The ETS factor TEL2 is a hematopoietic oncoprotein

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TEL2/ETV7 is highly homologous to the ETS transcription factor TEL/ETV6, a frequent target of chromosome translocation in human leukemia. Although both proteins are transcriptional inhibitors binding similar DNA recognition sequences, they have opposite biologic effects: TEL inhibits proliferation while TEL2 promotes it. In addition, forced expression of TEL2 but not TEL blocks vitamin D3--induced differentiation of U937 and HL60 myeloid cells. TEL2 is expressed in the hematopoietic system, and its expression is up-regulated in bone marrow samples of some patients with leukemia, suggesting a role in oncogenesis. Recently we also showed that TEL2 cooperates with Myc in B lymphomagenesis in mice. Here we show that forced expression of TEL2 alone in mouse bone marrow causes a myeloproliferative disease with a long latency period but with high penetrance. This suggested that secondary mutations are necessary for disease development. Treating mice receiving transplants with TEL2-expressing bone marrow with the chemical carcinogen 2-ethyl-N-ethyl-N-nitrosourea (ENU) resulted in significantly accelerated disease onset. Although the mice developed a GFP-positive myeloid disease with 30% of the mice showing elevated white blood counts, they all died of T-cell lymphoma, which was GFP negative. Together our data identify TEL2 as a bona fide oncogene, but leukemic transformation is dependent on secondary mutations. (Blood. 2006;107:1124-1132)

Introduction

The ETS (E26 transformation specific) proteins belong to a large family of eukaryotic transcription factors (TFs) unique to the metazoan lineage and highly conserved throughout evolution. Some ETS proteins are expressed ubiquitously while others show a restricted tissue distribution. All ETS proteins possess a highly conserved 85--amino acid (aa) ETS domain that binds a purine-rich GGAA/T core motif present in promoters and enhancers of different target genes. A subgroup of ETS TFs, including the TEL proteins, also contains a conserved pointed (PNT) protein-protein interaction domain, which mediates the formation of homodimers/oligomers and heterodimers/oligomers. In TEL this domain is involved in transcriptional repression.

Vertebrate ETS TFs are implicated in many aspects of normal development and differentiation, including that of the hematopoietic system. For example, Ets-1, Fli-1, and Erg are expressed early during mouse development in the blood islands of the yolk sac where hemangioblasts are present. A high level of Pu-1 expression in CD34+ hematopoietic progenitors directs their differentiation toward the macrophage lineage, while low Pu-1 expression induces B-cell differentiation. The expression of IL-7Ra, a receptor essential for pro-B-cell development, is directly regulated by Pu-1 in lymphoid progenitors. The ubiquitously expressed TEL/ETV6 is essential for normal embryonic yolk sac angiogenesis and is important for the maintenance of the adult bone marrow microenvironment.

Several ETS genes, including ETS1, ETS2, PU1, FLI1, TEL/ETV6, and ERG, possess oncogenic properties. Human TEL/ETV6 is a frequent target of chromosome translocations both in hematopoietic malignancies as well as in some solid tumors. TEL/ETV6 translocations mostly encode oncogenic fusion proteins with some exceptions. In addition, TEL might also have tumor suppressor activity because the gene is often deleted during later stages of t(12;21) childhood pre--B-cell acute lymphoblastic leukemia (pre--B-ALL). This hypothesis is supported by the observation that TEL expression inhibits Ras-induced transformation of NIH3T3 fibroblasts.

Recently, a novel ETS gene highly related to TEL1 was isolated and coined TEL2. TEL2 is expressed predominantly in human hematopoietic tissues, contains a PNT domain and an ETS DNA binding domain, and localizes to the nucleus. TEL2 self-associates via its PNT domain but can also form heterodimers/oligomers with TEL1, suggesting that these proteins might affect each other’s function in vivo. Although both proteins function as transcriptional repressors in transient transfection assays, their biologic effects appear distinct. For example, TEL1 inhibits colony formation of Ras-transformed NIH3T3 fibroblasts, while TEL2 slightly stimulates colony formation. During vitamin D3--induced differentiation of the promonocytic cell line U937, TEL2 but not TEL1 expression is down-regulated, and forced expression of TEL2 blocks differentiation. In contrast, overexpression of TEL2 mutants

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The online version of the article contains a data supplement.

