Identification of Genes That Are Downregulated in the Absence of the POU Domain Transcription Factor pou3f1 (Oct-6, Tst-1, SCIP) in Sciatic Nerve

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Despite the importance of myelinating Schwann cells in health and disease, little is known about the genetic mechanisms underlying their development. The POU domain transcription factor pou3f1 (Tst-1, SCIP, Oct-6) is required for the normal differentiation of myelinating Schwann cells, but its precise role requires identification of the genes that it regulates. Here we report the isolation of six genes whose expression is reduced in the absence of pou3f1. Only one of these genes, the fatty acid transport protein P2, was known previously to be expressed in Schwann cells. The LIM domain proteins cysteine-rich protein-1 (CRP1) and CRP2 are expressed in sciatic nerve and induced by forskolin in cultured Schwann cells, but only CRP2 requires pou3f1 for normal expression. pou3f1 appears to require the claw paw gene product for activation of at least some of its downstream effector genes. Expression of the novel Schwann cell genes after nerve injury suggests that they are myelin related. One of the genes, tramtorin1, encodes a novel amino acid transport protein that is localized to paranodes and incisures. Our results suggest that pou3f1 functions to activate gene expression in the differentiation of myelinating Schwann cells.

Key words: myelin; pou3f1; Oct-6; Tst-1; SCIP; claw paw; Schwann cells; tramtorin; dendrin; CRP1; CRP2; P2; representational difference analysis

Schwann cells generate myelin in the peripheral nervous system. One myelinating Schwann cell forms a single myelin sheath around a single axon, whereas nonmyelinating Schwann cells typically ensheathe several unmyelinated axons without myelinating any of them. Axons are thought to regulate the differentiation of Schwann cells; presumptive myelinated axons signal Schwann cells into the myelinating lineage, whereas the axons lacking this signal remain associated with nonmyelinating Schwann cells (Mirsksy and Jessen, 1996, 1999; Taylor and Suter, 1997; Garbay et al., 2000).

pou3f1 [also known as octomer-6 (Oct-6), Testes-1 (Tst-1), suppressed cAMP-inducible POU (SCIP), Octamer transcription factor-6 (Otf6)] is a member of the POU domain family of transcription factors (for review, see Ryan and Rosenfeld, 1997). pou3f1 is expressed transiently during Schwann cell development (Monuki et al., 1989), specifically in promyelinating Schwann cells, which have formed a 1:1 relationship with axons but have not yet formed a myelin sheath (Zorick et al., 1996; Arroyo et al., 1998). In the absence of pou3f1, the differentiation of myelinating Schwann cells is delayed, transiently arrested at the promyelinating stage (Bermingham et al., 1996; Jaegle et al., 1996). The role of pou3f1 in Schwann cell differentiation is unknown. It has often been considered to be a repressor of Schwann cell gene expression and/or differentiation. Cotransfection experiments suggest that pou3f1 inhibits the expression of the P0 (Mpz) myelin basic protein (Mbp), and p75 (low-affinity neurotrophin receptor) genes (Monuki et al., 1990, 1993; He et al., 1991). Furthermore, an N-terminal deletion of pou3f1 (ΔSCIP) acts as a dominant-negative inhibitor of pou3f1 repression of a 1.1 kb rat Mpz (P0) promoter in cotransfection experiments. In transgenic mice, ΔSCIP causes precocious PNS myelination, taken as evidence that pou3f1 represses the progression from promyelinating Schwann cells to myelinating Schwann cells (Weinstein et al., 1995). However, Schwann cells in pou3f1-null mice show delayed differentiation, and their expression of myelin-related mRNAs, including those of Mpz and Mbp, was not higher than in wild-type mice (Bermingham et al., 1996; Jaegle et al., 1996). These results contradict the idea that pou3f1 represses the expression of these genes in Schwann cells. To reconcile these observations, pou3f1 has been hypothesized to activate genes required for Schwann cell differentiation but to repress terminal differentiation (Mirsksy and Jessen, 1996; Zorick and Lemke, 1996; Jaegle and Meijer, 1998). In this scheme, pou3f1-null mice lack both functions, whereas ΔSCIP inhibits only the repression function.
Our approach sought to identify target genes of pou3f1 in Schwann cells to better understand the mechanisms by which it controls their differentiation. Representational difference analysis (RDA), a PCR-based technique that permits the isolation of DNA fragments that are present in one DNA sample but absent in another (Lisitsyn and Wigler, 1993; Lisitsyn, 1995), was used with a carrier RNA facilitating the isolation of differentially expressed genes (Erkman et al., 2000; Bermingham et al., 2001). We found six genes that are misexpressed in the sciatic nerves of pou3f1-null mice, and the expression of five of these genes correlates with the expression of other myelin-related genes after nerve injury. Because all of these mRNAs are expressed at lower levels in the sciatic nerves of pou3f1-null mice, pou3f1 appears to be an activator of gene expression in Schwann cells. The identification of these genes indicates that pou3f1 functions to induce membrane biogenesis, cytoskeletal rearrangement, and amino acid transport during the differentiation of myelinating Schwann cells.

MATERIALS AND METHODS

Isolation of RNA from cultured Schwann cells. Schwann cells were isolated from 3-d-old rat pups (Brookes et al., 1979) and expanded on 10 cm plates coated with poly-l-lysine in DMEM supplemented with 10% FCS, 2 µM forskolin (Porter et al., 1987), and 10 ng/ml recombinant human secreted β isofrom of neuregulin (glial growth factor-2). The cells were passaged six times, grown to confluency, and then maintained for 3 d in DMEM and 10% FCS. The cells were maintained for an additional 3 d in DMEM and 10% FCS alone or supplemented with 20 µM forskolin. All of the cultures used in these experiments were essentially free of fibroblasts. RNA was isolated from rat sciatic nerves and Schwann cells by CsCl2 gradient centrifugation (Chirgwin et al., 1979).

Isolation of RNA from sciatic nerves from lesioned rat sciatic nerves. Adult Sprague Dawley rats were anesthetized with 50 mg/kg pentobarbital. Sciatic nerves were dissected from hindquarters (Bermingham et al., 1996) for use on genomic RDA with Pou3f1. RPMI 1640 was supplemented with 20% FCS and 1 mM NaHCO3. All samples were maintained at 37°C in an incubator for 3 hr.

Sequence analysis and rapid amplification of cDNA ends. Twenty-four clones were sequenced for each experiment (96 total). Some clones contained multiple inserts. The sequences were compared with the GenBank nonredundant and expressed sequence tag (EST) databases using the National Center for Biotechnology Information (NCBI) basic local alignment search tool (BLAST) (blastn) program (www.ncbi.nlm.nih.gov/blast) (Altschul et al., 1997). Similar queries were performed using the Celera mouse genome assembly (www.celera.com). The sequence indicated which cDNAs were derived from the same gene and which were duplicates. Based on sequence, some inserts were not analyzed further, such as those corresponding to ribosomal RNAs (rRNAs) that were found in both of the samples. Clones 18 and 138 contained overlapping cDNAs of a novel gene, 18-138. 5′ and 3′ rapid amplification of cDNA ends (RACE) was performed with the SMARTrace kit (Clontech, Cambridge, UK) using cDNA that had been synthesized using Superscript II polymerase (Invitrogen) and P5-P2 Sprague Dawley rat sciatic nerve RNA.

