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Enteric Neural Cells From Hirschsprung Disease Patients Form Ganglia in Autologous Aneuronal Colon

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**Publication Details**

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Abstract

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Methods: Cells were obtained and cryopreserved from 31 HSCR patients from the proximal resection margin of colon, and ENS cells were isolated using flow cytometry for the NC marker p75 (nine patients). Aneuronal colon tissue was obtained from the distal resection margin (23 patients). ENS cells were assessed for NC markers immunohistologically and by quantitative reverse-transcription polymerase chain reaction, and mitosis was detected by ethynyl-2'-deoxyuridine labeling. The ability of human HSCR postnatal ENS-derived cells to colonize the embryonic intestine was demonstrated by organ coculture with avian embryo gut, and the ability of human postnatal HSCR aneuronal colon muscle to support ENS formation was tested by organ coculture with embryonic mouse ENS cells. Finally, the ability of HSCR patient ENS cells to colonize autologous aneuronal colon muscle tissue was assessed.

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SUMMARY

We show that enteric neural cells isolated from Hirschsprung disease patients can colonize autologous colon tissue to generate neurons and glia. Our findings establish the therapeutic potential of using patient’s own neural cells to form an enteric nervous system in autologous tissue.

BACKGROUND & AIMS: Hirschsprung disease (HSCR) is caused by failure of cells derived from the neural crest (NC) to colonize the distal bowel in early embryogenesis, resulting in absence of the enteric nervous system (ENS) and failure of intestinal transit postnatally. Treatment is by distal bowel resection, but neural cell replacement may be an alternative. We tested whether aneuronal (aganglionic) colon tissue from patients may be colonized by autologous ENS-derived cells.

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CONCLUSIONS: NC-lineage cells can be obtained from HSCR patient colon and can form ENS-like structures in aneuronal colonic muscle from the same patient. (Cell Mol Gastroenterol Hepatol 2016;2:92–109; http://dx.doi.org/10.1016/j.jcmgh.2015.09.007)

Keywords: Hirschsprung Disease; Aganglionosis; Cell Therapy; Megacolon; Enteric Nervous System.
enteric neural crest (ENC) cells migrate along the entire tract in a distal-directed wave over the 4th to 7th weeks of gestation in humans.8–10 Experiments in animal models indicate that the sacral level of the NC also provides neurons to the gut,11,12 but these are numerically less capable13 and cannot compensate for a lack of vagal NC-derived cells.14,15 Recently a late-arising population of ENS neurons has been identified in mice derived from Schwann cell intermediaries; these are a minority source and give a restricted range of neuron types.16 HSCR results when this wave does not complete colonization of the entire gut. Cell-replacement therapy postnatally after HSCR diagnosis for a defect of early embryonic origin faces difficulties. ENC stem or progenitor cells would have to be obtained and expanded before transplantation; then they would have to be introduced into the aneuronal gut without immune rejection. They must then migrate, colonize, differentiate, and establish connections to mediate bowel function at later stages and in a far larger tissue than would occur during normal ENS development.

Rodent and avian embryonic ENC progenitor cells can colonize embryonic aneural gut and form ENS in vitro.17–20 Further, rodent embryo NC cells can colonize the aneural gut and differentiate into neurons and glia of postnatal HSCR model mice in vivo.21,22 In addition, it has been shown that postnatal mouse ENC cells surgically placed in the wall of the colon of postnatal normal and HSCR-model syngeneic mice are able to migrate, proliferate, assemble into correctly placed ganglia and differentiate as both glia and neurons that extend axons, make and receive synapses, and show electrical activity.23 This is a key step to answering the therapeutic questions of whether ENS formation can be accomplished when both ENS-lineage donors and colon tissue recipients are postnatal. A further question is, can this be accomplished when both ENC and colon are not only of postnatal human origin, but when (to avoid immune rejection) they are of autologous HSCR patient origin?

We obtained human ENS-derived cells from the proximal margin of HSCR patient colon using flow cytometry for the NC marker p75. These were assessed for additional NC markers by antibody labeling and quantitative reverse-transcription polymerase chain reaction (qRT-PCR), and for proliferation with ethynyl-2′-deoxuryridine (EdU). We tested the ability of these cells to migrate in embryonic aneural gut in organ culture by combining the human ENS-derived cells with quail embryo intestine,24 a tissue known to support cell colonization. We then ascertained the ability of human postnatal HSCR aneural distal colon muscle tissue to support ENS formation by, in organ culture, transplanting into it genetically labeled ENC-derived cells from embryonic mice, cells with proven ENS-forming ability.25 We then labeled human p75-sorted cells using fluorescent tags and implanted them into distal colonic muscle tissue from the same HSCR patients and maintained the combinations in organ culture. The results show that NC-lineage ENS cells can be isolated from neuronal regions of HSCR patient postnatal colon, and that these ENS-derived cells can spread, proliferate, and differentiate in receptive intestinal tissues including aneural HSCR colon muscle from the same patient.