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containing an impaired ETS or PNT domain had no effect on vitamin D3–induced differentiation of U937 cells.29 Also TEL2, but not its PNT or ETS versions, cooperates with Myc in B-lymphomagenesis in mice,30 and its expression is up-regulated in some adult human leukemia samples29 and in more than 30% of pediatric ALL patients.30 Moreover, TEL2 is expressed in many human tumor cell lines.31 Together these data suggest a role for TEL2 in hematopoietic malignancy. Herein we show that expression of TEL2 alone in the mouse hematopoietic system causes a nontransplantable myeloproliferative disease (MPD) after a long latency period. This suggests that secondary mutations are necessary for disease development. Indeed, treatment with the DNA-damaging agent N-ethyl-N-nitrosourea (ENU) in mice receiving transplants greatly accelerated tumorigenesis.

Materials and methods

**Plasmid constructs, retroviral production, and viral titer determination**

TEL2 cDNA was cloned upstream of the IRES element into the unique EcoRI site of the murine stem cell virus (MSCV) IRES- GFP vector.29,32-35 Retroviral constructs were used to generate replication-incompetent retroviral stocks by transient transfection of the Ecotropic Phoenix 293T packaging cells line36 using Fugene 6 (Roche, Indianapolis, IN). Viral titers were determined by infection of NIH3T3 cells in the presence of Polybrene (Sigma Chemical, St. Louis, MO) (8 μg/mL) and varied between 4 × 10^3 to 1 × 10^5 CFU/mL.

**Bone marrow extraction, Lin− isolation, and retroviral transduction**

Bone marrow (BM) cells were harvested by flushing the femurs and tibiae of FVB-J male donor mice 3 days after intraperitoneal injection of 150 μg/g body weight 5-fluorouracil (SFU) (Sigma Chemical). Lineage-negative (Lin−) cells were purified by immunodepletion of cells presenting myeloid, erythroid, and lymphoid differentiation markers using biotinylated mouse antibodies Ly-6G (Pharmingen, San Diego, CA), Cd11b (Pharmingen), Cd45R/B220 (Pharmingen), CDS 5 (Pharmingen) and TER-119 (Pharmingen) and streptavidin-coated beads (Dynabeads M-280 streptavidin; Brown Deer, WI). The resulting progenitor-enriched population was prestimulated for 48 hours in Iscove medium (Gibco, Carlsbad, CA), 20% fetal bovine serum (Hyclone, South Logan, UT), supplemented with interleukin-3 (IL-3; 20 ng/mL) (Preprotech, London, United Kingdom), interleukin-6 (IL-6; 30 ng/mL) (Peprotech), interleukin-7 (IL-7; 10 ng/mL) (Peprotech), and stem cell factor (SCF; 50 ng/mL) (R&D Systems, Minneapolis, MN) in nontissue culture grade plastic Petri dishes. After prestimulation the cells were plated onto Retronectin (Takara, Otsu, Japan)—coated plates and incubated with the retroviral supernatants twice daily for 2 days in the presence of cytokines.

** Colony-forming unit assays**

Transduced and normal Lin− cells were plated in semisolid methylcellulose medium (MC1) plus cytokines (10 μg/mL insulin, 200 ng/mL human transferrin, 50 ng/mL SCF, 10 ng/mL IL-3, 10 ng/mL IL-6, 3 U/mL erythropoietin [Epo], StemCell Technologies 03434, Victoria, BC, Canada) at a concentration of 1 × 10^5 cells per dish Erythroid colony-forming unit (CFU-E), granulocyte erythrocyte monocyte macrophage colony-forming unit (CFU-GEMM), and granulocyte macrophage colony-forming unit (CFU-GM) colonies were scored 10 to 14 days later, pooled, and reseeded at a density of 1 × 10^5 cells per dish into secondary methylcellulose (MC2) cultures. Part of the cells were used to determine the fraction of GFP-positive cells using a FACSCalibur (Becton Dickinson, Franklin Lakes, NJ). We continued MC replating until the cultures were exhausted (usually 3 to 4 consecutive rounds of plating) due to terminal differentiation of the cells.

