5-Lipoxygenase Is a Candidate Target for Therapeutic Management of Stem Cell–like Cells in Acute Myeloid Leukemia

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Abstract
Nonsteroidal anti-inflammatory drugs such as sulindac inhibit Wnt signaling, which is critical to maintain cancer stem cell–like cells (CSC), but they also suppress the activity of 5-lipoxygenase (5-LO) at clinically feasible concentrations. Recently, 5-LO was shown to be critical to maintain CSC in a model of chronic myeloid leukemia. For these reasons, we hypothesized that 5-LO may offer a therapeutic target to improve the management of acute myeloid leukemia (AML), an aggressive disease driven by CSCs. Pharmacologic and genetic approaches were used to evaluate the effects of 5-LO blockade in a PML/RARα-positive model of AML. As CSC models, we used Sca-1+/lin– murine hematopoietic stem and progenitor cells (HSPC), which were retrovirally transduced with PML/RARα. We found that pharmacologic inhibition of 5-LO interfered strongly with the aberrant stem cell capacity of PML/RARα-expressing HSPC. Through small-molecule inhibitor studies and genetic disruption of 5-LO, we also found that Wnt and CSC inhibition is mediated by the enzymatically inactive form of 5-LO, which hinders nuclear translocation of β-catenin. Overall, our findings revealed that 5-LO inhibitors also inhibit Wnt signaling, not due to the interruption of 5-LO–mediated lipid signaling but rather due to the generation of a catalytically inactive form of 5-LO, which assumes a new function. Given the evidence that CSCs mediate AML relapse after remission, eradication of CSCs in this setting by 5-LO inhibition may offer a new clinical approach for immediate evaluation in patients with AML.

Cancer Res; 74(18); 5244–55. © 2014 AACR.

Introduction
Actually, one of the major clinical challenges in oncology is to target the cancer stem cell–like cells (CSC). Current therapeutic strategies, regardless of conventional cytotoxic or novel molecular approaches, often fail to eradicate cancer completely because they are unable to efficiently target CSCs. In this respect, a key signaling pathway for the maintenance of CSCs in many cancers, including colorectal, breast, and pancreatic cancer, is the Wnt signaling pathway (1–4).

In acute and chronic myeloid leukemias (AML and CML) the role of the Wnt signaling activation for the pathogenesis and the maintenance of the CSCs is well understood (5–7). The fact that in leukemia inhibition of the Wnt signaling directly targets the CSCs (leukemic stem cell, LSC) means that compounds and related signaling pathways that are able to inhibit Wnt signaling offer considerable potential within the setting of maintenance therapy of leukemia. Deregulated activation of the Wnt signaling leads to aberrant self-renewal of the LSCs in AML and CML and is fundamental to its maintenance (7–9). The activation of Wnt signaling in leukemia is mainly due to either a direct deregulation of key effectors such as APC, Axin, β-catenin, γ-catenin, or TCF/LeF by, i.e., mutations, or functional inhibition of their key regulators by leukemia-associated fusion proteins (LAFP), including PML/RARα, AML-1/ETO, MLL/AF9, or BCR/ABL. These LAFPs are generated by chromosomal translocations such as t(15;17) (PML/RARα), t(8;21) (AML-1/ETO), t(9;11)(MLL/AF9), or t(9;22) (BCR/ABL; refs. 6, 7, 10–12). The aberrant self-renewal of the LSCs depends on the aberrant activation of Wnt signaling by the LAFPs (6, 7, 10). The overexpression of γ-catenin, one of the key players in Wnt signaling in HSPCs, leads to the induction of leukemia in vivo (12).

Nonsteroidal anti-inflammatory drugs (NSAID) developed as COX-1/2 inhibitors are efficient Wnt signaling inhibitors.
Cell lines and chemicals

Meglomavirus (pRL-CMV) vectors were already described (12). pGL3- (Promega) have also been described previously (19). The identity of these cells was routinely controlled by immunodetection (immuno (Gibco). The identity of these cells was routinely controlled by fluorescence or Western blotting (unpublished observations). The spleens were embedded in paraffin blocks according to a conventional tissue processing procedure. Immunohistochemistry was performed on 5 μm sections of each paraffin block. The sections were deparaffinized with xylol (Sigma) and rehydrated through graded alcohol series (100%, 95%, 80%, 70%, 50%, 40%, 30%, 20%, and 10%, H2O). Antigen unmasking was achieved through heating with 0.01% sodium citrate buffer, pH 6.0, for 4 minutes. Endogenous peroxidase activity was blocked using 3% hydrogen peroxide.

Materials and Methods

Plasmids

The retroviral vectors, PINCO-PML/RARα and PINCO-HA-β-cateninS33A, as well as the expression vector, pCDNA3.1-5LO, have been described elsewhere (16). The TopFlash/FopFlash system and the Renilla luciferase pRL-TK construct (Promega) have also been described previously (19). The pGL3-γ-catenin construct and the pGL3basic and pRL-cytomegalovirus (pRL-CMV) vectors were already described (12).

