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Transplanted progenitors generate functional enteric neurons in the postnatal colon

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Cell therapy has the potential to treat gastrointestinal motility disorders caused by diseases of the enteric nervous system. Many studies have demonstrated that various stem/progenitor cells can give rise to functional neurons in the embryonic gut; however, it is not yet known whether transplanted neural progenitor cells can migrate, proliferate, and generate functional neurons in the postnatal bowel in vivo. We transplanted neurospheres generated from fetal and postnatal intestinal neural crest–derived cells into the colon of postnatal mice. The neurosphere-derived cells migrated, proliferated, and generated neurons and glial cells that formed ganglion-like clusters within the recipient colon. Graft-derived neurons exhibited morphological, neurochemical, and electrophysiological characteristics similar to those of enteric neurons; they received synaptic inputs; and their neurites projected to muscle layers and the enteric ganglia of the recipient mice. These findings show that transplanted enteric neural progenitor cells can generate functional enteric neurons in the postnatal bowel and advances the notion that cell therapy is a promising strategy for enteric neuropathies.

Introduction

The enteric nervous system (ENS) plays an important role in regulating a number of gut functions including motility (1, 2). Enteric neuropathies, which result from diseased, damaged, or congenitally absent enteric neurons, cause motility disorders, most of which are poorly managed by current treatments (3). Cell-based therapies have potential for the treatment of enteric neuropathies by replacing diseased neurons (for example, in gastroparesis or achalasia) or by generating enteric neurons in regions that entirely lack an ENS due to developmental defects (as in Hirschsprung disease) (4–12). Cell-based therapies also hold promise for the treatment of the injured or diseased CNS, but cell therapy for enteric neuropathies is likely to be more complicated because of accessibility and the potential of expanding stem/progenitor cells from healthy regions of the intestine for transplantation into diseased regions of the same patient (13).

Many studies have demonstrated the ability of a variety of sources of stem/progenitor cells to give rise to enteric neurons in the embryonic gut (14–18). For example, enteric neural stem/progenitor cells isolated from postnatal human bowel migrate within the embryonic chick or mouse gut and differentiate into neurons and glial cells (13, 19). However, it is essential that cell therapy to treat enteric neuropathies be carried out postnatally in infants, children, or adults, as diagnosis only occurs after birth. During development, the structure of the gut wall changes dramatically from undifferentiated mesenchyme to a highly organized, concentric-layered structure of differentiated cells (20–24). It is unknown whether the fully differentiated gut wall is permissive for migration of neural progenitor cells. Furthermore, molecules produced by the gut mesenchyme are essential for the normal development of the ENS (12, 25–27), but it is unclear whether these factors are expressed at sufficient levels in the postnatal bowel to permit the development of enteric neurons from progenitors. Previous studies have transplanted CNS neural stem cells, ENS stem/progenitor cells, or ENS cell lines into the gut of postnatal animals in vivo (4, 28–33) or grown cocultures between stem/progenitor cells and the muscle of postnatal human gut (13), but the extent of migration, and whether the graft-derived neurons have the electrophysiological properties of enteric neurons and are incorporated into the neuronal circuitry, have not been determined.

In the present study, we generated neurospheres (NSs) from enteric neural crest–derived progenitors isolated from the fetal and postnatal gut and transplanted them into the postnatal mouse colon in vivo. Although there are a number of possible sources of enteric neurons (4, 5, 7, 8, 10, 11, 13, 16, 34–39), enteric neural crest–derived ENS progenitors were chosen, as they are likely to be the most clinically relevant source of cells, are readily accessible (13), and can give rise to enteric neurons in the embryonic gut or when cocultured with colonic muscle from infants (13, 14, 18). We showed that after transplantation into the colon of postnatal mice, ENS progenitors proliferated; migrated extensively and differentiated into neurons with the neurochemical, morphological, and electrophysiological characteristics of enteric neurons; and received synaptic inputs.

