Dynamic Pattern of Reg-2 Expression in Rat Sensory Neurons after Peripheral Nerve Injury

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The 16 kDa pancreatitis-associated protein Reg-2 has recently been shown to facilitate the regeneration of motor and sensory neurons after peripheral nerve injury in the adult rat. Reg-2 has also been shown to be a neurotrophic factor that is an essential intermediate in the pathways through which CNTF supports the survival of motor neurons during development. Here we report the dynamic expression of Reg-2 in rat sensory neurons after peripheral nerve injury. Reg-2 is normally not expressed by dorsal root ganglion (DRG) cells, but we show, using immunocytochemistry, that Reg-2 is rapidly upregulated in DRG cells after sciatic nerve transaction and after 24 hr recovery is expressed almost exclusively in small-diameter neurons that bind the lectin Griffonia simplicifolia IB4 and express the purinoreceptor P2X₃. However, by 7 d after axotomy, Reg-2 is expressed in medium to large neurons and coexists partly with the neuropoietic cytokines. Reg-2 is no longer expressed in small neurons, and there is no colocalization with IB4 binding neurons, demonstrating a shift in Reg-2 expression from one subset of DRG neurons to another. We also show by double labeling for activating transcription factor 3, a transcription factor that is upregulated after nerve injury, that Reg-2 expression occurs predominantly in axotomized DRG cells but that a small percentage of uninjured DRG cells also upregulate Reg-2. The selective expression within IB4/P2X₃ cells, and the dynamic shift from small to large cells, is unique among DRG peptides and suggests that Reg-2 has a distinctive role in the injury response.

Key words: regeneration; axotomy; dorsal root ganglia; neuropoietic cytokine; peripheral nerve injury; Reg-2

Reg-2 (also known as PAP1 in rat, RegIIβ in mouse, and HIP/PAP in humans) is a 16 kDa secretory protein that has recently been shown to have preregenerative properties in motor and sensory neurons after peripheral nerve injury in the rat (Livesey et al., 1997). Reg-2 is massively upregulated in subsets of sensory neurons and in all regenerating α motor neurons after sciatic nerve injury (Livesey et al., 1997). In vitro, Reg-2 has a mitogenic effect on Schwann cells, and direct injection of Reg-2 antibody into the crushed nerve retarded the regeneration of the relevant subsets of sensory and motor neurons. This results strongly imply a novel principle: neurons do not simply grow passively through a permissive environment, but they can actively secrete factors that can change the environment through which they are regenerating. Reg-2 is also constitutively expressed in subpopulations of motor neurons during development, and this expression is driven by cytokines of the interleukin-6 (IL-6) family, which includes ciliary neurotrophic factor (CNTF), leukemia inhibitory factor (LIF), and cardiotrophin (CT-1) (Livesey et al., 1997). These cytokines have been shown to prevent motor neuron cell death that follows neonatal axotomy (Sendtner et al., 1990; Cheema et al., 1994; Pennica et al., 1996) and to prolong motor neuron survival in strains of mice that show spontaneous motor neuron cell death (Sendtner et al., 1992; Mitsumoto et al., 1994; Lindsay, 1996; Winter et al., 1996; Bordet et al., 1999).

More recently, Nishimune and colleagues (2000) have shown that during development Reg-2 is a neurotrophic factor that is an essential intermediate in the pathways through which CNTF supports the survival of motor neurons (Nishimune et al., 2000).

The role of Reg-2 expression in sensory neurons of the dorsal root ganglion (DRG) has yet to be elucidated but is clearly of interest because peripheral sensory neurons have the capacity to regenerate, and neuropoietic cytokines play a role in the maintenance of sensory neurons after peripheral nerve injury (Simon et al., 1995; Thompson et al., 1998; Thier et al., 1999). In a previous study (Livesey et al., 1997) Reg-2 was shown to be upregulated in a subpopulation of DRG cells after sciatic nerve crush, but the DRG cell type was not characterized. Here we have performed a detailed analysis of Reg-2 expression in lumbar DRG cells and their central projections at various time points after sciatic nerve crush and transection. Reg-2 expression is dynamic, appearing within distinct populations of sensory neurons at different times after axon damage.