Cell lines and chemicals

293T, NB4, and THP-1 cells were obtained from the American Type Culture Collection (ATCC). 293T cells were cultured in DMEM (Gibco), supplemented with 10% fetal calf serum (FCS; Gibco). NB4, THP-1, and U937-PR9 cells were maintained in RPMI medium supplemented with 10% FCS (Sigma). The identity of these cells was routinely controlled by immunodetection (immunofluorescence or Western blotting) of PML/RARα (NB4 and U937-PR9) or the induction of 5-LO activity (THP-1) and the response to all-trans retinoic acid-induced granulocytic differentiation (NB4), respectively. CJ-13,610 was synthesized in-house (see Supplementary Materials and Methods) and calcium ionophore A23187, arachidonic acid, zileuton, indomethacin, and diclofenac were purchased from Sigma-Aldrich. The sEH inhibitor trans-4-[3-adamantan-1-ylureido] cyclohexyloxyl]-benzoic acid (t-AUCB) I was kindly provided by Bruce Hammock (University of California at Davis, Davis, CA).

Isolation of Sca1+/lin - hematopoietic stem and progenitor cells

Sca1+/lin - HSPCs were isolated from either 8- to 12-week-old female C57BL/6N mice (Janvier) or male and female B6.129S2-Alox5tm1Fun/J (Jackson Laboratory) and B6.129X1-Phtk2tm1Gona/J mice (provided by Frank Gonzalez, NIH, Bethesda, MD) after killing by CO2 asphyxiation. The isolation of Sca1+/lin - cells was performed as described previously (20). Two days after the isolation cells were retrovirally transfected.

Retroviral infection

Ectropic Phoenix packaging cells were transfected with PINCO-PML/RARα or empty vector by calcium phosphate precipitation according to widely established protocols. Retroviral supernatant was collected at days 2 and 3 after transfection. Infection of the target cells was performed as described previously (12). The minimal accepted infection efficiency was 70%, as assessed by the detection of GFP-positive cells by FACS. Differences in the infection efficiency between samples did not exceed 10%.

Determination of 5-LO product formation in THP-1 and Sca1+/lin - cells

To induce 5-LO expression, differentiation of THP-1 or transduced Sca1+/lin - cells were either induced by the addition of TGFβ1 (1 ng/mL) and 1, 25-dihydroxyvitamin D3 (50 nmol/L) or by addition of GM-CSF (20 ng/mL, 60 ng/mL; Cell Concepts) for 4 days. Determination of 5-LO product formation was performed as previously described (21). For Sca1+/lin - cells, leukotriene formation in the supernatant was measured with a LTB4-ELISA, according to the manufacturer’s instructions (Enzo Life Science). 5-LO product formations of CJ-13,610-, zileuton-, and sulindac-treated THP-1 cells were analyzed by high-performance liquid chromatography (HPLC) as described previously (22). For indomethacin, leukotriene levels in the supernatant were analyzed by liquid chromatography/tandem mass spectrometry (LC/MS-MS) as described previously (23). 5-LO product formation included 5-HETE for CJ-13,610 and sulindac and LTB4 for indomethacin.

Immunohistochemistry

The spleens were embedded in paraffin blocks according to a conventional tissue processing procedure. Immunohistochemistry was performed on 5 μm sections of each paraffin block. The sections were deparaffinized with xylol (Sigma) and rehydrated through graded alcohol series (100%, 95%, 80%, 50%, H2O). Antigen unmasking was achieved through heating (95°C) with 0.25 mmol/L EDTA buffer for 50 minutes. The endogenous peroxidase activity was blocked using 3% hydrogen peroxide.
hydrogen peroxide (5 minutes, room temperature), followed by preincubation with antibody mix (TBS, 2% BSA, 2% normal goat serum, 0.02% Tween 20) for 20 minutes at room temperature. The primary antibody (mouse monoclonal anti-β-catenin; BD Biosciences) and biotinylated isoelectric B4 (dilution 1/25; Sigma) were diluted in the antibody mix. This antibody mixture was applied and samples were then incubated for 2 hours at room temperature. After washing with TBS, the secondary antibody (DAKO Envision Dual-Link System HRP; DAKO) was added and samples were further incubated for 30 minutes at room temperature. DAKO AEC-high sensitivity substrate chromogen was applied for 15 minutes followed by intense washing with H2O, slides were counterstained with hematoxylin (Sigma) for 15 seconds and washed under running H2O. Immunohistomount (Santa Cruz Biotechnology Inc.) was used for mounting.

Competitive repopulation assay
The competitive repopulation assay (CRA) was based on the CD45.1-CD45.2 chimeraism of C57BL/6N mice, as described previously (24). CD45.2+/Sca1+/lin- were retrovirally transduced and cultivated in DMEM supplemented with 10% FCS (Gibco), mIL6, mSCF, and mIL3 upon exposure of either 0.001% DMSO, 0.3 μmol/L, and 3 μmol/L CJ-13,610 for 7 days. A total of 1 × 10⁵ CD45.2-/+Sca1+/lin+ cells were injected into lethally (11 Gy) irradiated CD45.1- and CD45.2- female recipients (B6DJLF1/JRj; Janvier) together with 5 × 10⁵ normal Ly5.1+ bone marrow (BM) cells. The proportion of CD45.2+ donor cell-derived hematopoietic cells was determined by FACS analysis of cells from the peripheral blood or spleen. The cells were stained with conjugated monoclonal antibodies specific for CD45.1 and CD45.2 or unstained control (Miltenyi Biotec).

