The p75NTR-interacting protein SC1 inhibits cell cycle progression by transcriptional repression of cyclin E

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Schwann cell factor 1 (SC1), a p75 neurotrophin receptor–interacting protein, is a member of the positive regulatory/suppressor of variegation, enhancer of zeste, trithorax (PR/SET) domain-containing zinc finger protein family, and it has been shown to be regulated by serum and neurotrophins. SC1 shows a differential cytoplasmic and nuclear distribution, and its presence in the nucleus correlates strongly with the absence of bromodeoxyuridine (BrdU) in these nuclei. Here, we investigated potential transcriptional activities of SC1 and analyzed the function of its various domains. We show that SC1 acts as a transcriptional repressor when it is tethered to Gal4 DNA-binding domain. The repressive activity requires a trichostatin A–sensitive histone deacetylase (HDAC) activity, and SC1 is found in a complex with HDACs 1, 2, and 3. Transcriptional repression exerted by SC1 requires the presence of its zinc finger domains and the PR domain. Additionally, these two domains are involved in the efficient block of BrdU incorporation by SC1. The zinc finger domains are also necessary to direct SC1's nuclear localization. Lastly, SC1 represses the promoter of a promitotic gene, cyclin E, suggesting a mechanism for how growth arrest is regulated by SC1.

Introduction

Schwann cell factor 1 (SC1) is a p75 neurotrophin receptor (NTR)–interacting protein (Chittka and Chao, 1999). It belongs to a small group of proteins referred to as the retinoblastoma-interacting zinc finger (RIZ) family of transcription factors. These proteins are characterized by the presence of zinc finger and positive regulatory/suppressor of variegation, enhancer of zeste, trithorax (PR/SET) domains, and are involved in cell differentiation and tumorigenesis (for review see Huang et al., 1998). Proteins containing the PR/SET domain include the lymphocyte PRDI-BF1/BLIMP protein (Keller and Maniatis, 1991), RIZ (Buyse et al., 1995), and the Evi1 gene (Fears et al., 1996). The presence of the PR/SET domain makes these proteins distinct from a multitude of other zinc finger motif-bearing proteins. The PR/SET domain is a modified SET domain. SET domain proteins possess protein methyltransferase activity and have been shown to methylate histones, thus regulating chromatin structure and transcription (Rea et al., 2000; Strahl et al., 2002). However, close inspection of the amino acid sequences of the PR domains reveals striking differences in the highly conserved residues necessary for the methyltransferase activity of the SET domain (Kouzarides, 2002). The zinc finger domains and the PR/SET domain of SC1 imply that this protein may mediate transcriptional activities. Thus, transcriptional activities of repressive and activating nature have been described for RIZ (Xie et al., 1997; Abbondanza et al., 2000; Steele-Perkins et al., 2001), Blimp-1/PRDI-BF1 (Lin et al., 1997; Ren et al., 1999; Yu et al., 2000), and Evi-1 (Kurokawa et al., 1998; Izutsu et al., 2001; Palmer et al., 2001). SC1 is the first member of this family that is implicated in the neurotrophin signaling.

Abbreviations used in this paper: DBD, DNA-binding domain; HDAC, histone deacetylase; NTR, neurotrophin receptor; PR/SET, positive regulatory/suppressor of variegation, enhancer of zeste, trithorax; RIZ, retinoblastoma-interacting zinc finger; SC1, Schwann cell factor 1; siRNA, small interfering RNA; TSA, trichostatin A.
Neurotrophins, such as NGF, BDNF, and NT-3, are important survival and differentiation factors. The neurotrophins interact with two distinct classes of receptors, members of the Trk tyrosine kinase receptor subfamily and p75NTR, a member of the TNF receptor superfamily (Chao and Hempstead, 1995). There is evidence that p75NTR can function as a cell death receptor (Casaccia-Bonnefil et al., 1996; Frade et al., 1996; Bamji et al., 1998; Frade and Barde, 1998). Several proteins that interact with p75NTR have been shown to transduce apoptotic signaling. These include NRIF (Casademunt et al., 1999), NRAGE (Salehi et al., 2000), and NADE (Mukai et al., 2000). Interestingly, p75NTR is expressed very early during development (Yan and Johnson, 1987, 1988), before differentiation of many precursor cells of the nervous system and before the onset of programmed neuronal cell death. This observation makes it a candidate for a receptor functioning in the control of precursor cell proliferation and differentiation. Supporting this idea is the observation that several proteins which interact with p75NTR induce growth arrest. Expression of SC1, for example, was previously found to be correlated with a decrease in BrdU incorporation (Chittka and Chao, 1999).

