Isolation of Human Colon Stem Cells Using Surface Expression of PTK7

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SUMMARY

Insertion of reporter cassettes into the Lgr5 locus has enabled the characterization of mouse intestinal stem cells (ISCs). However, low cell surface abundance of LGR5 protein and lack of high-affinity anti-LGR5 antibodies represent a roadblock to efficiently isolate human colonic stem cells (hCoSCs). We set out to identify stem cell markers that would allow for purification of hCoSCs. In an unbiased approach, membrane-enriched protein fractions derived from in vitro human colonic organoids were analyzed by quantitative mass spectrometry. Protein tyrosine pseudokinase PTK7 specified a cell population within human colonic organoids characterized by highest self-renewal and re-seeding capacity. Antibodies recognizing the extracellular domain of PTK7 allowed us to isolate and expand hCoSCs directly from patient-derived mucosa samples. Human PTK7+ cells display features of canonical Lgr5+ ISCs and include a fraction of cells that undergo differentiation toward enteroendocrine lineage that resemble crypt label retaining cells (LRCs).

INTRODUCTION

The epithelium that lines the small intestine and the colon is in constant self-renewal with a life cycle of 3–4 days (Clevers, 2013a). Regeneration occurs from units embedded within the submucosa known as crypts, each containing about 6–10 intestinal stem cells (ISCs) (Clevers, 2013a). Differentiated epithelial cells reside at the surface. The biology of mouse ISCs has been intensively studied in recent years. They express Lgr5, a receptor for the WNT pathway agonist R-Spondin 1 (R-SPO) (Barker et al., 2007; de Lau et al., 2011; Leushacke and Barker, 2014). Mouse Lgr5+ cells are rapidly dividing multipotent stem cells that reside at the bottommost positions of the small intestine and colonic crypts. Under culturing conditions that simulate the stem cell niche, mouse Lgr5+ cells form 3D structures in vitro, known as organoids (Sato et al., 2009).

The progeny of ISCs, transit amplifying cells or progenitors, will eventually give rise to the majority of differentiated cells as they migrate upward along the crypt axis. Recently it was shown that a fraction of Lgr5+ cells remains relatively slow proliferating and differentiates directly into less abundant differentiated lineages such as Paneth or enteroendocrine cells. This Lgr5+ cell subpopulation can be experimentally recognized by the retention of molecules that label DNA and therefore has been known as label retaining cells (LRCs) (Clevers, 2013b; Potten et al., 1974). They are characterized by co-expression of ISC-specific genes and markers of Paneth cells and enteroendocrine lineages (Buczacki et al., 2013; Grün et al., 2015). LRCs are fated to differentiate; nevertheless, under certain conditions, such as damage of the cycling ISC pool, they act as facultative stem cells and contribute to the regeneration of the intestinal epithelium (Buczacki et al., 2013). Mouse LRCs also expand as in vitro organoids (Buczacki et al., 2013).

There is a growing interest in understanding the biology of human colonic stem cells (hCoSCs), as it is widely believed that they play key roles in various human intestinal disorders (Barker et al., 2009). Furthermore, a better understanding of this cell population is key to develop therapeutic approaches to regenerate disease-affected tissue (Yui et al., 2012). We recently showed that in media containing WNT3a, R-SPO, EGF, and Noggin (WREN media) human colonic crypts form organoids that expand as undifferentiated 3D cell cultures (Jung et al., 2011). However, only around 5%–6% of the cells within these cultures remain as multipotent stem cells. We searched for surface markers that identify bona fide hCoSCs from human organoid cultures. We characterize protein tyrosine pseudokinase 7 (PTK7) as a hCoSC marker that allows purification and optimized ex vivo expansion of self-renewing multipotent hCoSCs.

RESULTS

Quantitative Mass Spectrometry Analyses of Human Colon Organoid-Derived Membrane Proteins

We searched for surface markers that would enable discrimination of distinct cell populations in human colon organoids (Jung et al., 2011). By quantitative mass spectrometry
Figure 1. Quantitative Mass Spectrometry Reveals PTK7 Is Enriched in Ex Vivo Cultured hCoSCs and Specifies Organoid Cells with Highest Self-Renewal Capacity

(A) Experimental scheme: Human colonic organoids were cultured under stem cell (WREN, STEM) or differentiation (EN, DIFF) conditions. Differentiation was induced for 72 hr. After isolation of membrane-enriched protein fractions, samples were subjected to quantitative mass spectrometry (LC-MS-MS).

