

Review

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Posted Date: 6 February 2024

doi: 10.20944/preprints202402.0346.v1

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Review

Subepithelial Stromal Cells: Their Roles and Interactions with Intestinal Epithelial Cells during Gut Mucosal Homeostasis and Regeneration

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Abstracts: Intestinal epithelial cell activities during homeostasis and regeneration are well described, but their potential interactions with stromal cells remain unresolved. Exploring the functions of these heterogeneous intestinal mesenchymal stromal cells (iMSCs) remains challenging due to the lack of specific markers for most functionally homogenous subpopulations. In recent years, however, novel clustering techniques such as single-cell RNA sequencing (scRNA-seq), fluorescence-activated cell sorting (FACS), confocal microscope, and computational remodeling of intestinal anatomy have helped identify and characterize some specific iMSC subsets. These methods help researchers learn more about the localization and functions of iMSC populations during intestinal morphogenic and homeostatic conditions. Furthermore, it is imperative to understand their cellular pathways for activation and how they interact with surrounding cellular components, particularly during intestinal epithelial regeneration that follows mucosal injury. This review provides insights into the spatial distribution and functions of some identified iMSC subtypes in intestinal morphogenesis, homeostasis, and regeneration. We reviewed related signaling mechanisms implicated during epithelial and subepithelial stromal cell crosstalk. Future research should focus on elucidating the molecular intermediates of these regulatory pathways to open a new frontier for potential therapeutic targets that can alleviate intestinal mucosa-related injuries.

Keywords: intestinal stem cell niche; mesenchymal stromal cells; sub-epithelial gradient factors; epithelial mesenchymal interactions

1. Introduction

The continuous self-renewal of the intestinal stem cells (ISC) underpins the high turnover rate of differentiated epithelial cells [1-4]. Over a decade ago, Hans and colleagues demonstrated how ISCs self-renew and proliferate into progenitor cell populations before differentiating into specialized epithelium cells [5,6]. From their study, we understand that stem cell division is confined to the crypt. In contrast, proliferating and differentiated cells form single-layered monolayers of different cell types arranged from the crypt base toward the apical villi region [5,7,8]. The proliferating progenitor cells in the transit amplifying (TA) zone undergo asymmetrical cell divisions, such that the half-daughter cells differentiate into specialized lineages, such as the secretory (Paneth, tuft, goblet, and enteroendocrine cells) or the absorptive enterocytes that migrate and form the single epithelial layer of the villi compartment [9]. This crypt-villus axis is characterized by distinct cellular interaction and molecular signatures that maintain the homeostasis [10-12]. For example, previous studies demonstrated that Paneth cells and subcryptal stromal cells produce canonical Wnt ligands to maintain the stem cell niche [13-17]. Likewise, stromal cells close to the sub-villi have been shown to communicate with the differentiated cells by providing BMPs and non-canonical Wnt ligands [18-20].

During regeneration after mucosa injury, reserve stem cells are activated, and specialized cells dedifferentiate into active stem cell populations that proliferate to support epithelial regeneration in

the damaged site [21]. Previous studies have suggested that injured intestinal epithelial cells activate subepithelial fibroblast via TGF β to support its proliferation and migration. [22-24] However, the specific functions of various subepithelial stromal cell populations to epithelial regeneration remain unresolved [25,26]. On this account, this paper reviews the recently characterized subpopulations of iMSCs in mice and their interactions with epithelium, highlighting their role in intestinal epithelial morphogenesis, homeostasis, and regeneration.

