Notch2 is involved in the overexpression of CD23 in B-cell chronic lymphocytic leukemia

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Members of the Notch family encode transmembrane receptors that modulate differentiation, proliferation, and apoptotic programs of many precursor cells, including hematopoietic progenitors. Stimulation of Notch causes cleavage followed by translocation of the intracellular domain (NotchIC) to the nucleus, where it activates transcription of CBF1 responsive genes. The aim of this study was to elucidate the mechanisms leading to the overexpression of CD23, a striking feature of B-cell chronic lymphocytic leukemia (B-CLL) cells. By electrophoretic mobility shift assays, we identified a transcription factor complex (C1) that binds sequence specific to one known and 4 newly identified putative CBF1 recognition sites in the CD23a core promoter region. With the use of Epstein-Barr virus (EBV)-infected B cells as a model for CBF1 mediated CD23a expression, C1 was found to be EBV inducible. Super-shift assays revealed that the nuclear form of Notch2 is a component of C1 in B-CLL cells, supporting a model in which NotchIC activates transcription by binding to CBF1 tethered to DNA. Transient transfection of REH pre-B cells with an activated form of Notch2 induced endogenous CD23a, confirming that CD23a is a target gene of Notch2 signaling. Finally, reverse transcription-polymerase chain reaction and kinetic analysis demonstrated that the Notch oncogene is not only overexpressed in B-CLL cells but might also be related to the failure of apoptosis characteristic for this disease. In conclusion, these data suggest that deregulation of Notch2 signaling is involved in the aberrant expression of CD23 in B-CLL. (Blood. 2002;99:3742-3747)

Introduction

B-cell chronic lymphocytic leukemia (B-CLL) is characterized by the relentless accumulation of monoclonal anergic B cells that coexpress CD5 and CD19 with faint to virtually undetectable amounts of surface immunoglobulins. The pathomechanisms underlying the disease primarily involve defects that prevent cell turnover because of apoptosis, rather than alterations in cell cycle regulation.

One of the hallmarks of B-CLL cells is the overexpression of the transmembrane glycoprotein CD23, which undergoes spontaneous proteolysis, giving rise to soluble CD23 (sCD23) molecules. The serum concentration of sCD23 can be several hundred-fold higher than in healthy individuals and parallels the clinical stage of the disease.

Two isoforms of CD23 exist, CD23a and CD23b, which are expressed from 2 different promoters. Expression of CD23a is restricted to B lymphocytes, whereas CD23b is found on a number of different hematopoietic cell types, predominantly after interleukin 4 (IL-4) treatment. In B-CLL cells, selective expression of CD23a has been found to be concurrent to a state of cell survival, thereby providing a link between CD23a and the regulation of apoptosis.

CD23 is also closely associated with Epstein-Barr virus (EBV)-mediated B-cell immortalization, because a naturally occurring EBV mutant (P3HR1), carrying a deletion of the EBNA2 gene, has lost its ability to induce CD23 expression and to transform primary B cells in vitro. EBNA2 activates the CD23a gene through a CBF1 repressor site located in the CD23a proximal promoter. Several lines of evidence strongly suggest that EBNA2 mimics Notch signaling by acting as a transcriptional activator after binding and masking the repression domain of CBF1.

The Notch gene family encodes transmembrane receptors that modulate differentiation, proliferation, and apoptotic programs in response to extracellular ligands expressed on neighboring cells. Ligand-mediated stimulation of Notch causes the proteolytic release of the intracellular domain (NotchIC), which then passes into the nucleus where it activates transcription of CBF1 responsive genes. Deregulation of this pathway by overexpression of a constitutively activated form of Notch not only diverts cell fate decisions but is also tumorigenic. For example, truncation of Notch1 found in a subset of human T-cell leukemias leads to the expression of a ligand-independent oncogenic protein lacking the extracellular domain. The truncated Notch proteins consist primarily of the intracellular domain and localize predominantly in the nucleus. Enforced expression of Notch1IC in bone marrow stem cells causes T-cell leukemia in mice, indicating a causative role for Notch1 in T-cell oncogenesis.