Transactivation assays
pCDNA3 or pc3-5LO expression plasmids were either cotransfected with pRT-LK, pGL3-OT, or pGL3-OF or pGL3-γ-catenin, pGL3-basic, and pRT-CMV into U937-PR9 cells by nucleofection according to the manufacturer’s instructions (Nucleofector Kit C, Lonza). Two hours later, transgene expression was induced by the exposure of the cells to 100 μmol/L ZnSO4 (Sigma) as described (25). Twenty-four hours after Zn²⁺ treatment, luciferase activity was determined using the Dual-Luciferase Reporter Assay System (Promega) accordingly and all assays were normalized to cotransfected Renilla activity.

Immunofluorescence
Cells were applied on positively charged coverslips (Menzel-Gläser), washed with TBS (10 mmol/L Tris-HCl pH 8, 150 mmol/L NaCl) and fixed in 4% paraformaldehyde (Applichem) for 30 minutes followed by blocking and permeabilization with 5% (w/v) nonfat dry milk (Roth) and 0.2% Triton X-100 (Roth) for 30 to 45 minutes. Cells were incubated with monoclonal anti-5-LO (BD Biosciences), polyclonal anti-β-catenin (clone H102; Santa Cruz Biotechnology Inc.), and polyclonal antIRARα (Santa Cruz Biotechnology Inc.) antibodies, respectively. After extensive washing in TBS, cells were stained with Alexa Fluor 488-conjugated goat anti-mouse or Alexa Fluor 594-conjugated goat anti-rabbit Ig antibodies (Invitrogen). Nucleus staining was obtained using TO-PRO 3 (Invitrogen). The coverslips were mounted with Moviol (Sigma). Images were acquired by a Leica TCS-SP5 confocal microscope (Leica) under identical conditions for pinhole opening, laser power, photomultiplier tension, and layer number. During data elaboration by Fiji software (www.fiji.sc), identical parameters were applied for all samples.

Statistical analysis
Statistical significance was determined using one-way ANOVA with Bonferroni posttest using GraphPad Prism 5.0 (GraphPad).

Results
5-LO is inhibited by NSAIDs and is expressed in normal and malignant hematopoietic stem cell compartments
Some NSAIDs, such as sulindac, indomethacin, or celecoxib, are able to inhibit Wnt signaling, but only at high concentrations, exceeding those for the suppression of COX-1/2 (Table 1; refs. 6, 7, 13–16, 26). It has recently been shown that sulindac sulfide (active metabolite of sulindac) is able to inhibit 5-LO at clinically relevant concentrations (17). Therefore, we investigated whether other NSAIDs such as indomethacin are able to inhibit 5-LO. Monocytic THP-1 cells, known to express

<table>
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<th>NSAIDS</th>
<th>COX-1 IC₅₀ values (μmol/L)</th>
<th>COX-2 IC₅₀ values (μmol/L)</th>
<th>Concentrations used for Wnt signaling inhibition (μmol/L)</th>
<th>Literature</th>
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<tr>
<td>Sulindac sulfide³</td>
<td>1.02</td>
<td>10.43</td>
<td>50–200</td>
<td>(13, 15, 16, 26)</td>
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<tr>
<td>Indomethacin</td>
<td>0.16</td>
<td>0.46</td>
<td>40–600</td>
<td>(5, 6, 14, 15, 26, 28)</td>
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<tr>
<td>Diclofenac</td>
<td>0.14</td>
<td>0.05</td>
<td>&gt;100</td>
<td>(15, 26)</td>
</tr>
<tr>
<td>Ibuprofen</td>
<td>4.75</td>
<td>&gt;30</td>
<td>1,000</td>
<td>(26, 28)</td>
</tr>
<tr>
<td>Celecoxib</td>
<td>21.9</td>
<td>0.2</td>
<td>60–100</td>
<td>(27, 29)</td>
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³In human whole-blood assay (endotoxin-induced PGE₂ synthesis).
⁴In different cell systems (see literature).
⁵Active metabolite of sulindac.
5-LO and to have inducible 5-LO activity (30), were preincubated with inhibitors. 5-LO product formation was induced by exposure to Ca^{2+}-ionophore A23187 (21). CJ-13,610, a selective 5-LO inhibitor, was used as a positive control. We found that indomethacin was able to suppress 5-LO product formation in a concentration-dependent manner (Fig. 1A). Indomethacin suppressed 5-LO activity at higher concentration as needed for COX inhibition (Fig. 1A). Noteworthy for both inhibitors, indomethacin and sulindac sulfide, the 5-LO–inhibitory concentrations were similar to those required for suppression of Wnt signaling and the aberrant self-renewal of LSCs (16, 17). Therefore, we investigated the subpopulation of cells that should be targeted in a “stem cell therapy” approach. We studied 5-LO product formation in a highly enriched Sca1^{+}/lin^- population to disclose whether 5-LO is expressed in HSPCs. Differentiation was induced by GM/G-CSF. 5-LO activity was assessed by the formation of LTBl, by ELISA. Here, we show for the first time that the HSPC compartment not only expressed 5-LO, but also exhibited 5-LO enzyme activity, which was only slightly induced by the GM-CSF–induced differentiation process (Fig. 1B). These findings were confirmed by Western blot analysis of Sca1^{+}/lin^- HSPCs (data not shown).