2. Intestinal stem cell regulation

Neighboring epithelial and subepithelial cells secrete gradient factors that influence ISC fate. These cellular interactions maintain ISC stemness at the base of the crypt and aid intestinal cell proliferation and differentiation activities along the crypt-villus axis [4,27]. Though the specificity of active stem cells is debatable, they are marked by LGR5⁺, OLFM4, ASCL2, RNF43, SOX9, MSI1, and SMOC2, and are capable of self-renewal and replenish differentiated epithelial cells [9,11,27,28]. To regulate these ISC activities, Paneth cells are adjacently interspersed between ISCs and produce growth factors (Wnt, EGF, and Notch-associated ligands). For instance, Paneth cells as described by Sato, et al. [29] improved organoid formations. Yilmaz, et al. [30] also reported that caloric restrictions preserved ISC self-renewal, and their adaptation is likely coordinated by mTORC1 signaling in the neighboring Paneth cells. Despite their regulatory support to the ISC niche, debates remain on the contribution of these non-stem cell populations in the ISC niche. Paneth cell depletion did not significantly alter ISC fate in vivo [31], and their functions can be substituted with exogenous Wnt supplement in ex vivo enteroid culture [32]. Additionally, other epithelial cell populations also help maintain intestinal epithelial balance by providing cues, which are highlighted in Table 1. Specifically, non-active stem cells, found in +4 positions, were reported to replenish the active stem cell pool during injury recovery through YAP1-dependent transient expansion [33]. Likewise, Paneth cells in the small intestine or *Reg4*⁺ expressing cells in the colon secrete canonical Wnt factors (such as *Wnt3*, *Wnt9b*, and *Wnt11*) and growth factors (EGF) to support epithelial regenerations via Notch signaling activation [16,34-37]. Similarly, recent studies have shown that iMSCs play critical and indispensable roles in regulating intestinal morphogenesis, homeostasis, and regeneration by providing regulatory factors [38-40]. It is therefore vital to address how iMSCs regulate Wnt, EGF, and bone morphogenetic protein (BMP) signaling pathways that confine active stem cell renewal at the crypt base and differentiate progenitors into specialized cells [2].

Epithelial cells	Cell Markers	Ligands	Functions	Signaling pathways	References
Paneth cells	<i>Defa4</i> expressing cell	<i>Wnt3a</i> , <i>Wnt9b</i> , <i>Wnt11</i>	Support regeneration	Notch	[35]
	pS6 ⁺	<i>Notum</i>	Wnt inhibitor	Non-canonical Wnt/mTORC1	[41]
Progenitor cells	Dll ⁺	<i>Dll1</i> , <i>dll4</i>	Support regeneration	Notch	[42]
Secretory lineage	Unknown	<i>Egf</i> , <i>Tgfa</i> ,	Promote IEC homeostasis	EGFR/RAS	[34]
Colonic Paneth cells	REG4 ⁺	<i>Dll1</i> , <i>Egf</i>	Promote stemness	Notch	[43]
+4 quiescent cells	BMI1, HOPX, MTERT	Unknown	Support regeneration	Hippo	[21,33]
Tuft cells	DCLK1	<i>Dll1</i>	Promote regeneration	Notch	[37]

2.1. Signaling pathways

2.1.1. Wnt pathway

ISC stemness, proliferation, and differentiation fate largely depend on ligand-mediated signaling processes. As demonstrated in previous studies, canonical Wnt (or Wnt/ β -catenin) signaling is crucial for ISC self-renewal maintenance [44,45]. For example, knocking out *Porcupine* and *Wntless*, essential mediators of Wnt secretion, greatly reduced the ISC population [14,43,46]. Similarly, the adenoviral delivery of the Wnt signaling antagonist, Dkk1, inhibited intestinal proliferative marker, Ki67, and Wnt/ β -catenin target genes, CD44 and EpcA2 [47]. These studies confirmed the critical roles of Wnt signaling in intestinal homeostasis [2].

Notably, the canonical Wnt/ β -catenin activation, regarded as a major promoter of ISC stemness maintenance, is triggered by ligands (e.g., Wnt2b, Wnt3, etc.) secreted from the ISC's neighboring cells. First, the Porcupine protein synthesized in the endoplasmic reticulum of the neighboring cells enables Wnt ligand secretions that bind on the Frizzled receptors of the ISC plasma membrane to simultaneously attach to the LRP5/6 co-receptors [48]. This Wnt ligand-receptor binding initiates Dsh protein recruitment, leading to LRP receptor phosphorylation and the disruption of the Axin-GSK3-CK1 complex that orchestrates the β -catenin ubiquitination [2,11]. As a result, β -catenins are stabilized and accumulated in the cytoplasm. They subsequently translocate into the ISC's nucleus to interact with LEF/TCF factors, thereby promoting canonical Wnt target genes transcription as illustrated in Figure 1. Blocking Wnt ligands may compromise Lgr5⁺ stemness and induce premature lineage differentiation [49]. This is partly due to the essential role of Wnt in ISC stemness maintenance. In contrast, the expressions of Wnt signal-related promoters such as Wnt2b and Rspodins (Rsops), and Wnt target genes increased the Lgr5⁺ population and ISC diminished as the proliferating cells migrated apically in the transit-amplifying region [38].