### Materials and Methods

#### Human Tissues

Colon tissue was obtained immediately after surgery for HSCR at the Royal Children’s Hospital (RCH). In most cases (24 of 31), the patients were younger than 4 months at surgery, the length of resected bowel was less than 12 cm (27 of 31), and no other clinical conditions were identified (Table 1). Familial forms of HSCR (patients with first- and second-degree relatives affected) were reported in a small number of cases (4 of 31). After obtaining tissue from the proximal (neuronal) and distal (aneuronal) ends, the remainder of resected bowel tissue was histologically examined at RCH Anatomical Pathology. Samples of the muscle layers of both ends of all specimens were fixed in 4% paraformaldehyde (PFA), embedded in optimal cutting temperature (OCT) compound (Tissue-Tek, Sakura Finetek Europe, Alphen aan den Rijn, Netherlands), frozen, and sectioned (CM1900 cryostat; Leica Microsystems, Wetzlar, Germany) for immunolabeling. The procedures for resected tissues are described here, and a flow chart is shown in Figure 1.

Human tissue was collected under approval from the Royal Children’s Hospital (Parkville, Melbourne, Australia) Human Research Ethics Committee (HREC 30014A). Parent/guardian consent was obtained for all participants. Animal tissues were obtained under the conditions of the Murdoch Children’s Research Institute Animal Ethics Committee (AEC651).

#### Human Colon Tissue Dissociation

To obtain human ENS cells from myenteric plexuses between proximal colon muscle layers, the mucosa was removed by scraping, and the muscle layer was cut into 1–2 mm³ fragments. Dissociation was achieved in Ham’s F-12 media (GIBCO/Invitrogen, Grand Island, NY) supplemented with 0.5% w/v Dispase II (Roche, Basel, Switzerland) and...
0.1% w/v CLSAFA collagenase (Worthington Biochemical, Lakewood, NJ) at 37°C with orbital shaking and mechanical trituration. After 1 hour, the initial cell suspension was decanted, and the remaining tissue pieces were dissociated for an additional hour. We added 1 mg/mL hyaluronidase Type I-S (Sigma-Aldrich, St. Louis, MO), 0.1 mg/mL DNAase I (Roche), and 1 mM EDTA (Sigma-Aldrich Australia, Sydney, Australia) to the pooled cell suspension for 15 minutes. The cell suspension was centrifuged at 400 g in a bench centrifuge for 5 minutes, the supernatant was removed, and the pellet was resuspended. The cell suspension was washed in Ham’s F-12 medium with 5% v/v heat-inactivated fetal bovine serum (FBS) (Thermo Fisher Scientific, Waltham, MA) and filtered through a 40-μm cell-strainer (Falcon/BD Biosciences, Bedford, MA).

Culture of Human Hirschsprung Disease Colon Cells

Dissociated cells from the muscle layers, including ENS cells, were grown on tissue culture plates (Fisher Scientific, Roskilde, Denmark) pretreated with 20 μg/mL human fibronectin (Roche, Switzerland). To maximize spontaneous aggregation, fibronectin was reduced or omitted in some cultures. Cells were grown in a 1:1 mixture of Dulbecco’s modified Eagle medium (Thermo Fisher Scientific) and Ham’s F-12 (GIBCO/Invitrogen) with l-glutamine, B27 and N2 (GIBCO/Invitrogen), 10 ng/mL human recombinant fibroblast growth factor 2 (basic) (FGF2) and 10 ng/mL human epidermal growth factor (both R&D Systems, Minneapolis, MN), and 10 U/mL penicillin, 100 μg/mL streptomycin (GIBCO/Invitrogen), 50 μg/mL gentamicin (Sigma-Aldrich Australia), and 50 μg/mL metronidazole (Sigma-Aldrich Australia), this is referred to as nTCM. In some cases nTCM was supplemented with 3 μM glycogen synthase kinase 3 (GSK3) inhibitor CHIR-99021 (6-[2-[[4-(2,4-dichlorophenyl)-5-(5-methyl-1H-imidazol-2-yl)pyrimidine-2-yl]amino]ethylamino]pyridine-3-carbonitrile; Tocris Bioscience/R&D Systems, Minneapolis, MN), which simulates Wnt signaling.25

Cell Sorting and Labeling of Human Enteric Nervous System-Derived Cells

After 1-7 days of culture, adherent cells were lifted with 0.025% w/v trypsin (Sigma-Aldrich) and stained for 45
minutes with mouse p75/CD271:Alexa 647 antibody (BD Biosciences/Pharmingen, San Diego, CA) (see Figure 1). Cells were washed in 5% FBS in Ham’s F-12, and 10 μg/mL propidium iodide (Sigma-Aldrich) was added to detect dead cells. Cells staining positive for p75 (647: bandpass filter 660/20) and negative for propidium iodide (bandpass filter 692/40) were isolated by flow cytometry using a 1-drop pure mode (MoFlo, Beckman Coulter, Brea, CA; or Influx, BD Biosciences, San Jose, CA) and a sheath pressure of 22 psi with a 100-μm diameter nozzle and Dulbecco’s phosphate-buffered saline (PBS) sheath fluid (prepared in house).

To observe human cells in human colon explants, we incubated the p75+ cells for 20 minutes in 10 nM MitoTracker Red (MTR) CMXRos (Life Technologies, Carlsbad, CA) at 37°C, then washed twice in nTCM. Microscopy confirmed that all cells contained MTR. In some cases the p75+ cells were transduced with lentiviral particles carrying a constitutive enhanced green fluorescent protein (eGFP) reporter construct (PLL3; CMG-eGFP).