**Targeting 5-LO by selective inhibitors reduces the stem cell capacity of PML/RARalpha-positive HSPCs**

Because NSAID concentrations that are able to inhibit aberrant LSC capacity lie within the range needed to suppress 5-LO, we speculated whether the effects of the NSAIDs on LSCs are mediated by inhibition of 5-LO. Therefore, we studied the effects of CJ-13,610, a potent and selective inhibitor of 5-LO, on LSC capacity.

The effects of 5-LO inhibition on aberrant LSC capacity were addressed in serial replating efficiency assays in PML/RARalpha-positive Sca1^{+}/lin^- HSPCs. Serial replating revealed the combined effect of PML/RARalpha on proliferation, differentiation, and self-renewal of HSPCs (12, 31). Therefore, we retrovirally expressed PML/RARalpha in Sca1^{+}/lin^- murine HSPCs and plated them in semisolid medium supplemented with mIL3, mIL6, and mSCF in the presence/absence of 0.3 and 3 μmol/L of CJ-13,610 (Fig. 2A). Empty vector–transduced cells were used as controls. Serial replatings were performed as graphically described in Fig. 2A.

CJ-13,610 abolished the aberrant serial replating capacity of PML/RARalpha-positive HSPCs at a concentration of 0.3 μmol/L, starting from the second plating (Fig. 2B). An effect of the treatment was already seen at the first plating, as revealed by the modifications in the morphology of the colonies in the presence of 3 μmol/L CJ-13,610 (Fig. 2C). The rare colonies in the controls also disappeared upon treatment (Fig. 2A).

In addition to the 5-LO and COX pathway, there is also a third arachidonic acid pathway, CYP (cytochrome P450)/sEH (soluble epoxide hydrolase) pathway, known to be involved in Wnt signaling, proliferation, and mobilization of HSC (32). To investigate the role of CYP/sEH in the effects of stem cell suppression and to confirm that only 5-LO inhibition is responsible for the observed effects, we extended the treatment of PML/RARalpha-positive Sca1^{+}/lin^- cells, to sulindac sulfide, in COX-inhibitory concentrations, as well as to tAUCB, an sEH inhibitor (Supplementary Fig. S1; ref. 33). In fact, treatment with neither sulindac sulfide or tAUCB alone nor in combination was able to suppress the replating efficiency of PML/RARalpha. Only the presence of CJ-13,610 led to a complete inhibition of the aberrant replating efficiency of PML/RARalpha (Supplementary Fig. S1). Also, the genetic inhibition of sEH, using murine Sca1^{+}/lin^- cells with a sEH^/-^ background,
showed no effect on the replating efficiency of PML/RARα-expressing HSPCs (Supplementary Fig. S2).

In summary, these data indicate that the pharmacologic inhibition of 5-LO is sufficient to interfere with the aberrant replating capacity of PML/RARα-positive HSPCs.

The abolition of the stem cell capacity of PML/RARα-positive HSPCs by CJ-13,610 is accompanied by the inhibition of Wnt signaling

CJ-13,610 not only inhibited serial replating capacity of PML/RARα-positive but also inhibited colony formation of empty vector–transduced HSPCs. Thus, we investigated whether the inhibition of 5-LO generally suppresses stem cells or only LSCs. First, we studied the effects of CJ-13,610 on short-term hematopoietic stem cells (ST-HSC) and early immature progenitors in a colony-forming unit spleen day 12 (CFU-S12) assay (34). Therefore, Sca1+/lin− HSPCs were retrovirally transduced with PML/RARα or with empty vector and exposed in liquid culture to 0.3 or 3 μmol/L CJ-13,610. After 7 days, cells were counted, harvested, and inoculated into lethally irradiated mice (Fig. 3A). On day 12, the mice were sacrificed, spleens were fixed, and the spleen colonies were counted. As reported in Fig. 3B, the exposure to CJ-13,610 led to a concentration-dependent and significant reduction of the number of spleen colonies expressing PML/RARα. The exposure to 3 μmol/L CJ-13,610 almost completely abolished these colonies (Fig. 3B and C). Interestingly, 3 μmol/L CJ-13,610 allowed the formation of a few colonies in the controls, indicating that CJ-13,610 did not exert stem cell toxicity (Fig. 3B).