To elucidate the mechanism by which SC1 transduces the neurotrophin signaling, we began to analyze its potential transcriptional activity. In a reporter gene assay, SC1 acted as a transcriptional repressor. Both the zinc finger domains as well as the PR/SET domain are necessary for SC1 to act as a repressor, and are needed to effectively block BrdU incorporation into cell nuclei. The repression exerted by SC1 required the presence of trichostatin A (TSA)–sensitive histone deacetylases (HDACs), and SC1 was found in the complex with HDAC 1, 2, and 3. Further, we show that SC1 behaves as a transcriptional repressor upon NGF application to the cells transfected with both SC1 and either p75NTR or TrkA. We found the zinc finger domains of SC1 are necessary for its nuclear localization. Finally, our analysis of genes transcriptionally regulated by SC1 revealed that SC1 down-regulates the expression of a promitotic gene, cyclin E, consistent with its ability to block DNA replication as measured by BrdU incorporation. These results implicate SC1 as a potential transcriptional mediator of NGF signaling that may be involved in modifying the chromatin structure during differentiation.

**Results**

**Regulation of the subcellular distribution of SC1**

SC1 is localized to both the nucleus and the cytoplasm of COS1 and Schwann cells (Chittka and Chao, 1999). We sought to determine which domains of SC1 were responsible for this differential distribution. To address this question, we generated a series of deletion mutants of SC1 fused to GFP. These include full-length SC1 or deletions lacking either the acidic COOH terminus, the zinc finger domains, or the PR/SET domain (see Fig. 1 for a schematic representation of the constructs used in these experiments). These constructs were then transfected into COS1 cells, and the distribution of each protein was investigated. The results of these experiments are shown in Fig. 1 (a–e). GFP alone distributed to the cytoplasm and nucleus (Fig. 1 a). Full-length SC1 as well as SC1ΔPR and SC1ΔC were found predominantly in the nucleus (Fig. 1, b, c, and e, respectively), as can be visualized by their overlapping distribution with the Hoechst stain in the nuclei. Strikingly, the deletion of zinc finger domains resulted in a cytoplasmic distribution of SC1 (Fig. 1 d). Therefore, we concluded that zinc finger domains are necessary for nuclear localization of SC1.

**SC1 is an HDAC-dependent transcriptional repressor**

The localization of SC1 is highly regulated, and translocation of the protein to the nucleus was closely correlated with a loss of BrdU incorporation (Chittka and Chao, 1999). These results suggest that SC1 may be involved in transcriptional events associated with growth arrest. To assess whether SC1 possessed the ability to modulate gene transcription, we fused the coding sequence of SC1 to the Gal4 DNA-binding domain (DBD; see Fig. 2 B for the schematic representation of the effector and reporter constructs). Fusion of these truncated SC1 proteins with Gal4-DBD, which contains an endogenous NLS, ensured nuclear localization of the proteins (see Fig. 5). A luciferase reporter under transcriptional control of Gal4 upstream activating sequences was used for measuring transcription relative to a control effector without SC1 (i.e., Gal4 alone). Gal4-SC1 or Gal4 and the reporter DNAs were cotransfected into HEK293 cells, and luciferase activity was measured 48 h later.

A reduction in luciferase activity was consistently observed when SC1 was fused to the Gal4-DBD and cotransfected with the reporter as compared with the luciferase activity measured in the presence of Gal4-DBD only (Fig. 2 A, lanes 1 and 2). We used a reporter construct where the luciferase gene was under the control of the TATA element only to verify the specificity of the results observed when luciferase is
SC1 is a transcriptional cell cycle inhibitor driven by the Gal4 upstream activating sequence. No repression was measured when SC1 was compared with Gal4 alone in these reporter assays (Fig. 2 A, lanes 3 and 4), supporting the conclusion that SC1 acts as a transcriptional repressor with a responsive promoter.

To define the domains of SC1 responsible for the repressive activity, we created truncated fusion proteins of NH₂-terminally Flag-tagged SC1 with the Gal4-DBD and used them for transfection of HEK293 cells in our reporter assays. Three truncated proteins were made: (1) Δ754–798 lacks the extreme COOH terminus, which contains a highly acidic domain; (2) Δ583–798 lacks the six zinc finger domains in addition to the COOH-terminal domain; and (3) Δ404–798 lacks the PR domain and the entire COOH terminus. (see Fig. 2 B for the schematic representation of the fusion proteins). Expression of these proteins was verified by transfection of HEK293 cells and subsequent immunoprecipitation of the proteins using anti-Flag antibodies (Fig. 2 C).

We observed that deletion of the zinc finger domains from SC1 leads to a loss of its repressive activity (Fig. 2 B, compare lane Δ583–798 lacking the zinc fingers with Gal4 alone), whereas deletion of the acidic-rich COOH terminus had no influence on the repression by SC1 (Fig. 2 B, compare lane Δ754–798 lacking the COOH-terminus with Gal4 alone). Interestingly, deletion of the PR domain rendered SC1 a transcriptional activator, as can be seen from an increase in luciferase activity over the control levels (Fig. 2 B, compare lane Δ404–798 with Gal4). Such behavior has been observed for Blimp/PRDI-BF1 (Ren et al., 1999; Yu et al., 2000). We noticed that the truncations of the zinc fingers and the PR/SET domain led to a reduction of the total protein levels detected. To exclude the possibility that the increase in the luciferase activity observed upon the truncations of SC1 is due to the decrease of the protein level, we performed a series of experiments where an increasing amount of DNA encoding these proteins was transfected and luciferase measurement was performed.

