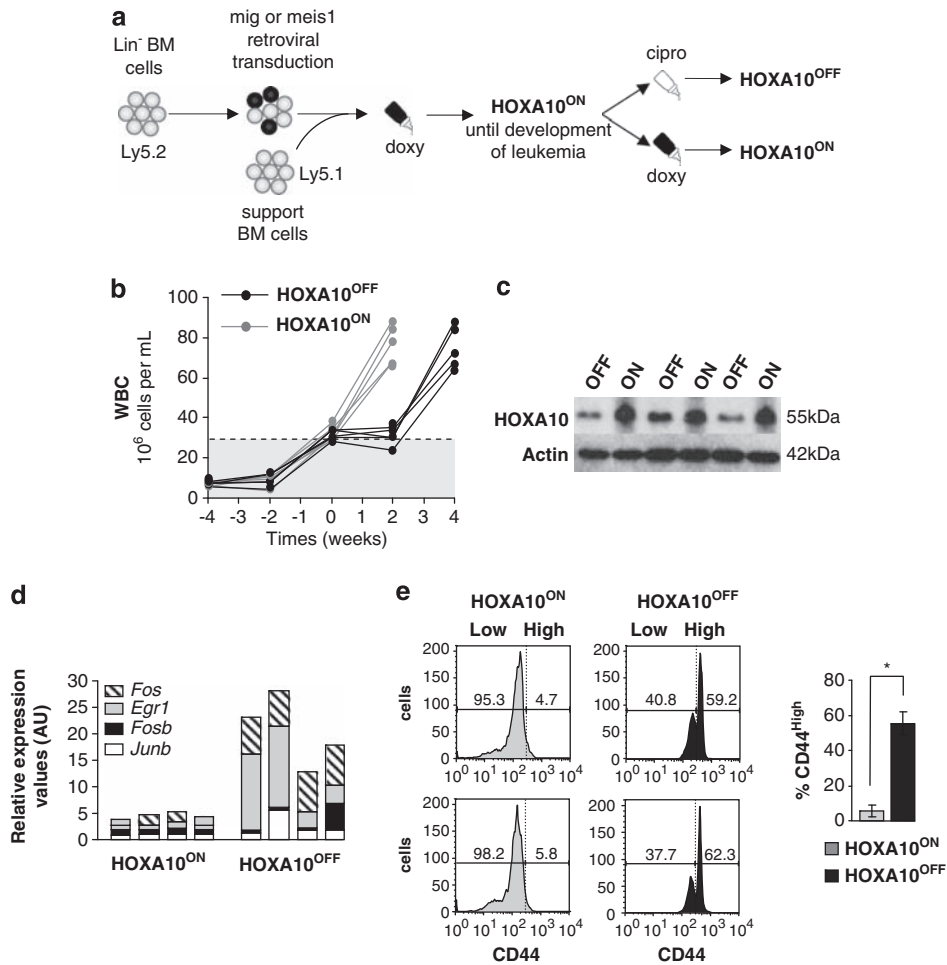


Figure 6 Leukemic cells expressing high levels of CD44 contribute toward rendering resistance to inactivation of the initial transformation event in primary recipients. (a) Illustration of the method used to analyze how withdrawal of *HOXA10* overexpression impacts on the maintenance of leukemia in primary recipient mice. (b) Modulation of the WBC level over time for mice under continuous administration of doxycycline ($n = 5$) or for mice where the *HOXA10*-induced expression was eliminated ($n = 5$). (c) Western blot of BM cells showing induced expression of the *HOXA10* protein in moribund mice receiving doxycycline continuously (ON; $n = 3$), whereas mice suffering from leukemia 4 weeks after withdrawal of doxycycline exhibit a clear downregulation of *HOXA10* in BM cells (OFF; $n = 3$). (d) Transcriptional activation of several proto-oncogenes in relapsed leukemic cells compared with leukemic cells where doxycycline was maintained ($n = 4$ mice). Data are expressed in arbitrary units (AU): real-time PCRs from samples of *HOXA10*^{OFF} ($n = 4$ mice) are normalized to the first sample of *HOXA10*^{ON} ($n = 4$ mice). (e) FACS analysis showing high levels of the CD44 protein on the surface of relapsed leukemic cells from the *HOXA10*^{OFF} mice compared with leukemic cells where doxycycline was maintained. Data show means \pm s.e.m. of five mice. * $P < 0.01$, as measured by unpaired *t*-test.

expression of several other surface markers expressed on leukemic cells (Supplementary Figure 6).