**Long-term culture-initiating cell (LTC-IC) assays**

Transduced and normal BM cells (5 × 10^3 cells per dish) were plated onto irradiated M2-10B4 stromal layers37 in Myelocult media (StemCell Technologies M5300) to which freshly prepared 0.5 μg/mL hydrocortisone (hydrocortisone 21-hemisuccinate; Sigma Chemical) was added. The cells were cultured for 4 weeks, while replacing half the medium every 7 days. At the end of this period the cells were harvested, analyzed for GFP expression by fluorescence-activated cell sorting (FACS), and plated in methylcellulose media. We scored the number and type of colonies 2 weeks later. Serial MC replating and concomitant GFP analyses were performed for 6 to 8 weeks.

**Bone marrow transplantation (BMT)**

Female 6- to 12-week-old FVB mice were lethally irradiated (single dose of 950 cGy) and 24 hours later received transplants by tail vein injection with 300 μL PBS plus heparin (30 U/mL) containing 3 × 10^6 to 5 × 10^6 Lin− cells transduced with retrovirus or not. After BMT, the animals were evaluated daily for possible signs of disease. Peripheral blood was obtained monthly by retro-orbital phlebotomy and was analyzed by FACS to determine the percentage of white blood cells (WBCs), red blood cells (RBCs), and platelets expressing GFP. Complete blood counts (CBCs) were performed with a Hemavet 3700 (Drew Scientific, Cumbria, United Kingdom), and Giemsa-stained blood smears were analyzed to verify the presence of abnormal cells.

**ENU mutagenesis**

**Bone marrow transplantation (BMT),” but 5 weeks after transplantation the recipients received intraperitoneal injections twice, at a 48-hour interval, with 100 mg/kg ENU. The health status of the animals was followed as described in “Bone marrow transplantation (BMT).”**

**Analysis of diseased mice and tissue preparation**

All the following animal procedures were conducted in accordance with the U.S. Public Health Service Policy on the Humane Care and Use of Laboratory Animals. Peripheral blood of moribund animals was isolated for GFP analysis, blood smears, and CBCs. The animals were killed using CO2 asphyxiation, and the weight of the spleen and liver was recorded. All organs were recovered, fixed in 10% neutral-buffered formalin, processed and embedded in paraffin, sectioned at 4 to 5 μm, and stained with hematoxylin and eosin (H&E) for routine histologic examination. Sternum and hind limbs were subjected to an additional decalcification step before tissue processing. Select tissues were also processed for immunohistochemical analysis with antibodies to the hematopoietic markers CD3 (Dako, Carpinteria, CA), CD45R/B220 (Pharmingen, San Diego, CA), terminal deoxynucleotidyl transferase (TdT; SuperTeches, Bethesda, MD), myeloperoxidase (MPO; Dako), TER-119 (Pharmingen, San Diego, CA), GATA1 (Santa Cruz Biotechnology, Santa Cruz, CA), and green fluorescent protein (GFP; Clontech, Palo Alto, CA).

Single-cell suspensions were prepared from bone marrow, spleen, and liver for analysis of GFP expression by FACS and for the preparation of cytospin slides (5 × 10^4 cells per slide) for morphologic examination after May-Grünewald-Giemsa (MGG) staining. After lysis of the red cells, the leukocytes were analyzed immediately for surface marker expression or frozen in fetal calf serum (FCS)/10% DMSO for later analysis. Images of tissue sections and cytospins were obtained using a BX51 microscope equipped with a Uplan FL 40 ×0.75 numeric aperture (NA) or a 100 ×1.30 NA objective (Olympus, Tokyo, Japan). Images were acquired using a SPOT camera and SPOT Advanced imaging software (Diagnostic Instruments, Sterling Heights, MI). Original magnification for tissue sections and cytospins was 400 × and 1000 ×, respectively.
Flow cytometric analysis

Single-cell suspensions of BM, spleen, and liver were washed and incubated for 30 minutes on ice in staining medium (SM: DMEM supplemented with 10% fetal bovine serum) containing human γ-globulin (100 mg/mL; Sigma Chemical) to block Fc receptors. After washing, cells were incubated with monoclonal antibodies (CD3c, CD4, CD8, CD11b/Mac1, CD19, CD34, B220, TER-119, Gr1, Sca1, c-Kit, Flt3, all from Pharmingen, San Jose, CA; anti-mouse IgM from Southern Biotechnology Associates, Birmingham, AL) on ice for 30 minutes. After a final washing step, cells were resuspended in SM and analyzed using a BD Biosciences FACS Calibur flow cytometer (BD Biosciences, San Jose, CA), selecting single cells by gating on forward versus side light scatter. Wild-type and TEL2-expressing bone marrow cells were cultured for 2 weeks, and apoptotic cells were identified by FACS after annexin V–FITC staining. Propidium iodine staining was used to exclude dead cells.