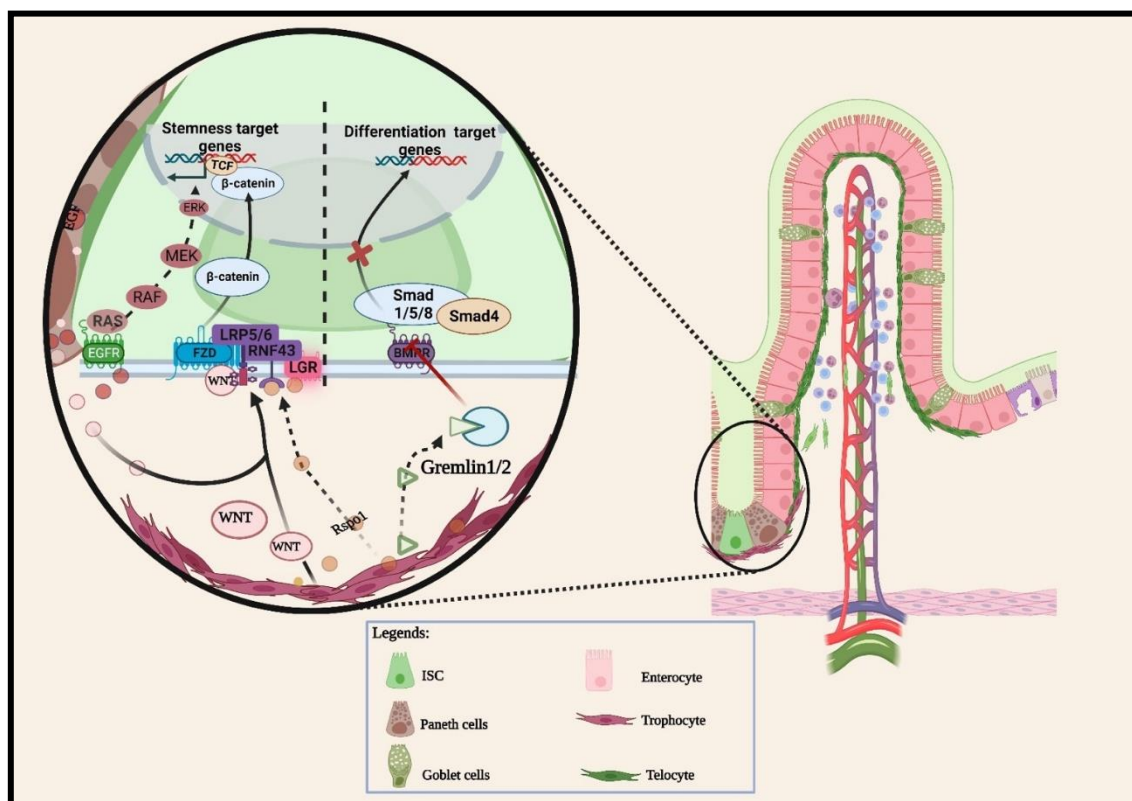


Figure 1. Multiple signaling pathways regulating intestinal epithelial-mesenchymal crosstalk in the crypt. The diagram provides an overview of the intricate mechanisms involved in the interactions between crypt-based epithelial cells and the neighboring iMSCs. This interplay controls intestinal stem cell (ISC) homeostasis and epithelial differentiation. The activation of Wnt-promoting pathways and inhibition of Bmp/Bmpr

binding orchestrate ISC stemness in the crypt base. Paneth cells secrete Wnt, Notch, and epithelial growth factors (EGF) ligands that induce Wnt target gene transcription in ISC. Wnt factors secreted by subcryptal iMSCs bind FZD and LRP6 co-receptors and stabilize β -catenin in the Wnt signaling cascade to promote ISC-related gene transcription. Rsp100 binding LGR family receptors stabilize FZD expressions, contributing to WNT pathway activation. In addition, subcryptal iMSCs secrete Bmp antagonists such as *Gremlin* to maintain Wnt activities in the ISC niche.