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cultures maintained in either WREN or EN media. Negative fold changes represent organoid stem cell-enriched proteins (see Table S1 for all analyses of these cultures revealed heterogeneous surface expression of PTK7 (Figure 1C, upper). We found that cells with high PTK7 surface abundance exhibited a higher re-seeding capacity when compared with PTK7-low, or PTK7-negative cells (Figures 1C, 2B, S1B, and S2B). To determine the frequency of organoid re-initiating cells (OIC) of each PTK7 population, we plated different cell numbers (40, 150, and 400 viable cells) and counted the number of organoids after 1 week. We then calculated OIC frequencies by applying the maximum-likelihood estimation method for limiting dilutions assay (Hu and Smyth, 2009). Simply re-plating the total of viable cells yielded a frequency of one OIC every 34 cells (3%), in line with our previous observations (Jung et al., 2011). We calculated an OIC of 1 in 6 for PTK7-high (16.7%), 1 in 23 for PTK7-low (4.3%), and 1 in 118 (0.85%) for PTK7-negative cells (Figure 1D). Furthermore, PTK7-high cells expressed elevated levels of bona fide ISC marker genes such as LGR5 and ASCL2 (Figure 1E, left), whereas PTK7-low and PTK7-negative WREN-cultured organoid cells expressed relatively higher levels of intestinal differentiation genes, including ANPEP, CEACAM7, and CA1 (Figure 1E, right). Upon removal of growth factors, secondary organoid cultures derived from PTK7-high cells underwent multilineage differentiation (Figure 1F).

We and others have previously identified surface markers for purification of mouse and human adult stem cells from intestinal samples. These include EPHB2 (Jung et al., 2011; Merlos-Suárez et al., 2011), CD44 (Gracz et al., 2013; Wang et al., 2013), and Prominin-1/CD133 (Hou et al., 2011; Zhu et al., 2009). These markers showed heterogeneous surface expression in human colon organoid cultures maintained in WREN media (Figures 2A and S2A). We thus assessed organoid-forming capacity of cells isolated using these surface markers by fluorescence-activated cell sorting (FACS). We compared the 10% more positive versus 20% more negative population for each of these markers within the same organoid culture. This experiment demonstrated

**High PTK7 Expression Levels Segregate with High Self-Renewal Capacity in Ex Vivo Cultured Human Colonic Crypts**

In a previous study, we observed that only around 5%–6% of viable cells within human colonic organoids cultured in WREN media hold the capacity to reform organoids upon serial passaging (Jung et al., 2011). Our data suggested that only a subpopulation of organoid cells remains as bona fide stem cells (Jung et al., 2011). Flow-cytometry analysis of these cultures revealed heterogeneous surface expression of PTK7 (Figure 1C, upper). We found that cells with high PTK7 surface abundance exhibited a higher re-seeding capacity when compared with PTK7-low, or PTK7-negative cells (Figures 1C, 2B, S1B, and S2B). To determine the frequency of organoid re-initiating cells (OIC) of each PTK7 population, we plated different cell numbers (40, 150, and 400 viable cells) and counted the number of organoids after 1 week. We then calculated OIC frequencies by applying the maximum-likelihood estimation method for limiting dilutions assay (Hu and Smyth, 2009). Simply re-plating the total of viable cells yielded a frequency of one OIC every 34 cells (3%), in line with our previous observations (Jung et al., 2011). We calculated an OIC of 1 in 6 for PTK7-high (16.7%), 1 in 23 for PTK7-low (4.3%), and 1 in 118 (0.85%) for PTK7-negative cells (Figure 1D). Furthermore, PTK7-high cells expressed elevated levels of bona fide ISC marker genes such as LGR5 and ASCL2 (Figure 1E, left), whereas PTK7-low and PTK7-negative WREN-cultured organoid cells expressed relatively higher levels of intestinal differentiation genes, including ANPEP, CEACAM7, and CA1 (Figure 1E, right). Upon removal of growth factors, secondary organoid cultures derived from PTK7-high cells underwent multilineage differentiation (Figure 1F).

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PTK7-high cells contained significantly more organoid forming cells than equivalent populations purified with the other ISC markers (Figure 2B). In addition, PTK7-high cells displayed higher expression levels of ISC markers genes (LGR5, SMOC2) and decreased expression of differentiation markers (KRT20, FABP2) compared with EPHB2-high, CD44-high, and CD133-high cells (Figure 2C). We obtained equivalent results using organoids derived from a different individual (Figure S2). Our data indicate that human colonic organoids in WREN media contain phenotypically distinct cell populations and that high PTK7 surface levels specify bona fide stem cells in these cultures.