Western blotting

For Western blotting, protein extracts were prepared from spleen samples of diseased TEL2 and healthy control mice, killed at the same time, using TRI Reagent (Sigma Chemical), following the manufacturer’s instructions. After quantification using the BCA Protein Assay Reagent (Pierce Chemical, Rockford, IL), 30 μg total protein was separated on 10% SDS-PAGE (sodium dodecyl sulfate–polyacrylamide gel electrophoresis) gels under reducing conditions and transferred onto a polyvinylidene difluoride membrane (Millipore, Billerica, MA). The membranes were incubated with the antibodies specific for Bcl-xL (Transduction Laboratories, Lexington, KY), recognizing both Bcl-xl and Bcl-xxs, Bcl2 antibody (554218; Pharmingen, San Diego, CA), and TEL2 antibody.30

Secondary transplantsations

Lethally irradiated (9.5 Gy [950 rad]) recipient mice received injections in the tail vein with 1 ¥ 106 freshly isolated bone marrow or spleen cells from moribund primary recipients after red cell lysis. Mice receiving transplants were followed and killed as described for the primary recipients.

Affymetrix GeneChip analysis

BM from 5FU-treated mice was either mock transduced (Un-BM) or transduced with MSCV-ires-GFP (GFP-BM) or MSCV-TEL2-ires-GFP (TEL2-BM) retrovirus as described in “Bone marrow transplantation (BMT).” After transduction, cells were allowed to recover in culture for 48 hours and were sorted for GFP expression by FACS. RNA of the GFP-positive cells and mock-transduced cells was isolated using Trizol (Invitrogen, Carlsbad, CA) following the manufacturer’s recommendations.

RNA quality was confirmed by UV spectrophotometry and by analysis on an Agilent 2100 Bioanalyzer (Agilent Technologies, Foster City, CA). Ten micrograms of total RNA was processed in the St Jude Hartwell Center Core Facility according to the standard Affymetrix (Santa Clara, CA) protocol18 and analyzed on the Affymetrix MOE-430A GeneChip array. Signal values, detection calls, and pairwise expression analyses were performed using the default parameters within the statistical algorithm of the Affymetrix GCOS software version 1.2. Signal values were scaled to a 2% trimmed mean target value of 500. Stringent selection criteria were applied to identify differential expression associated with TEL2. The following pairwise comparisons were performed: TEL2-BM versus GFP-BM, TEL2-BM versus Un-BM, and GFP-BM versus Un-BM. Initial selection was based on differential expression between TEL2-BM and GFP-BM. Probe sets with a log2 ratio greater than one (more than 2-fold change) plus a “change call” (P < .006, Wilcoxon signed rank test) were selected for further evaluation. To best identify TEL2-associated changes, the initial selection was filtered to exclude probe sets with less than 2-fold change in TEL2-BM versus Un-BM and those with more than 2-fold change in GFP-BM versus Un-BM. Probe set annotations were obtained from the Affymetrix website.39 Functional classification of genes was performed using the Affymetrix Gene Ontology browser. Biologic processes significantly enriched within the TEL2-associated gene list were identified using the χ2 test (P < .001).

Results

TEL2 expression affects the colony-formation potential of primitive myeloid progenitors in vitro

To test the effect of forced TEL2 expression in murine hematopoietic cells, Lin− cells of 5-FU treated donor mice were transduced with MSCV-ires-GFP or MSCV-TEL2-ires-GFP retrovirus (Figure 1A). FACS analysis 4 days later showed that 10% to 50% of the
cells expressed GFP (not shown), and were used for in vitro colony-forming assays and in vivo bone marrow reconstitution of lethally irradiated recipient mice.