MATERIALS AND METHODS

Materials and antibodies. All chemicals and materials were obtained from Sigma-Aldrich (Poole, UK) or Merck-BDH (Lutterworth, UK) unless stated otherwise. In this study, the following primary antibodies were used: anti-Reg-2 polyclonal antibody (Livesey et al., 1997) was raised in rabbit against whole recombinant protein and used in these studies at 1:20,000 for immunoperoxidase, 1:8,000 for indirect labeled immunofluorescence, and 1:120,000 with tyramide signal amplification (TSA; see below). Anti-trkA rabbit polyclonal antibody (Upstate Biotechnology, Lake Placid, NY) was used at 1:10,000 (TSA procedure); rabbit anti-neuropoietic Y (NPY) and anti-galanin antisera (both Affiniti, Exeter, UK) were used at 1:2,000, and guinea pig anti-P2X3 antiserum (Neuromics, Minneapolis, MN) was used at 1:1,500,000 (TSA procedure). Anti-activating transcription factor 3 (ATF3) rabbit polyclonal antibody (Santa Cruz, CA) was used at 1:200. Isolectin Griffonia simplicifolia IB4 FITC conjugate (Sigma-Aldrich, Dorset, UK) was used at a dilution of 1:100.
Animals and surgery. For all experiments, male Sprague Dawley or Wistar rats of ~150–200 gm were used, and preliminary experiments revealed no difference in Reg-2 expression, or upregulation, between these two strains. Unilateral sciatic nerve crush (15 Wistar, 4 Sprague Dawley) or transection (12 Wistar, 16 Sprague Dawley) was performed at mid-thigh level under deep anesthesia [4% (v/v) halothane for induction and maintained with 2% (v/v) during surgery]. For transection the sciatic nerve was first ligated and then cut distal to the ligature. Animals were allowed to recover for 24 hr, 5 d, 7 d, 30 d, or 8 weeks, at which time tissue was removed for immunocytochemical analysis. In some experiments (n = 4), the sciatic nerve was injected with 5 μl of 5% (w/v) Fast Blue at the time of transection to retrogradely label the axotomized sciatic afferents. In six other animals, axonal transport was studied by ligating the L4/L5 lumbar dorsal roots and proximal portion of sciatic nerve (two animals) or by ligating the sciatic nerve (two animals) or saphenous nerve (two animals) at mid-thigh level 3 d before perfusion fixation.

Tissue processing. Rats were deeply anesthetized with pentobarbitone (60 mg/kg, i.p.) and transcardially perfused with 100 ml sterile saline con-

Table 1. The percentage of DRG cells that express Reg-2, IB4, trkA, galanin (GAL), or ATF3 immunoreactivities at various time points after sciatic nerve transection

<table>
<thead>
<tr>
<th>Time point</th>
<th>Reg-2</th>
<th>IB4</th>
<th>trkA</th>
<th>GAL</th>
<th>ATF3</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>0.8 ± 0.4</td>
<td>49.7 ± 2.0</td>
<td>44.0 ± 0.6</td>
<td>8.7 (2)</td>
<td>0.8 ± 0.4</td>
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<tr>
<td>24 hr</td>
<td>14.1 ± 1.0</td>
<td>44.3 ± 2.4</td>
<td>47.5 ± 3.2</td>
<td>22.5 (2)</td>
<td>72.9 ± 4.4 (5)</td>
</tr>
<tr>
<td>7 d</td>
<td>10.0 ± 1.5 (4)</td>
<td>35.5 ± 0.4</td>
<td>31.2 ± 0.9</td>
<td>40.4 ± 3.2</td>
<td>58.7 ± 6.6</td>
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<tr>
<td>30 d</td>
<td>10.9 ± 0.5</td>
<td>32.2 ± 5.3</td>
<td>39.0 ± 3.2</td>
<td>39.3 ± 7.4</td>
<td>67.0 ± 5.0</td>
</tr>
</tbody>
</table>

The numbers shown are mean ± SEM and are based on counts from three animals unless indicated otherwise.