Adapters/primers. The J, N, and R oligonucleotides were designed by Lisitsyn and Wigler (1995) for use on genomic RDA with Pou3f1 and are used here with DpnII. The K, O, and S primers are the corresponding adapters/primers designed to be used with NlaIII: K Sph 24 (RDA with NlaIII), 5′-GAC AAC CGA CGT CGA CTA TGC ATG-3′; K Sph 12 (RDA with NlaIII), 5′-CAT CGA CGT CGA CTA TGC ATG-3′; S Sph 24 (RDA with NlaIII), 5′-AGG CAA CGT TGC TAT CCG AGC ATG-3′; O Sph 12 (RDA with NlaIII), 5′-CTC GGA TAG CAC-3′; S Sph 24 (RDA with NlaIII), 5′-GGC ACT CTC CAG CCT CTC AGC ATG-3′ and S Sph 12 (RDA with NlaIII), 5′-CTG AGA AGC TGG 3′.

Sequence analysis and rapid amplification of cDNA ends. Twenty-four clones were sequenced for each experiment (96 total). Some clones contained multiple inserts. The sequences were compared with the GenBank nonredundant and expressed sequence tag (EST) databases using the National Center for Biotechnology Information (NCBI) basic local alignment search tool (BLAST) (blastn) program (www.ncbi.nlm.nih.gov/blast) (Altschul et al., 1997). Similar queries were performed using the Celera mouse genome assembly (www.celera.com). The sequence indicated which cDNAs were derived from the same gene and which were duplicates. Based on sequence, some inserts were not analyzed further, such as those corresponding to ribosomal RNAs (rRNAs) that were found in both of the samples. Clones 18 and 138 contained overlapping cDNAs of a novel gene, 18-138. 5′ and 3′ rapid amplification of cDNA ends (RACE) was performed with the SMARTrace kit (Clontech, Cambridge, UK) using cDNA that had been synthesized using Superscript II polymerase (Invitrogen) and P5-P2 Sprague Dawley rat sciatic nerve RNA.
was exponential, and parallel PCRs were pooled to minimize variation from individual samples and obtain sufficient material. Driver DNA (0.5 or 1 μg) was electrophoresed through 3% NuSieve 3:1 agarose (FMC Bioproducts, Rockland, ME) or 4% acrylamide gels and then transferred by conventional blotting or electrophoretic transfer onto Hybond N membranes (Amersham Biosciences, Arlington Heights, IL).

In situ hybridization. Mice were anesthetized and perfused with formalin. Hindquarters were dissected to expose the sciatic nerve and then stored at 4°C in formalin. Tissues were frozen in a 1:1 mixture of optimal cutting temperature compound (O.C.T.) and Aquamount and sectioned at 20 μm. In situ hybridization was performed as described by Simmons et al. (1989). Single-stranded antisense transcriptions were prepared from the following clones: pou3f1, AP700 (Bermingham et al., 1996); LpZ, JBSN3, which consists of 423 bp of sequence starting at the DnHY site at 2344 in V00296; Po, rat cDNA as described by Bermingham et al. (1996); cysteine-rich protein-2 (CRP2), JBSN61B, which contains 231 bp starting 361 bp between nucleotides 2856 and 3203 of X96589; tramdorin (tran3), 343 bp between nucleotides 1549 and 1892 of AF512429; and 21-70, a 673 bp fragment of a mouse RACE clone that contains the 3′ 222 nucleotides of JBSN70, all of JBSN21, and 114 nt of additional 3′ untranslated region sequence. For 18-138, a 185 bp DnHY RDA fragment (clone JBSN70) was used as a probe at 5× MOPS, pH 7.0, 1% SDS, and 0.017 mg/ml phenylmethylsulfonyl fluoride (Sigma), followed by a brief sonication on ice with a dismembrator (Fisher Scientific, Houston, TX). Protein concentration was determined using the Bio-Rad (Richmond, CA) kit according to the manufacturer’s instructions. For each sample, after a 5–15 min incubation in loading buffer at room temperature, 100 μg of protein lysate was loaded onto a 12% SDS-polyacrylamide gel, electrophoresed, and transferred to an Immobilon-polyvinylidene fluoride membrane (Millipore) over a period of 1 hr, using a semidry transfer unit (Fisher Scientific). The blots were blocked (5% powdered skim milk and 0.5% Tween 20 in Tris-buffered saline) overnight at 4°C and incubated for 24 hr at 4°C in a rabbit antiserum against tramdorin (diluted 1:1000). After washing in blocking solution and Tris-buffered saline containing 0.5% Tween 20, blots were visualized by enhanced chemiluminescence (Amersham Biosciences) according to the manufacturer’s protocols.

Immunostaining. Transfected cells were plated onto four chamber glass slides (Nalge; Nunc, Roskilde, Denmark) and incubated for 2–3 d to ~60% confluency. The cells were washed in PBS, fixed in acetone at −20°C for 10 min, and then blocked with 5% fish-skim gelatin in PBS containing 0.1% Triton X-100 for 1 hr at room temperature. Cells were labeled with rabbit anti-tramdorin1 (1:500) and processed as described below. Unfixed rat sciatic nerves were embedded in O.C.T. and immediately frozen in a bath of liquid nitrogen. Sections were thaw mounted on SuperFrost Plus glass slides (Fisher Scientific) and stored at −20°C. Teased nerve fibers were prepared from adult rat sciatic nerves, dried on SuperFrost Plus glass slides overnight at room temperature, and stored at −20°C. Sections and teased fibers were postfixed and permeabilized by immersion in −20°C acetone for 10 min, blocked at room temperature for 1 hr in 5% fish-skim gelatin containing 0.5% Triton X-100 in PBS, and incubated 16–48 hr at 4°C with various combinations of primary antibodies: rabbit anti-tramdorin1 (1:500), mouse anti-rat myelin-associated glycoprotein (MAG) (clone 513, 1:100; Boehringer Mannheim, Indianapolis, IN), and mouse anti-lysosome-associated membrane protein (LAMP) (1:10; Developmental Hydriddoma Bank). After incubating with the primary antibodies, the slides were washed and incubated with the appropriate fluorescein- and rhodamine-conjugated donkey cross-affinity-purified secondary antibodies (diluted 1:100; Jackson ImmunoResearch, West Grove, PA). Slides were mounted with Vectashield (Vector Laboratories, Burlingame, CA), examined by epifluorescence with tetramethylrhodamine isothiocyanate (TRITC) and fluorescein- and rhodamine- labeled secondary antibodies, and photographed with a cooled Hamamatsu (Tokyo, Japan) camera or followed by image manipulation with Adobe Systems (San Jose, CA) Photoshop.