To generate lentiviral particles, human embryonic kidney 293 (HEK293) cells were transfected by Fugene HD (Promega, Madison, WI) with constructs pBR8.91 and pMDG and pLL3. The supernatant containing viral particles was collected after 2 days and then applied to p75+ cells for 24 hours. We found that the addition of 10% fetal calf serum (Thermo Fisher Scientific) to the cell culture medium aided in the survival of p75+ cells after lentiviral infection. After 4 days of culture, the cells that were positive for both p75+ and eGFP (p75+/eGFP+) were isolated by flow cytometry as described earlier (eGFP: bandpass filter 530/40). Both p75+/MTR and p75+/eGFP+ cells were spin aggregated, as we will describe herein.

**Cell Sorting for Mouse Enteric Nervous System–Derived Cells**

Mouse ENS Kikume+ cells were obtained by flow cytometry (Kikume: bandpass filter 530/40) from embryonic day E13.5–E14.5 Ednrbkil mice on a C57Bl/6 background.23,26 In these mice, all ENS cells express the fluorescent protein Kikume under the control of an enteric-specific region of the Ednrb promoter. Matings were conducted as described elsewhere.23 The gut from the stomach to the anus was dissected and screened by fluorescence microscopy to confirm the presence of labeled ENS. To collect mouse ENS cells, the gut was dissociated in 0.1% trypsin/EDTA (GIBCO/Invitrogen) at 37°C for 20 minutes, with gentle pipetting.

**Generation of Spun Enteric Nervous System Cell Aggregates**

After flow cytometry ENS cells (human: 5000; mouse: 10,000) were deposited into round-bottom, low cell-adherence plastic 96-well plates (Corning Life Sciences, Tewksbury, MA). The cells were aggregated by centrifugation at 400g for 5 minutes, and maintained for 1–2 days in normal culture conditions before use in coculture.
experiments. In most cases ENS cells formed spheres after overnight culture in low-adherence conditions.

**Avian Aneural Intestinal Organ Culture**

Fertilized quail (Coturnix japonica) eggs were obtained from Lago Game Supplies (Victoria, Australia) and were incubated at 38°C. Postumbilical gut was dissected from 4-day embryos to act as recipients for ENS cell immigration in organ culture. At this stage, this gut region is not invaded by ENC cells and is therefore aneural. Human ENS p75+ cell aggregates were applied to the proximal end of the gut, and the ensemble was maintained in culture in Ham’s F-12 medium supplemented with 10% FBS, L-glutamine, and 10 U penicillin and 100 µg/mL streptomycin. After 4 days, the explants were fixed and underwent immunohistochemical staining as whole mounts.

**Human Aneuronal Colon Explant Coculture With Enteric Nervous System Cells**

Human aneuronal colon muscle was obtained from patients as already described, but from the distal end of the resected bowel. The submucosa was removed, and the muscle was cut into 2–8 mm³ fragments containing both longitudinal and circular muscle layers; these layers were clearly distinguishable with a dissecting microscope (Leica MZ6; Leica Microsystems). These could be maintained for at least 12 days in culture in nTCM supplemented with 5% FBS at 37°C and 5% CO2 without alterations in cell nuclear morphology or in mesodermal smooth muscle, detected with Hoechst 33342 (5 µg/mL) and smooth muscle actin (SMA) antibody (see below), respectively. For combination explants, these human muscle tissues were cocultured with either EdnrbKik mouse or human ENS cell aggregates. These aggregates were inserted in a pocket created between the two muscle layers. These were maintained in culture for up to 7 days and monitored for Kikume expression for mouse/human combinations and MTR or eGFP for human/human combinations. Explants were fixed for 1 hour at 4°C in 4% PFA for whole-mount immunolabeling or tissue sectioning.

**Ethynyl-2’-Deoxyuridine Labeling and Cell Proliferation Analysis**

Human colon monolayer cultures and p75-sorted ENS cells before aggregation were treated with 5 µM EdU

<table>
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<tr>
<th>Cell Target</th>
<th>Identity</th>
<th>Type</th>
<th>Source</th>
<th>Product No.</th>
<th>Research Resource Identifiers (RRID)</th>
</tr>
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<tr>
<td>NC cell surface</td>
<td>p75</td>
<td>Rabbit IgG</td>
<td>Promega</td>
<td>G3231</td>
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</tr>
<tr>
<td>NC cell surface/FACS</td>
<td>p75: Alexa 647, clone c40-1457</td>
<td>Mouse IgG</td>
<td>BD Pharmaningen</td>
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<td>AB_2033986</td>
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<tr>
<td>NC cell surface</td>
<td>HNK1</td>
<td>Mouse IgM</td>
<td>MCRI</td>
<td>NA</td>
<td>NA</td>
</tr>
<tr>
<td>NC/glial cell nucleus</td>
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<td>Goat IgG</td>
<td>R&amp;D Systems</td>
<td>AF2864</td>
<td>AB_442208</td>
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<tr>
<td>Glial cytoplasm</td>
<td>GFAP</td>
<td>Rabbit IgG</td>
<td>Dako, Denmark</td>
<td>Z0334</td>
<td>AB_2314535</td>
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<tr>
<td>Glial cytoplasm</td>
<td>S100β, clone SH-B1</td>
<td>Mouse IgG</td>
<td>Sigma-Aldrich, Australia</td>
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<td>AB_477499</td>
</tr>
<tr>
<td>Neuron nucleus/cytoplasm</td>
<td>Hu</td>
<td>Human IgG</td>
<td>Vanda Lennon, Rochester, MN</td>
<td>NA</td>
<td>NA</td>
</tr>
<tr>
<td>Nitrergic neuron</td>
<td>nNOS</td>
<td>Sheep IgG</td>
<td>EMSON</td>
<td>H212</td>
<td>AB_2314957</td>
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<td>Nerve fiber</td>
<td>Neuronal class III β-tubulin clone (TUJ1)</td>
<td>Mouse IgG</td>
<td>Biolegend/Covance</td>
<td>MMS-435P</td>
<td>AB_2315514</td>
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<td>MitoTracker CMXRos</td>
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<td>Life Technologies</td>
<td>M7512</td>
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</table>

