Fluorescent Proteins Expressed in Mouse Transgenic Lines Mark Subsets of Glia, Neurons, Macrophages, and Dendritic Cells for Vital Examination


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To enable vital observation of glia at the neuromuscular junction, transgenic mice were generated that express proteins of the green fluorescent protein family under control of transcriptional regulatory sequences of the human S100B gene. Terminal Schwann cells were imaged repetitively in living animals of one of the transgenic lines to show that, except for extension and retraction of short processes, the glial coverings of the adult neuromuscular synapse are stable. In other lines, subsets of Schwann cells were labeled. The distribution of label suggests that Schwann cells at individual synapses are clonally related, a finding with implications for how these cells might be sorted during postnatal development. Other labeling patterns, some present in unique lines, included astrocytes, microglia, and subsets of cerebellar Bergmann glia, spinal motor neurons, macrophages, and dendritic cells. We show that lines with labeled macrophages can be used to follow the accumulation of these cells at sites of injury.

Key words: neuromuscular junction; Schwann cell; microglia; Langerhans cell; astrocyte; S100β; S100B; motor neuron; variegation; Bergmann glia

Introduction

Fluorescent proteins of the green fluorescent protein (GFP) family have revolutionized our ability to image cells in living organisms (Chalfie et al., 1994). By coupling nucleotide sequences encoding fluorescent proteins to cell-specific promoters, transgenes have been constructed that make specific cell types in living organisms fluorescent. As a consequence of incomplete promoters or random integration, expression of these transgenes can be limited to subsets of a particular cell type (cf. Feng et al., 2000) or occur in unexpected cell types. Because this variegation is generally consistent within animals of the same transgenic line, a small number of labeled cells of a particular type can be examined in each line. For example, variegation in the expression of fluorescent proteins driven by the thy 1.1 promoter has enabled repeated visualization of individual neurons in the cortex of mice and provided information about the dynamism of synapses in the adult brain (Grunzendorfer et al., 2002; Trachtenberg et al., 2002; Mizrahi et al., 2004). Similarly, expression of different colored fluorescent proteins in subsets of motor neurons has enabled detailed study of the competition occurring at neuromuscular junctions (NMJs) during neonatal synapse elimination (Walsh and Lichtman, 2003).

Terminal Schwann cells (TSCs), glial cells that cap the nerve terminal at the NMJ, play roles in maintaining and modulating this synapse (Robitaille, 1998; Reddy et al., 2003) and facilitate its repair after nerve injury (Son and Thompson, 1995a,b; Koizala et al., 2000). Vital imaging of nerve terminals and postsynaptic receptors in muscles has been successfully used to obtain new insights into the stability and plasticity of NMJs (Rich and Lichtman, 1989; Balice-Gordon and Lichtman, 1990, 1993, 1994; Walsh and Lichtman, 2003), and we wished to expand such observations to include TSCs. Other investigators have used antibodies or lectins that bind to cell surface components of Schwann cells and/or the extracellular matrix they deposit (Astrow et al., 1998) or have labeled Schwann cells with AM esters of fluorescent compounds that, for unknown reasons, give selective labeling of Schwann cells and/or the extracellular matrix they deposit (Astrow et al., 1998) or have labeled Schwann cells with AM esters of fluorescent compounds that, for unknown reasons, give selective labeling of Schwann cells and/or the extracellular matrix they deposit. To circumvent these problems, we...
prepared transgenic mice expressing fluorescent proteins in their Schwann cells. The protein S100B, previously known as S100\(\beta\), is a commonly used marker for glial cells (cf. Ludwin et al., 1976; Stefanson et al., 1982), and elements of its promoter have been characterized and shown to drive expression in stably transfected glial cells \textit{in vitro} (Alloire et al., 1990; liang et al., 1993; Castets et al., 1997). We coupled human-S100B upstream sequences to sequences coding for fluorescent proteins and prepared transgenic mice. A number of stable lines of mice were generated that selectively label TSCs. We show here that we can repeatedly image these cells in living, adult mice and that TSCs, like the nerve terminals they cover, are mostly stable. In addition, some of the lines showed labeling of subsets of Schwann cells. We used these subset lines to examine the mix of labeled and unlabeled Schwann cells at individual junctions. The nonrandom distribution of labeled cells suggests how Schwann cells come to populate junctions during development. Last, we have examined other cells labeled in these mice. We find that some of the expression patterns are predicted by the endogenous expression of S100B, whereas others are not. These transgenic lines offer the opportunity to study in a vital manner particular subsets of glia, neurons, and antigen-presenting cells of the immune system.

Materials and Methods

\textbf{Construction of transgenes and generation of transgenic mice.} Using a combination of restriction endonuclease fragment isolation and PCR amplification, 9.4 kb of human S100B regulatory sequence was inserted upstream of the enhanced GFP (EGFP) coding region in pEGFP-1 (Clontech, Cambridge, UK; BD Biosciences, Franklin Lakes, NJ) or enhanced yellow fluorescent protein (EYFP) coding region in pEYFP-1 (Clontech). In both cases the vectors were modified by alteration of the AflII site at the end of the SV40 polyadenylation module to a Pmel site. The human sequences include 6.6 kb of 5′ flanking sequence, the first noncoding exon of 71 nt and 2.7 kb of intron 1, including the 3′ splice acceptor site. The \(\sim 10.4 \) kb transgene was isolated from vector backbone sequences by digestion with restriction endonucleases SacII and Pmel and gel-purified. Transgenic mice were generated using standard techniques previously described (Hogan et al., 1994). Briefly, C57BL/6 X DBA (F2) zygotes were injected with the transgenes and subsequently transferred into the oviducts of pseudopregnant CD1 females. After genotyping (or phenotyping in the case of the YFP mice), resulting founders were bred to C57BL/6 X DBA (F1) mice to establish and maintain lines.