Figure 2. SC1 is a repressor that requires its zinc finger domains and the PR/SET domain for repression. (A) Histogram of representative relative luciferase units after a cotransfection of Gal4 or Gal4SC1 with the Gal4-luc reporter (lanes 1 and 2, respectively) or TATA-luc reporter (lanes 3 and 4, respectively) in HEK293 cells. Whole-cell extracts were used in the luciferase assays and β-galactosidase measurements. The basal promoter activity was measured in the presence of Gal4 alone (black bar), and SC1’s regulation of the basal promoter was compared with it (open bar). A schematic map of the reporter construct is represented on the top. The graph shows the result of one experiment performed in triplicate that was reproduced in several independent experiments. (B) Left: histogram of relative luciferase units measured after cotransfection of various truncated mutants of SC1 fused to the Gal4-DBD represented schematically on the right. The numbers on the schematic drawings specify the amino acids of SC1. PR, PR/SET domain; ZF, zinc finger domain; Ac, an acidic COOH-terminal portion of SC1 protein. Black bar represents relative luciferase units in a control cotransfection experiment using Gal4 alone as an effector; white bars represent the results of cotransfection with various Gal4SC1 mutants. (C) Western blot showing the expression of Gal4-SC1 fusion proteins in HEK293 cells, as detected by anti-Flag antibody. Fusion proteins were immunoprecipitated from whole-cell lysates with anti-Flag antibody and separated on a 7.5% SDS-PAGE before blotting and detection with anti-Flag antibodies. The graph below demonstrates the results of the control transfections where increased amount of cDNA coding for the deleted SC1 proteins with lower expression levels were tested for their ability to influence luciferase measurements. The amounts of DNA used are shown below the appropriate construct (compared with the 0.3 μg normally used in these experiments).
formed. We obtained similar results in these experiments (Fig. 2 C, bottom graph).

As many transcriptional repressors recruit HDACs and need this activity to exert their repression (Grunstein, 1997; Pazin and Kadonaga, 1997; Wolffe, 1997), we tested whether TSA, a potent inhibitor of HDACs, influences the repressive activity of SC1 in the transcriptional reporter assays. Fig. 3 A shows that the repressive activity of SC1 was completely abolished by the addition of 50 ng/ml TSA, indicating that SC1’s repression relies on the activity of TSA-sensitive HDACs. To directly test the interaction between SC1 and HDACs, we cotransfected Flag-tagged SC1 with HA-tagged HDAC 1, 2, or 3 into HEK293 cells and performed a coimmunoprecipitation assay. Anti-Flag Sepharose was used to bring down the Flag-tagged SC1. Expression of the fusion proteins was monitored by Western blot analysis. Cotransfection of Flag-vector with HDACs was used as a negative control. The results of these experiments are presented in Fig. 3 B, which demonstrated that SC1 can be found in a complex containing all the tested HDACs. Thus, SC1 associates with the class I HDACs and is likely to exert its repression by the recruitment of these proteins to the appropriate promoter sites.

The zinc finger domains and the PR/SET domain of SC1 are necessary to block BrdU incorporation

The zinc finger domains of SC1 are required for both nuclear localization and transcriptional repression, whereas the PR/SET domain is involved in the modulation of the repressive activity by SC1. In this work, we investigated whether the deletion of zinc finger domains or the PR/SET domain would influence BrdU incorporation. Full-length and truncated forms of SC1 fused to either GFP or Gal4-DBD were used for this investigation. Fig. 4 summarizes the results from transfections of COS1 cells and Schwann cells with the GFP-SC1 fusion proteins. The data were normalized with respect to the BrdU incorporation of cells expressing GFP alone (Fig. 4, A and B). The quantification of nuclei that have incorporated BrdU upon transfection with the various SC1 truncated proteins revealed that the deletion of zinc finger or the PR/SET domains resulted in an inability to block BrdU incorporation. Fig. 4 A contains the results for ΔPR and ΔZF in COS1 cells. Fig. 4 B contains the results for ΔPR and ΔZF in Schwann cells and the respective quantifications. Overexpression of both the full-length and COOH-terminally truncated SC1 protein led to a block of DNA synthesis to the same extent (Fig. 4, A and B). These experiments verified that the zinc fingers are necessary for nuclear entry of SC1. On the other hand, deletion of the PR/SET domain led to a loss of BrdU incorporation, even though the SC1ΔPR protein remained in the nucleus. Hence, the PR/SET domain is involved in the regulation of BrdU incorporation by SC1.