Results

Formation of NSs from dissociated fetal and postnatal gut. Previous studies have shown that all neural crest–derived cells in the gut express KikGR in embryonic EdnrbKik mice (40) and EGFP in embryonic RetTGM mice (41). Although KikGR is a photoconvertible protein that can be converted from green to red by the presence of UV light, we did not exploit this property in the current study, and the native green fluorescence was used to identify neural crest–derived cells. EdnrbKik– or RetTGM–positive cells were isolated by FACS from freshly dissociated gut of E13.5/E14.5 or P4 EdnrbKik or RetTGM

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mice, aggregated by gentle centrifugation and then cultured. After 7 days, NS-like bodies up to 250 μm in diameter had formed (Figure 1E, inset). To characterize NSs derived from dissociated fetal gut (fNSs; E13.5/E14.5) or postnatal gut (pNSs; P4) in vitro, NSs were grown on fibronectin-coated coverslips for 2 days, fixed, and processed for immunohistochemistry. Many cells emigrated from the fNS and pNS, and most of the cells within and surrounding the explanted NSs showed immunoreactivity for the neural crest cell marker Sox10 (Figure 1, A, B, and D). A subpopulation of cells (arrows) expressed the neuronal marker Tuj1 (C and D). (E) Composite image of low-magnification views of a whole-mount preparation of distal colon, showing graft-derived cells and fibers 4 weeks after transplantation of 2 fNSs (generated from the gut of E14.5 EdnrbKik mice). There was extensive migration of graft-derived cells away from the original transplantation sites (asterisks). Some of the graft-derived cells formed ganglion-like clusters (arrows). A NS at the same scale as the whole-mount colon preparation is shown in the inset. (F) Area occupied by graft-derived cells (left) and fibers (right) at the indicated times after transplantation of NSs generated from the gut of fetal and postnatal mice into the distal colon. Scale bars: 50 μm (A–D); 1 mm (E).

Figure 1
Enteric neural crest–derived NSs in vitro and after transplantation into the postnatal colon in vivo. (A–D) In vitro characterization of a pNS (derived from the gut of a P4 EdnrbKik mouse). After 2 days of culture on fibronectin, many EdnrbKik-positive cells had emigrated from the NS (A), and most showed immunoreactivity to the neural crest cell marker Sox10 (B and D). A subpopulation of cells (arrows) expressed the neuronal marker Tuj1 (C and D). (E) Composite image of low-magnification views of a whole-mount preparation of distal colon, showing graft-derived cells and fibers 4 weeks after transplantation of 2 fNSs (generated from the gut of E14.5 EdnrbKik mice). There was extensive migration of graft-derived cells away from the original transplantation sites (asterisks). Some of the graft-derived cells formed ganglion-like clusters (arrows). A NS at the same scale as the whole-mount colon preparation is shown in the inset. (F) Area occupied by graft-derived cells (left) and fibers (right) at the indicated times after transplantation of NSs generated from the gut of fetal and postnatal mice into the distal colon. Scale bars: 50 μm (A–D); 1 mm (E).
transplantation, graft-derived Hu+ and S100β marker Hu and the glial marker S100β were processed for immunohistochemistry using the pan-neuronal markers in the postnatal mouse colon. Many of the graft-derived cells that had migrated away from the transplantation site formed clusters resembling enteric ganglia (Figure 1E and Figure 2A). These data showed that cells derived from both fNSs and pNSs survived, resembling enteric ganglia (Figure 1E and Figure 2A). These data showed that graft-derived Hu+ neurons were also sometimes observed (Figure 2, A and D), and glial cells from the recipient were present at the periphery of most clusters (Figure 2A). Some graft-derived cells did not express Hu or S100β (arrowhead), which suggests that they had not yet differentiated. Scale bar: 20 μm.

Transplanted fNS- and pNS-derived cells differentiate into neurons and glial cells. To determine whether graft-derived neurons and glial cells were present, whole-mount preparations from recipient mice were processed for immunohistochemistry using the pan-neuronal marker Hu and the glial marker S100β. At 1–16 weeks after transplantation, graft-derived Hu+ and S100β+ cells were present in the group of cells that remained at the original transplantation site and in the ganglion-like clusters of graft-derived cells surrounding the transplant site (Figure 2). The ganglion-like clusters also contained some graft-derived cells that expressed neither Hu nor S100β (Figure 2, A and D), and glial cells from the recipient were present at the periphery of most clusters (Figure 2, C and D). Clusters containing a mixture of recipient and graft-derived Hu+ neurons were also sometimes observed (Supplemental Figure 2). These data showed that transplanted fNSs and pNSs gave rise to ganglion-like clusters of cells expressing neuronal and glial markers in the postnatal mouse colon.