Our first attempts used an S100-GFP transgene. By Southern blotting and PCR, six potential founders were identified from a total of 40 animals injected as embryos, and five of these were found to have germline integration of the transgene. Animals of one of these five F1 lines showed no detectable expression of GFP, and this line was discarded. Offspring of the remaining four founders showed GFP expression in Schwann cells located in the intramuscular nerves and overlying NMJs. Of these, the two lines showing the most intense labeling of Schwann cells were selected. Because these two lines gave identical labeling of the brain and periphery, one of them, designated “Kosmos,” was used for further experiments, and all the other S100-GFP lines were discarded.

A second set of S100-YFP transgenic animals was prepared, using exactly the same construct but with YFP substituted for GFP. Nine of 45 injected embryos brought to term were judged to be potential founders on the basis of Southern blotting and PCR. All nine of these animals had germline integration. Two of the lines were discarded on the basis of dim YFP expression in Schwann cells in muscles. We have analyzed the remaining seven lines and designated them YA, YB, YD, YE, YF, YG, and YH. With the exception of one line (YD), all lines have been bred to homozygosity.

\textbf{Screening of transgenics.} Transgenic founders from the initial series of injections were initially identified by Southern blot analysis of tail DNA with radiolabeled Klenow fragments for GFP sequences and by PCR using primers specific for the EGFP sequence (forward, ATG GTG AGC AAG GGC GAG GAG CT; reverse, TCG TTT GGG TCT TGT CTC AGG GC, yielding a 641 bp product). Founders from a second series of injections of the S100-YFP transgene were identified by the fluorescence detectable in the lens of the eye and in skin (Fig. 1A). The number of transgenes integrated in the four S100-GFP lines, estimated using Southern blotting, was six (Kosmos), four, 12, and three.
Repetitive in vivo imaging. Vital imaging of 2- to 4-month-old Kosmos mice was conducted using the methods of Lichtman et al. (1987). Briefly, each animal was anesthetized with ketamine and xylazine (0.10 – 0.15 ml of a 0.9% NaCl solution containing 17.4 mg/ml ketamine and 2.6 mg/ml xylazine). A ventral midline incision in the neck exposed the sternomastoid muscle. Acetylcholine receptors (AChRs) were labeled by application of Alexa-594- or rhodamine-conjugated α-bungarotoxin (Molecular Probes, Eugene, OR), 5 μg/ml in sterile, lactated Ringer’s solution applied for 5 min. The animal was intubated endotracheally, and its respiration was driven by a small rodent respirator (model 683; Harvard Apparatus, Holliston, MA). A number 0 coverslip held orthogonal to the optical axis of the microscope was applied to the top surface of the muscle and used to depress slightly the muscle surface to minimize movement and to flatten the field. Images were collected using a Zeiss (Oberkochen, Germany) Axiopt microscope equipped with a 100 W mercury lamp, 40× [numerical aperture (NA) 0.75] or 63× (NA 0.9) water immersion objectives, and a MicroMAX 5 MHz cooled CCD camera attached to a Macintosh computer running IP Lab software. Throughout the examination, the intensity of illumination reaching the preparation was reduced 50–90% using neutral density filters. Filter cubes were inserted into the light path to visualize GFP and rhodamine in turn. An Endow GFP filter set (Chroma Technology, Brattleboro, VT) was used for GFP. Images of the superficial junctions were captured using 2 × 2 binning. A similar procedure was followed to image junctions in the soleus muscle. In these cases, the soleus was exposed by incision through the lateral surface of the shank and retraction of the gastrocnemius muscle. Intubation was unnecessary for these experiments. After imaging, the incision was closed with 6–0 silk suture, and the animal was allowed to recover from anesthesia under a heat lamp and then returned to its cage.

Immunostaining of whole mounts and sections. After anesthesia, animals were perfused transcardially with PBS. For examination of the brain, the animal was also perfused with 4% paraformaldehyde. Brains were dissected and postfixed overnight. For examination of peripheral tissues, these tissues [sternomastoid, extensor digitorum longus (EDL), soleus, diaphragm, and triangularis sterni muscles; sciatic, peroneal, and tibial nerves; intestinal, thymus, spleen, and skin] were pinned into Sylgard-coated dishes and fixed in the same fixative for 30 min. For microscopic observation and immunostaining, tissues were observed as whole mounts (procedures are described in Love and Thompson, 1998) or as ~10 μm cryostat sections, prepared after cryoprotection (20% sucrose) and embedding in OCT (TissueTek, Miles, Elkhart, IN).