Secondary antibodies and probes

| Human IgG | Texas Red | Donkey IgG | Jackson ImmunoResearch | 709 075 098 | AB_2532064 |
| Mouse IgM | Texas Red | Donkey IgG | Jackson ImmunoResearch | 715 065 140 | AB_2340783 |
| Mouse IgG | Alexa 594 | Donkey IgG | Jackson ImmunoResearch | A21203 | AB_10563558 |
| Mouse IgG | Alexa 487 | Donkey IgG | Jackson ImmunoResearch | A21235 | AB_10562370 |
| Goat IgG | Alexa 488 | Donkey IgG | Jackson ImmunoResearch | A11015 | AB_10561557 |
| Goat IgG | Alexa 594 | Donkey IgG | Jackson ImmunoResearch | A11058 | AB_142540 |
| Goat IgG | Biotin | Donkey IgG | Jackson ImmunoResearch | 705 065 147 | AB_2340397 |
| Biotin | AMCA | Streptavidin | Jackson ImmunoResearch | 016 130 084 | AB_2337243 |

AMCA, methyl N-(4’-(9-acridinylamino)-phenyl) carbamate hydrochloride; GFAP, glial fibrillary acidic protein; HNK1, human natural killer-1; NA, not applicable; NC, neural crest; nNOS, neuronal nitric oxide synthase; SMA, smooth muscle actin; SOX10, sex-determining region Y-box 10.
(GIBCO/Invitrogen) for 12 hours at 37°C and washed twice in PBS. To quantify the proliferation of cells positive for sex-determining region Y-box 10 (SOX10) in the nTCM and CHIR-99021 groups, the EdU-labeled human colon monolayer cultures were fixed in 4% PFA for 10 minutes and stained for SOX10 (see below), and EdU-labeled cells were detected by the Click-iT EdU Imaging Kit (GIBCO/Invitrogen) according to the manufacturer’s instructions. Ten microscopic fields were visualized from each experimental group (nTCM with and without CHIR-99021 with cells from two HSCR patients), and the proportion of SOX10+/EdU+ cells was expressed as the ratio of total SOX10+ cells counted in each field. Human p75-sorted cell aggregates were used for coculture with human aneuronal colon, where incorporated EdU also served as a cell tracker. Mouse ENS-derived cell aggregates (Kikume+) in coculture with human aneuronal colon muscle explants were pretreated for 8 hours with 5 μM EdU during days 1–2 of the culture period.

**Gene Expression Analysis**

Total RNA was isolated from cells using RNeasy mini kit (Qiagen, Valencia, CA), and contaminant genomic DNA was removed with DNA-free reagents (Ambion/Life Technologies, Austin, TX). Primer sequences were designed using Primer3 (http://frodo.wi.mit.edu/primer3/) and are listed in Table 2.
Where possible, the primers sets were designed to span introns to discriminate between amplification of genomic DNA and cDNA. For qRT-PCR, oligo-dT primed cDNA was synthesized from 200 ng of total RNA using Murine Moloney Leukaemia Virus reverse transcriptase (Promega). We performed the qRT-PCR on an ABI Prism 7500 Real-Time PCR System using SYBR green master mix (Applied Biosystems, Foster City, CA) according to the manufacturer’s protocols. Relative gene expression values were obtained by normalization to the reference gene L32 using the $-2^{\Delta\Delta Ct}$ method, where $-2^{\Delta\Delta Ct} = \Delta Ct$ sample $- \Delta Ct$ calibrator as described elsewhere.28
Immunolabeling

Cells and tissues were fixed in 4% PFA for 5 and 60 minutes, respectively. The monolayer cultures and whole mounts were permeabilized and blocked in 0.1% Triton X-100/1% horse serum in PBS for 15 and 60 minutes, respectively. Tissue sections were blocked in 1% horse serum in PBS for 30 minutes. For immunolabeling, primary antibodies recognized NC cell markers (p75, HNK1 [human natural killer-1], SOX10), glial markers (GFAP [gliarial fibrillary acidic protein], S100β, SOX10), the neuronal marker (Hu), neuronal nitric oxide synthase (nNOS), nerve fibers (TUJ1, neuron-specific class III β-tubulin), and the muscle marker SMA. Secondary antibodies appropriate for these were coupled to AMCA (methyl N-(4′-9-acridinylamino)-phenyl) carbamate hydrochloride), Alexa 647, 594, 488, and Texas Red. A complete list of these antibodies is in Table 3. For monolayer culture and tissue sections, antibodies were incubated for 2 hours. For whole mount, antibodies were incubated overnight. Cell nuclei were counterstained with 5 μg/mL Hoechst 33342 (Invitrogen/Life Technologies, Carlsbad, CA), which labels all cell nuclei. Staining was visualized using a Leica SP2 confocal microscope (Leica Microsystems) or LSM 780 confocal microscope (Carl Zeiss Light Microscopy, Göttingen, Germany). Post-acquisition image analysis was performed by LAS AF light software (Leica Microsystems) or ZEN Blue software (Carl Zeiss Light Microscopy).