We furthermore inhibited CD44 with high concentration of anti-CD44 antibody preceding transplantation of secondary mice and provided evidence that targeting CD44 dramatically affect leukemogenesis of *HOXA10*^{OFF} cells (Figure 5d). In conclusion, CD44 represents the major driving force of the growth of leukemic cells upon withdrawal of the initial transforming events.

Increased expression of several proto-oncogenes and high levels of CD44 on leukemic cells correlate with relapsed leukemias in primary transplant recipients after withdrawal of *HOXA10* overexpression

We next asked whether upregulation of *HOXA10* is required for maintenance of the leukemia in primary transplant recipients, or alternatively, whether the presence of $\sim 20\%$ of resistant LICs among the pool of LICs could affect relapse to leukemia after

withdrawal of the primary oncogene. In primary transplantation settings, doxycycline administration was stopped immediately after increase of the white blood cell count above 30×10^6 cells per ml. Peripheral blood was subsequently monitored to test whether abolition of *HOXA10* might lead to a reversal of the leukemia (Figure 6a). In the positive control group, all mice were subject to continuous administration of the doxycycline, resulting in a dramatic increase of total white blood cells characteristic of acute leukemia (Figure 6b). In the group of mice where *HOXA10* expression was turned off, a stabilization of the number of leukemic cells in the blood was observed 2 weeks after withdrawal of the doxycycline, followed by relapse that occurs 2 weeks later with a substantial increase of myeloid cells in peripheral blood. Recipient mice with relapsed leukemias 4 weeks after withdrawal of doxycycline exhibit a clear downregulation of *HOXA10* in BM cells (Figure 6c). Activation of several proto-oncogenes was frequently observed in relapsed samples after removal of the *HOXA10* expression (Figure 6d). Furthermore, relapsed leukemic cells were also restricted to a

population of cells expressing high levels of the CD44 adhesion protein on the surface (Figure 6e). This finding suggests that complete remission cannot be achieved in the primary transplant mice despite withdrawal of *HOXA10*, because secondary mutations in additional proto-oncogenes and overproduction of the adhesion molecule CD44 contribute toward rendering the leukemic cells resistant to inactivation of the initial transformation event.

Discussion

Retrovirus-mediated overexpression of *HOXA10* in murine BM cells followed by transplantation has been shown to induce AML in a significant proportion of mice, with a latency of 19–50 weeks.¹¹ However, induced expression of *HOXA10* mediated by a tet-operator system is not leukemogenic neither for the original transgenic animals²⁹ nor when BM from these animals is transplanted into recipients.²⁰ There is experimental evidence that retroviral integration induces genetic hits that have an important contribution in leukemogenesis of AML in mice,^{34–36} but we never observed the development of leukemia in mice receiving transplants of inducible *HOXA10* BM cells transduced with the empty control vector. It is therefore likely that the oncoprotein levels may be critical for the development of the disease phenotype. In whole BM cells from *in vivo* induced *Rosa26-rtTA-nls/tetO-HOXA10* mice receiving doxycycline in the drinking water, the *HOXA10* transgene transcription was induced ~6-fold compared with uninduced control recipients receiving ciprofloxacin,²⁰ whereas the retroviral approach is well known to induce a constant activation of the transgene, at several proviral insertion sites in DNA, leading to very high expression levels of the transgene, often >20-fold.³⁷ In agreement with this, retroviral vector-mediated overexpression of *HOXA10* in murine BM cells induces AML *in vivo* with outgrowth of clones expressing extremely high levels of the *HOXA10* transgene.¹¹ Therefore, our model has many advantages since the expression can be induced upon demand, although the absolute expression levels are somewhat lower.