Figure 1. Expression of Reg-2 in DRG after peripheral nerve injury. A–D show Reg-2 immunoreactivity in ipsilateral (A–C) and naive control (D) lumbar DRG 1 d (A), 7 d (B), and 30 d (C) after sciatic nerve transection. Reg-2 is upregulated after sciatic transection, but expression is dynamic, appearing predominantly in small cells at 1 d (A) but in medium and large cells at 7 d (B). E–H show the disposition of Reg-2-immunoreactive axons in an L4 ganglion (DRG) with attached ventral root (VR), dorsal root (DR), and spinal nerve (SN) at 1 d after sciatic transection. The labeling in E shows the areas that were sampled for the high-magnification images in F–H. Immunoreactive axons (arrows) can be observed within and on the peripheral side of the ganglion (F) as well as within the spinal nerve (G). However, very few axons were present on the central side of the ganglion or within the attached dorsal root (H) or ventral root. Scale bars: A–D, 100 μm; E, 200 μm; F–H, 50 μm.
frozen on dry ice, and sectioning was performed on either a freezing microtome (Leica, Hemel Hempstead, UK) or a cryostat (Leica).

Microtome sections were cut at 20 or 40 μm into 5% (w/v) sucrose in 0.1 M PB containing 0.02% (w/v) sodium azide and were processed for Reg-2 immunoreactivity as free-floating sections. Tissue for cryostat sectioning was embedded in OCT, cut at 6 μm, and thaw-mounted onto Superfrost plus microscope slides.

**Immunocytochemistry.** To determine localization of expression in the DRG for cell size distribution analysis, free-floating microtome sections were first rinsed in 0.1 M PB, followed by a 30 min incubation in 0.1 M PB containing 0.6% (w/v) hydrogen peroxide at room temperature to block any endogenous peroxidase activity. Sections were then transferred into 0.1 M PB containing 3% (w/v) normal goat serum, 0.25% (v/v) Triton X-100, and 0.02% (w/v) sodium azide (PBT) and incubated for 1 hr at room temperature. For incubation in primary antibody, sections were transferred into fresh PBT containing Reg-2 polyclonal antiserum and incubated at 4°C for 2 d. After washes in 0.1 M PB, sections were incubated in biotinylated goat anti-rabbit IgG (Vector Laboratories, Peterborough, UK; 1:400 in PBT) for 1 hr at room temperature. After further washes, sections were incubated in avidin–biotin complex (Vector Laboratories; 1:200 in 0.1 M PB) that had been premixed 30 min previously. Finally, sections were washed again in 0.1 M PB followed by a brief wash in 0.15 M Tris-HCl, pH 7.4. Sections were then transferred into 0.15 M Tris-HCl, pH 7.4, containing 0.25 mg/ml diaminobenzidine, 2 mg/ml nickel sulfate, and 0.003% (w/v) hydrogen peroxide to induce color reaction. Tissue sections were washed in 0.1 M PB to stop the color development. Mounted sections were allowed to air-dry overnight. They were then dehydrated through increasing alcohol concentrations and placed in Histoclear. Slides were then coverslipped using DPX as mountant.