Generation of neurons after transplantation of fNS and pNS. To examine whether any neurons were generated from graft-derived cells after transplantation, the thymidine analog 5-ethyl-2′-deoxyuridine (EdU) was injected into recipient mice daily for the first 5 days after transplantation of fNSs or pNSs into the colon, and the mice were killed 4 weeks later. A subpopulation of graft-derived Hu+ cells arising from both fNSs (n = 3) and pNSs (n = 3) had incorporated EdU (Figure 3), which showed that neurons were generated in vivo from proliferating transplanted progenitors. Graft-derived cells that had incorporated EdU but were Hu+ were also commonly observed; these may be glial cells or neuronal or glial precursors.

fNS- and pNS-derived cells colonize the appropriate gut layers, and graft-derived nerve fibers associate with enteric neurons of the recipient. To identify the layers in the gut wall occupied by graft-derived cells and fibers, transverse frozen sections and whole-mount preparations from postsurgical mice were examined by confocal microscopy. The longitudinal muscle layer was thickened at the transplantation site. At the center of the transplantation site, graft-derived cells expressing the pan-neuronal markers Hu or PGP9.5 or the glial markers S100β or GFAP were present throughout the longitudinal and circular muscle layers; however, beyond the original transplantation site, graft-derived neurons and glia were most common in the myenteric plexus region (Figure 4, A–C). Graft-derived, varicose neurites were present in the longitudinal and circular muscle layers and also formed close associations with graft-derived neurons and myenteric neurons of the recipient (Figure 4, D–F). The location of graft-derived cell bodies and neurites within the different gut layers was similar to that of the endogenous neurons of the recipient, except that graft-derived neurons were not observed in the submucosa, and graft-derived neurites were rarely observed in the mucosa (Figure 4, A–C). Some of the varicosities of graft-derived neurites showed immunoreactivity to the synaptic vesicle protein synaptophysin (Figure 4, G–I). Our results showed that graft-derived cells and neurites colonized appropriate layers and cell types within the gut wall; future studies are required to determine whether graft-derived neurons form the appropriate synaptic connections with neurons and muscle within these layers.

fNS- and pNS-derived cells differentiate into neurons with appropriate enteric neuron subtype phenotypes. The ENS in mice and other mammals contains many different subtypes of neurons that can be identified by the expression of markers and morphology (2, 42, 43). We used markers that identify the majority of myenteric neurons in the mouse (43). Subpopulations of graft-derived neurons...
and fibers showed immunoreactivity to neuronal NOS (Figure 5, A–F), choline acetyltransferase (ChAT; Figure 5, G–I), vesicular acetylcholine transporter (VACHT; data not shown), and calretinin and calbindin (Supplemental Figure 3). Moreover, some of the graft-derived NOS neurons possessed flattened, lamellar dendrites (Figure 5, A–C), similar to the dendritic morphology of NOS enteric neurons in the mature intestine (43). The proportions of Hu+ neurons derived from fNSs that expressed ChAT or NOS were determined in 3 recipients 4 weeks after transplantation. ChAT+ and NOS+ neurons accounted for 52.5% ± 2.4% and 40.9% ± 15.0% of graft-derived Hu+ neurons, respectively, similar to the proportions of ChAT+ and NOS+ endogenous myenteric neurons in the distal colon adjacent to the transplant site (54.6% ± 4.8% and 38.2% ± 2.3%, respectively). These findings showed that, within the environment of postnatal colon, fNS- and pNS-derived cells differentiated into neurons with appropriate neurochemical and morphological phenotypes, and in the correct proportions.

No evidence of inappropriate differentiation or tumor formation. Interstitial cells of Cajal (ICCs) are found closely associated with enteric neurons (44–46), but do not arise from the neural crest (47, 48). ICCs can be identified by the expression of the receptor tyrosine kinase Kit (49, 50). We examined whether any fNS-derived cells expressed Kit. Endogenous Kit+ ICCs of the recipient were revealed by immunoreactivity to Hu, (G–I) Whole-mount preparation of graft-derived fibers. Some of the graft-derived varicosities (arrows) showed immunoreactivity to the synaptic vesicle protein synaptophysin. Scale bars: 50 μm (A–C); 20 μm (D–F); 10 μm (G–I).