These findings show that moderate levels of *HOXA10* have a restricted capacity to transform cells. To develop a mouse model for leukemia, we had to transduce *HOXA10*-inducible BM cells with the cofactor *MEIS1*,^{9,13,16} which generated AML in 70% of transplanted recipients and showed that *MEIS1* can potentiate the leukemogenesis induced by moderate expression of *HOXA10*. The long latency period of 15–30 weeks to induce leukemia suggests that the cooperation with *MEIS1* is not sufficient to transform cells. Rather, additional hits may occur later during the leukemogenesis *in vivo*. Combination of specific genetic events is involved in the multi-step evolution of premalignant cells to full-blown AML. In several AML mouse models, sequential genetic hits have been shown to collaborate to induce aggressive leukemias.^{25–28} This concept is supported by clinical evidence as well since multiple genetic hits are often diagnosed in AML patients.^{21–24}

The oncoproteins *BCR-ABL* and *MLL-ENL* have been investigated as chronic myeloid leukemia³⁸ and AML¹⁸ models, respectively, using conditional expression in mice. In both cases, the leukemic cells remained dependent on oncoprotein expression, whereas ablation of the transgene resulted in remission. Complete reversion of leukemias upon inactivation of the primary event suggests important implications for the design of therapy.^{18,38} It is however important to note that both *BCR-ABL* and *MLL-ENL* are potent oncogenes that can probably generate leukemia by themselves. The *HOXA10* oncogene is

less potent and needs additional genetic alterations for transformation. The reversal of a ‘weak’ initiating oncogene leads only to a partial relapse with a short period of remission because the clone is genetically unstable and has acquired many other alterations, whereas removal of a potent oncogene leads to elimination of the malignant clone because it is merely depending on that oncogene.^{18,38} The *MLL-ENL* fusion gene is one of the few genetic alterations that probably need very few secondary events to initiate AML, as the mouse model shows that the disease penetrance is high and the latency short, arguing that very few additional hits are needed.¹⁸ Not all models of *MLL* fusion-induced leukemia have short latencies, whereas the latency period required for the onset of acute leukemia *in vivo* is variable and partner protein dependent.³⁹ For example, a mouse model with *AML1-ETO* overexpression does not generate AML when the oncogene is expressed alone,⁴⁰ but AML develops when combined with dysregulation of additional oncogenes.^{25,27,28}

Our findings show that the majority of LICs are sensitive to ablation of the primary oncogenic event (*HOXA10* overexpression), whereas ~20% of the LICs cause leukemic relapse after withdrawal of *HOXA10* overexpression in a secondary transplantation setting. Activation of several oncogenes may compensate for reduction in *HOXA10* overexpression and confer additional advantage to a subpopulation of LICs, as the disease evolves and relapses. The set of commonly dysregulated proto-oncogenes (*Jun*, *Fos* and *Egr1*) we observed in relapsed leukemias have been previously described to occur in murine AML,⁴¹ and are found significantly activated in relapsing AML patients.⁴² Our findings show that activation of these proto-oncogenes is not owing to subchromosomal changes or retroviral vector integration in DNA flanking or within the proto-oncogenes. In most cases, we observed dysregulation of additional oncogenes in relapsed leukemias.

By performing proteomic analysis using mass spectrometry, we detected an important increase in the levels of specific proteins involved in invasiveness of LICs in the microenvironmental BM niche. By transplanting leukemic cells expressing different levels of the adhesion protein CD44, we clearly provide mechanistic evidence that relapse of the leukemia may be niche dependent, as only leukemic cells expressing high levels of CD44 cause leukemias upon withdrawal of the *HOXA10* oncogene. Although the bulk of the tumor failed to grow upon inactivation of the initiating *HOXA10* oncogene, our findings show that LICs are not totally eliminated by withdrawal of *HOXA10* overexpression. Rather, it seems that a population of LICs are resistant because proteins involved in cell–cell and cell–matrix interactions are highly expressed on the cell surface, modulating homing, motility and invasive properties of the leukemic cells. Among the proteins that we found increased in leukemic cells resistant upon withdrawal of *HOXA10* were MMP9 and CD44. The MMP9 protein is an important factor for tumor-associated tissue remodeling and metastasis,^{43,44} and several MMPs have been linked to leukemic survival rates.^{45,46} CD44 correlates with poor prognosis in AML, facilitating motility and tumor growth,^{31,32} and may be a good target to eradicate human AML stem cells.⁴⁷ Here we show the highly aggressive nature of leukemic cells expressing high levels of CD44 and conversely show the impaired outgrowth of leukemic cells expressing low levels of this surface marker.