The procedures for immunostaining have been described (cf. Love and Thompson, 1998). The following primary antibodies were used at the indicated dilutions: rabbit anti-cow S100B (1: 500; Dako, High Wycombe, UK); mouse monoclonal to neurofilament protein and synaptic vesicle protein (2H3 and SV2, respectively; 1:500; Developmental Studies Hybridoma Bank, University of Iowa, Iowa City, IA); anti-glial fibrillary acidic protein (GFAP; G8398; 1:100; Sigma, St. Louis, MO); anti-NeuN (MAB377; 1:100; Chemicon, Temecula, CA); Rip antibody (1:50; Developmental Studies Hybridoma Bank); F/4/80 rat monoclonal (MCA497R; 1:10–1:50; Serotec, Oxford, UK); FA-11 rat monoclonal (anti-macrosialin-CD68; MCA1957S; 1:10; Serotec); and hamster biotinylated monoclonal anti-CD11c and anti-CD3ε (553800 and 553239, respectively; 1:100; PharMingen, San Diego, CA). The following secondary antibodies and fluorophore reagents were used at the indicated dilutions: FITC-conjugated goat anti-rabbit (55671; 1:400; Cappel, Cochranville, PA); Alexa-594-conjugated goat anti-rat, anti-rabbit, and anti-mouse (A-11007, A-11037, and A-11020, respectively; 1:200 – 500; Molecular Probes); Cy5-conjugated goat anti-rabbit (111-175-144; 1:200; Jackson ImmunoResearch, West Grove, PA); tetramethylrhodamine isothiocyanate (TRITC)-conjugated goat anti-mouse (1070-03; 1:200; Southern Biotechnology); Texas Red donkey anti-rat and anti-rabbit (712-076-153 and 711-075-152, respectively; 1:100; Jackson ImmunoResearch); and Texas Red–streptavidin (SA100; 1:100; Caltag, San Francisco, CA).

AChRs were visualized using α-bungarotoxin conjugated to rhodamine, Alexa-594, or Alexa-647 (all obtained from Molecular Probes).

Nonvital imaging. Images were collected using a Leica epifluorescence microscope equipped with 40× (NA 1.0) and 63× (NA 1.32) objectives, filter cubes for 4’,6’-diamino-2-phenylindole (DAPI), GFP, YFP, TRITC, calcium crimson, and Cy5 (Chroma Technology), and an integrating CCD camera (Cool Snap HQ; Roper Scientific). Confocal imaging was performed using a Leica TCS-4. The AChR area within individual junctions was computed from an image of the bungarotoxin label as described previously (Love and Thompson, 1998).

Physiology. Soleus muscles and nerves were dissected and tested as described previously (cf. Lubischer and Thompson, 1999).

Averages in this manuscript are reported as ± SEM.

Results

A 9.4 kb fragment of genomic DNA encompassing the promoter region and first intron of the human S100B gene was inserted upstream of the coding region for a fluorescent protein (FP, either EGFP or EYFP) and the SV40 polyadenylation module. The constructs were used to generate 13 lines of S100-GFP and S100-YFP mice, eight of which have been retained and analyzed (see Materials and Methods). One line expresses GFP and was designated Kosmos; the remaining seven express YFP and were designated YA, YB, YE, YF, YG, and YH. In the course of screening these animals, we found that those bearing the transgene could be easily distinguished from nontransgenic littermates by examination under a fluorescence dissecting scope (Fig. 1 A).

GFP is a reliable marker of Schwann cells in the Kosmos transgenic line

As expected from the endogenous expression of S100B, myelinating Schwann cells were easily detected by their GFP fluorescence in intramuscular nerves (Fig. 1 B) or in small bundles of axons teased from the sciatic nerve. To examine GFP expression in TSCs at the NMJ, sternomastoid muscles were dissected from the Kosmos line and labeled with an antibody to S100 that marks Schwann cells (Fig. 1 E) and with α-bungarotoxin to identify the AChRs concentrated at each junction (Fig. 1 B). Almost without exception, every Schwann cell labeled by the S100 antibody was also labeled with GFP (Fig. 1, compare D, E). Of 197 junctions examined in sternomastoid muscles from four animals, there were only two junctions (one cell in each) in which there was any doubt of colabeling. The brightly fluorescent ovals revealed by anti-S100 and GFP were shown to be cell bodies by using a nuclear marker (DAPI; data not shown). Processes extending from these cell bodies covered the nerve terminal (identified by labeling with antibodies to neurofilament and synaptic vesicle proteins; data not shown). These processes could be visualized as well by their GFP fluorescence as by their S100 labeling. In addition to TSCs, GFP labeled the myelinating Schwann cells (Fig. 1 D, dou-
ble arrowhead) wrapping the motor axon (data not shown) that entered each junction. Identical GFP labeling was obtained regardless of whether the sternomastoid, soleus, EDL, diaphragm or triangularis sterni muscles were examined. We conclude that, in the Kosmos line, GFP is a reliable marker of Schwann cells. Such GFP labeling was present at the time of birth (data not shown).

**Junctions with fluorescent Schwann cells have normal morphology and physiology**

Transgenic expression of GFP in mammals is generally nontoxic (cf. Feng et al., 2000), but other studies have identified detrimental effects (cf. Huang et al., 2000). Although mice of the Kosmos line are seemingly normal in appearance, behavior and fertility, even when homozygous for the transgene, we were particularly interested in whether there were changes in the physiology and morphology of their NMJs.

To examine the morphology of transgenic NMJs, muscles were dissected from three transgenic Kosmos animals and three nontransgenic littermates at 2–3 months of age, fixed, and AChR and Schwann cells were labeled with different fluorophores. There were differences among the muscles (sternomastoid, soleus, and EDL) in the total area of their AChR-rich synaptic sites (490 ± 170 μm² in EDL to 695 ± 288 μm² in sternomastoid) and the number of Schwann cells per site (2.3 ± 0.8 in EDL to 3.3 ± 0.5 in sternomastoid). However, there were no significant differences in these parameters between transgenic and nontransgenic animals. At both transgenic and nontransgenic NMJs, TSCs precisely covered the nerve terminal.