For double-labeling experiments, standard immunofluorescence procedures were used using either indirect labeled immunofluorescence or a TSA kit (NEN Life Science Products, Hounslow, UK (Averill et al., 1995; Michael et al., 1997). Incubations were performed at room temperature and consisted of 1 hr in 10% (v/v) normal serum followed by 18–36 hr in each set of primary antisera and 3 hr in the developing secondary antisera. The two sets of antisera were applied sequentially, and this normally involved Reg-2 TSA followed by indirect-labeled immunofluo-

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**Figure 2.** Size distribution of Reg-2-immunoreactive sensory neurons in L4/5 DRG after sciatic nerve transection. Note that at 24 hr many small-diameter sensory neurons are immunoreactive, but by 5 d predominantly medium-diameter cells show Reg-2 expression. At 5 d after transection, some neurons are large, as indicated by the long tail seen on the size distribution graph, which is not seen by 8 weeks after transection.

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**Figure 3.** Analysis of Reg-2 immunoreactivity in the spinal cord and brain stem. A–D show the lumbar spinal cord 7 d after sciatic nerve transection. Reg-2 immunoreactivity is expressed in the ventral horn (VH) in axotomized motoneurons (A, arrows) but is absent from the dorsal horn (DH), indicating that it is not present in the central terminals of axotomized primary afferents. B–D show the superficial dorsal horn at high magnification, stained for IB4, Reg-2, or CGRP. Asterisks indicate the central terminal field of the axotomized sciatic nerve, which has downregulated IB4 (B). However, Reg-2 immunoreactivity in this region is not above background staining (C), although CGRP staining confirms that primary afferent terminals are present (D). E and F show the dorsal medulla 30 d after sciatic nerve transection. Ipsilateral to the transection, neuropeptide Y (NPY) immunoreactivity is upregulated within the gracile nucleus (E) in the central terminal fields of the axotomized primary afferents (asterisk). However, there is no indication of Reg-2 immunoreactivity in that region (F). Scale bars: A, 200 μm; B–F, 100 μm.
rescence. Tetramethylrhodamine isothiocyanate (TRITC)-labeled anti-rabbit IgG was used for indirect immunofluorescence (Jackson ImmunoResearch, West Grove, PA; 1:400 dilution). TSA labeling was performed using biotinylated goat anti-rabbit IgG (1:400; Vector Laboratories) and Vectastain Elite peroxidase reagent (Vector Laboratories) followed by biotin tyramide (NEN Life Science Products, Hounslow, UK; TSA-indirect kit) and ExtrAvidin-FITC (1:500, Sigma-Aldrich, Dorset, UK). After incubation in secondary reagents, sections were washed briefly in PBS and then mounted in PBS/glycerol (1:3) containing 2.5% (w/v) 1,4 diazobicyclo (2,2,2) octane (DABCO; anti-fading agent). Controls for double labeling included reversing the order of the primary antisera, as well as omitting the first or second primary antiserum.

Image analysis. For cell size distribution and fast blue experiments, images were obtained using a Leica DMR microscope and either a JVC KY-F50 color video camera (for DAB-labeled sections) or a Hamamatsu C5985 CCD camera (for immunofluorescence). Images were grabbed using VisionExplorer software, and cell diameters were measured using Leica Qwin (v2.2) image analysis software. For cell size distribution, we measured the diameters of Reg-2-positive cells from at least 16 sections taken from three animals at each time point. Only cells that displayed a distinct nucleus were measured. For immunofluorescence sections, quantitation of the proportion of Reg-2 expressing DRG cells was determined by counting the number of immunoreactive and non-immunoreactive neuronal profiles. In double-labeled sections, the percentage of Reg-2-expressing cells expressing a second marker was assessed by switching between FITC and TRITC filter blocks. At least 250 labeled DRG cells were examined for each marker and counted on randomly chosen sections. Photographs were taken using a Hamamatsu C4742-95 digital camera, and plates were assembled using Adobe Photoshop.