Unlike the canonical Wnt ligands that support intestinal stemness via β -catenin stabilization by orchestrating Wnt target gene transcriptions, non-canonical Wnt pathways are β -catenin independent [50]. Non-canonical Wnt pathways regulate the Wnt/PCP (planar cell polarity) and Wnt/ Ca^{2+} dependent signaling cascades to modulate the ISC fate [51-53]. Despite the extensive knowledge surrounding canonical Wnt signaling, the function of non-canonical Wnt signaling has not been thoroughly studied. Non-canonical Wnt signaling, triggered by non-canonical Wnt ligands, promotes cellular differentiation and regeneration to facilitate ISC migration. Non-canonical Wnt ligands, including Wnt4, Wnt5, Wnt5b, and Wnt16 were remarkably expressed along the villus axis to the tip [20,38]. Their presence indicates that some Wnt ligands support villus differentiation and homeostasis, which might complement BMPs that promote intestinal maturation [20,54]. Recent scRNA sequencings have identified some iMSCs clusters, that are abundant in the villus sub-region, express *Wnt4*, *Wnt5*, *Wnt 16*, and *Egf* [20,54]. This paradoxical revelation contradicts the classical roles of Wnt in maintaining the stemness [11,55]. For instance, ablation of villus tip subepithelial *Lgr5*⁺ iMSCs, which expressed high *Wnt5a* and *Bmp4*, resulted in enterocyte loss [54,56]. Additionally, the Wnt/ Ca^{2+} pathway can independently stimulate stem cell proliferation when Wnt5a/Fzd6 interactions increase intracellular Ca^{2+} levels in the cytosol during gastric cancer. This leads to PKC and CAMK-mediated downstream cascades to promote cell migration [53,57,58]. In a related pattern, Wnt ligands could bind and activate Rho receptor complexes (Ror/Ryk) to trigger c-Jun N-terminal kinase, initiating JNK target gene transcription [57,59]. JNK target genes participate in cellular proliferation, migration, and regeneration by targeting exoskeleton proteins and adhesion molecules such as actin and Rho GTPases. These suggest that non-canonical Wnt signaling downstream could be a promising target for pharmaceutical drugs to improve intestinal injury recovery [60].

2.1.2. BMP pathways

The bone morphogenetic protein (BMP) pathway is an important signal opposing the Wnt mechanism that stimulates epithelial differentiation. Contrary to canonical Wnt signaling, BMP/Smad activities increase toward the apical region, indicating repressed stem cell activities along the villi [61-63]. An extensive review has addressed BMP/Smad regulation of intestinal epithelial differentiation and homeostasis [63]. The focus here provides insight into how Bmp factors mediate the interaction between the intestinal epithelium and iMSCs. Of note, BMP4, a ligand stimulating Bmp signaling, was reported as a transcriptional target gene of hedgehog-responsive iMSCs [61,63-65]. The interaction between iMSCs and epithelium has complex roles in establishing the distinct identities of the crypt and villi by supplying Wnt and Bmp gradients [62,65-67]. A recent study found that populations of PDGFR α ⁺ iMSCs promote Wnt signaling, which can stimulate the expression of epithelial hedgehog ligands (SHH and IHH) during morphogenesis. [20]. In turn, epithelial-derived SHH ligands directly activate iMSC target gene transcriptions, including BMPs, that shape the developing intestinal villus formation [20,44,64,66,68]. Interestingly, sub-epithelial iMSC provides essential ligands that inhibit BMP activities in the crypt base where stem cell renewal and proliferation are at their peak. For instance, sub-cryptal iMSCs secrete Noggin and Gremlin1 to block BMP signaling, thereby supporting the ISC niche [20,69]. This means that epithelial-mesenchymal communications create a feedback loop to maintain intestinal homeostasis along the crypt-villus axis.

2.1.3. Other cellular signaling pathways

Subsequent to active ISC proliferation, cell lineage specialization fates are intricately regulated by the Notch pathway. Notch signaling actively commits progenitor cells to transform into differentiated specialized cells [36,70]. Moreover, increased Notch ligands from neighboring Paneth cells redirected proliferating cells toward enterocyte lineage by repressing *Math1* transcriptions [7,71]. Paneth cells, which is a secretory lineage, could trigger negative feedback to limit secretory cell lineage commitment, thereby supporting the importance of stringent regulations of ISC activities to maintain intestinal epithelial homeostasis. Notwithstanding, other mechanistic pathways, such as EGF, contribute to ISC maintenance and are well documented in previous studies and reviews [2,11,27].