To examine neuromuscular transmission and the organization of motor units, soleus muscles were dissected with their nerves from transgenic (n = 5) and nontransgenic (n = 2) animals, pinned into a dish, and superfused with oxygenated Ringer’s solution. Each muscle was connected to a tension transducer, and nerve stimulation, tetanic tension generated in response to stimulation of the nerve at 50 or 100 Hz, or tension produced by direct stimulation of the muscle (data not shown).

By these criteria, there are no detectable differences in the innervation of muscles of Kosmos despite the expression of soluble GFP in their Schwann cells.

**Vital imaging of Kosmos neuromuscular junctions shows the stability of the glial components of this synapse**

To assess the feasibility of vitally imaging Schwann cells at living NMJs, we exposed the sternomastoid muscle in anesthetized mice of the Kosmos line, labeled the AChR with nonblocking doses of rhodamine-conjugated bungarotoxin, and used an epifluorescence microscope to visualize receptor areas and the brightly labeled Schwann cells. We collected an initial set of images of 30 junctions that could be viewed en face in five animals at 2 months of age. Thirty days later we attempted to reimage these same junctions, identifying them on the basis of the unique, pretzel-like shape of their AChR (Fig. 2). We were successful in 25 cases; the remaining five had become situated on the sides of their muscle fibers. Most of the reimaged junctions (18) showed no major change in the position or number of Schwann cell bodies. Five of the junctions had added a Schwann cell body, and two had apparently lost a Schwann cell body. The synaptic AChRs were unchanged in all but three cases where there was a minor (<5%) loss of receptor area (Fig. 2, compare A, D).

The lost receptor areas were located where the motor axon entered the junction. Such losses of receptor areas have been previously observed by vital imaging (Balice-Gordon and Lichtman, 1993) and were suggested to result from loss of a functioning synaptic contact at these sites, perhaps because of encroachment of myelinating glia into previously synaptic areas. These experiments show the feasibility of performing repeated, vital imaging of Schwann cells at neuromuscular synapses. Moreover, they add to previous demonstrations that adult, mammalian NMJs have stable receptor areas and nerve terminals (Lichtman et al., 1987) by showing that this stability also applies to the associated glial cells.

**Vital imaging shows terminal Schwann cells extend and retract short processes**

During repeated, vital imaging of junctions in the sternomastoid and soleus muscles, we encountered TSCs that had extended short processes (Fig. 2G) (average length = 1.1 ± 0.1 μm, 66 processes) beyond the edge of the AChR (data not shown). These processes were present at approximately half of the junctions (42 of 90 in 32 sternomastoid muscles; 56 of 98 in 16 soleus muscles) and were much shorter than the numerous, longer extensions found after denervation (Son and Thompson, 1995a). To determine whether these short extensions might have been induced by the surgery or imaging, we examined sternomastoid and soleus muscles taken acutely from Kosmos or from nontransgenic animals, and we identified Schwann cells in the latter case by labeling...
We examined more fully the labeling of Schwann cell subsets in Schwann cells were labeled in some of the lines (Table 1, Fig. 3). Cells could be detected at birth (see below). Motor neurons, whereas other muscles in the same animals had labeled, although the size of the subset varied (Table 1). In addition, the labeling of motor neurons appeared to be minimal in some lines, in other lines only small subsets of motor neurons were labeled, although the size of the subset varied (Table 1). We first consider the labeling encountered at the first viewing. These observations suggest that Schwann cells extend and retract short processes in the environment around the synaptic site.

In some lines, subsets of Schwann cells are labeled and nonrandomly distributed to NMJs

Other transgenic lines we generated had labeling in some unexpected cell types (Table 1). We first consider the labeling encountered in the peripheral nerves and muscles, then other organs, and finally the CNS. Mice of each of the YFP lines were found to have some labeled motor axons and motor nerve terminals, either with or without labeling of Schwann cells (Fig. 3A, B). In these cases, the labeling of motor neurons could be confirmed in sections of the spinal cord (data not shown). Because spinal motor neurons are not labeled with antibodies to S100B, this YFP-labelling is an artifact of transgenesis in these lines. Although the majority of motor neurons appeared to be labeled in some lines, in other lines only small subsets of motor neurons were labeled, although the size of the subset varied (Table 1). In addition, muscles of mice from some of the lines had many labeled motor neurons, whereas other muscles in the same animals had none. The labeling of motor axons appeared late in postnatal development [postnatal day 7 (P7)], although expression in glial cells could be detected at birth (see below).

Besides the unexpected labeling of motoneurons, subsets of Schwann cells were labeled in some of the lines (Table 1, Fig. 3B). We examined more fully the labeling of Schwann cell subsets in.