To determine which stem cell compartment is targeted by CJ-13,610, we studied its effects on the frequency of short-term (ST) and long-term (LT) HSC in a CRA. In this assay, the ST-HSC population was detected at 12 weeks (Fig. 3F, left) and the LT-HSC population after 6 months (Fig. 3F, right). Therefore, Sca1+/lin− HSPCs were retrovirally transduced with PML/RARα or with empty vector and exposed in liquid culture to 0.3 or 3 μmol/L CJ-13,610. After 7 days, cells were counted, harvested, and inoculated into lethally irradiated mice (Fig. 3A). On day 12, the mice were sacrificed, spleens were fixed, and the spleen colonies were counted. As reported in Fig. 3B, the exposure to CJ-13,610 led to a concentration-dependent and significant reduction of the number of spleen colonies expressing PML/RARα. The exposure to 3 μmol/L CJ-13,610 almost completely abolished these colonies (Fig. 3B and C). Interestingly, 3 μmol/L CJ-13,610 allowed the formation of a few colonies in the controls, indicating that CJ-13,610 did not exert stem cell toxicity (Fig. 3B).

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Interestingly, the effects of CJ-13,610 were not related to the induction of apoptosis, excluding a cytotoxic effect of the 5-LO inhibitor (data not shown).

The effects of NSAIDs on cancer stem cells are mainly attributed to an inhibitory activity on β-catenin and the Wnt signaling. Therefore, we investigated whether suppression of the PML/RARα-induced aberrant stem cell capacity by CJ-13,610 was related to an inhibitory effect on β-catenin and Wnt signaling. Thus, we performed immunohistochemical staining of β-catenin in CFU-S12 spleen slices. Empty vector control and PML/RARα-expressing spleens of a CFU-S12 experiment in which the Sca1+/lin HSPCs were retrovirally transduced and treated with 3 μmol/L CJ-13,610. Slides were stained with either an anti-β-catenin and/or only secondary antibody (unstained), E, experimental strategy for studying the influence of CJ-13,610 on ST- and LT-stem cell capacity of PML/RARα-expressing murine HSCs (CRA). F, CRA. Sca1+/lin BM cells from CD45.2 mice were retrovirally infected with empty vector control or PML/RARα. Cells were treated with 3 μmol/L CJ-13,610. After 7 days in liquid culture, cells were cotransplanted with CD45.1 BM cells into lethally (11 Gy) irradiated CD45.1+2 recipient mice (4–5 mice/group). An analysis of donor chimerism was performed at 12 weeks and 6 months after transplantation. The plots show a representative donor-derived chimerism in (left) peripheral blood after 12 weeks and (right) spleen after 6 months from an individual mouse.
Loss of 5-LO expression does not recapitulate the inhibitory effects of the inhibitors on the leukemic phenotype by PML/RARα

To further investigate the role of 5-LO in the maintenance of normal and leukemic stem cells, we investigated the effect of targeted genetic inhibition of 5-LO on leukemogenic potential and the aberrant stem cell capacity induced by PML/RARα in Alox5+/− mice (35). PML/RARα was retrovirally expressed in Sca1+/lin−, Alox5+/− and Alox5+/+ HSPCs, respectively. First, we performed serial replating assays. As shown in Fig. 4A, the loss of 5-LO expression not only failed to inhibit the replating efficiency of PML/RARα-expressing HSPCs, but also led to increased proliferation of colony-forming cells, as revealed by an increased number of colonies in comparison with Alox5+/+ controls expressing PML/RARα (Fig. 4 A and B).

Next, we tested the LT-HSC capacity of Alox5+/− and Alox5+/+ HSPCs expressing PML/RARα in a CRA. Alox5+/− and Alox5+/+ HSPCs did not show significant differences in the frequency of the PML/RARα-positive LT stem cell population (CD45.1+), as revealed 9 months after transplantation into lethally irradiated recipient mice (CD45.1 and 2 population; Fig. 4C).

Taken together, these data show that loss of 5-LO expression does not interfere with the replating efficiency and LT stem cell capacity of PML/RARα-positive HSPCs. The data strongly suggest that stem cell inhibition may be mediated by the inhibited/inactive form of 5-LO, indicating a pathophysiological relevant difference between enzymatically active and inactive 5-LO.

Coexpression of catalytically inactive 5-LO inhibits the PML/RARα-induced activation of Wnt signaling in human leukemic U937 cells

Assuming that the catalytically inactive form of 5-LO is responsible for stem cell inactivation, it should, like the NSAIDs, also inhibit Wnt signaling. Therefore, we investigated activation of Wnt signaling in a cellular model in the absence and presence of 5-LO. We took advantage of the fact that PR9 cells, promonocytic U937 cells expressing PML/RARα under the control of the Zn2+-inducible metallothionein1 (MT-1) promoter do not express endogenous 5-LO due to the methylation of the 5-LO promoter (36). PML/RARα activates Wnt signaling in these cells through the activation of β-catenin and γ-catenin (10, 12). The lack of 5-LO expression in both control and P/R9 cells was confirmed by Western blotting and by PCR (data not shown). Noteworthy, transfected 5-LO is unable to synthesize leukotrienes in undifferentiated U937 cells (37). This may relate to cell type–specific features governing 5-LO activity, such as the redox state, the phosphorylation status of 5-LO, the activity of intracellular tyrosine kinases, structural