RESULTS

RDA using pou3f1 +/+ and −/− sciatic nerve

To identify target genes of pou3f1 in peripheral nerves, sciatic nerves were isolated from newborn [postnatal day 0 (P0)] pups. Not only do most pou3f1-null mice die at this time (Bermingham et al., 1996; Jaegle et al., 1996), but P0 is near the peak of pou3f1 target genes expression of myelin gene. However, because the maturation of myelinating Schwann cell is delayed in pou3f1-null mice (Bermingham et al., 1996; Jaegle et al., 1996), comparing gene expression after the onset of myelin gene expression would be complicated by the expression of myelin-related genes (i.e., genes whose function or expression correlates with myelination) in the wild-type nerves.

RDA on pou3f1 −/− and +/+ sciatic nerves was performed as

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described previously (Erkman et al., 2000; Bermingham et al., 2001). Nerves were dissected from newborn pups of heterozygous pou3f1 intercrosses and saved separately for each animal. After the pups were genotyped, the nerves from 7 to 10 pou3f1 +/- and +/- pups were pooled separately, and total RNA was isolated from each pool. DNA was also isolated from the pooled nerves and used to confirm their genotypes (data not shown). Poly(A⁺) RNA was isolated from a mixture of 1 μg of total RNA and 150 ng of carrier RNA and used to generate cDNA. cDNAs from +/- and +/- nerves were digested separately with DpnII or NlaIII, ligated to adaptors, and amplified by PCR to generate driver DNAs (Fig. 1A). From each driver DNA, tester DNAs were derived by replacing adaptor sequences, after which four RDA experiments were performed. For each enzyme, +/- cDNA served as driver and +/- cDNA as tester, to isolate putative pou3f1-activated genes. Separately, +/- cDNA was used as driver and +/- cDNA as tester to isolate putative pou3f1-repressed genes. Figure 1A shows +/- and +/- drivers, as well as the first and second difference products, DP1 and DP2, generated from one and two rounds of the RDA procedure, respectively. Amplified DP2 DNA was cloned into plasmid vectors and, after transformation, 24 random colonies from each experiment (96 total) were sequenced.

The sequences were compared to the NCBI nonredundant and EST databases. This analysis revealed, consistent with our previous RDA results (Bermingham et al., 2001), that some clones contained multiple cDNAs, some cDNAs were isolated multiple times, some cDNAs represented different parts of the same gene, and some RDA fragments were homologous to EST clones that correspond to two different genes. A total of 114 RDA cDNA fragments were sequenced. Of these, 23 RDA fragments corresponded to 18S and 28S ribosomal RNA genes; these were found as perinatal myosin heavy chain and troponin C, probably introduced by contaminating muscle tissue in the +/- nerves, also were eliminated. We were left with 78 RDA fragments that corresponded to 51 different candidate pou3f1 target genes. To determine whether these candidate genes were differentially expressed, we performed Southern blot hybridization of 44 of these genes to blots of PCR-amplified cDNA from wild-type/pou3f1 +/- and mutant/pou3f1 +/- sciatic nerves (Table 1). In our hands, these Snorthern blots have proved to be reliable indicators of differential expression (Bermingham et al., 2001) (our unpublished observations).

As summarized in Figure 1B, 11 RDA fragments from eight genes were confirmed to be differentially expressed by Snorthern blots. Hybridization to JBSN18 was seen only in +/- cDNA; an identical result was seen with JBSN138 (data not shown), which is a different fragment of the same gene (see below). JBSN21 hybridized to a differentially expressed cDNA and, to a lesser degree, with two cDNAs that are not differentially expressed. JBSN61B (CRP2), JBSN64 (dendrin), and JBSN67 (myelin P0) are all differentially expressed, as is JBSN70. Clone JBSN125 contains two inserts, a 215 bp unknown sequence, which is not differentially expressed and serves as an internal control, and a differentially expressed cDNA fragment that is a fragment of tramdorin1 (see below). A subclone (125-10) that contains only the smaller cDNA hybridizes only to the differentially expressed fragment (data not shown). pou3f1 itself was not identified in the experiments presented here, presumably because of an insufficient number of clones that were analyzed. However, hybridization of a Snorthern blot to a mouse pou3f1 RDA fragment from brain demonstrates that pou3f1 is present in +/- but not in +/- cDNA, providing further confirmation of the genotype of the tissue from which the cDNA is derived. P0 and peripheral myelin protein 22 kDa (PMP-22) cDNA fragments were isolated but were not differentially expressed on Snorthern blots, consistent with their apparent normal mRNA expression in pou3f1-null mice (Bermingham et al., 1996). In summary, six genes, representing nine RDA fragments, were expressed at lower levels in the absence of pou3f1 and thus are putatively activated by pou3f1. Two genes, LacZ (Fig. 1B) and Neo (data not shown), were expressed at higher levels in pou3f1-null sciatic nerve. These genes replace pou3f1 in the knock-out mutation and demonstrate that we could have detected repressed genes; the absence of other such genes suggests that pou3f1 is not a repressor in Schwann cells. Finally, these results demonstrate that misexpressed genes can be isolated from small amounts of mutant tissue using an artificial mRNA as a carrier.

Of the six activated genes, only one was known previously to be expressed by Schwann cells: the fatty acid transport protein myelin P0 (Narayan et al., 1991). Two genes had been identified previously but were not known to be expressed by Schwann cells: the LIM-domain protein CRP2 [smooth muscle LIM (smLIM)] (Weiskirchen et al., 1995; Jain et al., 1996), which binds to components of the actin cytoskeleton (Louis et al., 1997), and dendrin, a protein of unknown function found in dendrites of CNS neurons (Neuner-Jehle et al., 1996; Herb et al., 1997). The subclone 125-10 (from JBSN125) corresponds to numerous EST clones, including AI786604 (I.M.A.G.E. clone 1920302), AI035402 (1380757), and AI005767 (1363993). As described below, we have named this gene tramd1. Gene 18-138 is represented by clone JBSN18 (DpnII-derived) and clone JBSN138 (NlaIII-derived) fragments that overlap; the full characterization of this gene will be published elsewhere. RDA clones JBSN21 and JBSN70, which showed limited or no homology to sequences currently in the public databases, have been shown by RACE experiments to correspond to adjacent DpnII fragments of a novel putative extracellular matrix gene, 21-70 (data not shown).

**Schwann cells express putative pou3f1 target genes**

To determine whether the cDNAs that appeared to be differentially expressed on Snorthern blots are in fact misexpressed in vivo, in situ hybridization was performed with antisense cRNA probes. Figure 2A shows adjacent sections of P0 sciatic nerves from pou3f1 +/- and +/- mice. As expected, pou3f1 mRNA was expressed in +/- but not in +/- nerves. LacZ mRNA was expressed in the opposite pattern, and P0 mRNA was abundantly expressed in both genotypes (Bermingham et al., 1996). In contrast to P0 mRNA, P2 mRNA was more abundant in +/- nerves than in +/- nerves. Dendrin, 18-138, and CRP2 mRNA were all present in +/- nerve, but the signals in +/- nerve were not above background levels. Minimal expression was detected for 21-70, and no signal was observed using a sense cRNA probe from clone 18-138 or for tramdorin1 (data not shown).