Figure 4
Location of transplanted cells and their neurites within the wall of the recipient colon. (A–C) Graft-derived cells in a frozen transverse section through the distal colon of a mouse in which an fNS had been transplanted 8 weeks previously. This field of view was away from the original transplant site. Graft-derived cell bodies were present at the level of the myenteric plexus, and graft-derived fibers were present in the muscle layers, but not in the mucosa (A and C). PGP9.5+ cell bodies were present between the longitudinal muscle (LM) and circular muscle (CM) layers, at the level of the myenteric plexus (arrows), and PGP9.5+ fibers were present in the muscle layers and the mucosa (B and C). Note that the PGP9.5 staining showed both the recipient’s endogenous neurons as well as graft-derived neurons. (D–F) Whole-mount preparation of colon after transplantation of a fNS. This region containing graft-derived (Ednrbβ-S-positive) fibers was beyond the extent of graft-derived cell bodies. Graft-derived nerve fibers formed close associations with myenteric neurons of the recipient, revealed by immunoreactivity to Hu. (G–I) Whole-mount preparation of graft-derived fibers. Some of the graft-derived varicosities (arrows) showed immunoreactivity to the synaptic vesicle protein synaptophysin. Scale bars: 50 μm (A–C); 20 μm (D–F); 10 μm (G–I).
Together, these results showed that ENS progenitor cells isolated from the fetal and postnatal mouse gut differentiated into neurons with appropriate electrophysiological properties within the environment of postnatal gut.

Discussion

If cell therapy is to be used to treat gastrointestinal motility disorders, it is crucial to determine whether neural stem/progenitors can migrate and differentiate into neurons with the appropriate neurochemical and electrophysiological properties after transplantation into the postnatal gut in vivo. In the present study, we showed that NSs generated from neural crest–derived cells from both the fetal and the postnatal gut survived, proliferated, migrated, and differentiated into glial cells and a range of neuron subtypes that exhibited neurochemical, morphological, and electrophysiological characteristics similar to those of resident enteric neurons.

The wall of the embryonic gut consists largely of undifferentiated mesenchyme at the time of migration of neural crest–derived cells. The present study showed that NSs generated from neural crest–derived cells from both the fetal and the postnatal gut survived, proliferated, migrated, and differentiated into glial cells and a range of neuron subtypes that exhibited neurochemical, morphological, and electrophysiological characteristics similar to those of resident enteric neurons.

The neural circuits controlling motility consist of a variety of subtypes of enteric neurons, including intrinsic sensory neurons, interneurons, cholinergic motor neurons that mediate contraction of the gut wall, and inhibitory motor neurons that mediate relaxation and express NOS (56, 57). For cell therapy for motility disorders, all neuron subtypes will need to be generated from transplanted progenitors and form appropriate connections with each other and with muscle cells. In the present study, grafted ENS progenitor cells gave rise to neurons possessing neurochemical and morphological features similar to those of normal enteric neurons. For example, graft-derived neurons expressed a number

(Figure 6A). Together, these results showed that ENS progenitor cells isolated from the fetal and postnatal mouse gut differentiated into neurons with appropriate electrophysiological properties within the environment of postnatal gut.

Discussion

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The wall of the embryonic gut consists largely of undifferentiated mesenchyme at the time of migration of neural crest–derived cells. The present study showed that both fNS-derived cells and more clinically relevant pNS-derived cells were capable of migrating substantial distances in the gut wall of postnatal mice, consisting of concentric layers of differentiated cells. 4 weeks after transplantation of NSs about 0.25 mm in diameter, fNS- and pNS-derived cells occupied an average of about 9 mm² and 7 mm², respectively, and graft-derived neurites occupied an area around twice the size of the graft-derived cell bodies. Previous studies reported very limited migration of CNS stem cells when transplanted into the stomach of adult mice (4), or when enteric progenitor cells were cocultured with the hindgut of chick or mouse embryos after the gut had been fully colonized with neural crest–derived cells (24, 54). The differences in the extent of migration between the current study and earlier studies might reflect differences between the migratory abilities of CNS- and ENS-derived progenitor cells within the postnatal gut, differences between gut regions in their ability to permit migration of stem/progenitor cells, differences between the environment of the late embryonic gut versus the postnatal gut, and/or differences of the environment of the gut in vivo and in culture. Although we showed that the ENS progenitor cells migrated extensively in the postnatal mouse gut, stem/progenitor cells will need to be introduced in greater numbers and at multiple sites to treat humans with enteric neuropathies due to the larger size.

Previous studies have shown that ENS NSs, like their CNS counterparts, are composed of a mixture of progenitors and differentiated progeny (neurons and glial cells) (55). To examine whether some of the graft-derived neurons we detected were the progeny of transplanted cells, we injected EdU into the recipient mice immediately after transplantation of the NSs. 4 weeks later, Hu+ neurons that had incorporated EdU were readily encountered, which showed that at least some neurons were generated in vivo from transplanted proliferating progenitors.
of markers characteristic of subtypes of enteric neurons in mice, humans, and other species, including ChAT, VACHT, NOS, calbindin, and calretinin (43, 58–60). These markers are not, however, expressed exclusively by neurons in the ENS, but are also expressed by some classes of neurons elsewhere in the nervous system. Importantly, most of the graft-derived NOS neurons possessed lamellar dendrites, which is notable because enteric NOS neurons possess lamellar dendrites that are the sites of many of their synaptic inputs (61, 62). The proportions of graft-derived neurons expressing NOS and ChAT were similar to those of myenteric neurons in the neighboring region of distal colon. This is reassuring for the generation of an ENS in the aganglionic region of patients with Hirschsprung disease, in which all neuron subtypes will need to be generated (63); however, for enteric neuropathies such as achalasia, in which there is degeneration of specific classes of enteric neurons (3, 56), manipulation of the cells prior to transplantation is likely to be required to bias the differentiation of cells to particular neuron subtypes.