Figure 4. Effect of 5-LO+/− (Alox5+/−) on replating efficiency and long-term stem cell capacity of PML/RARα-expressing HSCs. A, Sca1+/lin− BM cells (wild-type and Alox5−/−) were retrovirally infected with empty vector control and PML/RARα and plated in methylcellulose medium supplemented with mIL3, mIL6, and mSCF to assess primary colony formation. The colony numbers were counted on day 10. After determining the colony number, the cells were harvested and serially replated every 10 days. B, colony morphology of platings I–VI. Data are shown from a single representative experiment of three that yielded similar results. C, CRA. Sca1+/lin− BM cells from wild-type and Alox5+/− CD45.2+ mice were retrovirally infected with empty vector control and PML/RARα. After 7 days in liquid culture, cells were cotransplanted with CD45.1+ BM cells into lethally (11Gy) irradiated CD45.1 and 2+ recipient mice (7–8 mice/group). An analysis of donor chimerism was performed at 9 months after transplantation to determine the LT-stem cell capacity. The graph shows a donor-derived chimerism at 9 months in peripheral blood. Statistical significance was tested using the Student t test (**+, P < 0.001). ns, nonsignificant.
characteristics in the cytoskeleton, and the individual expression of further enzymes involved in leukotriene biosynthesis (38). Notably, in myeloid cells, the activity of 5-LO was already reported by us to depend on the differentiation status and the activity of cellular peroxidases, which may also be true in 5-LO–transfected U937 cells (22). We expressed 5-LO in these cells under the control of a CMV promoter. Activation of Wnt signaling by PML/RARα was addressed by the transactivation of the “Topflash/Fopflash” reporter system, in which the activation of the reporter is under the control of a TCF/LEF responsive element (19). Here we show that the expression of 5-LO prevented the activation of the Wnt target promoter by PML/RARα (Fig. 5A).

To further understand the mechanism by which 5-LO suppresses Wnt signaling, we investigated the effect of 5-LO on γ-catenin–dependent transactivation. β- and γ-catenin are closely related and are both key mediators of Wnt signaling. In contrast to β-catenin, which exhibits transforming potential only upon constitutive stabilization by mutations, the over-expression of γ-catenin alone is able to transform cells (39). We used a construct in which the luciferase reporter was controlled by the γ-catenin promoter, which is activated by γ-catenin itself (12). As reported in Fig. 5B, only a slight reduction of the transactivation of γ-catenin promoter by PML/RARα was seen in the presence of 5-LO (Fig. 5B). In summary, these results suggest that 5-LO interferes with the PML/RARα-induced activation of Wnt signaling by inhibiting β-catenin- and not γ-catenin–dependent transactivation. This ability is mediated by inactive 5-LO.

5-LO and β-catenin show a direct interaction

To test the possibility of a direct influence of 5-LO on Wnt signaling, we investigated the interaction between 5-LO and β-catenin. Therefore, we coexpressed 5-LO in 293T cells together with a HA-tagged constitutive active β-catenin mutant, S33A, which lacks the phosphorylation sites for proteasomal degradation (Supplementary Fig. S5, right). Coimmunoprecipitation using anti–5-LO or high affinity anti-HA antibodies coupled to magnetic beads, followed by Western blotting, revealed a direct interaction between 5-LO and active β-catenin (Supplementary Fig. S5, left, IP 5-LO and IP HA). The slight bands of 5-LO in β-catenin in the respective anti–β-catenin or anti–5-LO precipitates were most likely due to specific binding of endogenous proteins. 293T cells expressed both endogenous β-catenin and 5-LO (data not shown). Nevertheless, upon overexpression of both proteins the amount of precipitated 5-LO or β-catenin, respectively, was clearly higher than that seen in the controls showing a direct interaction between the two proteins.

In summary, our data show a direct interaction between 5-LO and β-catenin, the key regulator of the Wnt signaling pathway, which may account for the inhibitory effect of inactive 5-LO on Wnt signaling.

5-LO colocalizes with β-catenin and hinders β-catenin from entering the nucleus

To confirm the biological relevance of the interaction between 5-LO and β-catenin revealed in vitro, we sought to disclose the mechanism by which this interaction interferes with Wnt signaling. Therefore, we studied the influence of the expression of 5-LO on the localization of β-catenin in PR9 cells by indirect immunofluorescence. Thus, we expressed 5-LO in PR9 cells and induced them to express PML/RARα by Zn2+ treatment. To investigate the colocalization of 5-LO and β-catenin, we performed double anti–5-LO/anti–β-catenin staining. As controls, we used empty vector–transfected P/R9 cells.