To elucidate the mechanisms leading to the up-regulation of CD23a in B-CLL cells, we analyzed the CD23a proximal promoter for sequence-specific DNA-protein interactions. By electrophoretic mobility shift assays (EMSAs), we detected a transcription factor
complex with low CD23 expression (referred to as B-CLL CD23 
KCl; 1.5 mM MgCl 2 ; 20 mM HEPES, pH 7.9; 0.2 mM EDTA; 25%
were collected after 1, 2, 3, and 5 days and processed for
signaling is involved in the pathogenesis of this disease.

Materials and methods

Cell lines and culture

Peripheral blood mononuclear cells (PBMCs) from 6 patients with typical
CD23a

Sequence analysis of the CD23a promoter

Potential transcription factor binding sites in the
that is based on the transcription factor database TRANSFAC 4.0

Preparation of nuclear extracts and EMSA

Nuclear extracts were prepared as described with minor modifications.30
briefly, 15 × 106 cells were lysed in 1 mL hypotonic buffer (10 mM
HEPES, pH 7.9; 1.5 mM MgCl2; 10 mM KCl) containing 0.15% NP-40 at
4°C for 10 minutes. The nuclear proteins were extracted from the nuclear
fraction by resuspending the nuclei in 600 μL extraction buffer (300 mM
KCl; 1.5 mM MgCl2; 20 mM HEPES, pH 7.9; 0.2 mM EDTA; 25%
glycerin) at 4°C for 20 minutes with constant agitation. After removing the
pellet by centrifugation, the supernatant was dialyzed against dialysis buffer
(20 mM HEPES, pH 7.9; 100 mM KCl, 0.2 mM EDTA, 20% glycerol) for
20 minutes with constant agitation. After removing the

Transient transfection experiments

REH pre-B cells were transiently transfected by electroporation. Cells
(1-2 × 106) were resuspended in 500 μL serum-free medium and were
mixed with 10 μg plasmid DNA (pSg5-nR2IC and pCMV-green fluores-
icient protein [GFP], respectively) in 0.4-cm electroporation cuvettes.
Electroporation was carried out after 5 minutes of incubation on ice on a
BioRad gene pulser set at 950 μF and 280 V. Immediately after pulsing,
cells were transferred to 4 mL culture medium and were incubated for 48
hours. Transfection efficiency was determined by quantification of GFP-

Determination of sCD23 concentration

Blood was obtained by venipuncture. After clotting, samples were centri-
fuged, and serum was stored at −20°C. The levels of sCD23 from sera and
culture supernatants were measured by using the Cellfree CD23 enzyme-
linked immunosorbent assay kit (Endogen, Woburn, MA).

RT-PCR analysis

Total RNA was extracted by using the RNeasy isolation system (CINNA/
MRC) according to the manufacturer’s instructions. One-step RT-PCR
reactions (Titan; Roche, Mannheim, Germany) were performed on total
RNA (50 ng per reaction) by using primer sequences as follows: Notch2,
forward 5′-TCGGAATGCGAGGTGCAAGGCACTGGAGG-3′ and reverse 5′-
GCGAGGAAACACCAATCTTTCACCTTAT-3′ (25 cycles; annealing tem-

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RNA used for detection of the target genes was accordingly adjusted. PCR products were sequenced to verify specificity of amplified DNA.

Results

**CD23a promoter sequence analysis**

Figure 1 depicts the sequence upstream of the transcriptional initiation site of the CD23a gene that has been shown to be sufficient for an efficient expression of CD23a in B cells. This region corresponds also to a nuclelease hypersensitive site found in B cells but not in T cells. In most cases such hypersensitive sites reflect the binding of transcription factors to specific DNA sequences, resulting in nucleosome displacement. Instead of a canonical TATA box, 2 putative Pax5 (BSAP) binding sites between −29 bp and −90 bp could be predicted. The upper Pax5 site is situated close to a previously described NF-κB-like element (−89 to −98) that has been shown to cooperate with a STAT6 site (−117 to −126) in CD40/IL-4–mediated induction of the highly homologous murine CD23 promoter. In addition to one known CBF1 site (CBF1.1 in Figure 1), we were able to identify 4 putative CBF1 recognition sites matching the consensus (GGTG/AGAA) in at least 6 of 7 positions. Furthermore, by using the matrix search program MatInspector V2.2 (Research Group Bioinformatics, Braunschweig, Germany), several potential binding sites for Ikaros (IK1–2), GATA1-3, LMO2, and MZF1 could be predicted.