In conclusion, our murine model of inducible *HOXA10* expression recapitulates many of the features of human AML and is helpful in analyzing the oncogene addiction and unravel the basic mechanisms involved in the initiation and maintenance of leukemia, and to study whether adhesion molecules

expressed on the surface of leukemic cells are important factors for leukemic relapse in the microenvironmental niches of the BM. Our findings support the notion that cell intrinsic genetic events are not the only factors causing leukemic relapse, but suggest that host-dependent extrinsic factors in the BM niche may also play a fundamental role in the mechanism mediating leukemic relapse.

Conflict of interest

The authors declare no conflict of interest.

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References

- Lawrence HJ, Sauvageau G, Humphries RK, Largman C. The role of HOX homeobox genes in normal and leukemic hematopoiesis. *Stem cells (Dayton, OH)* 1996; **14**: 281–291.
- Magli MC, Barba P, Celetti A, De Vita G, Cillo C, Boncinelli E. Coordinate regulation of HOX genes in human hematopoietic cells. *Proc Natl Acad Sci USA* 1991; **88**: 6348–6352.
- Magli MC, Largman C, Lawrence HJ. Effects of HOX homeobox genes in blood cell differentiation. *J Cell Physiol* 1997; **173**: 168–177.
- Andreeff M, Ruvolo V, Gadgil S, Zeng C, Coombes K, Chen W et al. HOX expression patterns identify a common signature for favorable AML. *Leukemia* 2008; **22**: 2041–2047.
- Golub TR, Slonim DK, Tamayo P, Huard C, Gaasenbeek M, Mesirov JP et al. Molecular classification of cancer: class discovery and class prediction by gene expression monitoring. *Science (New York, NY)* 1999; **286**: 531–537.
- Kavagoe H, Humphries RK, Blair A, Sutherland HJ, Hogge DE. Expression of HOX genes, HOX cofactors, and MLL in phenotypically and functionally defined subpopulations of leukemic and normal human hematopoietic cells. *Leukemia* 1999; **13**: 687–698.
- Borrow J, Shearman AM, Stanton Jr VP, Becher R, Collins T, Williams AJ et al. The t(7;11)(p15;p15) translocation in acute myeloid leukaemia fuses the genes for nucleoporin NUP98 and class I homeoprotein HOXA9. *Nat Genet* 1996; **12**: 159–167.
- Nakamura T, Largaespada DA, Lee MP, Johnson LA, Ohyashiki K, Toyama K et al. Fusion of the nucleoporin gene NUP98 to HOXA9 by the chromosome translocation t(7;11)(p15;p15) in human myeloid leukaemia. *Nat Genet* 1996; **12**: 154–158.
- Kroon E, Kros J, Thorsteinsdottir U, Baban S, Buchberg AM, Sauvageau G. Hoxa9 transforms primary bone marrow cells through specific collaboration with Meis1a but not Pbx1b. *EMBO J* 1998; **17**: 3714–3725.
- Thorsteinsdottir U, Kros J, Kroon E, Haman A, Hoang T, Sauvageau G. The oncoprotein E2A-Pbx1a collaborates with Hoxa9 to acutely transform primary bone marrow cells. *Mol Cell Biol* 1999; **19**: 6355–6366.
- Thorsteinsdottir U, Sauvageau G, Hough MR, Dragowska W, Lansdorp PM, Lawrence HJ et al. Overexpression of HOXA10 in murine hematopoietic cells perturbs both myeloid and lymphoid differentiation and leads to acute myeloid leukemia. *Mol Cell Biol* 1997; **17**: 495–505.
- Kroon E, Thorsteinsdottir U, Mayotte N, Nakamura T, Sauvageau G. NUP98-HOXA9 expression in hemopoietic stem cells induces chronic and acute myeloid leukemias in mice. *EMBO J* 2001; **20**: 350–361.
- Pineault N, Abramovich C, Ohta H, Humphries RK. Differential and common leukemogenic potentials of multiple NUP98-Hox fusion proteins alone or with Meis1. *Mol Cell Biol* 2004; **24**: 1907–1917.