To characterize the transcriptional activity of SC1 further, we used the Gal4-SC1 constructs used previously for the reporter gene assays. In contrast to the GFP fusion proteins used for experiments with COS1 and Schwann cells, SC1 deletion proteins were tethered to the Gal4-DBD with an endogenous NLS (see Fig. 5, A and C, for immunocytochemical detection of overexpressed proteins). This allowed us to establish a correlation between SC1’s repressive activity and its ability to influence BrdU incorporation. As described in the previous paragraph, we normalized the BrdU incorporation data with respect to Gal4-DBD–transfected cells. Again, we observed a loss of SC1’s capacity to block BrdU incorporation in the absence of its zinc finger domains (Fig. 5, B and D, lane SC1Δ583–798 for the respective quantification of BrdU incorporation in NIH3T3 and HEK293 cells, respectively). Overexpression of both full-length SC1 (Fig. 5, B and D, lane SC1FL for NIH3T3 and HEK293 cells, respectively) and a COOH-terminally truncated SC1 (Fig. 5, B and D, lane SC1Δ754–798 for NIH3T3 and HEK293 cells, respectively) resulted in much
lower BrdU incorporation in both cell lines. Thus, SC1’s repressive transcriptional activity could be involved in the implementation of the growth arrest in these cells.

The influence of NGF on the transcriptional activity of SC1
SC1 was originally isolated as a protein that interacts with the cytoplasmic domain of p75NTR, and its subcellular distribution was shown to be regulated by NGF. Therefore, we sought to investigate whether its transcriptional activity is regulated by the addition of NGF. To investigate this, we transfected HEK293 cells with Gal4-SC1 fusion protein as well as either TrkA or p75NTR and luciferase cDNAs under the control of Gal4 promoter as a reporter. Expression of p75NTR or TrkA was monitored by Western blotting after transfection (unpublished data). NGF was applied for either 1 or 5 h, and the luciferase activity was measured afterwards. We observed that coexpression of TrkA with SC1 led to a potentiation of the repressive activity of SC1 (see Fig. 6 A for the graphic representation of the relative luciferase units measured in these experiments). To verify the specificity of TrkA’s influence on the repressive activity of SC1, we used K252a, a potent inhibitor of Trk signaling, and SC1’s transcriptional output was then evaluated. Addition of K252a led to a loss of the potentiation of SC1’s repressive activity as measured by the reporter assay (refer to Fig. 6 A for the graphic representation of these results). This observation implicates TrkA in the regulation of SC1’s transcriptional activity. In the cotransfection experiments with p75NTR, SC1’s repressive activity also correlated with the addition of NGF to the transfected cells. Thus, the addition of NGF for 1 h activated the repressive activity of SC1, and this activity increased over time as can be seen from the 5-h measurement (see Fig. 6 B for the graphic representation of SC1’s transcriptional activity in the presence of p75NTR and NGF). Hence, SC1 is a transducer of both TrkA and...
SC1 represses transcription of genes that drive cell cycle progression

The fact that SC1’s transcriptional activity influenced BrdU incorporation led us to test the possibility that SC1 may repress the expression of promitotic genes. To this end, we used a reporter gene system where the expression of cell cycle–related genes was assayed using a luciferase reporter assay. We used cyclin E, cyclin A, and cyclin B promoters for this analysis. The cyclin E promoter contained 1.4 kb of the sequence upstream of the transcription start site, which has been shown to regulate its transcription during G1 phase (Geng et al., 1996); the cyclin A promoter contained 3.2 kb of the sequence upstream of the transcription start site known to regulate its transcription during S phase (Yoshizumi et al., 1995); and the cyclin B1 promoter contained 3.8 bp upstream of the transcription start site. This sequence has been shown to enhance its transcription during G2 phase (Cogswell et al., 1995). Flag-tagged full-length SC1 was used as an effector, and the cyclin promoter-specific driven luciferase constructs were used as reporters. NIH3T3 cells were synchronized by serum withdrawal (see the Materials and methods section) for 48 h and released from growth arrest by the addition of serum-containing medium.

