Conservation and divergence of NF-Y transcriptional activation function

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ABSTRACT

The CCAAT-binding protein NF-Y is involved in the regulation of a variety of eukaryotic genes and is formed in higher eukaryotes by three subunits NF-YA/B/C. We have characterized NF-Y of the trematode parasite Schistosoma mansoni and studied the structure and the function of the SMNF-YA subunit. In this work, we present the cloning and sequence analysis of the B subunit of the parasite factor. SMNF-YB contains the conserved HAP-3 homology domain but the remaining part of the protein was found to be highly divergent from all other species. We demonstrated by transfections of GAL4 fusion constructs, that mouse NF-YB does not contain activation domains while the C-terminal part of SMNF-YB has transcriptional activation potential. On the other hand, the N-terminal parts of SMNF-YA and mouse NF-YA were shown to mediate transactivation; the integrity of a large 160 amino acid glutamine-rich domain of NF-YA was required for this function and an adjacent serine- and threonine-rich domain was necessary for full activity in HepG2, but redundant in other cell types. Transactivation domains identified in SMNF-YB are also rich in serine and threonine residues. Our results indicate that serine/threonine-rich sequences from helminth parasites potenti ate transcription and that such structures have diverged during evolution within the same transcription factor.

INTRODUCTION

NF-Y (also called CBF, CP1) is a ubiquitous heteromeric complex that binds to the widespread CCAAT sequence present in the promoters and enhancers of a variety of eukaryotic genes (1,2). NF-Y has an absolute requirement for the CCAAT box as well as a strong preference for specific flanking sequences (3,4).

NF-Y从 higher eukaryotes is formed by three subunits, NF-YA, NF-YB and NF-YC (5,6). All subunits are required for DNA binding (6). Saccharomyces cerevisiae also has a CCAAT-binding activity displayed by the HAP-2/3/5 complex that is specifically required to activate transcription when cells are grown on a non fermentable carbon source (7–9). NF-YA/B/C subunits present a significant degree of identity with yeast HAP-2, HAP-3 and HAP-5 factors, respectively, in conserved regions called the HAP-2/3/5 homology domains (6,7,10). The yeast HAP-4 protein is not needed for DNA-binding but contains an acidic domain that is essential to promote transactivation when associated with the HAP-2/3/5 complex (11).

NF-YB and NF-YC form a dimer which in turn interacts with NF-YA allowing the whole complex to bind DNA (6). Interestingly, the conserved HAP-3/5 homology domains within NF-YB–NF-YC have sequence similarities to the histone fold motifs (HFM) of the H2B and H2A histones respectively, and to the archaeabacterial histone-like protein Hmf-2 (12). A previous mutational analysis of NF-YA indicated that the HAP-2 homology domain contains a segment for interaction with other NF-Y subunits and a domain necessary for DNA binding (11,13,14). NF-YA does not present similarity to known interfaces and motifs and therefore constitutes an unusual heteromeric DNA-binding factor.

Transcriptional activation functions of transacting factors generally lie on domains localized outside of the DNA-binding regions. Several transcription factors are classified according to their motifs capable of mediating transcriptional activation such as acidic, glutamine-rich and proline-rich (15,16). In higher eukaryotes, NF-Y transcriptional activation has been shown to be essentially displayed by the NF-YA and NF-YC subunits. NF-YA transactivation domains were predominantly localized in the N-terminal region of the subunit which is highly rich in glutamine (35%) and flanked on its C-terminal side by a stretch of serine/threonine residues. On the other hand, the NF-YC transactivation activity is contained in the C-terminal part of the protein, which is very hydrophobic and also highly rich in glutamine residues (17,18).

We have already characterized the NF-YA subunit of the human blood fluke Schistosoma mansoni (SMNF-YA). We have shown that the HAP-2 homology domain is highly conserved in SMNF-YA, that is able to associate with the mammalian NF-YB/C subunits to form a complex with affinity to Y boxes derived from the 5′ flanking region of the Sm28GST parasite gene (19,20). These results are in line with the functionality of the schistosome HAP-2 homology domain and the existence of NF-YB/C subunits in this parasite.
In this paper, we present and analyse the complete sequence of the B subunit from the SMNF-Y complex (SMNF-YB). As expected, SMNF-YB shows a high degree of conservation in the HAP-3 homology domain, containing the HFM and the TATA binding protein (TBP)-binding domain (21). By contrast, the remaining part of the protein is completely divergent from the mammalian and yeast counterparts. No glutamine stretches are found in the SMNF-YB N-terminus, while a large C-terminal region is rich in serine and threonine. Interestingly, the parasite NF-YB, unlike its eukaryotic counterpart, is herein shown to mediate transactivation via its serine/threonine-rich domain, suggesting the ability of such a structure to potentiate transactivation in helminth parasites.