**B-CLL–specific DNA-protein interactions in the CD23a proximal promoter**

To identify transcription factors implicated in the up-regulation of the CD23a gene in B-CLL cells, EMSAs were performed with 4 oligonucleotide probes derived from the CD23a core promoter region (Figure 1). Nuclear extracts were prepared from freshly isolated B-CLL cells (n = 6, Table 1), from PBMCs (n = 3), and from tonsillar B cells (n = 1). The Burkitt lymphoma cell line BL41 infected with the EBV strain B95-8 served as a positive control for CBF1-mediated CD23a expression. Nuclear proteins isolated from Th cells (n = 3) were included as a negative control for B-cell– and EBV-specific transcription factors.

<table>
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<th>Table 1. Serum and supernatant levels and membrane expression of CD23</th>
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<tr>
<td>CD23a (sCD23)</td>
</tr>
<tr>
<td>BL41</td>
</tr>
<tr>
<td>BL41/B95-8</td>
</tr>
<tr>
<td>PBMC HD</td>
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<tr>
<td>CLL (n = 3)</td>
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<tr>
<td>CLL</td>
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The soluble CD23 (sCD23) concentrations were determined by enzyme-linked immunosorbent assay in serum obtained from 3 representative healthy donors (HDs); data represent mean ± SEM. One patient with B-CLL had a low level of sCD23 in the absence of detectable levels of sCD23. The percentage of CD19+ and CD23+ cells from total cell populations were determined by using flow cytometry.

As demonstrated in Figure 2, EMSA with probe A to D, encompassing one known (Figure 1, probe A) and 3 newly identified putative CBF1 sites (Figure 1, probes B to D), visualized a prominent, slow-migrating DNA-protein complex designated as complex 1 (C1, Figure 2A-D). The significance of this complex was underlined by the fact that in all B-cell samples the intensity of C1 correlated with their respective levels of CD23a transcription. Most notable, C1 was found to be EBV inducible (Figure 2A-D, lanes 4 and 5), indicating that this complex is involved in CBF1-mediated CD23a expression. To support this finding, EMSA was performed, including an unlabeled competitor oligonucleotide (CBF1.Cp) spanning a well-characterized CBF1 site derived from the EBV C promoter. C1 was completely abolished upon the addition of the competitor (Figure 3A), confirming that C1 binds sequence specific to the CBF1 recognition sites. An additional putative CBF1 site matching the consensus at 7 of 7 positions (CBF1 site 2; GTGAGAA, −137 to −143) was identified in inverted orientation closely juxtaposed to a putative CAAT-box (Figure 1). A synthetic 20-bp unlabeled oligonucleotide comprising CBF1 site 2 (CBF1.2; −130 to −149) inhibited binding of C1 to probe A, indicating that C1 is also capable of interacting with CBF1 site 2 (Figure 3B).

Because B-CLL cells do not express detectable amounts of the EBV-encoded CBF1 activator EBNA2, we conducted supershift assays, including antibodies raised against the nuclear forms of Notch1 and Notch2. 2 members of the Notch family known to be expressed in normal B cells. Whereas antibodies specific
for Notch1IC had no discernible effect on the formation or migration of C1 (data not shown), antibodies specific for Notch2IC led to a marked decrease in the intensity of C1 in Th cells and in B-CLL samples (Figure 3C). This result indicates that in B-CLL cells the nuclear form of Notch2 is a major part of the cellular activity that binds to the CBF1 recognition sites in the \( \text{CD}23 \text{a} \) core promoter region.