- Shen WF, Montgomery JC, Rozenfeld S, Moskow JJ, Lawrence HJ, Buchberg AM et al. AbdB-like Hox proteins stabilize DNA binding by the Meis1 homeodomain proteins. *Mol Cell Biol* 1997; **17**: 6448–6458.
- Shen WF, Rozenfeld S, Kwong A, Kom ves LG, Lawrence HJ, Largman C. HOXA9 forms triple complexes with PBX2 and MEIS1 in myeloid cells. *Mol Cell Biol* 1999; **19**: 3051–3061.
- Thorsteinsdottir U, Kroon E, Jerome L, Blasi F, Sauvageau G. Defining roles for HOX and MEIS1 genes in induction of acute myeloid leukemia. *Mol Cell Biol* 2001; **21**: 224–234.
- Faber J, Krivtsov AV, Stubbs MC, Wright R, Davis TN, van den Heuvel-Eibrink M et al. HOXA9 is required for survival in human MLL-rearranged acute leukemias. *Blood* 2009; **113**: 2375–2385.
- Horton SJ, Walf-Vorderwulbecke V, Chatters SJ, Sebire NJ, de Boer J, Williams O. Acute myeloid leukemia induced by MLL-ENL is cured by oncogene ablation despite acquisition of complex genetic abnormalities. *Blood* 2008; **113**: 4922–4929.
- Buske C, Feuring-Buske M, Antonchuk J, Rosten P, Hogge DE, Eaves CJ et al. Overexpression of HOXA10 perturbs human lymphomyelopoiesis *in vitro* and *in vivo*. *Blood* 2001; **97**: 2286–2292.
- Magnusson M, Brun AC, Miyake N, Larsson J, Ehinger M, Bjornsson JM et al. HOXA10 is a critical regulator for hematopoietic stem cells and erythroid/megakaryocyte development. *Blood* 2007; **109**: 3687–3696.
- Roumier C, Lejeune-Dumoulin S, Renneville A, Goethgeluck AS, Philippe N, Fenaux P et al. Cooperation of activating Ras/rtk signal transduction pathway mutations and inactivating myeloid differentiation gene mutations in M0 AML: a study of 45 patients. *Leukemia* 2006; **20**: 433–436.
- Schneider F, Bohlender SK, Schneider S, Papadaki C, Kakadia P, Dufour A et al. AML1-ETO meets JAK2: clinical evidence for the two hit model of leukemogenesis from a myeloproliferative syndrome progressing to acute myeloid leukemia. *Leukemia* 2007; **21**: 2199–2201.
- Valk PJ, Bowen DT, Frew ME, Goodeve AC, Lowenberg B, Reilly JT. Second hit mutations in the RTK/RAS signaling pathway in acute myeloid leukemia with inv(16). *Haematologica* 2004; **89**: 106.
- Wang YY, Zhou GB, Yin T, Chen B, Shi JY, Liang WX et al. AML1-ETO and C-KIT mutation/overexpression in t(8;21) leukemia: implication in stepwise leukemogenesis and response to Gleevec. *Proc Natl Acad Sci USA* 2005; **102**: 1104–1109.
- Grisolano JL, O'Neal J, Cain J, Tomasson MH. An activated receptor tyrosine kinase, TEL/PDGFBetaR, cooperates with AML1/ETO to induce acute myeloid leukemia in mice. *Proc Natl Acad Sci USA* 2003; **100**: 9506–9511.
- Palmqvist L, Argiropoulos B, Pineault N, Abramovich C, Sly LM, Krystal G et al. The FLT3 receptor tyrosine kinase collaborates with NUP98-HOX fusions in acute myeloid leukemia. *Blood* 2006; **108**: 1030–1036.
- Schessl C, Rawat VP, Cusan M, Deshpande A, Kohl TM, Rosten PM et al. The AML1-ETO fusion gene and the FLT3 length mutation collaborate in inducing acute leukemia in mice. *J Clin Invest* 2005; **115**: 2159–2168.
- Yuan Y, Zhou L, Miyamoto T, Iwasaki H, Harakawa N, Hetherington CJ et al. AML1-ETO expression is directly involved in the development of acute myeloid leukemia in the presence of additional mutations. *Proc Natl Acad Sci USA* 2001; **98**: 10398–10403.
- Bjornsson JM, Andersson E, Lundstrom P, Larsson N, Xu X, Repetowska E et al. Proliferation of primitive myeloid progenitors can be reversibly induced by HOXA10. *Blood* 2001; **98**: 3301–3308.