MATERIALS AND METHODS

Cloning of the S.mansoni NF-YB subunit

The following degenerated oligonucleotides were designed following comparison of the HAP-3 homology region from yeast and mammalian proteins: 5′-AA(AG)(GA)(TC)GCAA(AG)-GA(AG)-3′ and 5′-AT(AG)(GC)(CT)TCICC(GA)TTTAT-3′. These oligonucleotides were used in a touchdown PCR (22) on first strand adult S. mansoni cDNA. The touchdown PCR was performed using decreasing annealing temperatures as follows: 45°C for 5 cycles, 40°C for 5 cycles and 37°C for the last 25 cycles. Amplification products were cloned in the pCRII plasmid (Invitrogen), sequenced and used as probes for screening of a λgt10 adult S. mansoni cDNA library.

NF-YA and NF-YB transactivation regions—DBD Gal constructs

Different oligonucleotide pairs were synthesized in order to generate an SMNF-YA segment coding from the M2 to the G171 residue positions (SmYA10: 5′-AAAGGATCCATGGACGTTTATACACGGCT-3′; SmYA12: 5′-AAAATTAGTGGCCGGCAGAGAAGACCCG-3′) and the SMNF-YB C-terminal region from the S119 to the S242 residue position (SmYB3: 5′-AAAGGATCTTCATCATCTATTGAATC-3′; SmYB: 5′-AAAGCCGCCGCAAGATTGCTTACGTAAT-3′). We also generated fragments corresponding to halves of this SMNF-YB C-terminal region from the S119 to the M198 residue (SmYB3; 5′-AAAGGATCCTTATGATAATTGAATC-3′) and from the K192 to the S242 residue (SmYB4: 5′-AAAGGATCCAAATTGCGCCAGAATG-3′; SmYB2: 5′-AAAGCCGCCGCAAGATTGCTTACGTAAT-3′). All the oligonucleotides contain adequate restriction sites to allow further cloning steps. The amplification products were initially cloned in the pCRII plasmid and sequenced. The corresponding fragments were purified after digestion with restriction enzymes and ligated into the pGAP424 plasmid in frame with the DBD Gal domain (23).

All mouse NF-YA and NF-YB GAL4 fusions, and mutants thereof, were generated by PCR using oligonucleotides containing CluI and BglII (for YA), or BamHI (for YB) sites and cloned into the corresponding sites in the polylinker of the GAL4-1-147 fragment as a probe, led us to the isolation of several clones containing a unique inserted sequence of 1547 bp with an open reading frame encoding a polypeptide of 242 amino acids homologous to the yeast HAP-3 and human NF-YB (CBF-A) proteins, named SMNF-YB.

RESULTS AND DISCUSSION

The DNA-binding and subunit interaction (HAP-3) domain of NF-YB appeared to be highly conserved at the primary sequence level. Based on this observation, we considered the possibility of isolating the S. mansoni NF-YB counterpart by a PCR-based strategy using degenerate oligonucleotides similar to the flanking regions of the conserved HAP-3 homology domain. Amplification was performed on adult S. mansoni cDNA that generated a major product of 130 bp. The sequence of the PCR product was determined, showing a high identity with the HAP-3 homology domain from yeast and mammalian factors. Further screening of an adult S. mansoni cDNA library, using the 130 bp DNA fragment as a probe, led us to the isolation of several clones containing a unique inserted sequence of 1547 bp with an open reading frame encoding a polypeptide of 242 amino acids homologous to the yeast HAP-3 and human NF-YB (CBF-A) proteins, named SMNF-YB.
Figure 1. Sequence alignment of NF-YB subunits. Amino acid sequence of SMNF-YB (Sm) was compared to human (H) and Saccharomyces cerevisiae (Sc) proteins (37). Dashes indicate the residues shared between S. mansoni and human and/or yeast proteins. The HAP-3 homology domain is boxed. The HFM is underlined and the TBP-binding domain is in italics. Stars indicate the positions of β-sheet structures in SMNF-YB predicted by the Garnier–Robson algorithm. The nucleotide sequence of SMNF-YB has been deposited to DDBJ/EMBL/GenBank under the accession no. AF037602.