Another notable mechanism that regulates intestinal cellular interactions is the Hedgehog (HH). HH ligands secreted by epithelium initiate negative regulation of its receptor (*Ptch1*) on neighboring iMSCs [65,72]. In the absence of HH ligands from epithelial cells, *Ptch1* inhibits Smo signaling transduction, leading to Gli family phosphorylation in the cytoplasm by degradation complex including GSK3 β , CK1 α , and PKA. In contrast, when HH ligands bind on *Ptch1* receptors, they release Smo to activate STK36, thereby inhibiting the degradation complex assembly. STK36 also phosphorylates SUFU complex to stabilize Gli and allow nuclear accumulation, which subsequently stimulates Gli-dependent target gene transcriptions Figure 2. Despite the extensive review on Hedgehog regulations, especially their intricate functions in fetal villigenesis and regenerations [20,45,65,69], their roles in promoting crypt stemness during homeostatic conditions are just gaining attention in recent studies [9,66,73]. This was best described when hedgehog-responsive mesenchymal cells were revealed to constitute a key colonic stem cell niche [73]. Consistently, the mesenchymal subsets that are localized in sub – and peri-cryptal regions form the main sources of the Wnt, Rspo, and Grem niche that supports the crypt stemness [38,39]. For example, activated HH pathway upregulated stromal Wnt expression which contributes to increased OLFM4 positive stem cell pool [74]. Likewise, Current data also suggest HH/Wnt pathways crosstalk potentially promote intestinal regeneration. Upon irradiation-induced intestinal epithelial injury, SHH was significantly upregulated, leading to increased production of Wnt ligands (Wnt2b, Wnt4, and Wnt5a) in underlying stromal cells – resulting in enhanced regeneration [67]. Together, these signaling networks complementarily or in opposing efforts regulate intestinal cell fate along the crypt-villi topologies during intestinal homeostasis and injury repair processes.