To determine whether these changes persisted in older nerves, we also performed in situ hybridization on sciatic nerves from Pou3f1 +/- and +/- mice (Fig. 2B). As expected (Monuki et al., 1989), pou3f1 mRNA expression in Pou3f1 +/- sciatic nerves was...
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Figure 1. Isolation of genes that are misexpressed in sciatic nerve in the absence of pou3f1. A. Photograph of an ethidium bromide-stained acrylamide gel containing driver, DP1, and DP2 cDNA from P0 pou3f1 +/+ and −/− sciatic nerves. The sizes of the cDNAs can be estimated from the ∆X174 HaeIII digest (lane M). RDA was performed in parallel using cDNAs that were digested with either DpnII or NlaIII. Faint bands, presumably attributable to ribosomal RNA or other highly repetitive RNAs, can be seen in the lanes containing the driver cDNA. Note the absence of any band derived from carrier RNA (584 bp). After one round of RDA (DP1 lanes), specific bands, many of similar sizes, begin to appear in both the +/+ and −/− lanes, with either DpnII- or NlaIII-digested cDNA. After two rounds of RDA, the +/+ and −/− lanes contain distinct banding patterns (DP2 lanes). DP2 DNAs were cloned into pBSKII + for analysis. In DP3 cDNA, only a few bands that correspond to Neo and LacZ increase in intensity relative to DP2 cDNA (data not shown). 

B. Southern blots of PCR-amplified cDNA (Southern blots) from pou3f1 +/+ and −/− sciatic nerves after hybridization to different radiolabeled cDNA probes. 

Expression also is reduced in P1.5 clp −/− nerves, but to a lesser degree compared with P1. In wild-type nerves, CRP2 expression is robust at P1.5 but is greatly reduced at P13. In clp mutant nerves, CRP2 activation is reduced, but its expression at P13 is less obvious. In clp mutant nerves, CRP2 activation is reduced, but its expression at P13 is elevated, consistent with the delay in myelination. CRP2 expression is reduced in P1.5 clp −/− sciatic nerves but recovered somewhat by P13. Dendrin mRNA was detected in P9 +/+ nerves but not in −/− nerves. The mRNA level of 18-138 was increased in both +/+ and −/− nerves between P0 and P9, but the levels were lower in P9 −/− nerves. Expression of CRP2 mRNA was comparable in P9 +/+ and −/− nerves, but compared with its expression at P0, its expression had decreased in P9 +/+ nerve and increased in P9 −/− nerve (see below). Expression of 21-70 mRNA was faint but greater in P9 +/+ nerve than in P0 +/+ nerve and greater in +/+ than in −/− nerve. Myelin P2 and 18-138 expression is reduced in P1.5 clp −/− sciatic nerves, whereas LacZ mRNA is unaffected, although the nerves are thinner in clp mutant mice. These results demonstrate that delayed expression of the genes that we have identified correlates with delayed peripheral myelination in clp as well as pou3f1.
which multiple independent cDNAs were isolated for these genes (either the same or different sequences corresponding to that gene), their numbers are also indicated.

of genes and cDNA fragments that were isolated, tested, and confirmed are putatively repressed by pou3f1. Not all genes were analyzed (see Results), and for some genes, multiple cDNA fragments were isolated; for each experiment, the number of genes isolated by RDA are misexpressed in pou3f1 mutant nerves are shown after in situ hybridization with antisense probes for pou3f1, LacZ, P0 (Mpz), P2 (P2), dendrin, 18-138, CRP2, and 21-70 mRNA. A, P0 sciatic nerves. The left panels show wild-type nerves (+/+); the right panels show knock-out (−/−) nerves. pou3f1 hybridizes to +/+ but not to −/− nerves, whereas LacZ is the opposite; these results confirm the genotypes of the tissues. Hybridization to mRNA for P0, whose expression is independent of pou3f1, is observed in both +/+ and −/− sciatic nerves and serves to demarcate them. Myelin P2 and CRP2 are expressed in +/+ sciatic nerve, but in pou3f1 mutant nerve, little or no signal is observed. Dendrin and 18-138 hybridization produce faint signals in +/+ and no signals in −/−. At this age, 21-70 mRNAs were not detected. These results demonstrate that differences in expression can be detected by in situ hybridization, and that the genes isolated by RDA are misexpressed in pou3f1 mutant nerves in vivo. B, P9 sciatic nerves. The same probes that were hybridized to P0 sciatic nerves were used to assay expression in P9 sciatic nerves. pou3f1 and LacZ expression confirm the genotypes of the tissues, although by P9, pou3f1 expression is declining, whereas LacZ expression in mutants remains high, as has been observed previously (Jaegle and Meijer, 1998). Myelin P0 mRNA expression confirms no significant differences in P0 expression in +/+ and pou3f1 −/− mice. At P9, myelin P2 and 18-138 expression in mutant sciatic nerve is much stronger than at P0 but remains reduced relative to wild type. Dendrin expression is present in +/+ nerves, but there is little or no signal from mutant nerves. CRP2 expression appears equivalent in +/+ and −/− at this time, expression is declining in +/+ nerves but is probably increasing in −/− nerves. Faint 21-70 expression can be observed in +/+ nerves but not in −/− nerves. Scale bar, 0.5 mm.