To assess whether TEL2 expression conferred a proliferative advantage to hematopoietic cells in vitro, we compared the growth characteristics of Lin− cells transduced with empty retrovirus (MSCV-BM) or a TEL2 retrovirus (TEL2-BM) with that of uninfected Lin− cells (Un-BM). To assess the colony-forming capacity of committed progenitors we plated the cells into MC1 and 2 weeks later scored the cultures for the number and type of colony-forming units (CFU-E, CFU-GEMM, CFU-GM). After counting, 10^7 cells from the MC1 were replated into fresh MC and CFUs were scored, and this process was repeated until the CFU activity was exhausted (3 to 4 rounds). However, we observed no differences in the type, size, or number of the Un-BM, MSCV-BM, and TEL2-BM colonies (Figure 1B). We next tested whether TEL2 expression affected the growth of primitive progenitors and followed the clonogenic potential of TEL2-BM and MSCV-BM progenitors in myeloid MCs after long-term culture-initiating cell (LTC-IC) cultures. This experiment was repeated twice with both types of bone marrow containing 10% GFP-positive cells. After 4 weeks of LTC-IC, cells were collected and plated into serial MC assays. While the percentage of GFP-positive cells in MSCV-BM was similar before and after LTC-ICs, the percentage of GFP-positive cells in the MC1 of TEL2-BM LTC-ICs had increased to 60% and the number of colonies in the MC1 was 3-fold higher than that in the MC1 of MSCV-BM cells. Upon serial replating, the elevated percentage of GFP-positive cells in the TEL2 MCs remained, but the number of colonies rapidly dropped to a level similar to that of MSCV-BM cells (Figure 1C). Thus, TEL2-expressing primitive progenitors appeared to have a growth advantage in LTC-IC assays only. When transduced Lin− bone marrow cells used for the LTC-ICs were directly plated onto tissue culture plastic, they formed their own stromal layer. Under these conditions TEL2-BM cells grew faster than control MSCV-BM cells (not shown), and the percentage of GFP-positive cells again increased from 10% to 60% during 3 weeks of culture. This suggested that interaction with stromal cells, whether in LTC-IC cultures or in stroma-forming BM cultures, provided a growth advantage to TEL2-BM cells.

**Bone marrow–expressing TEL2 induces a myeloproliferative disease in FVB recipient mice**

We simultaneously tested whether TEL2 expression in reconstituted BM would affect normal hematopoiesis in vivo and transplanted Un-BM, MSCV-BM, and TEL2-BM cells into 7 lethally irradiated FVB recipients and assessed their peripheral blood parameters and the percentage of GFP-positive cells monthly. The same experiment was repeated 3 months later, and the combined results of these experiments are presented in Figure 1D-G. All animals receiving transplants engrafted, and the peripheral blood counts (PBCCs) remained stable for 6 months, followed by an increasing leukocytosis (Figure 1D) until the mice became moribund 2 to 3 months later. At the time the animals were killed, they showed an average peripheral WBC count of 75 × 10^9/L (75 × 10^9/μL) (Figure 1E), with all cells expressing GFP (Figure 1F). All TEL2-BM mice died within 8 to 14 months after transplantation (Figure 1G). Instead, mice receiving transplants with Un-BM or MSCV-BM cells showed normal blood parameters and remained healthy, although 2 MSCV-BM animals died of causes not related to hematopoietic disease.

Average CBCs of mice (Un-BM, MSCV-BM, and TEL2-BM) were determined at the time of death of the TEL2-BM mice (Table 1).