Quantification of Cell Types in Aggregates

Cell types present in spontaneously forming aggregates was detected by immunolabeling aggregates for SOX10, p75, and Hu and then counterstaining with Hoechst 33342. For each labeled sphere, a Z-stack confocal microscopy series was generated with optical sections taken every 10 μm. Three optical sections for each sphere were assessed and for cells staining positive for SOX10+/p75−,

Figure 4. Human enteric nervous system (ENS)–derived cells proliferate in vitro. (A) Incorporation of ethynyl-2′-deoxyuridine (EdU) (white arrowheads) into SOX10+ ENS cells with and without CHIR-99021. (B) Proliferation of SOX10+ cells is significantly increased by CHIR-99021, expressed as a ratio of EdU+/SOX+ cells to total SOX10+ cells. (C) Proliferation occurs in putative progenitor cells (EdU+/SOX10+/S100β-ve; white arrowheads), but was not detected in neurons (HU+/SOX10-ve; open arrowhead) during the culture period. Images acquired using confocal microscopy. Data derived from two Hirschsprung disease (HSCR) patient samples.
Hu\(^{+}\), or Hoechst 33342 were counted. Ten spheres were counted for each treatment group (nTCM/CHIR-99021, n = 2 patients). Data were represented as a percentage of SOX10\(^{+}\)/p75\(^{+}\) or Hu\(^{+}\) cells of all cells present (Hoechst 33342\(^{+}\)).

**Statistical Analysis**

Data were analyzed with GraphPad Prism version 5.1 software (GraphPad Software, San Diego, CA). Data were expressed as mean ± standard error of the mean (SEM). Statistical analysis was performed by one-way analysis of variance or Kruskal-Wallis test and the appropriate parametric (Student t test) or nonparametric (Mann-Whitney) post test. \(P < .05\) was considered statistically significant.

**Results**

**Enteric Nervous System Cells Can Be Identified in the Hirschsprung Disease Patient Colon and Persist in Organ Culture**

Serial sections of the distal aneuronal segment of HSCR colon from four HSCR patients were examined. These contained no Hu\(^{+}\) neurons, but p75\(^{+}\), SOX10\(^{+}\), and S100\(\beta\)\(^{+}\) cells were plentiful. The neural elements of the aneuronal segment consisted of large bundles of TUJ1\(^{+}\) fibers surrounded by p75\(^{+}\) cells as well as single TUJ1\(^{+}\) fibers (Figure 2A). This confirmed that the HSCR patient distal colon was aneuronal but not aneural. Aneuronal colon muscle maintained for up to 11 days in organ culture maintained expression of these markers, including the persistence of TUJ1\(^{+}\) nerve fibers (see Figure 2B). In sections of proximal neuronal colon from four HSCR patients, ganglia within the myenteric plexus could be identified containing the NC markers p75, SOX10 and HNK1 (not shown), neuronal (HU) and neurite (TUJ1; not shown) markers and glia markers (SOX10, GFAP; not shown S100\(\beta\)) (Figure 3A and B). Closely associated with ganglia were putative ENS progenitor cells, which were SOX10\(^{+}\) but lacked neuronal and glial markers, including GFAP (see Figure 3B).

**Human Enteric Nervous System and Mesoderm Cells Can Be Propagated in Vitro**

Cells dissociated from proximal colon (21 HSCR patients) produced monolayers on fibronectin; these showed multiple morphologically distinct cell types (see Figure 3C). Neurosphere-like bodies (see Figure 3D) also appeared in cell cultures, and omission or reduction of substrate fibronectin promoted this. Within the monolayer cultures, ENS cells could be identified by staining for NC surface antigens p75 and HNK1 (see Figure 3E) and by the pan-neuronal marker Hu and glia/progenitor cell markers SOX10 and S100\(\beta\) (see Figure 3E and F). The ENS-derived cells formed clusters of 5–30 cells in a carpet of fibroblast-like cells, many of which stained strongly for SMA (not shown) and were presumed to have originated from smooth muscle cells.
Fluorescence-activated cell sorting analysis (see below) indicated that ENS cells formed <5% of the total cells present in these cultures.

To test whether the proliferation rate of SOX10+ cells could be increased in monolayer cultures, we added the GSK3 inhibitor CHIR-99021 to the nTCM culture medium. This molecule was chosen because inhibition of the GSK3 pathway to mimic canonical Wnt signaling promotes proliferation of central nervous system progenitors in vitro. Additionally, Wnt in combination with bone morphogenetic proteins (BMP) signaling promotes the survival and proliferation of mouse NC progenitor cells. Proliferation of SOX10+ cells in monolayer cultures with nTCM alone was low but could be increased more than 4-fold by the addition of CHIR-99021 (Figure 4A and B). The fibroblast-like cells did not require CHIR-99021 for proliferation. Proliferating ENS cells in the presence of CHIR-99021 were either glia (SOX10+/S100β-) or putative progenitor (SOX10+/S100β+ve) cell types (Figure 4C). Neurons (SOX10-ve/Hu+) could not be identified as differentiating in culture from a proliferating (ie, EdU+) precursor labeled before the prior EdU exposure period (see Figure 4C). Primary colon monolayer cells were routinely passaged and cryopreserved without loss of the ENS cell moiety (not shown).