Table 1. Genes isolated by RDA of pou3f1 +/+ versus −/− sciatic nerve

<table>
<thead>
<tr>
<th>A. Increased expression in +/+</th>
<th>Isolated-tested-confirmed</th>
<th>Confirmed genes</th>
</tr>
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<tbody>
<tr>
<td>DpnII-digested cDNA</td>
<td>28-26-10 cDNAs</td>
<td>Unknown gene 18–138 (4 copies)</td>
</tr>
<tr>
<td></td>
<td>21-19-5 genes</td>
<td>Unknown gene 21–70 (3 copies, 2 different adjacent DpnII fragments)</td>
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<td></td>
<td></td>
<td>CRP2 (D88792)</td>
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<td></td>
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<td>Dendrin (X96589)</td>
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<td></td>
<td></td>
<td>P2 (S39508)</td>
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<tr>
<td>NlaIII-digested cDNA</td>
<td>20-13-4 cDNAs</td>
<td>Unknown gene 18–138 (3 copies)</td>
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<tr>
<td></td>
<td>12-8-2 genes</td>
<td>Unknown gene 125-10: Tran dorin 1</td>
</tr>
<tr>
<td>B. Increased expression in −/−</td>
<td>Isolated-tested-confirmed</td>
<td>Confirmed genes</td>
</tr>
<tr>
<td>DpnII-digested cDNA</td>
<td>29-20-12 cDNAs</td>
<td>LacZ V00296 (7 copies, 4 different fragments)</td>
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<tr>
<td></td>
<td>14-12-2 genes</td>
<td>pMC1neo U43612 (5 copies)</td>
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<tr>
<td>NlaIII-digested cDNA</td>
<td>38-21-6 cDNAs</td>
<td>LacZ V00296 (5 copies, 2 different fragments)</td>
</tr>
<tr>
<td></td>
<td>21-14-2 genes</td>
<td>pMC1neo U43612</td>
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The results of four independent experiments are summarized using cDNA from pou3f1 +/+ and −/− nerves, digested with either DpnII or NlaIII. Clones with increased expression in pou3f1 +/+ sciatic nerve refer to experiments in which tester cDNAs were derived from wild-type sciatic nerve; these genes are putatively upregulated by pou3f1. Likewise, clones with increased expression in pou3f1 −/− sciatic nerve refer to experiments in which tester cDNAs were derived from pou3f1 −/− sciatic nerve; these genes are putatively repressed by pou3f1. Not all genes were analyzed (see Results), and for some genes, multiple cDNA fragments were isolated; for each experiment, the number of genes and cDNA fragments that were isolated, tested, and confirmed is indicated. The genes that were shown to be differentially expressed are listed, and in the cases in which multiple independent cDNAs were isolated for these genes (either the same or different sequences corresponding to that gene), their numbers are also indicated.

Figure 2. Expression of novel Schwann cell genes in pou3f1 +/+ and −/− sciatic nerves at P0 and P9. Sections of sciatic nerves are shown after in situ hybridization with antisense probes for pou3f1, LacZ, P0 (Mpz), P2 (P2), dendrin, 18-138, CRP2, and 21-70 mRNA. A, P0 sciatic nerves. The left panels show wild-type nerves (+/+); the right panels show knock-out (−/−) nerves. pou3f1 hybridizes to +/+ but not to −/− nerves, whereas LacZ is the opposite; these results confirm the genotypes of the tissues. Hybridization to mRNA for P0, whose expression is independent of pou3f1, is observed in both +/+ and −/− sciatic nerves and serves to demarcate them. Myelin P2 and CRP2 are expressed in +/+ sciatic nerve, but in pou3f1 mutant nerve, little or no signal is observed. Dendrin and 18-138 hybridization produce faint signals in +/+ and no signals in −/−. At this age, 21-70 mRNAs were not detected. These results demonstrate that differences in expression can be detected by in situ hybridization, and that the genes isolated by RDA are misexpressed in pou3f1 mutant nerves in vivo. B, P9 sciatic nerves. The same probes that were hybridized to P0 sciatic nerves were used to assay expression in P9 sciatic nerves. pou3f1 and LacZ expression confirm the genotypes of the tissues, although by P9, pou3f1 expression is declining, whereas LacZ expression in mutants remains high, as has been observed previously (Jaegle and Meijer, 1998). Myelin P0 mRNA expression confirms no significant differences in P0 expression in +/+ and pou3f1 −/− mice. At P9, myelin P2 and 18-138 expression in mutant sciatic nerve is much stronger than at P0 but remains reduced relative to wild type. Dendrin expression is present in +/+ nerves, but there is little or no signal from mutant nerves. CRP2 expression appears equivalent in +/+ and −/− at this time, expression is declining in +/+ nerves but is probably increasing in −/− nerves. Faint 21-70 expression can be observed in +/+ nerves but not in −/− nerves. Scale bar, 0.5 mm.
mutant mice. In contrast, pou3f1 expression appears normal in P1.5 clp−/− nerves but fails to be downregulated at P13. Therefore, expression of pou3f1 is insufficient for activation of its putative target genes, indicating either that pou3f1 regulates these genes indirectly or that it requires a cofactor to activate them.

Axotomy results in a prompt decline in the levels of myelin-related mRNAs distal to the lesion, including those of P0, MBP, P2, connexin32, PMP-22, myelin-associated glycoprotein, and periaxin (Poduslo, 1993; Scherer et al., 2001). To determine whether CRP2, tramd1, 18-138, and dendrin are expressed in parallel with other myelin-related genes, we performed Northern blot analysis on rat sciatic nerves collected at various times after crush or transection (Fig. 4). For comparison, the blot was also hybridized for CRP1, pou3f1, P0, p75, and glyceraldehyde-3-phosphate dehydrogenase (GAPDH). The transections were performed to prevent axonal regeneration, whereas nerve crush allows axonal regeneration, and many of the regenerating axons are remyelinated in a proximal-to-distal manner. To better illustrate these points, the distal stumps of crush-injured nerves were divided into proximal (D1) and distal (D2) segments. Tramd1, 18-138, and dendrin mRNA levels paralleled that of P0; their mRNA levels fell after transection and did not increase thereafter, whereas their mRNA levels returned after crush, first in the D1 segment (at 24 d) and then in the D2 segment (at 58 d). The expression of CRP2 mRNA, in contrast, was remarkably similar to that of pou3f1, being low in unlesioned adult nerves and increasing after nerve crush but not transection. These data indicate that tramd1, 18-138, and dendrin are expressed in myelinating Schwann cells and are expressed as part of the program of myelin-related gene expression.

Expression of CRP genes in Schwann cells
CRP2 mRNA expression parallels that of pou3f1 both during development as well as after nerve injury. In situ hybridization of CRP2 mRNA at P0, P1.5, P9, and P13 (Figs. 2, 3) indicates that CRP2 expression is transient, like that of pou3f1 (Monuki et al., 1989). To evaluate the relationship between CRP2 and pou3f1 further, we examined CRP2 mRNA expression in cultured Schwann cells and observed that CRP2 is induced by forskolin (Fig. 5A), as is pou3f1 (Monuki et al., 1989). These results further indicate that CRP2 expression closely mimics that of pou3f1 and that CRP2, like pou3f1, may be expressed in promyelinating Schwann cells (Arroyo et al., 1998). To determine whether CRP2 performs an essential role in Schwann cell myelination, we examined sciatic nerves from P4 and P5 mice that lack the CRP2 gene (kindly provided by S.-F. Yet and M.-E. Lee, Harvard Medical School). We found no differences in myelin from these mice and their wild-type littermates in epoxy sections (data not shown), suggesting that CRP2 is not essential or that other CRP genes may compensate for the absence of CRP2. To test the latter possibility, we examined the expression of CRP1, which is closely related (Liebhaber et al., 1990; Crawford et al., 1994). CRP1 is also induced by forskolin, but unlike CRP2, its expression is not regulated by pou3f1 (Fig. 5B) or nerve injury (Fig. 4). As potential regulators of the actin cytoskeleton, the CRP proteins may act...
in the changes in Schwann cell morphology that accompany myelination.