**Figure 5.** Effect of 5-LO on PML/RARα–mediated Wnt pathway activation. Transactivation assay for Wnt signaling–related transactivation (A) β-catenin (TCF/LEF) or γ-catenin–dependent transactivation (B). The expression vectors pCDNA3.1 or pCDNA3.1-5-LO were cotransfected by nucleofection with either Topflash (OT) or Fopflash (OF) reporter constructs (A) or pGL3-γ-catenin or pGL3 basic constructs (B) into 5-LO negative U937 cells expressing PML/RARα under the control of a Zn2+–inducible metallothionein1 (MT-1) promoter. The transgene was induced by exposure to 100 μmol/L ZnSO4. Luciferase activity was measured 24 hours later and normalized with Renilla activity. The data are expressed as the means of two (A) and three (B) independent experiments with SD. Statistical analysis was performed by the Student t test (*, P < 0.05).
The expression of PML/RARα upon Zn²⁺ induction was confirmed by anti-RARα staining, which revealed the typical microspeckled pattern of PML/RARα (Supplementary Fig. S6; refs. 40, 41). The β-catenin expression in U937 cells showed a more scattered pattern than in other cell lines, most often fibroblasts, where the distribution of β-catenin is more diffuse (42–44). As shown in Fig. 6A, upon Zn²⁺ induction, the proportion of nuclear β-catenin (red fluorochrome) increased, confirming activation of Wnt signaling upon PML/RARα induction. In contrast, in the absence of PML/RARα, anti-β-catenin staining was seen exclusively in the cytoplasm. The expression of 5-LO in the PR9 cells led to perinuclear anti–5-LO staining (green fluorochrome), whereas empty vector–transfected cells did not reveal any specific anti–5-LO staining, further confirming that these cells do not express 5-LO. The anti–5-LO staining was not influenced by the presence of PML/RARα. In contrast, in the presence of 5-LO, no nuclear anti–β-catenin staining was seen. The superimposition of anti 5-LO (green fluorochrome) and anti–β-catenin (red fluorochrome) staining showed partial colocalization of 5-LO and β-catenin (yellow) (Fig. 6A). Four confocal sections (Fig. 6B, a–d) of the same cell taken on the horizontal axis of the microscope (0.25–0.33 μm interval) are shown in Fig. 6B. To view the colocalization in greater detail, a further electronic 4-fold magnification of the indicated area of each layer is shown (Fig. 6B, a’–d’). A three-dimensional reconstruction of all the layers of the cell (data not shown) illustrated the same colocalization seen in Fig. 6B. The apparently small amount of colocalizing 5-LO and β-catenin accords with the data obtained by coimmunoprecipitation.

To investigate whether CJ-13,610 is capable of inducing colocalization of 5-LO and β-catenin, human NB4 acute promyelocytic leukemia cells exhibiting well-defined cytosolic 5-LO expression were exposed to 3 μmol/L CJ-13,610. As can be seen treatment with CJ-13,610 led to partial colocalization of 5-LO and β-catenin (Supplementary Fig. S7) strengthening the hypothesis of a 5-LO–mediated suppression of Wnt signaling induced by CJ-13,610. Taken together, immunofluorescence studies could confirm the direct interaction of 5-LO with β-catenin, triggered by CJ-13,610, suggesting also a novel mechanism for Wnt signaling inhibition by which β-catenin is prevented by 5-LO from entering the nucleus.

**Discussion**

The aim of our study was to determine whether the inhibition of 5-LO may account for the effects of NSAIDs on the Wnt signaling pathway and to assess their capacity to suppress CSCs in leukemia. We could show that the pharmacologic inhibition of 5-LO at clinically feasible concentrations of selective inhibitors, CJ-13,610 or zileuton (45–47) suppresses the aberrant stem cell capacity of PML/RARα-positive HSPCs via inhibition of Wnt signaling. In addition, inhibitors of the other arachidonic acid metabolizing enzymes, COX and sEH, failed to suppress PML/RARα-positive HSPCs. Finally, a number of subsequent experiments substantiated the CJ-13,610–dependent suppressive action of ectopically expressed 5-LO.

![Figure 6. Colocalization between 5-LO and β-catenin in PR9 cells. Cells were transfected by nucleofection with either empty vector control or 5-LO. Two hours after nucleofection, transgene expression was induced by 100 μmol/L ZnSO₄. A, twenty-four hours later, cells were imaged in phase contrast (gray, a), stained with TO-PRO-3 (blue, b), and stained with anti–5-LO and anti–β-catenin antibody, respectively. Alexa Fluor 594–conjugated anti-rabbit-Ig (red, c) and Alexa Fluor 488, anti-mouse-Ig (green, d) were used for immunostaining. Merged images (e) were obtained by electronic overlapping of images recorded in a–d. B, cells, treated and stained as in A, were scanned in 30 layers (0.25–0.33 μm interval) on z-axis. Electronic overlapped merged images of phase contrast, TO-PRO-3 (nuclei), red fluorescence (β-catenin), and green fluorescence (5-LO) are shown in B. Left, four confocal sections (a–d) are shown. Colocalizations are indicated with arrows. A ×4 magnification (a’–d’) is reported on the right side of B.](cancerres.aacrjournals.org/content/74/18/5252/F6.large.jpg)
on Wnt signaling and leukemic cells. Thus, suppression of aberrant stem cell capacity should mainly result from the drug’s ability to specifically target 5-LO. However, pleiotropic off-target effects of CJ-13,610 cannot ultimately be excluded. Nevertheless, even the existence of such possible off-target effects, contributing to the antileukemic effect of the drug, would not detract from the documented key finding of a pivotal role of 5-LO in AML.