**RESULTS**

Reg-2 immunoreactivity was assessed in rat lumbar DRG cells at various times after sciatic nerve injury. By 24 hr after nerve transection, Reg-2 was expressed by ~14% of DRG cells (Table 1). Immunoreactivity was observed in predominantly small diameter (26.46 ± 4.65 μm) sensory neurons, which appeared evenly distributed throughout the DRG (Figs. 1A, 2). By 5–7 d after sciatic nerve section, a similar percentage of DRG cells were stained, but the immunoreactivity was now observed in predominantly medium to large diameter (44.47 ± 8.98 μm) cells with some cells having very large diameters (>60 μm) (Figs. 1B, 2). Immunoreactivity remained elevated at longer time points, but by 8 weeks the Reg-2-positive cells were predominantly of small to medium diameter with no very large cells observed (Fig. 2). Similar results were obtained with both nerve transection and crush at 1 and 7 d survival. Reg-2 immunoreactivity in contralateral lumbar DRG and in naïve control DRG was observed in just a few isolated profiles at each time point and omission of primary antisera resulted in a loss of immunoreactivity (not shown). In some material, light labeling of satellite glial cells was present (see Fig. 7C), but controls indicated that this was not specific. In addition to DRG cells, a few Reg-2-immunoreactive axons were visible within ganglia at all time points studied, and in well-stained preparations they could be observed running into and

**Figure 4.** Axonal transport of Reg-2. A, C, and E show anterograde accumulation proximal to a L4/L5 dorsal root ligature, and B, D, and F show anterograde accumulation proximal to a sciatic nerve ligature. Vertical arrows in A and B indicate the site of each ligature. CGRP shows a prominent accumulation in dorsal roots (A) and in sciatic nerve (B), whereas very little Reg-2 accumulation (C, D, arrows) is seen in dorsal roots compared with sciatic nerve. IB4 staining proximal to the ligatures is not as prominent as CGRP but is present in both dorsal roots (E) and sciatic nerve (F). Scale bars: A, C, E, 100 μm; B, D, F, 200 μm.
within the spinal nerve (Fig. 1E–G). However, only a few immunoreactive axons were present within dorsal roots (Fig. 1H), which suggests that Reg-2 protein from DRG cells is transported peripherally but mainly not centrally after nerve injury. This conclusion was supported by analysis of the central termination territory of DRG axons and of the effect of nerve ligation. Immunostaining of lumbar spinal cord after sciatic nerve transection revealed the previously described expression of Reg-2 in motor neurons (Livesey et al., 1997) that remained only for the period of regeneration. Despite upregulation of Reg-2 in the DRG, as described above, no staining was observed in the dorsal horn of the spinal cord (Fig. 3A–D) or in the dorsal column nuclei (Fig. 3E,F) at any time points studied (1, 7, and 30 d). Accumulation of Reg-2 immunoreactivity was observed proximal to a ligature of the sciatic nerve (Fig. 4B,D,F) or saphenous nerve (a purely sensory nerve), but very little was present proximal to a dorsal root ligature (Fig. 4A,C,E).

The population of small-diameter Reg-2-positive neurons observed in L5 lumbar DRG 24 hr after sciatic nerve injury coexists almost exclusively (>95%) with IB4 binding and purinoreceptor P2X3 immunoreactivity (Fig. 5A–D, Table 2). IB4 binding and P2X3 expression have been shown to coexist in that ~98% of P2X3-expressing sensory neurons are IB4-positive (Bradbury et al., 1998). At this same 24 hr time point, a proportion of the L5 DRG Reg-2-positive profiles (21% after transection and 25% after nerve crush) also colabel for the nerve growth factor (NGF) receptor trkA (Fig. 5E,F). Numerous DRG cells showed galanin immunoreactivity, but there was little coexistence with Reg-2 (Table 2).