Our study showed that transplanted ENS progenitor-derived cells migrate and settle in locations similar to those occupied by neural crest–derived cells during normal development. Furthermore, varicose, graft-derived neurites were present in the muscle layers and formed close associations with myenteric neurons of the recipient and with other graft-derived neurons. Thus, cues must exist in the postnatal gut that graft-derived neurites can use to navigate to specific targets. However, graft-derived fibers were

**Figure 6**
Morphology and electrophysiological properties of graft-derived neurites. (A–F) Impaled graft-derived neuron 3 weeks after transplantation of a FNS. (A) Low-magnification image showing a biocytin-filled neuron, which had a single long, circumferentially projecting, axon-like process (yellow arrow) that projected for about 0.6 mm in the plane of the myenteric plexus and finished in an expansion bulb, where the process had broken off during tissue preparation. (B) High-magnification image of the neuron in A, showing multiple filamentous (open arrows) and lamellar (filled arrows) dendrite-like processes. (C and D) Single optical section through neuron in A (asterisk), confirming that it expressed KikGR (D) and hence was graft-derived. (E) The neuron fired a single AP at the beginning of a 500-ms depolarizing step current. (F) fEPSPs occurred both spontaneously (open arrows) and were evoked by a single-pulse stimulus (0.6 mA; filled arrow). Membrane potential was held at –82 mV. (G–K) Impaled graft-derived neuron 4 weeks after transplantation of a pNS. (G) The neuron had lamellar dendrite-like processes (white arrows) and a single long, axon-like process (yellow arrow). (H and I) Single optical section through the neuron in G (asterisk), confirming that it expressed KikGR (I). (J) The neuron fired 3 single APs at the beginning of a 500-ms depolarizing step current. (K) A fEPSP was evoked by a single-pulse stimulus (1.4 mA; arrow). Membrane potential was held at –100 mV. Scale bars: 50 μm (A); 10 μm (B–D and G–I).
located in the submucosal plexus (66, 67). It is therefore highly unlikely that the neurons involved in the circuits controlling motility are intrinsic sensory neurons involved in the regulation of motility in small mammals in the myenteric plexus, some of the intrinsic sensory neurons involved in the circuits controlling motility are located in the submucosal plexus (66, 67). It is therefore highly likely that submucosal ganglia are also essential for a fully functioning bowel, not only for the control of motility, but also for water and electrolyte transport. Hence, techniques will need to be developed to generate submucosal ganglia that consist of appropriate functional types of neurons from transplanted cells.

This is the first study to report the electrophysiological properties of neurons generated from neural progenitor cells transplanted into the gut. During ENS development, some cells express pan-neuronal markers, but do not have the electrophysiological properties of neurons (68). However, graft-derived neurons in the postnatal colon had electrophysiological properties similar to those of mature, functional enteric neurons. Importantly, the presence of fEPSPs showed that the graft-derived neurons had integrated into the neuronal circuitry, although we were unable to determine whether the synaptic inputs arose from other graft-derived neurons, from the recipient’s neurons, or both. There are 2 main electrophysiological classes of myenteric neurons in the ENS, S neurons and AH neurons (53, 69). S neurons are uniaxonal and show monophasic repolarization after an AP and fEPSPs in response to fiber tract stimulation. In contrast, AH neurons show a biphasic repolarization and a slow afterhyperpolarizing potential after an AP.