On the basis of our findings that the 5-LO directly interacts with β-catenin, we conclude that the Wnt inhibitory effect of NSAIDs in our leukemia model is mediated not by inhibition of the COX-1/2 and prostaglandin E2 synthesis as observed in models of colorectal cancer (48). This is supported by the finding that the activation of Wnt signaling in LSC is not related to upregulation of COX, but is related to the presence of aberrant chimeric transcription factors resulting from chromosomal translocations such as t(15;17)-PML/RARα or t(8;21) AML-1/ETO fusion proteins (10, 12). The inhibitory effect on tumor stem cells of the inhibition of 5-LO is not limited to our leukemia model but has also been described in glioblastoma. Here, the selective 5-LO inhibitor, Nordy, was able to reduce sphere formation and the frequency of stem cells in xenograft tumors (49).

Both the genetic disruption and pharmacologic inhibition of 5-LO suppress the tumor-initiating cells of BCR/ABL-driven CML-like disease in mice (18). This is, at least partially, in contrast with our findings on the PML/RARα-driven AML model, in which the genetic targeting of 5-LO did not replicate the effects of pharmacologic inhibition on the stem cell capacity. This is in accordance with recent findings showing full leukemogenic potential of AML-1/ETO9x—expressing HSPCs in an Alox5−/− background although an impairment of the in vitro self-renewal of HSPCs expressing either AML-1/ETO9x or MLL/AF9 was seen (50). CML is a myeloproliferative disease characterized by proliferation of hematopoietic progenitors, which are still able to fully differentiate. PML/RARα induces an acute leukemia characterized by aberrant self-renewal of the HSCs and differentiation block. The stem cell compartment of these two diseases differ being more mature for the CML-like disease and very primitive HSPCs for PML/RARα (31, 51). Furthermore, the induction and the maintenance of AML-LSC depend on the activation of Wnt signaling, whereas for the induction of CML-like disease, Wnt signaling seems to be nonessential (5). Therefore, we hypothesize that the presence of the enzymatically inactive form of 5-LO is needed to inhibit Wnt signaling in the PML/RARα-positive HSPCs by blocking the transition of β-catenin to the nucleus. That 5-LO is able to alter nuclear trafficking of a protein, was shown for p53, as 5-LO antagonized genotoxic stress-induced apoptosis by inhibiting binding of p53 to the promyelocytic leukemia protein (PML) and its relocalization into PML-nuclear bodies (52).

The exact molecular mechanism by which inhibitor-bound and catalytically inactive 5-LO interacts with β-catenin and possibly other Wnt components remains unclear. Possibly, 5-LO inhibitors, such as CJ-13,610 and zileuton, induce certain conformational changes in the 5-LO enzyme, thereby allowing suppressive interactions of the enzyme with other proteins of the Wnt signaling pathway. Similarly, AA and competitive 5-LO inhibitors mimicking AA were observed to trigger changes in subcellular localization of 5-LO that may relate to the effects observed in this study with CJ-13,610 (53). As a consequence, cytosolic and perinuclear β-catenin bound to 5-LO may be trapped and hindered from entering the nucleus, suggesting that the protein cannot drive anymore the proleukemic transcription of Wnt target genes, as demonstrated in our study. Notably, in PR9 cells, ectopically transfected 5-LO is capable of suppressing Wnt signaling, whereas in other cell types, including the PML/RARα-positive HSPCs, the presence of CJ-13,610 is required to convert 5-LO to an active Wnt suppressor. We conclude that the environment of PR9 cells into which the ectopically transfected 5-LO is imbedded renders 5-LO catalytically inactive by still unknown mechanisms thereby allowing inhibition of Wnt signaling. Further biophysical studies on protein structure are required addressing in detail the possible conformational changes in 5-LO after binding to 5-LO inhibitors allowing interaction with β-catenin.

Our study demonstrated a crucial role for 5-LO in the maintenance of LSCs in AML, confirming recent data in murine models of AML (50). The fact that, in a subset of tumors, the inhibition of 5-LO is able to suppress the tumor stem cell renders feasible a stem cell therapy as a maintenance regimen after induction and consolidation therapy protocols. The use of 5-LO inhibitors, instead of COX inhibitors, would further avoid the adverse effects of COX inhibitors, such as gastrointestinal bleeding and ulcerations or cardiovascular side effects, including hypertension, myocardial infarction, and stroke. Taken together, our data establish pharmacologic inhibition of 5-LO as a novel approach of stem cell therapy in leukemia, by targeting Wnt signaling.

Disclosure of Potential Conflicts of Interest
No potential conflicts of interest were disclosed.

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Acknowledgments
The authors thank Prof. Dr. Mike Parnham for critically proofreading the article and Carlo Angioni for excellent technical assistance.

Grant Support
This work was funded by the German José Carreras Leukemia Foundation (DJKLS RF-11/15; M. Ruthardt and I. Fleming, by LOEWE (Landesinitiative zu DJCLS RF-11/15; M. Ruthardt and I. Fleming, by LOEWE (Landesinitiative zu...
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Received October 30, 2013; revised June 9, 2014; accepted June 21, 2014; published OnlineFirst July 31, 2014.

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