Neurosphere-like bodies were immunolabeled with SOX10, p75, and Hu (Figure 5) to determine the percentage of ENS and non-ENS cells present. From confocal sections, on average the neurosphere-like bodies were 14.11% (SEM = 3.20%) p75+/SOX10+ cells when grown in nTCM (total 4781 cells sampled) and 12.31% (SEM = 3.34%) when grown with the addition of CHIR-99021 (total 4126 cells sampled). Less than 1% of cells were Hu+ neurons in these neurosphere-like bodies after culture with or without CHIR-99021 (see Figure 5). Although there was slight enrichment for ENS cells in neurosphere-like bodies compared with primary cultures of monolayered colon cells, the major constituents (over 85%) were not ENS cells.

**Enrichment of Human Enteric Nervous System Cells by Flow Cytometry Cell Sorting**

We did not find sorting for p75+ ENS cells effective using cells directly dissociated from the proximal colon. Instead, proximal colon cells were cultured for 1–7 days in monolayer, and then ENS lineage cells were isolated by flow cytometric sorting of p75+ cells (see Figure 1). The average percentage of p75+ cells in colon cultures was 4.82% (SEM = 1.43, from six patients) (Figure 1A). Analysis by qRT-PCR (see Figure 6B) showed an increase of NC gene expression in p75-sorted cells. These were SOX10 (mean 407-fold enrichment, SEM = 205), FOXD3 (mean 165-fold, SEM = 32), AP2 (mean 176-fold, SEM = 90), and p75 (mean 204-fold, SEM = 126) when compared with cells negatively sorted for p75 (negative sort) (n = 4 patients). The p75+ve sorted cells showed enrichment of ENS markers when compared with unsorted cells on the day of tissue dissociation (precultured), or cultured cells on the day of flow cytometry (presort) (see Figure 6B). A comparison of precultured and presort groups revealed that no significant loss of ENS markers occurred during the cell culture period. The p75-sorted ENS cells on replating showed neuronal or multipolar morphologies (see Figure 6C). These cells formed aggregates after centrifugation into low-attachment plates (see Figure 6D). Immunologic staining of these p75+ spin-aggregates confirmed that almost all cells expressed either p75\SOX10 or Hu (see Figure 6D and E).

**Embryonic Avian Aneural Gut Supports Human Enteric Nervous System Cell Colonization**

The avian embryonic aneural gut favors ENC cell colonization, so we used this to test the colonization ability of the postnatal human p75+ ENS-derived cells. When p75+ cell aggregates were combined in explants with aneural quail guts, cells migrated as chains into the gut and expressed SOX10 (n = 4 quail intestines, ENS cells from two HSCR patients) (see Figure 6F). This was similar to migrating avian embryonic ENC cells, but the colonization distance was less; approximately 600 μm in 4 days, compared with >1500 μm for avian ENC cells in similar culture conditions.

**Postnatal Human Aneuronal Hirschsprung Disease Colonic Muscle Supports Mouse Enteric Neural Crest Cell Colonization**

To test whether postnatal human HSCR colon tissue could support colonization by ENC cells, we employed cells...
with demonstrated colonization potential: embryonic ENC cells from EdnrbKik mice. Aggregates of these cells were combined with human HSCR aneuronal colonic muscle in vitro (n = 2 patients, eight explants). The diameter of the mouse ENS-derived spheres were approximately 150 μm at the time of combination (Figure 7A); however, after 1 week the ENC cells had migrated greater than 1 mm radially over and into the surrounding human HSCR colon tissue (see Figure 7A, inset). Mouse ENS cells formed small ganglion-like clusters of neuronal (Hu+) and glial or progenitor cells (SOX10+) These ganglion-like clusters contained nitric oxide synthase positive (NOS+) neural subtypes (C) and also glia (GFAP+, glial fibrillary acidic protein positive) (D). (E) Proliferating EdnrbKik+ ENC cells differentiated into neurons (EdU+/Hu+) (white arrowheads) within human HSCR muscle. (F) Combination cultures of EdnrbKik cells with HSCR colon tissue for 5 days shows synaptophysin immunoreactivity closely associated with Kikume graft–derived nerve fibers. Results were from 8 explants from two HSCR patients. Images were acquired by confocal microscopy.

Postnatal Human Aneuronal Hirschsprung Disease Colon Muscle Supports Autologous Human Enteric Nervous System Cell Colonization

HSCR patient ENS-derived cell aggregates were cocultured with aneuronal colon muscle from the same patient. To investigate the degree of ENS cell spreading, eGFP+/p75+ ENS-derived cells were monitored in HSCR muscle from one patient, in two explants (Figure 8A).
SOX10+/eGFP+ cells extended with surprising speed, and by 4–5 days were spread, principally over the circular muscle layer of the explanted aneuronal muscle tissue (see Figure 8B). ENS aggregate cells (from six patients, 30 explants) were also tracked by MTR vital stain (Figure 9A) and by prior EdU-labeling.