Table 1

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<tr>
<th>Property</th>
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<th>pNS derived</th>
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<tr>
<td>Resting membrane potential (mV)</td>
<td>–27 ± 4</td>
<td>–36 (n = 1)</td>
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<tr>
<td>Input resistance (MΩ)</td>
<td>231 ± 30</td>
<td>316 ± 33</td>
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<tr>
<td>AP amplitude (mV)</td>
<td>43 ± 4</td>
<td>33 ± 4</td>
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<tr>
<td>AP duration (ms)</td>
<td>1.3 ± 0.1</td>
<td>1.4 ± 0.2</td>
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<tr>
<td>Maximum APs (no.)</td>
<td>3 ± 1</td>
<td>3 ± 1</td>
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<tr>
<td>AP threshold (pA)</td>
<td>123 ± 10</td>
<td>78 ± 12</td>
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<tr>
<td>Neurons with spontaneous fEPSPs (%)</td>
<td>74 (17 of 23)</td>
<td>82 (31 of 38)</td>
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<tr>
<td>Neurons with evoked fEPSPs (%)</td>
<td>100 (23 of 23)</td>
<td>100 (23 of 23)</td>
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<tr>
<td>Evoked fEPSP maximum amplitude (mV)</td>
<td>30 ± 2</td>
<td>27 ± 2</td>
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<tr>
<td>Evoked fEPSP total duration (ms)</td>
<td>57 ± 5</td>
<td>83 ± 13</td>
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For evoked fEPSPs, membrane potential was held at –100 mV. AP duration was calculated as time to decay from maximum amplitude to half amplitude (APD50).

Because AH neurons are not generated from transplanted progenitor cells in the postnatal colon, or because neurons do not develop AH characteristics until 3–4 weeks after transplantation, when all of the electrophysiological experiments were performed.

The neural circuitry controlling motility involves precise connectivity among different functional classes of neurons (26, 70). Although we showed that graft-derived neurons projected to the correct gut layers, it remains to be determined whether each of the different neurochemical types of graft-derived neurons formed appropriate synaptic connections. Also important is that graft-derived neurons do not make incorrect synaptic connections. However, while no gut motility studies were performed, the recipient animals survived and did not exhibit any obvious signs of motility defects, such as stool retention, which would be suggestive of inappropriate circuitry.

There were no significant differences in the behavior of cells derived from fNSs and pNSs. This is important because patient-derived cells are an accessible source of cells to treat congenital motility disorders such as Hirschprung disease (13), and their use will avoid immune responses and the ethical issues associated with some sources of stem/progenitor cells (10). It remains to be determined whether cells derived from NSs generated from the adult gut have post-transplantation properties similar to those of fNS- and pNS-derived cells.

Our data lay the foundation for studies in which ENS progenitors are transplanted into animal models of enteric neuropathies to determine whether graft-derived neurons ameliorate the motility defects. In preliminary studies, we transplanted genetically labeled (EdnrB<sup>cre</sup>) fNSs into the aganglionic region of colon of postnatal f<sup>/</sup>f<sup>/</sup> mice, a mouse model of Hirschsprung disease (71). However, very few transplanted cells survived beyond 1 week due to immunological rejection, as the EdnrB<sup>cre</sup> mice are on a different genetic background from f<sup>/</sup>f<sup>/</sup> mice. As a result, detailed studies

Table 2

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<tr>
<td>Calbindin</td>
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<sup>a</sup>V. Lennon, Mayo Clinic, Rochester, Minnesota (74).<sup>b</sup>P. Emson, Cambridge University, Cambridge, United Kingdom (75, 76).<sup>c</sup>C. Smith, Murdoch Childrens Research Institute, Melbourne, Australia (77).
have had to be postponed until the Ednrb<sup>Kik</sup> mice are backcrossed onto the same genetic background as the s/s<sup>2</sup> mice. Nonetheless, we performed some preliminary experiments in which NSs were generated from N4 backcrossed mice and implanted into the aganglionic region of s/s<sup>2</sup> mice. After 4 weeks, graft-derived cells were present, some of which had migrated away from the transplant site and formed clusters of Hu<sup>+</sup> cells (Supplemental Figure 4, A and B), and graft-derived neurites were abundant in the circular muscle layer. Furthermore, electrophysiological recordings from 2 briefly impaled graft-derived neurons revealed EPSPs (Supplemental Figure 4C). These preliminary data showed that cells transplanted into the aganglionic region survived and migrated in the absence of endogenous enteric neurons and that graft-derived neurons received synaptic inputs. Our findings of immunological rejection after transplantation of cells between mouse strains strongly suggest that patient-derived cells will be the best source of enteric neurons to transplant into patients with enteric neuropathies. Furthermore, although our data using postnatal donor and recipient mice support the idea that cell therapy might be used to treat pediatric enteric neuropathies, additional studies in which cells isolated from the adult mouse gut are transplanted into adult mice are required to demonstrate proof of principle that cell therapy might also be used to treat adult enteric neuropathies.