**Tissue-specific expression of putative pou3f1 target genes**

The expression of dendrin, 18-138, and tramdorin1 mRNAs was analyzed on a Northern blot of adult mouse tissues to determine their tissue distribution and the sizes of their transcripts. A dendrin probe hybridized to a 4 kb mRNA in adult cerebrum, as expected (Neuner-Jehle et al., 1996; Herb et al., 1997), and more weakly to a similarly sized mRNA from sciatic nerve (Fig. 6A). The 18-138 probe generated a strong hybridization signal with a 1.9 kb mRNA and a weaker signal with a 4.7 kb transcript (Fig. 6B). Of the tissues examined, 18-138 mRNA was detected only in sciatic nerve. The major tramdorin1 transcripts in sciatic nerve were 2.2 and 2.7 kb; the 2.7 kb species is the most prominent tramdorin1 mRNA in lung, adrenal gland, and thymus (Fig. 6C). No expression was observed in the spleen, but a faint band (>6.5 kb) was observed in cerebrum. These results indicate that EST cDNA 1920302, as well as the RACE cDNAs, define a nearly full-length tramd1 mRNA that likely corresponds to the larger 2.7 kb transcript, because they are ~2.45 kb without their poly(A) tails. Therefore, tramdorin1 expression is neither ubiquitous nor restricted to sciatic nerve. A similar Northern blot analysis of rat tissue showed expression in sciatic nerve but not in lung and thymus (data not shown), suggesting either variation in tramdorin1 expression among species or cross-hybridization of the tramd1 probe to related sequences.

**Chromosomal localization of putative pou3f1 target genes**

The chromosomal locations of all six putative pou3f1 target genes were examined to ascertain any linkage to known human or mouse peripheral neuropathy loci. Dendrin has been mapped to distal mouse chromosome 15 (Brady et al., 1997), and myelin P2 (Pmp2) has been mapped to proximal mouse chromosome 3 (Dietrich et al., 1996). The mouse chromosomal locations of the CRP2 (Csrp2), 18-138, 21-70, and tramd1 genes were determined by interspecific backcross analysis using progeny derived from [(C57BL/6J × Mus spretus)F1 × C57BL/6J] mice (N. A. Jenkins, D. Gilbert, and N. Copeland, unpublished observations). The location of Csrp2 in mice near Myf6 on chromosome 10 confirms its previous mapping to human to 12q21.1 (Weiskirchen et al., 1997) (data not shown). 18-138 resides near the Lag locus on distal mouse chromosome 4, in a region that corresponds to human 1q36. 21-70 is closely linked to the Atm and Csk loci on mouse chromosome 9; the corresponding human locus is on 15q21. The chromosomal locations of these genes were confirmed using Celera and public mouse and human genome databases. Currently, there are no peripheral neuropathy loci for...
which dendrin, myelin P₂, CRP2, 18-138, or 21-70 are candidate loci.

The results of backcross mapping of the tramdl locus confirm its localization to 11 exons residing between nucleotides 56,775,171 and 56,748,143 on mouse chromosome 11 in the Celera mouse chromosome database (May 5, 2002 update; our unpublished observations; data not shown). The proximal region of mouse chromosome 11 is syntenic with human chromosome 5q31–33, and the homology of mouse tramdorin1 coding exons to 5q sequences in the Celera and public human genome assemblies (Lander et al., 2001; Venter et al., 2001) confirms the mapping in mouse. Although no candidate mouse mutations reside near the tramdl locus, kindreds with an autosomal recessive form of inherited demyelinating neuropathy have been mapped to this region (LeGuern et al., 1996; Gabreels-Festen et al., 1999; Guilbot et al., 1999a,b), prompting us to characterize this gene further.

**Figure 6.** Tissue-specific expression of dendrin, 18-138, and tramdorin1 genes. Each lane contains an equal amount (10 μg) of total RNA isolated from the indicated mouse tissues. The blot was successively hybridized to a radiolabeled probe corresponding to dendrin (A), 18-138 (B), and tramdorin1 (C) and exposed to film for 8, 7, and 14 d, respectively, after each hybridization. The sizes of the transcripts are estimated according to the sizes of 28S and 18S rRNAs (4712 and 1869 nt, respectively) (Hassouna et al., 1984; Raynal et al., 1984). Sciatic N., Sciatic nerve.
cDNAs homologous to clone 125-10 encode a novel gene product, tramdorin1

Rat and mouse cDNAs that correspond to clone 125-10 were isolated by 5′ and 3′ RACE rat cDNAs, a 3′ RACE rat cDNA, and a mouse EST cDNA. Of these, the 2.45 kb EST cDNA 1920302 appears to include the entire open reading frame, which is depicted in black. This cDNA appears to be nearly full length, based on the calculated size of tramd mRNA (Figs. 3, 7). The organism and tissue of origin of each cDNA fragment are listed. The open reading frame commences with an ATG initiation codon (in bold) that closely matches the Kozak consensus (Kozak, 1987); nucleotides that are an exact match are underlined. 14dpc, 14 d postcoitum.

B. Amino acid sequences encoded by mouse tramdorin EST cDNA 1920302 and a hypothetical rat tramdorin1 cDNA, derived from 5′ and 3′ RACE sequences, are shown. These sequences have been assigned GenBank accession numbers AF512429 and AF512430, respectively. The mouse cDNA encodes a 478 aa protein, whereas the corresponding rat protein is 481 aa. The amino acid sequence of mouse EST cDNA 1920302 was analyzed using the transmembrane domain prediction programs Memsat2 (McGuffin et al., 2000) and TMHMM (Sonnhammer et al., 1998). Eleven putative transmembrane domains are numbered. Transmembrane domains predicted by TMHMM are marked with capital Ts. For Memsat2, transmembrane domains are marked as follows: O, Outside transmembrane helix cap; X, central transmembrane helix segment; I, inside transmembrane helix cap. For both Memsat2 and TMHMM, predicted cytoplasmic domains are marked by +, whereas noncytoplasmic domains are marked by −. Five consensus glycosylation sites within the protein sequence are shown in bold and are boxed. C. Diagram depicting the predicted topology of tramdorin1 with 11 transmembrane domains, as predicted by Memsat2 and TMHMM. The three extracellular/lumenal glycosylation sites are marked with branched structures. However, the electrophoretic mobility of tramdorin suggests that it is not glycosylated extensively (Fig. 8).