PTK7 Enriches in Self-Renewing, Multipotent Stem Cells of the Human Colon

Immunohistochemical analysis confirmed that PTK7 antibodies stained the bottom third of human colonic crypts with the highest intensity observed at the bottommost
PTK7 antibodies recognized the mouse protein, which precluded isolation of PTK7+ cells by flow cytometry from mouse tissues (data not shown). Therefore, we used Cyp1a1-H2B-YFP mice, which upon induction with beta-Naphthoflavone, express histone H2B-YFP fusion protein in intestinal cells (Buczacki et al., 2013). In pulse-chase experiments, around 8%–10% of CD24+Lectin− crypt cells retained the H2B-YFP label 10 days after induction (Figures 4B and 4C). Mouse SI LRCs (CD24+/H2B−YFP+) expressed Lgr5, Chga, and Rfx6 compared with non-LRCs (CD24+/H2B−YFP−). This experiment revealed that LRCs also expressed high levels of Ptk7 (Figure 4D).

To extend our analysis, we studied the expression of genes characteristic of mouse Lgr5+ ISCs (Muñoz et al., 2012) or of LRCs (Buczacki et al., 2013) in PTK7+ hCoSCs. We derived an ISC (Lgr5) signature comparing the transcriptomic profiles of Lgr5-GFP-high versus Lgr5-GFP-low crypt cells purified from Lgr5-GFP knockin mice (Muñoz et al., 2012) (Table S4). This gene set contained many Wnt targets, including the ISC marker genes Lgr5, Smoc2, Ascl2, and Apcdd1. To generate a specific LRC signature, we selected genes upregulated in H2B-YFP-retaining crypt cells (H2B−YFP−; CD24+) compared with crypt non-retaining cells (H2B−YFP−; CD24+) (Table S4; Supplemental Experimental Procedures for details) (Buczacki et al., 2013). Only 12 genes overlapped between the LGR5 and LRC gene sets (Figure 4E). Gene Set Enrichment Analysis (GSEA) showed that both signatures were highly enriched in PTK7-high hCoSCs (Figures 4F and S4D). We confirmed by qRT-PCR that genes specific of LRCs, such as Chga, Tph1, Rfx6, and Cacna1a, were upregulated in PTK7-high cells compared with PTK7-low cells purified from human colon samples (Figure 4G). In contrast, Ephb2-high or CD44-high cells purified from these samples were not enriched in LRC-specific genes (Figures S4E and S4F). These results indicate that the PTK7+ human colonic epithelial cell population expresses gene programs of both canonical ISCs and of LRCs.

**DISCUSSION**

Insertion of reporter cassettes into the Lgr5 locus has enabled characterization of mouse ISCs (Barker et al., 2007, 2010) and development of ex vivo organoid culture systems (Sato et al., 2009). Subsequent transplantation assays using fetal or adult ISCs have paved the way for regenerative therapies of intestinal injuries (Fordham et al., 2013; Yui et al., 2012). However, low cell surface abundance of the LGR5 protein and lack of high-affinity anti-LGR5 antibodies represent a roadblock to isolate human colon stem cells using this marker gene. Our data show that PTK7 is a surrogate of LGR5 in human colon. The methodology developed herein enables the isolation of a cell population...
Figure 3. PTK7 Specifies Self-Renewing, Multipotent Stem Cells of the Human Colon

(A–D) Immunohistochemistry analysis of (A) PTK7, (B) EPHB2, (C) FABP1, and (D) CK20 expression on serial sections of normal human colonic mucosa. Black arrows indicate specific staining. Scale bar represents 100 μm.

(E) FACS profile of single-cell suspensions from primary human colon stained with antibody to PTK7 (APC-conjugate) (upper). Only epithelial cells were included (EpCAM+) and non-epithelial cells were excluded (CD11+/CD31+/CD45-) from the analysis. Control staining was performed to define PTK7-negative fraction (lower). H, PTK7 high; M, PTK7 medium; L, PTK7 low; N, PTK7 negative.

(F) Representative images of organoid cultures 10 days after seeding equal cell numbers.

(G) Quantification of organoid-forming capacity of colon cells sorted in (E). Shown are data collected from three independent 3D cultures (biological replicates). t test, *p value of the average PTK7 high versus average medium lower than 0.05.

(H) qRT-PCR analysis of gene expression on primary human colon epithelial cells purified by FACS according to different PTK7 surface levels. Error bars indicate SD (n = 3 technical replicates, see Figure S3C for an additional supporting experiment).
**Figure 4.** PTK7+ Human CoSCs Show Mixed Features of Canonical CoSCs and LRCs

(A) Immunofluorescence staining on formalin-fixed paraffin sections of DAPI (blue), PTK7 (green), and CHGA (red). Scale bar size is indicated.