In conclusion, the ability of ENS stem/progenitor cells to proliferate, migrate extensively, differentiate into neurons of the appropriate phenotype, associate closely with endogenous enteric neurons, and incorporate into the neuronal circuitry in postnatal colon suggests that cell therapy to replace the diseased ENS in postnatal colon is a distinct possibility.

### Methods

**Animals.** Postnatal wild-type mice and E13.5/E14.5 and P4 Ednrb<sup>Kik</sup> and Ret<sup>TGM</sup> mice, all on a C57BL/6 background, were used. In Ednrb<sup>Kik</sup> mice, all neural crest–derived cells within the embryonic gut express the fluorescent photoconvertible protein Kikume, under the control of an enteric-specific region of the Ednrb promoter (40). Ret<sup>TGM</sup> mice have had cDNA encoding tau-GFP-myc (TGM) inserted into the first coding exon of the Ret gene (72), and all neural crest–derived cells in the embryonic gut express EGFP (41). The genotype of adult Ednrb<sup>Kik</sup> and Ret<sup>TGM</sup> mice were determined by PCR using primers and conditions reported previously (40, 72). Timed pregnant mice were killed by cervical dislocation. The morning on which a copulatory plug was observed was designated E0.5.

**Generation of Ednrb<sup>Kik</sup>- or Ret<sup>TGM</sup>-positive ENS NSs.** Wild-type female mice were plug-mated to Ednrb<sup>Kik</sup> or Ret<sup>TGM</sup> heterozygote males. The entire gut, from the stomach to the anus, of E13.5/E14.5 and P4 mice was dissected and screened. E13.5/E14.5 gut was dissociated in 0.1% trypsin/EDTA (GIBCO, Invitrogen) at 37°C for 20 minutes, with gentle pipetting. P4 gut was dissociated in 0.5% Dispase II (Roche Applied Science) and 0.05% Collagenase CLS AFA (Worthington Biochemical Corp.) at 37°C for 30 minutes, with gentle pipetting. Medium containing 10% fetal calf serum was then added, and the cell suspension was passed through a 40-μm cell strainer (BD Biosciences). The cell suspension was centrifuged at 850 g in a bench centrifuge for 2 minutes, the supernatant was removed, and the pellet was resuspended. The Ednrb<sup>Kik</sup>- or Ret<sup>TGM</sup>-positive cells were isolated by flow cytometry (MoFlo, Beckman Coulter) and sorted into round-bottomed, low-attachment, sterile 96-well plates (Corning, Costar) at a density of 10,000 cells/well (or 50,000 cells/ml). The cells were aggregated by centrifugation at 480 g for 3 minutes at 4°C, as described previously (73) and then were cultured in DMEM/F12 (GIBCO, Invitrogen) containing 1% l-glutamine (Sigma-Aldrich), 1% penicillin/streptomycin, 1x B-27 supplement (GIBCO, Invitrogen), 1x N-2 supplement (GIBCO, Invitrogen), and 20 ng/ml EGF and bFGF in a humidified incubator at 37°C and 5% CO<sub>2</sub> for 7 days to allow for NS formation.

**In vitro migration and differentiation of Ednrb<sup>Kik</sup>- and Ret<sup>TGM</sup>-positive ENS stem/progenitor cell NSs.** Ednrb<sup>Kik</sup>-positive NSs were transferred to glass-bottomed chamber slides coated with 20 μg/ml fibronectin (Sigma-Aldrich) and cultured for a further 2 days in NS growth medium before being fixed in 4% paraformaldehyde in 0.1M phosphate buffer (PB) and processed for immunohistochemistry.

**In vivo transplantation of Ednrb<sup>Kik</sup>- or Ret<sup>TGM</sup>-positive stem/progenitor cell NSs to the colon of postnatal mice.** Recipient wild-type mice (2–3 weeks of age) were anesthetized by subcutaneous injection of a mixture of 20 mg/kg xylazine (Troy Laboratories) and 100 mg/kg ketamine hydrochloride (Troy Laboratories). A midabdominal incision was made, and the distal colon was exposed. 2 or 3 NSs, dyed by brief exposure to 0.1% trypan blue in PB, were transplanted into the external muscle layer of the distal colon. At 1, 2, 4, 8, 12, or 16 weeks after surgery, recipient mice were killed by cervical dislocation, and the distal colon was removed.