At 7 d after sciatic nerve injury (both transection and crush), Reg-2 coexpression with IB4 binding was much reduced (Fig. 6A,B, Table 2), but a large proportion of the Reg-2-immunoreactive profiles showed immunoreactivity for galanin (Fig. 6C,D) and NPY (Fig. 6E,F). Both of these peptides have been shown to be upregulated in sensory neurons after sciatic nerve injury (Hokfelt et al., 1987; Villar et al., 1989; Wakisaka et

Figure 5. Colocalization of Reg-2-positive cells 24 hr after sciatic nerve transection. Immunofluorescent staining of single L5 DRG sections using Reg-2 (A, C, E) and IB4-FITC conjugate (B) or P2X3 (D) or trkA (F) polyclonal antibodies 24 hr after sciatic nerve transection. Arrows show double-labeled cells, and arrowheads show Reg-2-positive cells that are negative for the second marker. Note that many Reg-2-positive cells show IB4 or P2X3 labeling. Scale bar, 50 μm.
Reg-2 expression has a number of interesting features that make it quite unique and were not reported in the original study (Livesey et al., 1997). First, Reg-2 is one of a very small number of molecules that are rapidly upregulated after nerve injury. Second, Reg-2 is the first example of a molecule that is selectively upregulated in the IB4/P2X<sub>3</sub> GDNF-sensitive population of nociceptors. Third, the shift in Reg-2 expression from initially small-diameter to subsequently large-diameter neurons is unique and may reflect the selective expression of cytokines and growth factors adjacent to the site of damage or denervated target (see below). Fourth, Reg-2 is axonally transported predominantly peripherally and not centrally.

Reg-2 is a gene belonging to a larger family of Reg-related genes. The original member of the family, Reg, was a novel gene expressed in regenerating pancreatic islet cells (Terazono et al., 1988) and was found to code for a 16 kDa secretory protein. In subsequent years, a family of related genes has been described under a rather varied nomenclature and been broadly categorized into three groups (for review, see Okamoto, 1999). Type I Reg proteins include the original Reg-1 protein and have been shown to have a role in promoting regeneration and proliferation of insulin-producing β-cells of pancreatic islets (Terazono et al., 1988, 1990; Zenilman et al., 1997; Levine et al., 2000). The type II Reg gene has been described only in the mouse (Unno et al., 1993), and the biological function of this gene has not been determined. Type III Reg proteins, which include rat Reg-2 described here, have been described as growth factors in liver cells (Christa et al., 1996), as possible anti-apoptotic agents in pancreatic acinar cells (Ortiz et al., 1998), and as a novel motor and sensory neuron survival factor (Livesey et al., 1997; Nishimune et al., 2000). In addition, a regenerative role of Reg-2 as a Schwann cell mitogen released at the regrowing axon tip has been described (Livesey et al., 1997).

Sensory neurons of the DRG can be categorized into subpopulations according to their size and expression of various neurochemical markers (for review, see Snider and McMahon, 1998; Hunt and Mantyh, 2001). Small-diameter sensory afferents represent ~70% of the total lumbar DRG neuron population, have unmyelinated axons (C-fibers), and act mainly as nociceptors. The large-diameter afferents of the neuron population have myelinated axons (A-fibers), innervate mechanoreceptors peripherally, and mediate proprioceptive and tactile responses. They can be immunocytochemically identified using antibodies that recognize high molecular weight neurofilament protein. Small-diameter DRG neurons can be further characterized into a peptide-expressing [such as calcitonin-related gene product (CGRP) and substance P] and NGF-responsive subset and a nonpeptidergic, GDNF-responsive subset that bind isoclinet-B4 (IB4) and express the purinoreceptor P2X<sub>3</sub> (Averill et al., 1995; Bennett et al., 1998; Priestley et al., 2002).

Peripheral nerve injury induces dramatic changes in gene expression in DRG neurons. For instance, after peripheral nerve transection or crush, the transcription factors c-jun (Herdegen et al., 1992; Jenkins et al., 1993) and ATF3 (Tsujino et al., 2000) are expressed in all injured neurons within 24 hr, whereas the pattern and extent of the expression of neuropeptides such as galanin and NPY are dependent on the time after injury (Hokfelt et al., 1987; Wakisaka et al., 1991; Zhang et al., 1998; Landry et al., 2000).