Table 1. FP expression among the S100-FP lines

<table>
<thead>
<tr>
<th>S100-FP line</th>
<th>FP expression</th>
<th>Schwann cells</th>
<th>Motor axons</th>
<th>Muscle macrophages</th>
<th>CNS</th>
</tr>
</thead>
<tbody>
<tr>
<td>Kosmos</td>
<td>All, in all muscles examined</td>
<td>None</td>
<td>Very few</td>
<td>Astrocytes; Bergmann glia</td>
<td></td>
</tr>
<tr>
<td>YA</td>
<td>Subsets, + in so, ts, di; + + in st, EDL; weak expression in so</td>
<td>Subsets, none in di, so; + + in st, EDL, ts</td>
<td>Few</td>
<td>Microglia with few or weak astrocytes; Subset of Bergmann glia</td>
<td></td>
</tr>
<tr>
<td>YB</td>
<td>Weak expression in subsets. + in so, di; + + in st, EDL, ts</td>
<td>Subsets, + in so, di; + + in st, EDL, ts</td>
<td>Few–many</td>
<td>Bright microglia, dim astrocytes; Bergmann glia</td>
<td></td>
</tr>
<tr>
<td>YD</td>
<td>Subsets, + + in st, di, ts, so; + + + in EDL; weak expression in so and di</td>
<td>Subsets, + + in st, EDL, di; + + + in ts</td>
<td>Some–many</td>
<td>Not examined</td>
<td></td>
</tr>
<tr>
<td>YE</td>
<td>Subsets, + + in ts, di, st; + + + in so, EDL; weak expression in so and di</td>
<td>Subsets, None in ts, so, di; + + in st, EDL</td>
<td>Few–many</td>
<td>Bright microglia, dim astrocytes; Bergmann glia</td>
<td></td>
</tr>
<tr>
<td>YF</td>
<td>All, in all muscles examined</td>
<td>Subsets, None in ts, so, di; + + in st, EDL</td>
<td>Few</td>
<td>Bright microglia, dim astrocytes; Bergmann glia</td>
<td></td>
</tr>
<tr>
<td>YG</td>
<td>All, in all muscles examined</td>
<td>Subsets, None in di, so; + + in st, ts, EDL</td>
<td>Few</td>
<td>Bright microglia, dim astrocytes; Bergmann glia</td>
<td></td>
</tr>
<tr>
<td>YH</td>
<td>Weak expression in subsets. none in so, di; + + in st, EDL, ts</td>
<td>Subsets, + + in di, so; + + in st, EDL, ts</td>
<td>Some–many</td>
<td>Bright microglia, dim astrocytes; Bergmann glia</td>
<td></td>
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Sizes of subsets are indicated as + + + for most, + + for some, and + for few of cells present. For each line, two to eight muscles and one to four brains were examined. st, Sternomastoid; so, soleus; EDL, extensor digiforum longus; ts, triangularis sterni; di, diaphragm.
individual junctions was also observed in one YH sternomastoid (observed 75%, predicted 34%). The differences in observed versus predicted homogeneity in YA animals would be expected by chance in <5% of the cases (paired t test; n = 3 YA muscles), making them statistically significant. It is interesting to note that the YFP status (+ or −) of TSCs was apparently mirrored by the YFP status of the myelinating Schwann cell forming the last internode before the junction. A total of 125 junctions and their axons were analyzed in this manner in two YA sternomastoid muscles. For 50 of these junctions, YFP was present in both the Schwann cell forming the last internode and the TSCs; for 72 of them, YFP was present in neither; only three of them had no label in the Schwann cell on the last internode but some YFP+ TSCs.

One possible explanation for this nonrandom distribution of YFP+ and YFP− cells is that YFP is expressed in response to local influences exerted by the particular motor axon or muscle fiber with which TSCs associate. Bundles of axons teased from the sciatic or tibial nerves of YA animals showed that myelinating Schwann cells also had this variegation: some were YFP+, and some were YFP− (data not shown). Thus, the variegation occurs some distance from the peripheral targets of these axons, making it unlikely that a muscle-derived signal is controlling YFP expression. Moreover, it was possible in some cases to observe single axons teased from peripheral nerves over several internodes. In these cases, axons were observed that had one or more internodes wrapped by YFP+ Schwann cells while an adjacent internode was wrapped by a YFP− Schwann cell (Fig. 3E). Because the YFP expression along single axons can be nonuniform, it appears unlikely that axons directly YFP expression in Schwann cells.

YFP expression in Schwann cells could be stable, or it could change over time. We addressed this issue in two ways. First, we vitally imaged the TSCs at 39 NMJs in the sternomastoid muscles of four YA 3-month-old mice as described above. Of these 39 NMJs, 17 had YFP+ TSCs, and most of these also had YFP+ axons, whereas the remaining 22 junctions had only axonal labeling. Thirty days later we reimaged these same NMJs, and we found that all of the junctions retained their original labeling pattern (Fig. 5), with one exception. In this single case, a junction that had YFP+ TSCs on the initial viewing had added an additional YFP+ TSC by the second viewing. Thus, YFP expression in TSCs appears stable, at least over an interval of 30 d in adult animals. As a second approach, we attempted to determine whether there is an obvious switch in YFP expression during development. We observed that YFP+ and YFP− TSCs were present at sternomastoid NMJs at the time of birth (Fig. 3C–D). Therefore, the variegation seen in the adult does not emerge from uniformity present at birth. If Schwann cells do switch their expression pattern postnatally, this change must occur coordinately in groups of cells, otherwise mixtures of cell types would be more common.