- 30 Nielsen ML, Savitski MM, Zubarev RA. Improving protein identification using complementary fragmentation techniques in Fourier transform mass spectrometry. *Mol Cell Proteomics* 2005; **4**: 835–845.
- 31 Krause DS, Lazarides K, von Andrian UH, Van Etten RA. Requirement for CD44 in homing and engraftment of BCR-ABL-expressing leukemic stem cells. *Nat Med* 2006; **12**: 1175–1180.
- 32 Legras S, Gunthert U, Stauder R, Curt F, Oliferenko S, Kluin-Nelemans HC *et al*. A strong expression of CD44-6v correlates with shorter survival of patients with acute myeloid leukemia. *Blood* 1998; **91**: 3401–3413.
- 33 Bendall LJ, Bradstock KF, Gottlieb DJ. Expression of CD44 variant exons in acute myeloid leukemia is more common and more complex than that observed in normal blood, bone marrow or CD34+ cells. *Leukemia* 2000; **14**: 1239–1246.
- 34 Erkeland SJ, Verhaak RG, Valk PJ, Delwel R, Lowenberg B, Touw IP. Significance of murine retroviral mutagenesis for identification of disease genes in human acute myeloid leukemia. *Cancer Res* 2006; **66**: 622–626.
- 35 Yin B, Delwel R, Valk PJ, Wallace MR, Loh ML, Shannon KM *et al*. A retroviral mutagenesis screen reveals strong cooperation between Bcl11a overexpression and loss of the Nf1 tumor suppressor gene. *Blood* 2009; **113**: 1075–1085.
- 36 Touw IP, Erkeland SJ. Retroviral insertion mutagenesis in mice as a comparative oncogenomics tool to identify disease genes in human leukemia. *Mol Ther* 2007; **15**: 13–19.
- 37 Goverdhan S, Puntel M, Xiong W, Zirger JM, Barcia C, Curtin JF *et al*. Regulatable gene expression systems for gene therapy applications: progress and future challenges. *Mol Ther* 2005; **12**: 189–211.
- 38 Koschmieder S, Gottgens B, Zhang P, Iwasaki-Arai J, Akashi K, Kutok JL *et al*. Inducible chronic phase of myeloid leukemia with expansion of hematopoietic stem cells in a transgenic model of BCR-ABL leukemogenesis. *Blood* 2005; **105**: 324–334.
- 39 Ayton PM, Cleary ML. Molecular mechanisms of leukemogenesis mediated by MLL fusion proteins. *Oncogene* 2001; **20**: 5695–5707.
- 40 Rhoades KL, Hetherington CJ, Harakawa N, Yergeau DA, Zhou L, Liu LQ *et al*. Analysis of the role of AML1-ETO in leukemogenesis, using an inducible transgenic mouse model. *Blood* 2000; **96**: 2108–2115.
- 41 Yuan W, Payton JE, Holt MS, Link DC, Watson MA, DiPersio JF *et al*. Commonly dysregulated genes in murine APL cells. *Blood* 2007; **109**: 961–970.
- 42 Staber PB, Linkesch W, Zauner D, Beham-Schmid C, Guelly C, Schauer S *et al*. Common alterations in gene expression and increased proliferation in recurrent acute myeloid leukemia. *Oncogene* 2004; **23**: 894–904.
- 43 Somerville RP, Oblander SA, Apte SS. Matrix metalloproteinases: old dogs with new tricks. *Genome Biol* 2003; **4**: 216.
- 44 Lapidot T, Dar A, Kollet O. How do stem cells find their way home? *Blood* 2005; **106**: 1901–1910.
- 45 Stefanidakis M, Koivunen E. Cell–surface association between matrix metalloproteinases and integrins: role of the complexes in leukocyte migration and cancer progression. *Blood* 2006; **108**: 1441–1450.
- 46 Redondo-Munoz J, Ugarte-Berzal E, Terol MJ, Van den Steen PE, Hernandez Del Cerro M, Roderfeld M *et al*. Matrix metalloproteinase-9 promotes chronic lymphocytic leukemia B cell survival through its hemopexin domain. *Cancer Cell* 2010; **17**: 160–172.
- 47 Jin L, Hope KJ, Zhai Q, Smadja-Joffe F, Dick JE. Targeting of CD44 eradicates human acute myeloid leukemic stem cells. *Nat Med* 2006; **12**: 1167–1174.



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Supplementary Information accompanies the paper on the Leukemia website (<http://www.nature.com/leu>)