First, we determined the time course of expression of cyclins E, A, and B in these cultures by performing Northern blots at 0, 4, 8, 12, 16, 20, and 24 h after serum addition. Cyclin E could be detected at 12 h first and persisted until 24 h, cyclin A could first be detected at 20 h, and cyclin B was first detected at 24 h after serum addition of the growth-arrested cells (unpublished data). The cells were transfected with either SC1-bearing plasmid or a control plasmid, and lysates were collected at the following time points after serum addition: at 12 h for cyclin E measurement; at 20 h for cyclin A measurement; and at 24 h for cyclin B measurement. Subsequently, luciferase activity was measured in these lysates. We observed that cyclin E was down-regulated as measured by the luciferase activity, and the repression of the cyclin E promoter was comparable in magnitude to the repression exerted by SC1 when it is tethered to the Gal4 moiety (compare Fig. 7 A with Fig. 2 A). There was also a slight reduction of cyclin B expression as measured by the luciferase activity (Fig. 7 C). Expression of cyclin A, the major cyclin acting during S phase of the cell cycle, was not down-regulated by SC1 in these experiments (Fig. 7 B). To verify that the cyclin genes are expressed in these cells, we performed RT-PCR to monitor the endogenous expression of the cyclins (Fig. 7). Then, we investigated whether the levels of endogenous cyclin E protein were affected by the overexpression of SC1. To this end, we analyzed the protein lysates from NIH3T3 cells that had been transfected with either SC1 or a control Flag-bearing plasmid and treated as described above. Equal amounts of experiments are shown). C and D are the same as A and B for HEK293 cells. Arrows indicate the cells transfected with SC1 constructs and the corresponding BrdU incorporation into their nuclei. Bars, 20 μm.
total protein were loaded on the gels, which were then transferred for Western blot analysis. We used an anti-cyclin E antibody to detect the levels of total cyclin E protein in these lysates, and normalized these by probing the blots with an anti-actin antibody (Fig. 7 D). The images were analyzed using a densitometer to determine the relative levels of cyclin E and actin present in transfected and mock-transfected cells, and were corrected for the transfection efficiency of these cells (which was routinely between 30 and 40%). We calculated a reduction of the total cyclin E protein level by a factor of 2.5 in cells overexpressing SC1, supporting our previous observations on the repression of the cyclin E promoter in the reporter assays (see Fig. 7 A).

Our observations that SC1 represses the expression of cyclin E and that its activity is regulated by both TrkA and p75NTR led us to test the validity of these observations in a well-characterized cell line used in studying neurotrophin signaling, namely PC12 cells. Thus, the cells were synchronized using the method of Rudkin et al. (1989) by serum withdrawal, released from the cell cycle block by the addition of different factors, and samples to be analyzed were taken at \( \sim 50 \) h after transfection for luciferase measurement (expression of the endogenous cyclin E at this time was verified by performing RT-PCR; unpublished data). We tested the effects of NGF on the repression of the cyclin E promoter by SC1. The results of these experiments are summarized in Fig. 8. Application of NGF to these cells enhanced the repression of cyclin E promoter by SC1. These observations implicate SC1 in the control of cell cycle progression and make it an important component of NGF signaling.

Next, we investigated the role of the endogenous SC1 in regulating cyclin E expression in PC12 cells. To this end, we used the small interfering RNA (siRNA) method to block the expression of endogenous SC1 in PC12 cells, and measured the levels of cyclin E proteins in the lysates of transfected or mock-transfected cells. In parallel experiments with GFP expression plasmids, transfection rates in PC 12 cells were in the range of 30–50%, and thus resembled those in a recent paper (Rossoll et al., 2003). This corresponded to a reduction of SC1 by 40–60% in our experiments (Fig. 9). Down-regulation of SC1 leads to a significant increase in cyclin E expression, whereas the levels of cyclin A are not modified (Fig. 9), consistent with our observations in NIH3T3 cells as measured by luciferase reporter gene assays (Fig. 7, A and B). The quantification of the expression levels of SC1, cyclin E, and cyclin A proteins in PC12 cell lysates transfected with either SC1 siRNA or a scrambled siRNA construct is presented in Fig. 9 B. These experiments clearly show the involvement of endogenous SC1 in the regulation of transcription of cyclin E, but not cyclin A in PC12 cells.

**Discussion**

Transcriptional regulation through chromatin modification is emerging as one of the crucial ways to control differentiation during development. In this paper, we describe a function of SC1, a novel p75NTR-interacting zinc finger protein with a PR/SET domain. We present evidence that SC1 can act as a transcriptional repressor, it is found in a complex with HDACs 1, 2, and 3, and its activity is sensitive to TSA. Additionally, we demonstrate here that zinc finger domains are required for multiple functions of SC1 (i.e., nuclear localization, transcriptional repression, and SC1’s ability to negatively influence DNA replication). The PR/SET domain is involved in transcriptional repression and decreases BrdU incorporation. Further, we present evidence that SC1...
can down-regulate the expression of cyclin E, which is instrumental for the S-phase entry during cell cycle (Ewen, 2000). This down-regulation is observed in both NIH3T3 cells and in PC12 cells where the effect is enhanced by NGF.

Analysis of the transcriptional activity by SC1 revealed that it acts as a transcriptional repressor both in the context of being tethered to a DBD of another protein, Gal4, and by itself in transfection experiments. Our data indicate that the six zinc finger domains and the PR/SET domain are required for transcriptional repression by SC1. This implicates the zinc finger’s and PR/SET domain’s direct involvement in transcriptional activity either by allowing DNA binding of SC1 or by recruitment of other proteins that may modulate transcriptional repression. Consistent with the latter hypothesis, we observed that SC1 exerts its repressive activity by recruiting TSA-sensitive HDACs. Complexes involving HDACs have been implicated in silencing neuronal-specific genes (Naruse et al., 1999; Ballas et al., 2001). SC1 is an HDAC-dependent repressor, which may be involved in actively predisposing cells for differentiation by blocking their proliferative potential through down-regulation of promitotic genes.