Homogeneous YFP expression in the TSC population at individual NMJs could arise during development by sorting of SCs during the process of neonatal synapse elimination, because each junction loses synaptic inputs to become innervated by a single motor axon (Brown et al., 1976). We examined three sternomastoid muscles dissected from three YA animals on P6, using the same methods outlined above for adult muscles. Fewer TSCs were present at P6 junctions (an average of three, compared with approximately four in the adult), a finding consistent with reports that Schwann cells are added to junctions during postnatal development (Love and Thompson, 1998). The percentage of S100+ TSCs that were YFP+ in these muscles was 66, 60, and 58%. Based on a random assortment of Schwann cells according to their YFP phenotype, the expected frequencies of homogenous junctions were 33, 28, and 27% (Fig. 4B), significantly lower than the observed 90, 87, and 88% (< 0.001; n = 3; paired t test). Thus, TSCs at individual junctions tend to be of the same YFP status at P6, although half the junctions receive input from more than one motor axon at this time (Balice-Gordon and Lichtman, 1993). This homogeneity in Schwann cells and its possible interpretation is discussed below.
Resident macrophages in muscles and peripheral nerves express YFP and respond to injury in several lines

In addition to expression in Schwann cells and axons, the YA, YD, YE, YF, and YH lines had prominent YFP labeling within a population of cells scattered among muscle fibers, peripheral nerve fibers, and their connective tissues (Fig. 6A,B). These cells had short processes that were usually oriented along the axons or muscle fibers. These cells were not labeled by anti-S100 antibody (Fig. 6C,D), and thus represent another example of apparently aberrant expression of the transgene. Such labeled cells were particularly prominent in the triangularis sterni and diaphragm. Because these YFP + cells resemble “resident” macrophages known to be present within tissues (cf. Gordon et al., 1992), we immunostained sections of muscle and nerve with antibodies to the murine macrophage markers F4/80 and FA-11 (Gordon et al., 1992) (supplemental Fig. 1, available at www.jneurosci.org as supplemental material). In sections of the EDL muscles and plantar nerves of YH mice, >80% of the YFP + cells were labeled by these antibodies (data not shown). However, some cells positive for the macrophage markers were not YFP +. We conclude that the YFP label in some of our lines serves as a marker for a subset of resident macrophages in muscles and nerves, as it does for lymphoid tissues (see below). All the FP lines we generated have at least an occasional, rare FP + cell located outside the NMJ that is obviously not a Schwann cell, but the frequency of such labeled cells is dramatically higher in the YE and YH lines and moderately higher in the YA, YD, and YF lines. Thus, there is considerable line-to-line variability in the extent of labeling of macrophages, suggesting that macrophage subsets are labeled in the different lines.

Because macrophages are known to accumulate at sites of tissue damage (cf. Monaco et al., 1992), we tested whether damage to superficial muscle fibers or to the sciatic nerve would lead to accumulation of YFP + macrophages. One day after nerve crush, the density of YFP + cells at the crush site was at least five times as high as that at sites approximately a millimeter away (data not shown). Six days after resection of the sciatic nerve, YFP + cells had filled the nerve distal to the site of resection (data not shown). Such YFP + cells were still present 14 d later. Most of these cells appeared to be filled with phagocytic vacuoles (supplemental Fig. 1C,D, available at www.jneurosci.org as supplemental material). Two days after injury to muscle fibers (cutting a few superficial fibers with iridectomy scissors), YFP + cells accumulated at the injury site and along individual, degenerating fibers on each side of the injury site. Confocal imaging showed rounded, YFP + cells were present on the surface and inside these fibers (Fig. 6E,F). Thus, the YA, YE, and YH lines could be used for the vital imaging of macrophages at sites of tissue injury.
bright fluorescence, cells could also be seen in other internal organs (e.g., lungs, heart, liver). Fluorescent cells were also encountered in blood smears and were occasionally seen moving through capillaries during the vital imaging of NMJs. Examination of cryostat sections of thymus, spleen, and intestine from a mouse of the YF line showed that some dendritic cells (marked by anti-CD11c) (supplemental Fig. 2, available at www.jneurosci.org as supplemental material) and some macrophages (marked by antibody F4/80; data not shown) expressed YFP. YFP+ dendritic cells were encountered in the anticipated locations: in the cortex and medulla of the thymus (data not shown) and as interdigitating cells in T-cell-containing cryptopatches, B- and T-cell-containing Peyer’s patches, and the lamina propria of the small and large intestines (data not shown). B- and T-cells were YFP− in all locations (data not shown). Dendritic cells have been previously reported to express S100B (Cocchia et al., 1981, 1983; Michetti et al., 1983, 1985). In our material, however, many YFP+ dendritic cells were not labeled by a polyclonal antibody to S100B, suggesting that the relationship between expression of the endogenous gene and the transgene is an artifact of transgenesis.

Fluorescent proteins are expressed in astrocytes in the CNS

Because S100 has been commonly used as an astrocytic marker in the CNS (cf. Ghandour et al., 1981), our expectation was that FPs would be expressed in these cells in our transgenics. This expectation was met to varying extents in the different lines. In the more extensively examined Kosmos line, GFP+ cells that colabeled with anti-GFAP, an astrocytic marker (supplemental Fig. 4A–C, available at www.jneurosci.org as supplemental material), were present throughout the CNS. However, not all astrocytes (GFAP-positive cells) expressed GFP (data not shown). In hippocampus of these mice, GFP+ cell bodies were absent in the pyramidal and granule cell layers (supplemental Fig. 3A, available at www.jneurosci.org as supplemental material), but GFP-labeled processes outlined the cell bodies of many of the

Figure 5. Vital images of NMJs in the YA line show that YFP+ and YFP− TSCs retain their phenotype in adults. Images of YFP fluorescence obtained 30 d apart from two junctions in sternomastoid muscles. Images of AChR are not shown. A, B, NMJ in which TSCs express YFP. C, D, NMJ in which TSCs are YFP− but the motor nerve terminal is YFP+. Scale bar, 50 μm.