In all explants from all patients, within a week MTR-labeled cells spread from the initial graft, and were detected as p75+ cells in fixed specimens using confocal microscopy (see Figure 9B). Ganglion-like clusters of MTR−/p75+ cells within HSCR muscle stained for neuronal (HU), axonal (TUJ1) and glial (SOX10, S100b) markers (see Figure 9C and D). Neuronal differentiation from previously cycling ENS precursor cells was shown by the presence of some EdU+/HU+ double-positive cells (see Figure 9E). Some neurons showed the characteristic enteric neuron type marker nNOS (see Figure 9F). These p75+ ENS-derived neurons were often at and close to the implant site, but MTR+/p75+/SOX10+ cells occurred more distally from the implant site. In no cases were neurons detected in aneuronal explant tissue (six patients, 21 explants) without the prior engraftment of p75+ ENS cells.

**Discussion**

Delivery of neural progenitor cells to the postnatal aneuronal (aganglionic) colon is a promising therapeutic strategy for HSCR.23 Avoidance of immunologic rejection of such cells is obviously of prime importance, hence autologous cells would be preferred. Here we have shown that HSCR patients’ ENS-derived cells can form ENS in aneuronal bowel muscle tissue of the same HSCR patient. Our findings have fulfilled three key requirements: 1) human postnatal ENS-derived cells can be harvested from the colon of HSCR patients; 2) postnatal aneuronal human colonic tissue from HSCR patients is still permissive and supportive of de novo ENS formation, despite derivation from a much later developmental stage than is typical for this process; and 3) postnatal ENS-derived cells from a HSCR patient colon are still competent to colonize not only embryonic gut (when colonization occurs normally) but are also able colonize and differentiate in postnatal colon tissue of the same patient.

A number of sources of autologous cells for ENS reconstitution for the HSCR distal colon can be considered, but potentially favorable sources are NC-derived cells already associated with the ENS. The aneuronal HSCR distal colon has numerous sacral and trunk NC-derived cells associated mostly with nerve trunks, as we show here. These can be isolated and can form neurons in culture and in rodent embryonic aneuronal colon in organ coculture.32 In addition, normal enteric neurogenesis has recently been reported in postnatal mice, the source being trunk and/or sacral NC-derived Schwann cell precursors.16 However, animal model experiments suggest that neither of these has adequate ENS-forming ability in terms of cell numbers and neuron types and cannot replace the absent vagal

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**Figure 8.** Autologous transplant of human p75+/eGFP enteric nervous system (ENS)-derived cells in Hirschsprung disease (HSCR) muscle. (A) p75+/eGFP ENS aggregate combined with HSCR muscle (day 0) and after the culture period (day 5). During this period enhanced green fluorescent protein (eGFP) cells have spread throughout the muscle explant. Most eGFP+ cells spread on the circular muscle (CM) and at the interface (dotted line) with the longitudinal muscle (LM); this is shown (white arrowhead) in the side view of the lower explant at day 5. (B) Confocal whole-mount immunolabeling revealed that the majority of eGFP cells maintained expression of SOX10 and these cells were found extensively over muscle tissue (white arrowheads). SOX10+ cells not expressing eGFP were also detected and are likely of glial origin, which reside in the aneuronal tissue (see Fig. 2).
NC-derived ENS source. The most favorable cell population for ENS reconstitution may be that derived from NC with vagal positional identity that has experienced prior enteric inductive influences—that is, cells from the ENS of the upstream proximal ganglioneuronal colon of HSCR patients. An alternative route to autologous ENS-competent cells is via an iPSC cell route. This would solve problems of ENS progenitor cell number, but would require the iPSC cells to be converted to NC identity, and preferably to vagal and ENC identity. Techniques to achieve these higher-order specification steps are as yet not known.

It is important to use optimal methods for harvesting and maintaining ENS-competent cells. Spontaneous formation of neurosphere-like bodies, requiring one or more weeks in vitro, has been used for isolating ENS cells from dissociated gut. However, we found that spontaneously forming spheres consisted mostly of mesodermal cells, reflecting the low proportion of ENS cells in the intestinal wall. This means that spherogenesis is a slow and imperfect method for isolating ENS cells, as concluded by others. We obtained human ENS-derived cells rapidly by flow cytometry of dissociated and replated colon cells using the NC cell-surface marker antibody p75. This method produced NC-derived cells, shown by enrichment for NC markers detected by qRT-PCR (p75, FOXD3, SOX10, and AP2) and by immunolabeling (p75, SOX10, HNK1, GFAP, S100β, HU, TUJ1). NC markers were expressed similarly in cells isolated from freshly resected colon and in cells cultured for 1 week, suggesting that the in vitro environment was not, at least in the short term, detrimental to the maintenance of ENS cell types.

Proliferation of SOX10+ ENS cells in the mixed monolayer cell cultures was greatly increased by the GSK3 inhibitor CHIR-99021. Inhibition of the GSK3 pathway to mimic canonical Wnt signaling promotes proliferation of central nervous system progenitors in vitro. Activation of the Wnt pathway in part mimics the signals necessary for NC cell proliferation, which drives colonization of the gastrointestinal tract. Nevertheless, more work is required to optimize the culture conditions for human ENS cells.