Interestingly, truncation of the region of SC1 that contains the PR/SET domain transforms SC1 into a transcriptional activator, thus pointing to an involvement of this domain in repressive activity of SC1. Similar observations were made when Blimp-1’s PR domain was removed (Yu et al.,...
In this respect, it is noteworthy that two related proteins (RIZ and MDS1-EVI1) encode two different transcripts, which differ only in the presence of the PR/SET domain in the respective proteins (Bartholomew and Ihle, 1991; Liu et al., 1997). The absence of the PR/SET domain correlates with a weaker repressive activity in RIZ protein (Xie et al., 1997), suggesting a modulatory role of PR/SET domains during repression. Additionally, Blimp-1/PRDF-BF1 can exert repression through its PR/SET domain (Ghosh et al., 2001).

SET domains are found in chromosomal proteins that modulate gene expression and chromatin structure (for review see Jenuwein, 2001). Several proteins containing SET domains possess lysine histone methyltransferase activity (O’Carroll et al., 2000; Rea et al., 2000; Strahl et al., 2002). These histone modifications can lead to transcriptional repression (Firestein et al., 2000), but other enzymes are likely to be required, e.g., HDACs, to act together with methyltransferases. We do not know whether the PR/SET domain of SC1 possesses a protein methyltransferase activity. However, given SC1’s ability to repress transcription in an HDAC-dependent manner, it would make SC1 a prime candidate for developmental regulation and subsequent maintenance of differentiation. Such a role is further substantiated by the observation that SC1’s action is correlated with a block of BrdU incorporation, suggesting a role in cell differentiation and possible maintenance of the differentiated state.

One of the genes whose transcription is negatively regulated by SC1 is cyclin E, the major cyclin at G1–S phase transition. The down-regulation of cyclin E is consistent with the hypothesis that SC1 may play a major role at the G1–S decision making stage, and possibly during terminal mitosis as the cells prepare to enter a differentiative program. A down-regulation of cyclin E mRNA was observed by Tramtrack protein during glial development in Drosophila melanogaster, blocking entry into S phase and thus regulating glial cell proliferation (Badenhorst, 2001). Additionally, the activity of cyclin E together with CDK2, its binding partner, is instrumental for the regulation of cell cycle progression in neural and glial progenitor cells, and its block leads to an efficient cell cycle arrest (Casaccia-Bonnefil et al., 1999; Ferguson et al., 2000). Thus, SC1 may be one of the key regulators determining the decision of proliferating cells to exit cell cycle in the nervous system.

Several interesting points arise concerning the possible role of SC1 in neurotrophin signaling. It has been shown that in PC12 cells, the responsiveness to NGF is cell cycle stage-dependent (Rudkin et al., 1989), i.e., they will respond by differentiating when in G1 phase, but progress through the cell cycle when exposed to NGF during the other cell cycle phases (van Grunsven et al., 1996a,b). This responsive difference to NGF can be partially explained by the different temporal cell surface expression of TrkA and p75NTR (Urdiales et al., 1998). Thus, TrkA is observed on the cell surface during late M/early G1 phase, whereas p75NTR is highly expressed at the surface during the remaining time of cell cycle. Consistent with this temporal expression of the two NGF receptors, SC1 may act as a transducer of anti-proliferative signaling by NGF at an appropriate cell cycle stage, i.e., at G1–S transition as a blocker of further S phase entry. We observed that both TrkA and p75NTR enhanced the repressive activity of SC1 in our cotransfection experiments, implying the role of SC1
as the transducer of NGF signaling by these two receptors. This observation is particularly interesting in view of our data, showing that NGF enhances the repressive activity of SC1 at the cyclin E promoter in PC12 cells. As such, SC1 could play a decisive role during the differentiation of PC12 cells upon NGF addition. Specifically, as the decision to exit the cell cycle is made during G1 phase in these cells and both TrkA and p75NTR are expressed at this stage, they would be involved in mediating the anti-mitogenic response through SC1 as one of the key players in this process. The observations presented here offer novel venues for probing the molecular mechanism of NGF action as it is transduced by both TrkA and p75NTR through the activation of SC1.