Assessment of cell morphology of peripheral blood smears (Figure 2A) showed anemia, large numbers of mature and immature neutrophils, and the presence of some blastlike cells. BM cytopsins (Figure 2B) of affected mice revealed the presence of mostly myeloid cells with a similar abundance of mature and immature neutrophils, but also blastlike cells, monocytes, and promyelocytes. All TEL2-BM animals showed splenomegaly and hepatomegaly, due to extensive infiltration of a population of cells with a high mitotic index, which expressed GFP (Figure 2C-D). The spleen had completely lost its normal architecture (not shown), and the cell population in the red pulp consisted of erythroid precursors, megakaryocytes, and a vast excess of mature granulocytes and immature myeloid cells. Unlike the spleen, the liver infiltrate consisted of mature granulocytes and immature myeloid cells. The immature cells were the dominant constituent in both tissues, and most of them expressed MPO consistent with a granulocytic lineage (Figure 2E). Western blot analysis confirmed expression of TEL2 in the spleen of the diseased mice (Figure 2F). Given that the antipapoptotic effect of TEL2 on B-cell progenitors is associated with increased expression of Bcl2 but not Bcl-xl,27 we also tested the expression of these 2 antipapoptotic proteins in the same spleen samples. Compared with MSCV-BM spleen cells, 2 TEL2-BM tumor samples expressed an increased amount of Bcl-xl and Bcl-x, whereas Bcl2 expression was increased in 4 of 6 tumor samples. Two tumors samples showed no up-regulation of Bcl2 or Bcl-xl, indicating that the antipapoptotic proteins were not consistently up-regulated in all tumors. FACS analysis of BM (Figure 3) showed that the malignant cells were Mac1+ and Gr1+. BM also showed an increase in Sca1+ c-Kit+, and Thy1.1+ progenitors. Together these features are consistent with a diagnosis of a chronic myelomonocytic leukemia (CMML)–like disease,40 suggesting that TEL2 enlarges an early myeloid progenitor population.

Transplantation of the diseased bone marrow into 12 sublethally irradiated secondary recipients failed to reproduce disease. The percentage of GFP-positive cells in the peripheral blood of the secondary recipients ranged between 5% and 35% during the first 3 months after transplantation but diminished to almost undetectable levels thereafter. Only 1 of 12 secondary recipients showed a distinct myeloproliferation (93 × 10^9/L [93 × 10^9/μL]) associated with organ infiltration, but the percentage of GFP-positive cells in peripheral blood was 12% and the population of cells infiltrating the different organs was GFP negative.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>UN-BM</th>
<th>MSCV-BM</th>
<th>TEL2-BM*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Complete WBC count, × 10^9/L</td>
<td>8.6 ± 3.6</td>
<td>10.3 ± 4.9</td>
<td>76 ± 35.32</td>
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<tr>
<td>Neutrophil count, × 10^9/L</td>
<td>1.48 ± 0.52</td>
<td>2.36 ± 0.87</td>
<td>14.7 ± 13.94</td>
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<tr>
<td>Lymphocyte count, × 10^9/L</td>
<td>4.99 ± 1.68</td>
<td>4.64 ± 1.6</td>
<td>15.3 ± 14.05†</td>
</tr>
<tr>
<td>Monocyte count, × 10^9/L</td>
<td>0.55 ± 0.28</td>
<td>0.57 ± 0.39</td>
<td>3.2 ± 2.8</td>
</tr>
<tr>
<td>Eosinophil count, × 10^9/L</td>
<td>0.019 ± 0.008</td>
<td>0.013 ± 0.005</td>
<td>1.14 ± 0.96</td>
</tr>
<tr>
<td>Basophil count, × 10^9/L</td>
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<td>0 0</td>
<td>0.23 ± 0.19</td>
</tr>
<tr>
<td>MCH, pg</td>
<td>13.42 ± 1.45</td>
<td>15.19 ± 1.31</td>
<td>17.9 ± 4.4</td>
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<td>RBC count, × 10^12/L</td>
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<td>7.6 ± 0.811</td>
<td>4.66 ± 1.65</td>
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<td>MCV, fl</td>
<td>44.84 ± 3.14</td>
<td>49.23 ± 3.32</td>
<td>62.68 ± 14.29</td>
</tr>
<tr>
<td>Platelet count, × 10^12/L</td>
<td>17.17 ± 68.63</td>
<td>1406 ± 204</td>
<td>1094 ± 303</td>
</tr>
<tr>
<td>Hemoglobin level, g/L</td>
<td>127 ± 17 130 ± 17</td>
<td>84 ± 14</td>
<td></td>
</tr>
</tbody>
</table>

†The high number of lymphocytes in the Hemavet counts of the TEL2-BM mouse blood samples is a gross overestimate because the blast cells and partially differentiated myeloid cells have the same size as lymphocytes and are therefore scored as "lymphocytes." This was confirmed by morphologic analysis of the blood smears (Figure 2A).