As shown in Figure 1, a region highly similar to human and yeast proteins (80 and 61.5% identity, respectively) was observed between the P30 and the L104 residues of the schistosome protein. This region contained the HFM already described in the HAP-3 homology domain of mammalian transcription factors, are present in the S. mansoni NF-YB subunit. Interestingly, the C-terminal portion of SMNF-YB showed a high content of serine/threonine (30.3%) and proline (8.2%), and was also rich in valine (9.8%) residues. Analysis of this region using the Garnier–Robson algorithm, revealed that it contains a high proline content, and the enhancer binding proteins (37). Dashes indicate the residues shared between S. mansoni and human and/or yeast proteins. The HAP-3 homology domain is boxed. The HFM is underlined and the TBP-binding domain is in italics. Stars indicate the positions of β-sheet structures in SMNF-YB predicted by the Garnier–Robson algorithm. The nucleotide sequence of SMNF-YB has been deposited to DDBJ/EMBL/GenBank under the accession no. AF037602.

The N-terminal part of SMNF-YB was found to be shorter than that of other factors and very different at the protein sequence level. The remaining part of the protein was also completely different from that of already described NF-YBs. The SMNF-YB C-terminal region was larger than that of yeast and mammalian proteins and that of other factors and very different at the protein sequence level. In mammals, NF-Y-dependent activation processes are mainly mediated by NF-YA and NF-YC subunits (17,30) while in yeast, CCAAT-dependent transactivation activity has been attributed to the HAP-4 component (11). The SMNF-YA subunit does show the typical glutamine-rich transactivation domain found in the N-terminal part of NF-YA proteins from higher eukaryotes but contains also an elevated proportion of serine/threonine residue in its N-terminal region (19). A small group of transcription activators contain activation domains that have been already characterized as serine- and threonine-rich. These include the C-terminal activation domain of Oct-2 (31,32), which also possesses a high proline content, and the enhancer binding proteins (37). Dashes indicate the residues shared between S. mansoni and human and/or yeast proteins. The HAP-3 homology domain is boxed. The HFM is underlined and the TBP-binding domain is in italics. Stars indicate the positions of β-sheet structures in SMNF-YB predicted by the Garnier–Robson algorithm. The nucleotide sequence of SMNF-YB has been deposited to DDBJ/EMBL/GenBank under the accession no. AF037602.

As shown in Figure 1, a region highly similar to human and yeast proteins (80 and 61.5% identity, respectively) was observed between the P30 and the L104 residues of the schistosome protein. This region contained the HFM already described in the HAP-3 homology domain of the human and yeast proteins (12) and shown to be dependent on strict structural constraints for function. A conserved domain has also been described within the HAP-3 homology domain as a short segment adjacent to HFM and necessary for TBP-binding (21). A similar TBP-binding domain is also found at the same position (between N99 and R105 residues) in the SMNF-YB sequence.

The N-terminal part of SMNF-YB was found to be shorter than that of other factors and very different at the protein sequence level. The remaining part of the protein was also completely different from that of already described NF-YBs. The SMNF-YB C-terminal region was larger than that of yeast and mammalian proteins and does not show identity with any other sequences. In particular, neither glutamine nor proline stretches characteristic for transactivation domains in mammalian transcription factors, are present in the S. mansoni NF-YB subunit. Interestingly, the C-terminal portion of SMNF-YB showed a high content of serine/threonine (30.3%) and proline (8.2%), and was also rich in valine (9.8%) residues. Analysis of this region using the Garnier–Robson algorithm, revealed that it may adopt a structure formed by six β-sheets, each composed of 13–23 residues and homogeneously distributed from the end of the HAP-3 homology region to the C-terminus of the protein. The same algorithm analysis was not able to predict any similar structure in yeast or mammalian protein sequences.