Materials and methods

Plasmids

We used 0.75 μg 5xGal4 UAS-luciferase DNA (a gift of Al Fisher, Cornell University Medical College, New York, NY; Catron et al., 1995); 0.25 μg Gal4SC1 or other deletion mutants of SC1; and 0.05 μg pCMVβ-gal DNA for transfections of HEK293 cells. The same relative ratios and amounts of plasmid DNA were used for transfections with cyclin-specific luciferase constructs. Reporters were used as follows: pGL2-cyclin E-Luc containing 1.4 kb of cyclin E promoter; pGL2-cyclin B-luc containing 3.8 kb of the cyclin B promoter; and pGL2-cyclin A-Luc containing ~3.5 kb of cyclin A promoter (all these were gifts of E. Kerkhoff, University of Würzburg). As a control for transfection efficiency, ciferase activity was measured using a luminometer (Berthold). As an internal RT-PCR was performed using the SuperScript™ II kit (Invitrogen). Primers were used as follows: cyclin E, forward: 5’-GTGAAAGCGAGGATAGCAG-3’; reverse: 5’-TGTGTGATGCCATGTAACG-3’; cyclin B, forward: 5’-GCCACTGTAAAGCCCTACC-3’; reverse: 5’-GTCTGTGATGACCATC-3’; and cyclin A, forward: 5’-AAGGACCTTCCTTATAACG-3’; reverse: 5’-TCTCCACCTCAACCGC-3’. Cycling was performed as follows: for cyclin E and cyclin B; 5 min at 94°C for 1 cycle; 30 s at 94°C, 30 s at 53°C, 30 s at 72°C for 30 cycles; 10 min at 72°C for 1 cycle. For cyclin A, the annealing was performed at 50°C.

Co-immunoprecipitation experiments and Western blotting

Co-immunoprecipitations of Flag-SC1 and HDACs were performed as follows: HEK293 cells were transfected with either Flag-SC1 or Flag alone and HDAC 1, 2, or 3, and were harvested 48 h after transfection. Cells were lysed using RIPA buffer on ice and centrifuged to get rid of cell debris. Lysates were then precleared using protein A/G beads (Amersham Biosciences) and Flag-tagged proteins were precipitated using anti-Flag M2 Sepharose (Sigma-Aldrich). The beads were then washed with RIPA buffer 4–5 times and the immune complexes were separated by SDS-PAGE. Half of the reaction was run out on a 10% gel, blotted and probed for HDACs using anti-HA antibody (Santa Cruz Biotechnology, Inc.); another half was separated on a 7.5% gel, blotted and probed with anti-Flag M2 antibody (Sigma-Aldrich) to detect SC1. For Western blotting with anti-cyclin E antibody, we used a polyclonal antibody (Santa Cruz Biotechnology, Inc.); the anti-actin mouse monoclonal antibody was from Ambion. As a template for the in vitro transcription, a fragment of 1.2 kb corresponding to nucleotides 1–1200 of the rat SC1 sequence was subcloned in pCNA3 in both orientations. PC12 cells were transfected with SC1 siRNA or scrambled siRNA (0.5 μg/well) using LipofectAMINE™ 2000. 24 h later, cells were serum starved for 24 h. Serum-containing medium was added, and 24–48 h later, cells were collected directly in protein loading buffer and boiled for 5 min. Western blotting was performed to determine levels of SC1, cyclin E, cyclin A, and actin.

sRNA experiments

SC1 sRNA experiments was obtained using the Silencer™ siRNA Cocktail Kit (RNase II; Ambion). As a template for the in vitro transcription, a fragment of 1.2 kb corresponding to nucleotides 1–1200 of the rat SC1 sequence was subcloned in pCNA3 in both orientations. PC12 cells were transfected with SC1 siRNA or scrambled siRNA (0.5 μg/well) using LipofectAMINE™ 2000. 24 h later, cells were serum starved for 24 h. Serum-containing medium was added, and 24–48 h later, cells were collected directly in protein loading buffer and boiled for 5 min. Western blotting was performed to determine levels of SC1, cyclin E, cyclin A, and actin.

SC1 antibodies

Antibodies against SC1 were obtained from rabbits injected with the peptide GSTMTEGCRSMVSAWYSDELSLASHHC coupled to KLH. Antibodies were purified using an affinity column with the peptide coupled to CNBr Sepharose (Roche). Antibodies were used at 2 μg/ml for Western blotting.

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Immunocytochemistry

For the experiments using NIH3T3 and HEK293 cells, BrdU was purchased from Sigma-Aldrich and used according to a previously published protocol (Krek and DeCaprio, 1995). Cells were plated on sterile glass coverslips coated with poly-ornithine and were pulse for 2–3 h with 10 μM BrdU before fixation with cold 50:50 methanol/acetic for 2–3 min. Immunostaining was performed as suggested by the manufacturer of the anti-BrDU antibody (Zymogen). Immunostaining of labeled cells with a monoclonal anti-FLAG FITC-conjugated antibody (Upstate Biotechnology) to visualize SC1 expressing cells was performed according to the manufacturer’s instructions. Coverslips were embedded in Mowiol after washing. COS1 and Schwann cells were plated as above, except that poly-o-lysine was used for coating the coverslips. They were pulse for 24 h with 10 μM BrdU before processing for immunocytochemistry. The cells were fixed in 4% PFA in PBS. Primary anti-BrDU antibody from DakoCytomation was used, with subsequent visualization by application of anti-mouse rhodamine-conjugated secondary antibody from Jackson Immunoresearch Laboratories. Scans of stained cells were made using a confocal microscope (model TCS; Leica) for NIH3T3, COS1, and Schwann cells with identical settings for pinhole and voltage for any panel of analysis, and a fluorescent microscope (model IX70; Olympus) was used for HEK293 cells.