The protein encoded by mouse cDNA 1920302 was predicted by the Memsat2 (PSIPRED) (McGuffin et al., 2000) and transmembrane hidden Markov model (TMHMM) (Sonnhammer et al., 1998) protein structure prediction programs to consist of 11 transmembrane helices, with the N terminus facing the cytoplasm (Figs. 7B, C). Although other programs produced differing predictions (data not shown), the following considerations support this topology. First, those programs that predicted 11 transmembrane domains more accurately predicted the number and topology of the transmembrane helices of a set of transmembrane proteins of known structure (Tusnady and Simon, 2001). Second, a signal sequence prediction program (Nielsen et al., 1997) indicates that tramdorin1 does not appear to contain a cleaved N-terminal signal sequence that would be expected if the N terminus was lumenal/extracellular. Third, membrane proteins are glycosylated only on their extracytoplasmic faces, and the distribution of glycosylation consensus sites (NXS/T, where X is...
any amino acid except proline) within the protein supports the proposed topology. Five putative glycosylation sites were found, of which three are predicted to be extracytoplasmic in the structures predicted by Memsat2 and TMHMM. A fourth is not conserved in humans (our unpublished observations), and a fifth is buried in predicted transmembrane helix 5. Fourth, the protein displays homology to the amino acid and anxin permease superfamily of amino acid transport proteins, which are predicted to possess 11 transmembrane domains (Young et al., 1999). However, the related vesicular GABA transporters are thought to consist of 10 transmembrane domains that correspond to predicted transmembrane domains 2–11 (McIntire et al., 1997; Sagne et al., 1997). Regardless of their differing topological predictions, all of the programs indicated that cDNA 1920302 encodes a polytopic transmembrane protein. Therefore, we have named the protein the polytopic transmembrane protein. The tramdo1 gene has been assigned the human and mouse gene symbol tramd1. The known proteins that are most closely related to tramdo1 are two additional tramdoins: tramdo2 and tramdo3 (our unpublished observations). While this manuscript was in preparation, tramdo1 was isolated independently and shown to encode a proton-dependent amino acid transporter (Boll et al., 2002). A recently cloned rat lysosomal amino acid transporter (LYAAT), LYAAT-1 (Sagne et al., 2001), is the rat homolog of tramdo3; rat tramdo1 is 67% identical to this protein.

To characterize tramdo1, an antiserum was raised against a sequence near the N terminus that was divergent among the tramdo1 gene family (see Materials and Methods). In immunoblots of adult rat sciatic nerve, the antiserum binds to a single band close to the predicted molecular weight of unglycosylated tramdo1 (Fig. 8A). A band of similar molecular weight is observed on immunoblots of cells transfected with full-length mouse tramdo1 cDNA (Fig. 8B). Transfected cells were stained with the anti-tramdo1 antiserum, whereas parental cells were unstained (Fig. 8C,D). To localize tramdo1, we labeled unfixed myelinated fibers from rat sciatic nerves with this antiserum. Tramdo1 immunoreactivity was observed in the paranodes and in most incisures, as shown by colabeling with a monoclonal antibody against MAG (Fig. 9A–C). To determine whether tramdo1, like tramdo3/LYAAT-1, is associated with lysosomes, teased fibers were double labeled for tramdo1 and LAMP1, a lysosomal marker. Figure 9D–F shows that tramdo1 and LAMP1 are found in paranodes but do not colocalize. These observations indicate that tramdo1 is localized to noncompact myelin but is not a component of lysosomes.

**DISCUSSION**

To identify candidate target genes of pou3f1, we used RDA in conjunction with an artificial carrier mRNA to isolate genes that are differentially expressed in sciatic nerves from newborn pou3f1−/− pups compared with their +/+ littermates. The sequences of the RDA fragments led to the discovery of six candidate genes, each of which was verified to be misexpressed by Snorthern blots and *in situ* hybridization. Furthermore, the mRNA levels of five of six of these genes paralleled other myelin-related genes in nerve-injury paradigms. These genes are good candidates to fulfill important functions in generating or stabilizing the myelin sheath or in the metabolism of Schwann cell proteins.

**Figure 8.** Immunoblot analysis of tramdo1. A, B, Western blot analysis of anti-tramdo1 antiserum. Lysates (100 μg) from adult rat sciatic nerve (sciatic N; A) and 293 cells (B) transfected with tramdo1 expression vector (+) and untransfected 293 cells (−) were used for Western blot analysis of the tramdo1 antiserum (diluted 1:1000); the blots were developed with chemiluminescence. Arrowheads mark the location of immunoreactive tramdo1 protein. C, D, Micrographs showing transfected (C) or untransfected (D) 293 cells viewed by immunofluorescence using the anti-tramdo1 antiserum. Nuclei were visualized using 4',6'-diamidino-2-phenylindole.

**Misexpressed genes in pou3f1 mutant mice**

CRP2 is the only putative gene whose mRNA levels parallel those of pou3f1 in developing as well as lesioned adult peripheral nerves and in cultured Schwann cells treated with forskolin. Their parallel expression patterns strongly support the idea that CRP2 is a direct target of pou3f1. In keeping with this suggestion, the CRP2 promoter (Yet et al., 1998) has seven consensus pou3f1-binding sites within 4.8 kb of the transcription start site (data not shown). However, pou3f1 expression is insufficient to activate CRP2 in clp mutant mice, suggesting that if CRP2 is directly activated by pou3f1, this activation also requires the clp gene product. Although absence of CRP2 does not appear to affect Schwann cell myelination, it is a second gene whose expression marks the promyelinating stage of Schwann cell differentiation.
The related protein CRP1 is induced in Schwann cells by forskolin treatment, also expressed in adult sciatic nerve, but its expression is not modulated by nerve injury or dependent on pou3f1. CRP1 and CRP2 (smLIM) are members of the cysteine-rich protein subclass of LIM-only proteins, which consist of two LIM domains, each followed by a glycine-rich region (Liebhaber et al., 1990; Arber et al., 1994; Crawford et al., 1994; Jain et al., 1996). All three members of this subclass are associated with actin fibers, bind to the actin cytoskeletal proteins α-actinin and zyxin (Louis et al., 1997), and are thought to function in the differentiation of the cells that express them. CRP1 and CRP2 are downregulated in transformed cells (Weiskirchen and Bister, 1993; Weiskirchen et al., 1995), and mice that lack the third member of the family, CRP3/MLP, display a disruption of the cytoarchitecture of differentiated muscle cells (Arber et al., 1997). Thus, if CRP2 and CRP1 perform similar functions in Schwann cell differentiation, the loss of CRP2 may be compensated for by CRP1. These proteins may participate in cytoskeletal changes that accompany the transition from a promyelinating Schwann cell to a myelinating Schwann cell (Kidd et al., 1996; Fernandez-Valle et al., 1997; Bernier et al., 1998).

Myelin P2 is a member of the fatty acid binding protein (FABP) family of cytosolic lipid binding proteins (Narayan et al., 1988), which are thought to function by binding and transporting fatty acids and/or by modulating fatty acid-mediated signal transduction (Glatz et al., 1995; Coe and Bernlohr, 1998). Adipocyte P2 (aP2), the most closely related FABP to myelin P2, activates the nuclear receptor transcription factor peroxisome proliferator activated receptor (PPAR)γ, which activates the aP2 gene in a positive feedback loop that controls lipid metabolism in adipocytes (Coe and Bernlohr, 1998; Hertzel and Bernlohr, 1998). If myelin P2 performs an analogous function to aP2 in Schwann cells, then lipid metabolism and perhaps PPAR-mediated gene regulation may be compromised in pou3f1−/− Schwann cells. In addition, myelin P2 has been shown to bind retinoic acid and retinol (Uyemura et al., 1984), suggesting that it could modulate the activities of retinoid nuclear receptors. The myelin P2 promoter (Narayanan et al., 1991; Bharucha et al., 1993) contains four consensus pou3f1-binding sites (data not shown), located in sequences that are required for forskolin induction of the myelin P2 gene and high-level expression transient transfection assays. Together, these observations suggest that myelin P2 is a target of pou3f1, and that forskolin induction of myelin P2 (Monuki et al., 1989) may be mediated by pou3f1.