Endogenous expression of these peptides in normal sensory neurons is limited to a small number (~5%) of small-diameter cells, but after injury expression is robustly enhanced and maintained in neurons of all sizes.

Reg-2 expression is rarely seen in control ganglia, but both mRNA (Livesey et al., 1997) and protein (this study) are upregulated after nerve injury. Twenty-four hours after either sciatic nerve transection or crush, Reg-2 is transiently expressed within

### Table 2. The percentage of Reg-2-immunoreactive DRG cells that also express IB4, P2X3, trkA, galanin, NPY, or ATF3 at various time points after sciatic nerve transection

<table>
<thead>
<tr>
<th>Time point</th>
<th>Markers</th>
<th>% of Reg-2-expressing cells that also express the marker</th>
<th>% of marker-expressing cells that also express Reg-2</th>
</tr>
</thead>
<tbody>
<tr>
<td>24 hr</td>
<td>Reg-2 and IB4</td>
<td>95.3 ± 1.8</td>
<td>20.3 ± 3.3</td>
</tr>
<tr>
<td></td>
<td>Reg-2 and P2X3</td>
<td>94.7 ± 1.9</td>
<td>28.2 ± 2.0</td>
</tr>
<tr>
<td></td>
<td>Reg-2 and trkA</td>
<td>21.2 ± 2.0</td>
<td>5.0 ± 0.2</td>
</tr>
<tr>
<td></td>
<td>Reg-2 and galanin</td>
<td>2.2</td>
<td>1.3</td>
</tr>
<tr>
<td></td>
<td>Reg-2 and NPY</td>
<td>NA</td>
<td>NA</td>
</tr>
<tr>
<td></td>
<td>Reg-2 and ATF3</td>
<td>77.8 ± 1.8</td>
<td>7.2 ± 0.2</td>
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<tr>
<td>7 d</td>
<td>Reg-2 and IB4</td>
<td>3.2 ± 1.6</td>
<td>0.6 ± 0.3</td>
</tr>
<tr>
<td></td>
<td>Reg-2 and P2X3</td>
<td>20.3 ± 4.9</td>
<td>3.2 ± 1.3</td>
</tr>
<tr>
<td></td>
<td>Reg-2 and trkA</td>
<td>0.7 ± 0.7</td>
<td>0.1 ± 0.1</td>
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<tr>
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<td>Reg-2 and galanin</td>
<td>48.6 ± 3.5</td>
<td>9.4 ± 1.2</td>
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<td></td>
<td>Reg-2 and NPY</td>
<td>76 ± 1.7</td>
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<td>Reg-2 and ATF3</td>
<td>96.5 ± 2.6</td>
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<td>30 d</td>
<td>Reg-2 and IB4</td>
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<td>Reg-2 and NPY</td>
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<td></td>
<td>Reg-2 and ATF3</td>
<td>71.8 ± 9.5</td>
<td>7.1 ± 0.3</td>
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</table>