It is assumed that the most effective NC-derived cell for ENS reconstitution would be a progenitor cell, combining proliferative ability and multilineage differentiation potential. However, it is difficult to unequivocally assign progenitor status to p75+ sorted cells described here due to the lack of markers for bona fide ENS progenitors. For instance, SOX10, present in ENS progenitor cells, remains expressed in differentiated glial cells. However, we were able to identify potential progenitors as SOX10+ cells that were negative for glial markers. Nevertheless, glial cells have been reported to be capable of enteric neurogenesis after wounding at least in rodents. Moreover, recent analyses in mice of development of the related NC-derived parasympathetic ganglia suggest that some neurons are normally derived from cells that had expressed glial markers, while some late-originating calretinin+ ENS neurons in mice are of Schwann cell origin.

Experiments with embryonic mice on colonization of gut by ENS progenitors show that the stage of the ENC cells and of the recipient colon are influential, and a temporal mismatch between ENC cells and colon restricts innervation. For HSCR patients, temporal variation options are limited: only postnatal cells and tissues are available. Therefore, the study of enteric neurogenesis abilities used postnatal human ENS-derived cells and recipient colon from HSCR patients. Comparison with age-matched cells from healthy neonatal donor colons is difficult because the latter are so rarely obtainable, but it can be compared with that of equivalent postnatal ENS cells of normal rodent models.

We investigated whether postnatal human HSCR patient ENS-derived cells could migrate into receptive gut and whether postnatal human HSCR colon tissue was permissive for ENS formation, given that these processes are normally confined to embryonic development. To address the first question, we used the embryonic quail anorectal intestine. In this model, the human ENS-derived cells adopted the morphology of motile ENC cell chain, which are typical of NC migration in the embryonic gut, and expressed NC cell markers, but the rate of colonization of these human HSCR-derived ENS cells was slower than that of avian embryonic ENC cells. This may be because the rate of occupation is driven by proliferation, which may be expected to be lower in cells of older and/or human origin. However, this confirmed that p75-sorted human HSCR ENS cells are capable of enteric colonization in a permissive embryonic tissue. To answer the second question, we challenged human HSCR colon tissue with mouse embryonic ENC progenitors of known colonization ability. We conclude that postnatal human aneuronal HSCR colon muscle tissue is permissive of mouse embryonic ENC cell migration, proliferation, ganglion-like aggregation, ENS neuronal, and glial differentiation, and neurite extension.

Previously, neonatal and adult human bowel has been dissociated and cultured in vitro to form spontaneously neurosphere-like bodies for transplantation into aneural embryonic mouse hindgut in organ culture or into postnatal mouse colon in vivo. Neurons (Hu, PGP9.5) and glia (S100β, GFAP) were presumed to have arisen from the implanted human cells. Human bowel-derived neurosphere-like bodies have also been cocultured with postnatal aneuronal human gut, although analysis was...
limited to detection of TUJ1+ fibers.36 We have confirmed32 that the aneuronal tissue contains many single TUJ1+ nerve fibers and large nerve fiber bundles that persist after in vitro culture. We also observed TUJ1+/p75+ cells that were SOX10+ and lacked neuronal morphology in aneuronal (aganglionic) HSCR colon both in vivo and in explants. We concur that this antibody does not discriminate between neurons/axons and non-neuronal cells in human ENS as clearly as it does in mouse ENS.45 Therefore, it is difficult based on TUJ1 labeling alone to unequivocally identify whether neurosphere-derived donor cells had colonized and differentiated in the human gut tissue.

We prelabeled HSCR patient ENS-derived cells with eGFP by viral transfection and by vital staining with MTR to allow detection in HSCR muscle. Because of the rapidity with which we could obtain these cells, we could implant them into aneuronal muscle explants from the same individual. We conclude that postnatal human ENS-derived cells from HSCR colon are capable of migration, proliferation, ganglion-like aggregation, and neuronal and glial differentiation in autologous aneuronal colon muscle tissues.

HSCR is genetically complex, with numerous mutations in many gene(s) expressed in ENS cells,50 most frequently the gene for the tyrosine kinase receptor RET that mediates signaling from mesoderm-derived GDNF (glial-derived neurotrophic factor). It has been argued that, without gene correction, autologous cells may therefore be poor candidates for ENS replacement therapy. Yet in our studies all patient donor cells were able to colonize autologous smooth muscle explants. In the majority of HSCR patients a large portion of the ENS is present and is functional, so an alternative argument is that in HSCR ENC cells could form ENS if they could be delivered to the region lacking ENS.

In summary, we provide strong evidence that ENS-derived cells from HSCR patients can establish ENS in aneuronal (aganglionic) bowel of the same patients. Critical to this is our demonstration that postnatal human colon tissue remains permissive for colonization by a competent ENS source. Nevertheless, we caution against over-interpretation of our conclusions as there are still quantitative and distributional hurdles to surmount before cell therapy can become practicable in HSCR, or achieve the lesser goal of improving the outcome of current surgery by reinervating the remaining aneuronal muscular cuff.51 Generation of sufficient numbers of progenitor cells to be effective in the large postnatal aneuronal area is challenging. Even with adequate cell numbers, how to surgically transplant them to achieve coverage of the large area of HSCR colon in vivo is a problem. It is likely that the refinement of cell growth conditions combined with the use of elastic polymer substrates permissive for cell proliferation and cell delivery will assist in achieving this therapeutic goal.52

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Conflicts of interest
The authors disclose no conflicts.

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