Table 1. Average complete blood count (CBC) in peripheral blood of mice receiving transplants with Un-BM, MSCV-BM, and TEL2-BM, at the moment of death of the TEL2-BM mice

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We concluded that this unique case was caused by a genetic event not
related to overexpression of TEL2.

**Annexin V staining of FVB Lin\(^{-}\) BM cells**

Because TEL2 expression in pre-B cells inhibits their rate of apoptosis,\(^{30}\) we assessed whether a similar effect of TEL2 in myeloid cells might provide a proliferative advantage in the mice receiving transplants. We analyzed the percentage of apoptotic cells by annexin V staining in Lin\(^{-}\) Un-BM cells or in GFP-positive MSCV-BM and GFP-positive TEL2-BM cells after culture for 7 days in liquid media promoting myeloid proliferation. Compared with both Un-BM and GFP-positive MSCV-BM cells, GFP-positive TEL2-BM cells showed a consistent 50% decrease in the number of apoptotic cells (Figure 4). This suggested that the myeloproliferative disease caused by TEL2-BM cells in mice might in part be attributed to a reduced level of apoptosis of myeloid progenitors in vivo.

To obtain information about TEL2-induced changes in gene expression, we transduced FVB Lin\(^{-}\) cells with MSCV-IRES-GFP (MSCV-Lin\(^{-}\) cells) or MSCV-TEL2-IRES-GFP retrovirus (TEL2-Lin\(^{-}\) cells) and sorted GFP-positive cells 72 hours later by FACS. Relative RNA abundance in these 2 GFP-positive cell populations and in untransduced Lin\(^{-}\) cells was determined by Affymetrix microarray analysis (MOE-430A). We then compared gene expression differences between Un-Lin\(^{-}\) and MSCV-Lin\(^{-}\) and MSCV-Lin\(^{-}\) and TEL2-Lin\(^{-}\) cells (Figure 5). This 3-way comparison allowed elimination of genes whose expression was changed due to
GeneChip microarrays. Log2 ratio values from pairwise comparisons are plotted. (A) Relative expression of 195 probe sets identified as differentially expressed (see “Materials and methods”) in both TEL2 versus GFP (x-axis) and TEL2 versus uninfected (y-axis) bone marrow. (B) Relative expression of differentially expressed probe sets found enriched (P < .001) in the Gene Ontology categories of apoptosis (left) or differentiation (right). Gene symbols are indicated in the panels.

Discussion

TEL2, a new member of the ETS TFs, was identified by several groups a few years ago. The protein was coined TEL2 because of its high homology with TEL (38.2% identity). However, despite this homology, its function appears to be quite distinct from that of TEL. TEL has an antiproliferative effect whereas TEL2 has a mostly proliferative effect, although a direct assessment of the tumorigenic activity of TEL2 alone has not been determined. In our in vitro LTC-IC studies strongly suggested that TEL2 stimulated the growth of primitive progenitors, provided they are in contact with a feeder layer, a condition that mimics the bone marrow environment. This effect on myeloid cells is different from TEL2’s effect on B-cell progenitors, whose growth is stimulated in the absence of feeder cells.

TEL2 is specifically expressed in the human hematopoietic system, and its expression is up-regulated in some adult human leukemias and more than 30% of pediatric B-ALL. TEL2 also accelerated B lymphomagenesis in Eμ-Myc mice, a model for Burkitt lymphoma, and possibly cooperates with N-MYC/C-MYC in pediatric B-cell lymphoma. Therefore, we tested whether enforced TEL2 expression alone in mouse bone marrow predisposed the animals to hematopoietic malignancy. Our experiments indeed verified the oncogenicity of TEL2 in this setting, but the long latency of disease indicated that secondary mutations are necessary for disease development.