In mammals, NF-Y-dependent activation processes are mainly mediated by NF-YA and NF-YC subunits (17,30) while in yeast,
with an anti-GAL4 antibody: Figure 3B shows that mutants negative in transcription were present at equal levels compared to the functioning fusion proteins.

The same type of experiment was tried with mouse NF-YB–GAL4DBD fusions (Fig. 4): in this case, neither the wt NF-YB, nor any of the mutants showed any detectable degree of activation on the M2TA TA-CA T vector. To further prove that mouse NF-YB subunit is truly devoid of any activating potential in this assay and rule out that more than two GAL4 sites are required for NF-YB proteins, we tried the more sensitive cis-vector containing the TK promoter in front of two GAL4 sites. As shown in Figure 4, this vector was efficiently activated by the positive GAL4 control, but not by any of the NF-YB constructs: if anything, a small but reproducible degree of repression was observed. We also transfected NIH3T3 cells in parallel with SMNF-YB, mouse NF-YB and YA6, with the 5× GAL4 sites-containing vector used before: the results shown in Table 1 indicate that unlike SMNF-YB, mouse NF-YB does not activate. This experiment also allowed us to quantify the YA6 potential as being of similar magnitude with respect to the SMNF-YA and SMNF-YB. In these experiments, the expression efficiency of all mutants was controlled by western blot analysis, as described for NF-YA constructs (data not shown). Moreover, the mouse GAL4-NF-YB constructs do activate, provided that the two other subunits are cotransfected (A.di Silvio and R.Mantovani, in preparation). The conclusion of this set of experiments is that mouse and schistosome NF-YA are both able to activate transcription, while only the helminth NF-YB harbours such potential.

### Table 1. Comparative analysis of schistosome and mouse NF-YB transcriptional potential

<table>
<thead>
<tr>
<th>Plasmid</th>
<th>Luciferase activity (fold activation)</th>
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<tbody>
<tr>
<td>GAL4Poly</td>
<td>1</td>
</tr>
<tr>
<td>GAPYB (S.mansoni)</td>
<td>10.4</td>
</tr>
<tr>
<td>GALKYB (mouse)</td>
<td>1.6</td>
</tr>
<tr>
<td>GALY (mouse)</td>
<td>11.1</td>
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Luciferase activity was measured in NIH3T3 cells co-transfected with NF-Y constructs and pGSTATALuc plasmid.

NF-Y represents an ideal system to follow conservation/divergence of transcriptional activation domains. Unlike many general transcription factors (such as TBP, TFIIIB, TFIIA, etc.) which are highly conserved, but do not possess a distinct transcription activating domain, and unlike many DNA-binding factors, whose conservation in different phyla is much lower and restricted to the DNA-binding domain, NF-Y shows among the highest degree of conservation (67–73% identity, 85–90% similarity) in the subunit association and DNA-contacting parts, but also in recognizable activating domains. At this time, little is known about transcription factors in helminth parasites. Besides the A subunit of the SMNF-Y complex, a limited number of *Schistosoma* transcription factors have been described, that include the Heat Shock Factor (HSF) (35) and several nuclear receptors considered as transcriptional regulators (36). In all cases, a relatively high conservation of sequences involved in the binding...
to specific DNA targets was observed between schistosomes and higher eukaryotes. However, no significant identity was found when other regions, particularly activation domains of mammalian factors, were compared to their *Schistosoma* counterparts.

The NF-YA N-terminal domains of mammals, sea urchin, *S.mansoni* and *Aspergillus nidulans* contain clearly recognizable glutamine-rich domains (19,37 , R.Mantovani, unpublished). Similar in overall amino acid composition, hydrophobics predominate as well as glutamines, they are largely different in the primary structure. Many factors of different phylogenetic origins have been shown to activate transcription by their acidic- and/or proline-rich transactivation domains in the budding yeast (38–40). The *S.cerevisiae* acidic transcription factor GAL4 also activates transcription in mammals as well as in plants, suggesting a conserved and interchangeable mechanism of acidic transactivation factors throughout evolution (41,42). In contrast, glutamine-rich domains, such as those found in the human SP1 factor, failed to potentiate activation of transcription in *S.cerevisiae* (43), while they were functional in *Schizosaccharomyces pombe* (44). Results indicating that glutamine-rich sequences have different functions in human, *S.pombe* and *S.cerevisiae* could mean that such domains need additional transcription cofactors which are present in mammals and absent, or playing distinct functions, in *S.cerevisiae*. In keeping with this observation, a recent paper reported binding of NF-YA–NF-YC Q-rich activation domains to dTAFII110 (45), one of the few TBP-associated factors for which no *S.cerevisiae* counterpart has been found so far. In *S.cerevisiae*, a fourth subunit, HAP-4, containing