EMSA with probe C and D (Figure 2C,D), spanning 2 predicted Pax5 sites, revealed the appearance of an additional complex, designated C2. This complex appeared in all B–cell samples (Figure 2C, lanes 4–10, and Figure 2D, lanes 3–10) but was absent in the Th-cell control (Figure 2C, lane 3, and Figure 2D, lane 2). Incubation of nuclear extracts with Pax5 antibodies before the addition of probe C diminished the signal of C2 (Figure 3D), implying that this complex contains Pax5. A higher order complex was observed in all samples positive for both, C1 and C2 (Figure 3E), suggesting that these factors bind the shared DNA regions in a cooperative manner. The presence of 2 Pax5 binding sites in the promoter region, revealed the appearance of an additional complex, designated C2. This complex appeared in all B-cell samples (Figure 3C). This result indicates that in B-CLL cells the nuclear form of Notch2 is a major part of the cellular activity that binds to the CBF1 recognition sites in the \( \text{CD}23 \text{a} \) core promoter region.

**Notch2 is overexpressed in B-CLL cells**

To examine whether elevated levels of Notch2IC in nuclear extracts from B-CLL cells were caused by increased transcription of the corresponding Notch2 gene, RT-PCR analysis was performed. As demonstrated in Figure 4, Notch2 was found to be consistently overexpressed in B-CLL cells (6 of 6) as compared with resting peripheral B cells from healthy donors (Figure 4B, lane 7, representative for \( n = 3 \)) and to other B–non-Hodgkin lymphoma cell lines (Figure 4B, lanes 4 and 5). In total PBMCs from healthy donors, relative high levels of Notch2 transcription were observed in purified Th cells (Figure 4B, lane 3) and in peripheral blood B cells (Figure 4B, lane 7) that are in accordance with published data. No signals for Notch1 were detected under these conditions (data not shown). Remarkably, neither Notch1 (data not shown) nor Notch2 messages (Figure 4B, lane 6) were found in a rare case of B-CLL (B-CLL \( \text{CD}23 \text{a}^{+/-} \)) in which the leukemic cells expressed low amounts of surface \( \text{CD}23 \) (Table 1). Because no detectable amounts of \( \text{CD}23\text{a} \) messenger RNA (mRNA) were found in this sample (Figure 4B, lane 6), it is likely that the surface expression of \( \text{CD}23 \) on these cells resulted from \( \text{CD}23\text{b} \) that is differently regulated. Figure 4B shows also the induction of the \( \text{CD}23\text{a} \) gene through EBV (lanes 4 and 5).

**Notch2IC induces endogenous \( \text{CD}23\text{a} \) in B cells**

To directly demonstrate that \( \text{CD}23\text{a} \) is a target gene of Notch2 signaling in B lymphocytes, the \( \text{CD}23\text{a} \) pre–B–cell line REH was transiently transfected with a mammalian expression vector containing the complementary DNA coding for rat Notch2IC (kindly provided by Dr Diane Hayward). This vector leads to the expression of a ligand-independent constitutively-active Notch2IC protein lacking the extracellular domain. The transfection efficiency in the living cell population was 28% as determined by flow cytometry (not shown). After 2 days of incubation, RT-PCR analysis demonstrated that ectopic expression of Notch2IC induces \( \text{CD}23\text{a} \) transcription (Figure 5, lane 3), whereas a GFP-encoding control plasmid had no effect (Figure 5, lane 2).
Expression of CD23a correlates with B-CLL cell survival

CD23 is detectable on virtually all resting B cells that coexpress surface IgM and IgD. In vitro, freshly isolated normal B cells rapidly lose both, CD23 protein and mRNA, suggesting that the expression of CD23 on resting B cells is not constitutive. To compare the expression of CD23a in normal B cells with B-CLL cells, gene expression kinetic studies were performed. The results confirmed that after 24 hours of incubation purified normal B cells completely lose CD23a transcription (Figure 6D, lane 3). Down-regulation of CD23a was accompanied by a down-modulation of Notch2, the antiapoptotic gene bcl-2, and by a rapid increase in cell death by apoptosis as determined by FACS analysis (Figure 6B,C) and by RT-PCR (Figure 6D, lane 3). In contrast, CD23a, bcl-2, and Notch2 remained high in B-CLL cells, and this finding was associated with high cell viability.