**Fixation and tissue preparation.** For whole-mount preparations, the distal colon was opened along the mesenteric border, pinned, stretched on Sylgard-coated dishes, and fixed in 4% paraformaldehyde in 0.1M phosphate buffer (PB) overnight. The tissue was then washed and the mucosa removed. For cryosections, the distal colon was opened along the mesenteric border, loosely pinned to a Sylgard dish, and then fixed in 4% paraformaldehyde in 0.1M PB overnight. The tissue was washed and transferred to 5% sucrose in 0.1M PB, then 1:1 OCT/sucrose, and then transferred to cryomold containing OCT (TissueTek). Frozen sections (12 μm thick) were cut transversely on a cryostat.

**Immunohistochemistry.** Cells or tissues were fixed overnight in 4% paraformaldehyde in 0.1M PB at 4°C, exposed to 0.1% Triton X-100 for 30 minutes, and then exposed to primary and secondary antibodies (Tables 2 and 3). Note that an antibody was not used to reveal KikGR-expressing cells; only the native KikGR protein was observed. Preparations were viewed on a confocal microscope. Segments of control intestine (lacking transplanted cells) did not contain any fluorescent cells in the external muscle layers.

**EdU studies.** For 5 consecutive days after transplantation of NSs, mice were injected intraperitoneally with 50 mg/kg EdU. 4 weeks later, mice were killed by cervical dislocation, and the colon was dissected, pinned out, and
fixed overnight in 4% formaldehyde at 4°C. The mucosa was removed, and EdU was detected using the Click-iT EdU Imaging Kit (Invitrogen) according to the manufacturer’s instructions. The axon guide in this detection reaction, which covalently binds the alkyne group associated with the incorporated EdU, was coupled to Alex Fluor 647. The preparations were also processed for immunohistochemistry using a human anti-Hu.

**Measurement of area occupied by graft-derived cells.** To determine the area occupied by graft-derived cells plus fibers or by cells only, tile scans of whole-mount preparations of recipient colon were taken using 5× or 10× objectives on a confocal microscope. The total area occupied by graft-derived cells plus fibers, or cells only, in each preparation was measured using Image J software.

**Electrophysiology.** Segments of distal colon were removed and immediately placed in physiological saline (118 mM NaCl, 25 mM NaHCO₃, 11 mM tri-glucose, 5 μM KCl, 2.5 mM CaCl₂, 1.2 mM MgSO₄, 1.0 mM Na₂HPO₄, 25 μM nicardipine, and 1 μM hyoscine) bubbled with 95% O₂ and 5% CO₂. The region of distal colon containing EdnrB°-positive grafted cells was cut along the mesenteric border and pinned flat, mucosa side up, in an organ bath lined with a silicone elastomer (Sylgard 184; Dow Corning). The mucosa and submucosa were dissected and removed from the underlying smooth muscle and myenteric plexus layers. The preparation was continually superfused with physiological saline (33°C–34°C) and left to equilibrate for 1 hour.

Standard intracellular recording methods (52) were used to impale and record from EdnrB°-positive grafted cells. Intracellular microelectrodes (100–200 MΩ) containing 1M KCl and 2% biocytin (Sigma-Aldrich) were used. Electrical stimuli of a single pulse (0.4–1.8 mA) or trains of stimuli (3, 10, or 15 pulses) was applied via a focal stimulating electrode positioned on interganglionic fiber tracts about 200 μm oral to the impaled cell region to determine whether the grafted cells display synaptic potentials. The excitability of the grafted cells was examined by holding the membrane potential at ~60 mV and applying depolarizing current pulses (500 ms duration) in 10-pA increments over a range of 50–300 pA. Input resistance of the grafted cells was measured from hyperpolarizing current pulses (500 ms, 10-pA increments, 100–300 pA). After electrophysiology, the preparations were fixed in 4% formaldehyde and processed to reveal the impaled neurons (Streptavidin Alexa Fluor 594, 1:200; Invitrogen). Hyoscine and hexamethonium (Sigma-Aldrich) and were prepared as stock solutions dissolved in distilled water, and diluted to their final concentrations before usage in experiments. The amplitude of all the peaks (or maximum peak amplitude, where indicated) and total duration of the stimulated fEPSP complexes were measured. The number of APs triggered by depolarizing current pulses was counted. The amplitude of APs was measured, and the duration of an AP was measured as the time to decay from maximum amplitude to half amplitude.

**Statistics.** Data are displayed as mean ± SEM and were analyzed using 2-tailed t tests. A P value less than 0.05 was considered significant.

**Study approval.** All studies were approved by the Anatomy and Neuroscience, Pathology, Pharmacology, and Physiology Animal Ethics Committee of the University of Melbourne (ethics ID 0911131).

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