Figure 6. Some transgenic lines have prominent labeling of resident macrophages within nerve and muscle, and these cells accumulate at sites of injury. A, YFP fluorescence from the plantar branch of the sciatic nerve from an animal of the YH line. YFP+ cells are oriented mostly along the axis of axons that run horizontally in this image. B, YFP fluorescence from the triangularis sterni muscle of a YE animal. Near the top of the panel, YFP+ axons within the intramuscular nerve form NMJs (arrows). In addition, there are YFP+ macrophages (arrowheads) located along the fibers throughout the muscle that have no obvious relation to NMJs. The processes of these cells are commonly oriented along muscle fibers, i.e., at ~45° in this image. C, D, Higher-magnification image of a junction from a muscle like that in B, showing YFP labeling of the motor axon and terminal (C) and labeling of the Schwann cells with an anti-S100 antibody (D). YFP is present in two TSCs present at the junction that are labeled by anti-S100 as well (D, arrows). YFP is also present in a nearby cell with several processes (C, arrowhead) that is negative for S100 (D). E, F, In a sternomastoid muscle of a YE animal, YFP+ cells (E) accumulate within and around a muscle fiber 2 d after the fiber was injured with iridectomy scissors. The DIC image (F) shows that the sarcomeric organization clearly evident in adjoining fibers is absent from the damaged fiber. Scale bar: A, E, F, 50 μm; B, 35 μm; C, D, 25 μm.
neurons. Similar labeling was seen in the spinal cord and cerebral cortex (data not shown). In the cerebellum, GFP was expressed in astrocytic-like cells in the granule and Purkinje cell layers, but the most prominent labeling was in Bergmann glia, with the molecular layer being intensely fluorescent (supplemental Fig. 3A). Most prominent labeling was in Bergmann glia, with the astrocytic-like cells in the granule and Purkinje cell layers, but the cortex (data not shown). In the cerebellum, GFP was expressed in neurons. Similar labeling was seen in the spinal cord and cerebral cortex (data not shown). In the cerebellum, GFP was expressed in astrocytic-like cells in the granule and Purkinje cell layers, but the most prominent labeling was in Bergmann glia, with the molecular layer being intensely fluorescent (supplemental Fig. 3B, available at www.jneurosci.org as supplemental material). No GFP+ cells were colabeled with the Rip antibody to oligodendrocytes (Friedman et al., 1989), indicating that these glia do not express the transgene (data not shown). In the olfactory bulb, GFP was brightest in the olfactory nerve layer and sparse in areas of high neuronal cell body density (e.g., the mitral cell layer, data not shown). Despite this labeling of cells previously shown to express S100B, it is noteworthy that not all CNS cells that labeled with anti-S100 antibody expressed GFP (data not shown).

Microglia and subsets of Bergmann glia are labeled in some transgenic lines

In contrast to the expected astrocytic labeling described above in mice of the Kosmos line, some of the YFP lines labeled distinct populations of cells in the CNS. Two clear examples emerged from an examination of the brains of YA and YH mice. In YA animals subsets of Bergmann glial cells in the cerebellum were brightly labeled (Fig. 8A), in contrast to Kosmos, in which all Bergmann glia appeared to be labeled. Both YA and YH mice had bright labeling of ramified cells that were identified as microglia on the basis of their morphology, labeling with antibodies to the markers FA-11 and F4/80 (Ohmi et al., 2003) (supplemental Fig. 4D–F, available at www.jneurosci.org as supplemental material), and absence of labeling with GFAP (data not shown). In the cortex of YA mice, very little labeling was present that was not in microglia (Fig. 8B), whereas in YH the microglia were present on a background of weaker labeling of astrocytes (supplemental Fig. 4G, available at www.jneurosci.org as supplemental material). Last, in addition to the labeling of spinal motor neurons described above, some YFP lines showed labeling of neurons in the brainstem. These cells have been shown previously to express S100B (cf. Friend et al., 1992; Yang et al., 1996).

Discussion

We report here the generation of transgenic mice that express fluorescent proteins under the control of the human S100B promoter. In many of the lines, expression occurs in the same cells in which the S100B protein itself is reportedly present, i.e., in Schwann cells, dendritic cells of the immune system, astrocytes, microglia, adipocytes, and some neurons in the brainstem. These results mirror some findings in a previous report in which 4.8 kb of mouse S100B regulatory sequence was used to drive expression of GFP (Vives et al., 2003). They also complement findings in which other promoters were used to drive transgenic expression of FPs in glial cells (Zhuo et al., 1997; Fuss et al., 2000; Nolte et al., 2001). However, we also found considerable variegation in transgene expression among the transgenic lines we generated. In some lines, cells in which the S100B protein is not normally found were clearly FP-positive. A prominent example is expression in spinal motor neurons. A second example is the presence of YFP+, S100+ macrophages distributed throughout skeletal muscles. We also observed cases in which only some of the expected cell types were labeled. For example, in some of the lines microglial labeling was the most prominent labeling in the cortex. Still another type of variegation was labeling of subsets of a given cell type. Examples include Bergmann glia, peripheral Schwann cells, dendritic cells, macrophages, and α motor neurons. Thus, these lines present an array of labeling patterns.