Discussion

The overexpression of the transmembrane glycoprotein CD23 is one of the major characteristics of B-CLL cells. Besides the prognostic potential of its soluble cleavage product, sCD23, which reflects tumor load and progression of disease, selective expression of the CD23a isofrom is concurrent to a state of B-CLL cell survival, thereby providing a link between CD23a and the malfunction of apoptosis characteristic for this neoplastic B-cell type. The molecular basis underlying the up-regulation of CD23a on the gene level, however, has remained elusive.

In this report, we provide evidence that the Notch2 proto-oncogene plays a critical role in this process. Bandshift assays visualized a prominent transcription factor complex (C1) that binds sequence specific to one known and 4 newly identified putative CBF1 sites in the CD23a core promoter region. The significance of this complex was underlined because in all B-cell samples the intensity of C1 correlated with their respective levels of CD23a transcription. Moreover, by using EBV-infected BL41 cells as a positive control for CBF1-mediated CD23a expression, C1 was found to be EBV inducible. Supershift assays with antibodies raised against the nuclear forms of Notch1 and Notch2 pointed to the fact that Notch2IC is a component of C1 in B-CLL cells. Because transfection of REH pre-B cells with Notch2IC confirmed that CD23a is a target gene of Notch2 signaling, it is reasonable to conclude that Notch2 is involved in the overexpression of CD23a in B-CLL cells.

Considering that induction of CD23 through EBNA2 is an initiating event in EBV-driven B-cell immortalization, it is tempting to speculate that the Notch2 proto-oncogene plays an equally important role in the transformation process of B-CLL cells. In this context, Notch2 might not only account for B-CLL-specific CD23a expression but also for other phenotypic changes characteristic for this B-cell malignancy. In T cells, for instance, ectopic expression of Notch1IC results in the up-regulation of bcl-2 and renders thymomas resistant to glucocorticoid-induced apoptosis. Given these observations, the oncogenic nature of Notch receptors may reflect an inhibition of programmed cell death that would be consistent with the phenotype of B-CLL cells.

Another characteristic feature of B-CLL lymphocytes that might be explained by Notch2 signaling is the low expression of surface immunoglobulins. Recently, it has been demonstrated that activated Notch1, which shares many functions with Notch2, acts as transcriptional suppressor of the Igμ gene, indicating that IgM expression is negatively controlled by the Notch signal transduction pathway.

So far, however, the role of Notch2 signaling in B-cell differentiation and tumorigenesis is still not clear. In human fetal B cells, Notch2 expression is restricted to the late pre-B (CD19+ IgM+IgD-) compartment, suggesting that Notch2 confers an antiapoptotic signal when these cells undergo selection through the pre-B-cell receptor. In the current study, we show that Notch2 as well as CD23a are also expressed in normal peripheral blood B cells that served as control because they were clustered next to B-CLL cells by genomic scale gene expression profiling. Recirculating blood B lymphocytes comprise naïve and memory B cells and are characterized by their low proliferation rate and by their relative long life span. As compared with these cells, Notch2 was found to be consistently overexpressed in B-CLL cells, thereby explaining the up-regulation of CD23a. Another clear difference between normal B cells and B-CLL lymphocytes was observed by in vitro kinetic studies showing that in normal B-cells CD23a is
down-regulated within 24 hours, whereas in B-CLL cells CD23a levels remain high during 5 days of incubation. Interestingly, the CD23a levels correlate not only with Notch2 expression but also with the levels of the antipoptotic gene bel-2 together with enhanced cell viability pointing to a link between the Notch2/CD23a axis and the failure of apoptosis in B-CLL cells.

Several lines of evidence suggest that deregulation of Notch signaling locks cells into a less differentiated, proliferative state and inhibits apoptosis in certain leukemic cell lines. Therefore, a structure-function analysis together with approaches to determine Notch2 function in vivo should provide further insights into the molecular mechanisms underlying the regulation of Notch2 and its role in the pathophysiology of B-CLL.

Acknowledgments

We thank Drs M. Busslinger and T. Decker for critically reviewing the manuscript and Dr Diane Hayward for providing the pSG5-rN21C plasmid.

References
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