The disease was not transplantable in secondary recipients, although the proliferation of the myeloid cells appeared strictly cell autonomous given the tight correlation between GFP expression and the number of leukemic cells (Figure 1E). This indicated that...
progenitor cells expressing TEL2 were not immortalized and that the conditions in the reconstituted bone marrow were unable to support reconstitution by the MPD. Similar observations have been reported for MPD caused by NUP98-Hox A. \(^{57}\) Annexin V analysis of TEL2-expressing bone marrow after a short period (1 week) of in vitro culture showed that TEL2 suppressed the apoptotic index of these cells as it does inhibit apoptosis in Myc-overexpressing or wild-type B-cell progenitors. \(^{30}\)

Given the large excess of myeloid progenitors in the BM and the outcome of the Affimetrix array analysis of Lin\(^-\) cells 72 hours after transduction with TEL2 or vector retrovirus suggested that TEL2 also inhibits apoptosis of myeloid progenitors. Several proapoptotic genes were repressed, such as Trp53inp1, Stk17b, Dapk2, and Bmf, which might mediate this effect. In addition, 4 of 6 mice analyzed for Bcl2 or Bcl-x\(_l\) expression in myeloid cells, which invaded the spleen, showed up-regulation of Bcl2, Bcl-x\(_l\) and Bcl-x\(_s\). Similar to the effect of TEL2 in B cells, \(^{30}\) up-regulation of Bcl2 and Bcl-x is probably not at the transcriptional level, because Bcl2 and Bcl-x mRNA was not increased in the Affimetrix arrays of TEL2-transduced bone marrow. We believe that Bcl2 and Bcl-x overexpression contributes to the expansion of the myeloid cells, but currently we do not know which events substitute for Bcl2 and Bcl-x overexpression in the 2 tumor samples that did not overexpress these proteins. However, changes in expression of genes inhibiting (Btg1, Btg2, Cdkn1B, Rb1cc1) or stimulating (cKit) proliferation could have an additive effect on the enlargement of the myeloid progenitor pool. It remains to be determined whether any of these genes are direct transcriptional targets of TEL2, an issue that we will address in future studies.

It is reasonable to speculate that myeloid progenitors in the bone marrow of mice receiving transplants have a reduced apoptotic rate, but this does not lead to increased numbers of myeloid cells in the peripheral blood during the first 6 months after transplantation, a feature also apparent in mice expressing MLL fusion oncogenes and AML1-ETO. \(^{58,59}\) It is possible that compensatory mechanisms control the number of myeloid cells entering the periphery and that additional genetic changes, possibly those that affect cell proliferation, may override this check, resulting in accumulation of differentiating myeloid cells in the peripheral blood. In addition, expansion of the myeloid compartment during the first 6 months after transplantation would increase the frequency of secondary mutations and thereby promote development of disease. The extensive differentiation of the cells is the likely cause of the chronic character of the disease, which after initial detection took 2 to 3 months to kill the animal. The differentiation of the malignant cells in affected animals seems in conflict with the observation that TEL2 inhibits differentiation of HL60 and U937 myeloid cells. \(^{29}\) The reason for this is unknown but, unlike bone marrow, cell lines harbor many mutations, which might account for this difference in behavior.

Given the development of myeloproliferative disease in mice receiving transplants with TEL2, we expected that ENU mutagenesis would induce myeloid leukemia. This occurs in mice expressing RUNX1-ETO or CBFB-MYH11, \(^{59,60}\) indicating that these fusion transcription factors specifically affect myeloid cells despite the fact that conditional RUNX1-ETO knock-in mice also express the fusion protein in their T cells. \(^{61}\) Our experiments indicate that in combination with ENU all TEL2-BM mice developed distinct myeloid aberrations in the bone marrow within 3 to 6 months after ENU treatment, which is much more rapid than TEL2-BM mice only (Figure 1G). However, only 3 mice showed elevated numbers of dysplastic cells in the peripheral blood, indicating a possible development of myeloid leukemia. Surprisingly, all mice also developed GFP-negative T-cell lymphoma. We must conclude that changes caused by
TEL2 expression promoted T-cell lymphomagenesis, because none of the control ENU-treated mice developed T-cell lymphoma. It is possible that T-cell proliferation is stimulated indirectly by TEL2-expressing hematopoietic cells, which could contribute to increased genetic instability, making these cells more susceptible to ENU-mediated leukemic transformation. The lethal lymphoid disease might have prevented the myeloid preleukemic state of these mice to develop into AML.

Together our results suggest that TEL2 is a hematopoietic oncogene in the mouse, and its overexpression in some adult and 30% of pediatric ALL samples also argues for a causative role of TEL2 in human leukemogenesis.

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The ETS factor TEL2 is a hematopoietic oncoprotein

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