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largely enriched in CoSCs from human mucosa samples. Remarkably, the possibility of purifying the cell population with highest self-renewal capacity from organoid cultures through PTK7 could facilitate the development of therapeutic approaches based on CoSC transplantation.

Identification of a pool of slow cycling ISCs committed to differentiate toward Paneth and enteroendocrine lineages has clarified the long-lasting debate about the existence of quiescent stem cells in the intestine (Clevers, 2013b). One of the defining features of intestinal LRCs is the ability to gain multipotent growth capacity upon tissue injury (Buczacki et al., 2013). However, it remains unclear whether a similar cell type is also present in the human intestine. Expression profiling of PTK7-high cell population suggests the existence of LRC-like cells in the human colonic epithelium and implies phenotypic and functional heterogeneity within the hCoSC compartment similar to that present in mouse crypts. Importantly, Buczacki et al. (2013) demonstrated that mouse LRCs maintain the ability to form intestinal organoids when plated in the culture dish. However, under these conditions, LRCs lose their quiescent/slow-cycling character and revert into actively proliferating ISCs (Buczacki et al., 2013). Similarly, we were unable to identify LRCs in human colon organoids (data not shown). Future efforts should aim to optimize organoid culture conditions in order to promote the concomitant propagation of canonical stem cells and LRCs in vitro. The biology of these reserve stem cells in humans holds great interest as they might be responsible for regeneration of human colon mucosa upon certain insults such as chemotherapy.

EXPERIMENTAL PROCEDURES

Organoid Cultures from Colonic Crypts

Colonic tissue pieces were incubated in 10 mM DTT for 5 min at room temperature and transferred to 8 mM EDTA for 60 min at 4°C, and vigorous shaking yielded supernatants enriched in colonic crypts. Crypts were either directly cultured, or single-cell suspensions were obtained by incubation with 0.4 mg/ml Dispase (GIBCO). Crypts or single cells were mixed with 50 µl matrigel (BD Biosciences) and plated on 24-well culture dishes. Embedded cells were overlaid with WREN medium (Wnt3a-conditioned medium and ADF 50:50, Glutamax, 10 mM HEPES, N-2 [1×], B-27 without retinoic acid [1×], 10 mM Nicotinamide, 1 mM N-acetyl-L-cysteine, 1 µg/ml RSPO1, 50 ng/ml human epidermal growth factor [EGF], 100 ng/ml human Noggin, 1 µg/ml Gastrin, and 0.05 µM PGE2, 0.1 µM A83-01, 10 µM p38 inhibitor SB202190, 10 µM Y27632). Medium was replaced with fresh WREN medium every other day. Biological samples were obtained from individuals treated at the Hospital del Mar (Barcelona, Spain) or from Hospital Clinic (Barcelona, Spain) under informed consent. Experiments were approved by the ethics committee of IRB/Hospital Clinic (projectERC-208488/CRC programme).

ACCESSION NUMBERS

The accession number for the microarray dataset reported in this paper is GEO GSE68340.

SUPPLEMENTAL INFORMATION

Supplemental Information includes Supplemental Experimental Procedures, four figures, and four tables and can be found with this article online at http://dx.doi.org/10.1016/j.stemcr.2015.10.003.

AUTHOR CONTRIBUTIONS

P.J. designed, planned, and performed experiments and analyzed the results. C.S. and M.S. performed LC-MS-MS analysis and analyzed results. E.B. provided crucial assistance with experiments and analyzed data. S.J.B. and D.J.W. contributed transgenic animals and contributed crucial experimental protocols. X.H.-M. provided assistance with transgenic animals. M.S. provided assistance with immunohistochemistry. M.D.-F. and P.A. analyzed the results. C.S. and M.S. performed LC-MS-MS analysis and ADF 50:50. Glutamax, 10 mM HEPES, N-2 [1×], B-27 without retinoic acid [1×], 10 mM Nicotinamide, 1 mM N-acetyl-L-cysteine, 1 µg/ml RSPO1, 50 ng/ml human epidermal growth factor [EGF], 100 ng/ml human Noggin, 1 µg/ml Gastrin, and 0.05 µM PGE2, 0.1 µM A83-01, 10 µM p38 inhibitor SB202190, 10 µM Y27632). Medium was replaced with fresh WREN medium every other day. Biological samples were obtained from individuals treated at the Hospital del Mar (Barcelona, Spain) or from Hospital Clinic (Barcelona, Spain) under informed consent. Experiments were approved by the ethics committee of IRB/Hospital Clinic (projectERC-208488/CRC programme).

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