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**Figure 3.** Transcriptional activation of mouse NF-YA. (A) The scheme depicts the wt short (YA1) and long (YA13) forms of NF-YA proteins fused to the DNA-binding domain of GAL4. YA2 (amino acids 161–346), YA3 (amino acids 262–346), YA4 (amino acids 1–317), YA5 (amino acids 1–261), YA6 (amino acids 1–160), YA8 (amino acids 161–317), YA9 (amino acids 262–317). YA10, YA11, YA12 are equivalent to YA4, YA5, YA6, except that they contain the 28 amino acids of the optional exon B (46). YA14 (amino acids 1–70, short form), YA15 (amino acids 1–70 long form), YA16 (amino acids 71–160). The reporter CAT vector M2TATA-CAT contains two GAL4-binding 17mers in front of the AdML TATA box (24). (B) Western blot analysis of extracts derived from HeLa cells transfected with the indicated YA mutants using an anti-GAL4 monoclonal antibody.

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**Figure 4.** Transcriptional potential of mouse NF-YB. GAL4–NF-YB constructs were made essentially as described for NF-YA in Figure 3. YB-1 (amino acids 1–161), YB2 (amino acids 1–50), YB4 (amino acids 51–140), YB6 (amino acids 141–207). The N- and C-terminal domains of NF-YB are indicated by dotted boxes, while a grey box indicates the central conserved domain of NF-YB. Results of transfection of HeLa cells with M2TATA-CAT (left panel) or MX2-CAT (containing a tandem of GAL4-binding 17mers fused to the TK promoter) are presented.
a powerful highly acidic domain is required to activate the HAP-2/3/5 complex: this subunit is apparently absent in other phyla, where the activating information has been incorporated within the core subunits within complex, perhaps modulable domains. In NF-YA, a serine- and threonine-rich segment was described as being essential for full transactivation in concert with the glutamine-rich region in an in vitro system (17). The data of transfections in different cell lines presented here support the idea that this domain is necessary for full activation in some but not all cell lines, thus suggesting that activation is influenced by the cell-type context, probably reflecting the existence, or abundance, of intermediate cell-specific cofactors capable of mediating activation in a given cell type. It is worth noting that mouse NF-YA activation emerges only in GAL4 constructs devoid of the HAP-2 homology domain: this could be explained by supposing that such fusion proteins are complexed by the endogenous NF-YB–NF-YC homology domain: this could be explained by supposing that such NF-Y protein identity is restricted to the HAP-2/3/5 homology domains, the latter two containing putative histone fold domains and favour of a complex evolution of the structure/function of NF-Y factors as well as of other transcription factors.

Our study, performed in mammalian cells, represents the first indication that activation domains of a Schistosoma transcription factor, as divergent from human as it is from yeast, efficiently operates in a mammalian context. The results presented here also show, for the first time, the existence of a functional activation domain in the B subunit of an NF-Y complex. We found that, unlike NF-YA and NF-YC that mediate transactivation via their glutamine-rich domains, the mouse NF-YB subunit is completely unable to potentiate transcription in higher eukaryotes. Our experiments suggest a model whereby transactivation by NF-Y in helminth parasites could be mediated by the A and B subunits via glutamine- and serine/threonine-rich domains, with the latter being slightly more efficient. This observation could also signify the preference for serine/threonine-rich conformations as activators for the schistosome RNA PolII, depending on the context of the parasite promoters and the position of DNA-binding sites that govern responsiveness to different regulatory signals. The fact that the C-terminal region of SMNF-YA also contains serine and threonine residues in addition to glutamines, when compared with the mammalian NF-YA (15), is in line with this hypothesis. Further cloning and analysis of the third SMNF-YC subunit will certainly shed more light on the evolution of transcription activating function of this conserved transcription factor.

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