RT-PCR for cyclin-specific mRNA

RNA was extracted from the cells using TRIzol® (Invitrogen), according to the manufacturer’s instructions. RT-PCR was performed using the SuperScript™ II kit (Invitrogen). Primers were used as follows: cyclin E, forward: 5’-GGAGAGGCAGGATAGCAG-3’; reverse: 5’-GGTGTGATGCCATGTAACG-3’; cyclin B, forward: 5’-GCCACTGTAAAGCCCTACC-3’; reverse: 5’-GTCTGTGATGACCATC-3’; and cyclin A, forward: 5’-AAGGACCTTCCTTATAACG-3’; reverse: 5’-TCTCCACCTCAACCGC-3’. Cycling was performed as follows: for cyclin E and cyclin B; 5 min at 94°C for 1 cycle; 30 s at 94°C, 30 s at 53°C, 30 s at 72°C for 30 cycles; 10 min at 72°C for 1 cycle. For cyclin A, the annealing was performed at 50°C.

Plasmids

HEK293 cells and NIH3T3 cells were kept in DME supplemented with 10% FCS unless otherwise indicated. We used a subline of PC12 cells, PC6-3 (a gift of Jonathan Ham, Institute of Child Health, London, UK; Pittman et al., 1993). These were grown in DME with 10% horse serum and 5% FCS, except for the synchronization experiments and treatment with growth factors where low serum was used (1% horse serum and 0.5% FCS). Synchronization was done according to a published protocol (Rudkin et al., 1989) where cells were kept without serum for 3 d and then released from the cell cycle block by the addition of either 100 ng/ml NGF or serum. For synchronization of NIH3T3, we used a previously published method (Kerkhoff and Rapp, 1997). In brief, the cells were arrested by keeping them in low serum (0.5% FCS) for 48 h, and were released from growth arrest by the addition of full serum-containing medium. In the experiments with K252a, the latter was added to the cells at a final concentration of 0.2 μM for 30 min before NGF addition. Transfections were performed using the LipofectAMINE™ 2000 kit according to the manufacturer’s instructions (Invitrogen). 24-well plates were used for all the experiments, and the amount of DNA was kept constant, as suggested by the manufacturer. For TSA (Upstate Biotechnology) treatment of transfected cells, TSA was dissolved in DMSO and added directly to the transfected cells at a final concentration of 50 ng/ml. DMSO was added to mock-treated cells in control reactions. Cells were lysed in the luciferase assay buffer as suggested by the manufacturer (Promega), and luciferase activity was measured using a luminometer (Berthold). As an internal control for transfection efficiency, β-galactosidase-expressing plasmid (pCMVβ-gal) was cotransfected with the reporter and effector plasmids. The data were normalized to the β-galactosidase activity. Luciferase activity measurement was corrected for the transfection efficiency by calculating the ratio of luciferase units to β-galactosidase units, respectively as the relative luciferase units. Transfections of COS1 and primary Schwann cells were performed using the Effectene Kit (Qiagen) following the instructions of the vendor. Primary Schwann cells were isolated from P1 sciatic nerve according to a previously published protocol (Tikoo et al., 2000).

Cells lines, transfections, and reporter gene assays

Western Blotting

The expression levels of cyclin E and cyclin A were determined by Western blotting of cell lysates using anti-cyclin E antibody (Santa Cruz Biotechnology, Inc.) and anti-cyclin A antibody (Cell Signaling). As a loading control, an antibody recognition specific to actin was used (Zhao et al., 1999). After SDS-PAGE, proteins were transferred onto nitrocellulose membranes. The membranes were blocked for 30 min with 5% non-fat dry milk in TBS-Tween and probed with primary antibodies against cyclin E, cyclin A, or actin overnight at 4°C. The blots were then washed in TBS-Tween and incubated with horseradish peroxidase-conjugated secondary antibodies for 1 h at room temperature. Immunoreactive bands were visualized using the ECL plus detection system (Amersham Pharmacia Biotech). Densitometry analysis was performed using Molecular Dynamics software (Molecular Dynamics). Immunostaining of labeled cells with a monoclonal anti-FLAG antibody (Sigma-Aldrich) was detected using a polyclonal antibody (Sigma-Aldrich, Inc.); the anti-actin mouse monoclonal antibody was from DakoCytomation.
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