As its name suggests, dendrin was originally found in dendrites (Neuner-Jehle et al., 1996; Herb et al., 1997). It is a highly basic protein with putative phosphorylation sites for PKA and protein kinase C, suggesting that phosphorylation regulates its function. It is one of several proteins thought to be translated in dendrites using internal ribosome entry sites (Pinkstaff et al., 2001), suggesting that its translation could be localized to discrete locations in Schwann cells as well. Our data indicate that dendrin is expressed by myelinating Schwann cells, but its function both in dendrites and in Schwann cells remains to be determined.

Tramdorin1/mPAT2 has been shown to function as a proton-dependent transporter of small amino acids (Boll et al., 2002). It is related to yeast vacuolar and rat lysosomal amino acid transport proteins, but unlike tramdorin3/LYAAT-1, tramdorin1 does not appear to be a lysosomal protein. Tramdorin1 lacks a possible lysosomal targeting motif [an acidic residue located near the C terminus, −4 to −6 relative to a dileucine (LL[I,M,V]) (Sandoval et al., 2000) that is present in tramd3/LYAAT-1. Tramdorin1/ mPAT2 shows strong specificity for glycine, L-alanine, and L-proline, with greatest inward currents generated by glycine (Boll et al., 2002). Its restricted substrate specificity and localization suggest an unsuspected function of glycine in myelinating Schwann cells.

**Figure 9.** Localization of tramdorin1 in myelinating Schwann cells. Images of unfixed teased fibers from adult rat sciatic nerve, double labeled with a rabbit tramdorin1 antiserum (A, D) (TRITC) and a mouse monoclonal antibody against MAG (B) (FITC) or LAMP1 (E) (FITC) are shown. Tramdorin immunoreactivity is found at paranodes (arrows) and most incisures (arrowheads), colocalizing with MAG (A–C), and in puncta along the outer surface. The failure to detect tramdorin 1 in all incisures is more likely the result of technical problems of antibody penetration than of heterogeneity of expression. At paranodes (D–F), tramdorin1 immunostaining does not appear to colocalize with that of LAMP1, a lysosomal marker. Both markers possess cone-like expression patterns in the paranode, in which the tramdorin expression domain is nested within the LAMP1 expression domain.
Regulation of Schwann cell differentiation by pou3f1

Several observations indicate that the genes we have identified are not regulated exclusively by pou3f1. First, pou3f1 is expressed in P1.5 clp mutant sciatic nerves, whereas dendrin, myelin P2, CRP2, and 18-138 are not, suggesting that factor(s) in addition to pou3f1 are required for normal activation of those genes. The gene product that is altered by the clp mutation is unknown; directly or indirectly, this gene product could act in concert with pou3f1 or downstream of it. Second, only the expression of CRP2 appears to parallel that of pou3f1; the other genes normally are expressed after the decline of pou3f1 expression. It is possible that pou3f1 indirectly activates these genes through its activation of the distal enhancer of Krox-20 (Ghislain et al., 2002). In fact, two of the putative pou3f1 target genes are also activated by Krox-20. Dendrin is one of the most strongly activated genes by overexpression of Krox-20 in Schwann cells (Nagarajan et al., 2001), and myelin P2 expression is reduced or absent in krox-20-null mice (Topilko et al., 1994). However, the transient expression pattern of CRP2 suggests that this gene may be regulated independently of Krox-20. Third, the genes that we have identified appear to be activated slowly in the absence of pou3f1, in conjunction with the delayed onset of myelination (Bermingham et al., 1996; Jaegle et al., 1996). The cause of delayed maturation of myelinating Schwann cells in pou3f1-null mice is unknown, but compensation by another POU protein is possible. In skin, pou3f1 and a member of a different class of POU domain protein, Skn-1, compensate for one another in keratinocyte differentiation (Andersen et al., 1997), indicating that divergent POU domain transcription factors can regulate pou3f1 target genes. The POU domain transcription factor Brm5 is a good candidate for such a compensatory factor, because it is expressed in myelinated Schwann cells (Wu et al., 2001).

Our results suggest that pou3f1 functions primarily as an activator of gene expression in Schwann cells. All six genes that we identified were upregulated by pou3f1. The two repressed genes that we found, Neo and LacZ, indicate that such genes could have been found, and we were able to identify genes that were repressed by forskolin using the same approach (Bermingham et al., 2001). It is possible that repressed genes were missed or that pou3f1 acts as a repressor in Schwann cells at times other than P0 or in other tissues. However, the simplest explanation for our data, and for the phenotype of the pou3f1-null mice, is that pou3f1 is an activator. Because the POU domain of pou3f1 itself can function as an activator (Fyodorov and Denerer, 1996), we suggest that the ∆SCIP transgene (Weinstein et al., 1995) also functions as an activator in vivo, triggering precocious myelination by prematurely activating pou3f1 target genes.

The sample of RDA clones that we analyzed does not contain all of the genes that are misexpressed in the absence of pou3f1. By looking at P0, before most myelin-related genes are highly expressed, we may have reduced the number of genes that we identified. For example, pou3f1 is required for its own downregulation (Jaegle and Meijer, 1998) (Fig. 2), but this effect is not apparent at P0. The Krox-20 distal enhancer has been shown to be activated by pou3f1 (Ghislain et al., 2002), but we did not isolate any Krox-20 clones, perhaps because of confounding effects of its expression from the early, pou3f1-independent enhancer. We did not find other genes that are known or likely to be differentially expressed by myelinating Schwann cells, such as periaxin, dystroglycan-related protein-2, connexin32, MAG, neurofascin, β4 integrin, and E-cadherin, although most of their cDNAs have appropriately spaced restriction sites for DpnII and NheIII.

Either these genes were missed or their expression does not require pou3f1.

Only a subset of myelin-related genes appears to be downregulated in the absence of pou3f1. The mRNA levels of P0, MBP, PMP-22, and MAG do not appear to be altered significantly in the first 10 d after birth by the absence of pou3f1 (Bermingham et al., 1996) (our unpublished results). Of the genes that were known previously to be expressed by myelinating Schwann cells, only myelin P2 was identified as a putative pou3f1 target gene. The results presented here suggest that myelination involves the coordinated expression of more genes than has been appreciated previously. The use of RDA screens to identify genes that are downregulated in the absence of pou3f1 is a fruitful approach to identify novel genes that may perform important functions in myelination. Until microarrays that contain every mouse or human gene are available, RDA remains a useful way to identify novel genes from specialized tissues, such as sciatic nerve.

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