The numbers shown are mean ± SEM and are based on counts from three animals, with the exception of the 24 hr Reg-2 and galanin counts, which are based on two animals. NA, Not applicable.
a subpopulation of the purinoreceptor P2X₃- expressing, GDNF-sensitive, small-diameter sensory neurons. At this time point, all Reg-2-expressing cells show IB4 labeling, and the small percentage (20%) that show trkA labeling is consistent with the reported overlap between IB4 and trkA (Averill et al., 1995). Many molecules are downregulated after peripheral nerve injury, and the rapid upregulation that we have observed for Reg-2 has so far been reported only for galanin and for the transcription factors c-jun and ATF3. In addition, most molecules that are upregulated are expressed in either the small trkA-expressing cells or large-diameter neurons. Reg-2 is the only molecule, to date, that is selectively upregulated in the IB4-labeled, GDNF-sensitive population of cells. However, at longer time points, Reg-2 expression was restricted to medium- to large-diameter sensory neurons, although some expression in small neurons was observed at 30 d. Our double-labeling studies, cell size distribution data, and counts of total percentage of DRG cells that express Reg-2 all suggest that this change is caused by a shift in Reg-2 expression from the small- to medium/large-diameter neurons. This shift in Reg-2 expression is very unusual. Galanin, for example, is expressed in medium- and large-diameter neurons at longer time points but continues to be expressed in small-diameter neurons. After peripheral nerve axotomy, brain-derived neurotrophic factor (BDNF) expression is also upregulated in medium- to large-diameter sensory neurons (Cho et al., 1998; Michael et al., 1999), but this is the only cell group that shows upregulation. BDNF expression in small cells has been reported either to be downregulated (Cho et al., 1998) or to show no significant change (Michael et al., 1999). In addition, unlike Reg-2, crush injury can induce an increased expression of BDNF in all cell sizes (Cho et al., 1998). The pattern of expression seen for Reg-2 is thus unlike any other peptide, and because similar changes are seen after both transection and crush, it is likely that similar mechanisms for Reg-2 upregulation are activated in both injury models.

The dynamic changes in Reg-2 expression that we have observed may indicate that quite different factors control Reg-2 expression, and function, in the small and large DRG cells. For example, Reg-2 expression in the small-diameter IB4 cells may

Figure 6. Colocalization of Reg-2-positive cells 7 d after sciatic nerve transection. Immunofluorescent staining of single L4/5 DRG sections using anti-Reg-2 polyclonal antibody (A, C, E) and IB4-FITC conjugate (B) or galanin (D) or NPY (F) polyclonal antibodies, 7 d after sciatic nerve transection. Arrows show double-labeled cells, and arrowheads show Reg-2-positive cells that are negative for the second marker. Note that many Reg-2-positive cells at this time point show colocalization with NPY and galanin but not IB4. Scale bar, 50 μm.
have local functions. The transient expression of Reg-2 may result in the delivery of a bolus of the peptide to the site of injury, or Reg-2 could be released within the DRG itself and act on satellite glia and other neurons. This could account for the appearance of Reg-2 in neurons that were not axotomized in our experiments, as well as the lack of transport of the peptide into the dorsal horn. It has generally been observed that peptides, such as NPY, that are upregulated within the DRG are exported through the central axonal processes of the DRG to the dorsal horn and/or dorsal column nuclei. However, our studies indicate that Reg-2 is transported predominantly peripherally. A similar lack of central transport has recently been reported for the degenerin/epithelial sodium channel (DEG/ENaC) family member BNaC1 (Garcia-Anoveros et al., 2001). A local role for Reg-2 would be consistent with developmental studies in which Reg-2 appears to act in an autocrine/paracrine manner. After induction by a peripheral factor (possibly a cytokine of the LIF/CNTF family), Reg-2 can act on the parent cell or neighboring cells as an obligatory survival factor (Nishimune et al., 2000). The factor inducing Reg-2 in IB4 cells is not known but could be a member of the LIF/IL-6/CNTF family because IB4 cells are known to have binding sites for this family (Thompson et al., 1997). Whether locally released Reg-2 is responsible for its induction in large neurons is not known but worth further investigation. Such a role for the IB4 group of small-diameter neurons would also complement that of the peptidergic, small-diameter, NGF-responsive sensory fibers that have a well established role in the peripheral inflammatory response (McMahon, 1996).

Reg-2 expression in medium- to large-diameter sensory neurons 7 d after sciatic nerve section predominantly colocalized with NPY or galanin. The expression of these neuropeptides in uninjured DRG neurons is maintained at a low level, and it is thought that the upregulation in the same neurons after peripher-