Such variegation is not unusual in transgenic animals and has been attributed to variability in the site of transgene integration and methylation (silencing) as well as the absence in the transgene construct of some genomic sequences normally conferring control of the expression of the endogenous gene (Martin and Whitelaw, 1996). A transgenic mouse has been generated previously using a transgene containing 17.3 kb of sequence encoding human S100, including 9.0 kb of sequence upstream of the trans-
lation start site and some 4 kb of sequence 3′ to the polyadenylation signal (Friend et al., 1992). These transgenic animals were reported to overexpress S100 appropriately in all cell types examined. The differences in transgene expression reported here and in this overexpression study could be explained by the improved sensitivity of a FP reporter over in situ hybridization. Differences between human and mouse promoter sequences also might explain some of the unexpected expression of FP we encountered. However, the variation among lines seen in our study was obtained using a single transgene and is therefore more likely to be caused by position effects. Whether FP expression in subsets of cells in any given line represents intrinsic differences among cells or is attributable to some chance occurrence during development remains unclear (cf. Feng et al., 2000).

Other investigators preparing transgenic mice have noted variation like that reported here and have shown that this variation can enable imaging of a small number of cells against a background of cells of the same type having no labeling (Walsh and Lichtman, 2003). The mice described here could be used to examine subsets of axons in muscles, Bergmann glial cells in the cerebellum, microglial cells in the cortex, blood-derived antigen-presenting cells (macrophages and dendritic cells), and Schwann cells in nerves and at synapses. In this study we have made initial attempts to understand the variegation in Schwann cell labeling at individual NMJs.

**Variegation in Schwann cell labeling at neuromuscular junctions**

The YFP label in Schwann cells follows an intriguing pattern in some of the lines. There is a high probability that, for any particular junction, either all of the TSCs express YFP or none of them do. Based on the presence of a similar expression pattern in early development, the presence of variegation in myelinating Schwann cells, and the association of both YFP+ and YFP− Schwann cells with single axons, it seems unlikely that this pattern is dictated by motor neurons or muscle. Rather, the simplest explanation appears to be some type of random event during early development that results in Schwann cells or their progenitors becoming one of two types, YFP+ or YFP−, with this commitment retained by daughter cells. Because the first Schwann cell to reach developing NMJs almost certainly arrives there by migration along an innervating axon and because the number of TSCs increases subsequently, at least in part, by cell division (cf. Love and Thompson, 1998), the first Schwann cell could serve as the founder for all the other TSCs as well as the Schwann cells that generate the myelin wrappings of the axon near the junction. Because multiple axons innervate fetal junctions, it is possible that different axons could recruit different potential founder Schwann cells to the same junction. To achieve the homogeneity seen at adult NMJs, the axon that survives synapse elimination would have to retain one of the sets of Schwann cells, possibly the Schwann cells that associated with it during development. Alternatively, the same founder Schwann cell and its descendants could populate a bundle of axons leading to each junction and separate these axons later in development, in the same way that Schwann cells migrate between and separate axons within developing nerve trunks (Peters and Muir, 1959). Inspection of the data in Figure 4 suggests that Schwann cells are almost as homogeneous at day 6, when half the fibers in the muscle are polynervously innervated, as they are in the adult. However, at a stage when approximately half of the junctions are polynervously innervated and approximately half of the Schwann cells are YFP+, one would anticipate only a quarter more junctions with mixed Schwann cell labeling than seen in the adult animal. Repeated, vital imaging in the neonate could be used to determine the relationship between the Schwann cells and axons that remain at NMJs after synapse elimination.

**Vital imaging reveals the stability of synaptic glia**

The original objective of this study was to produce mice useful for vital observation of TSCs at the NMJ. At least two of the lines (Kosmos and YG) appear useful in this regard because they have strongly labeled myelinating and terminal Schwann cells but little or no labeling of other cell types in the muscle. We have shown the feasibility of using these mice for vital imaging. Because few changes were found in the glia at imaged junctions, we believe that the imaging itself does not change the junctions. This result contrasts with the marked changes in Schwann cells observed in transgenic animals after denervation of the muscle (L. Tian, H. Kang, and W. J. Thompson, unpublished results), changes that are also seen in nontransgenic animals (cf. Son and Thompson, 1995a).

Our observations extend to the glial components of the synapse the relative stability previously reported for the nerve terminal and postsynaptic apparatus at adult NMJs (Balice-Gordon and Lichtman, 1990). This finding contrasts with variability in glial cell numbers and positions observed during vital imaging of synapses in parasympathetic ganglia (Pomeroy and Purves, 1988), but these synapses are more dynamic than neuromuscular junctions.

We also found that TSCs send out short, unstable processes that explore the territory around the synapse (Robbins and Polak, 1988). These extensions may play an important role in matching TSC processes to nerve terminals and AChRs, should they happen to change. For example, as the junction grows during development, its glial coverings expand, with the addition of TSCs as required (Love and Thompson, 1998). The extension of longer TSC processes after denervation may represent an elaboration of these short, exploratory TSC processes. We are using these mice to follow the growth of these processes and to elucidate their role in nerve repair after full and partial denervation.

**References**


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