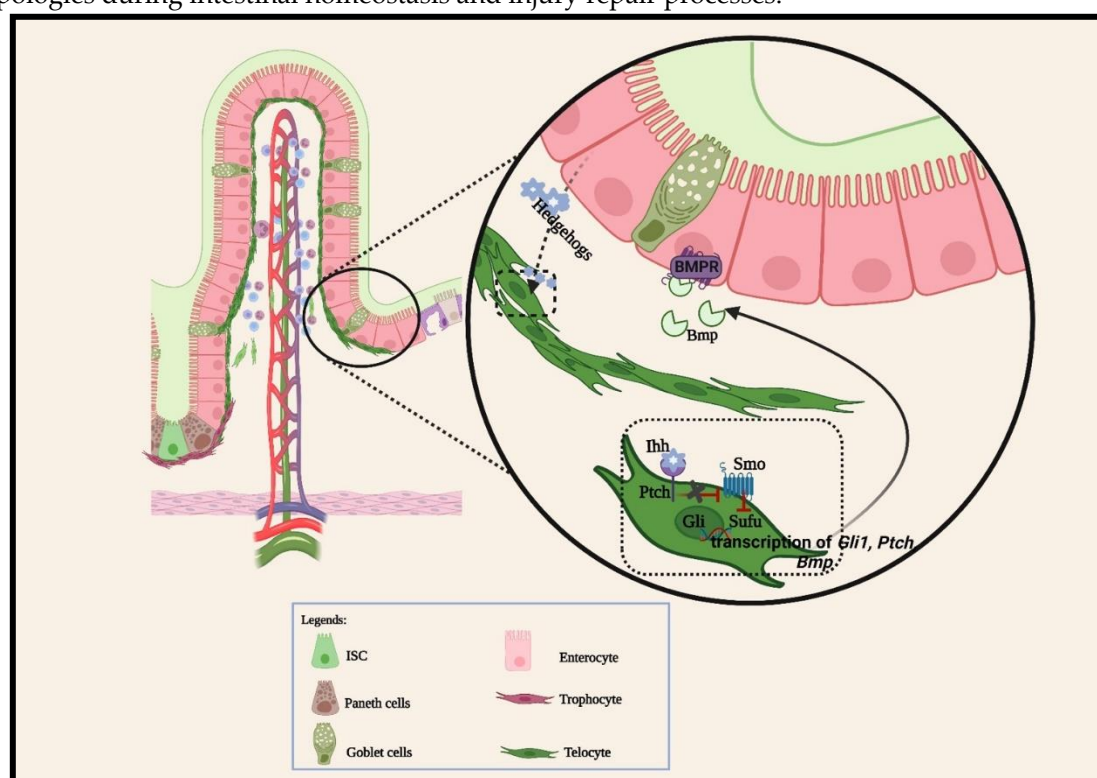


Figure 2. Intestinal villus base epithelial secreted ligands activate hedgehog signaling in neighboring mesenchymal cells. These epithelial–mesenchymal interactions at the villus base stimulate intestinal epithelial cell terminal differentiation and coordinate the migration of differentiated cells toward the villus tip. Hedgehog ligands produced by epithelial cells bind on the *Pitch1* receptor of PDGFR α^{hi} cells located at the villus base. This binding triggers the transcription of hedgehog target genes. The genes transcribed by the Gli family of PDGFR α^{hi} cells include *Bmps*. Secreted Bmp ligands from these iMSCs bind with the Bmp receptor (Bmpr) on adjacent epithelial cells. The Bmp and Bmpr binding subsequently phosphorylate the Smad family transcription factor, which in turn induces the differentiation of villus epithelial cells.

3. Intestinal cell plasticity and regeneration

The intestinal epithelium is a delicate but resilient organ. It undergoes rapid turnover of epithelial cells, predisposing it to luminal insults [75]. Epithelial cell exposure to luminal contents can cause severe damage induced by physical or pathogenic agents [21]. These perturbations necessitate addressing the concept of intestinal cell plasticity. While active Lgr5 $^{+}$ stem cells continuously proliferate to replenish epithelium during normal homeostatic conditions, the proliferating properties make them more susceptible to mucosal injuries, including irradiation or inflammation [76]. Studies showed that various mucosal injuries, such as inflammation, hypoxia, or irradiation caused Lgr5 $^{+}$ stem cell loss, thereby exacerbating intestinal epithelium regenerations [77-80]. Though the Lgr5 $^{+}$ stem cell pool is depleted during intestinal injury [78], another stem cell population (known as reserve stem cells) is activated to replenish the active stem cell pool [37,81,82]. This population of non-active Lgr5 $^{-}$ ISC plays a crucial role in regeneration following intestinal mucosal damage and helps maintain intestinal homeostasis [33,83-85]. For example, both DSS (Dextran sulfate sodium) and irradiation-induced Lgr5 $^{+}$ cell loss caused the proliferation of quiescent Clu $^{+}$ expressing cells to reconstitute columnar base Lgr5 $^{+}$ cells and replenish damaged epithelium during regenerations [33]. To buttress this idea, a previous study reported that the transitory loss of Lgr5 $^{+}$ stem cells did not disrupt intestinal architecture, and this could result from the reserved stem cells recruited following Lgr5 $^{+}$ stem cell loss [78,82]. This hypothesis corroborates the activation of putative reserve stem cells, including Bmi1, Hopx, and mTert, during Lgr5 $^{+}$ cell depletion [82,83,86,87]. It is suggested that quiescent cells are resistant to mucosal injury and play critical functions during the regeneration process [82,83]. Reserve stem cell populations are less proliferative and express high anti-apoptotic and DNA repair genes [33,88]. Though these properties make them a great candidate to replenish the active stem cell pool after mucosal injury [89,90], researchers still debate whether reserve stem cells at the +4 position of the crypt are responsible for regeneration [28].

Others proposed that several differentiated epithelial cells undergo fetal-like reprogramming and revert to active Lgr5 $^{+}$ cells during intestinal regeneration [23,24,67]. Secretary lineage seems to be the most prominent among the differentiated cell types that support intestinal regeneration during mucosal healing [91]. The dedifferentiation by these non-putative active stem cells to replenish the Lgr5 $^{+}$ pool is known as intestinal plasticity [28,82,92]. In the last decades, lineage tracing techniques using transgenic animals have provided the model to delineate cell migration [5,7]. This technique can track the specific cell lineage by editing the gene of interest. Van Es, et al. [42] explored this model to reveal that Dll1 $^{+}$, a secretory cell progenitor, can establish organoids containing Lgr5 $^{+}$ cells in *in vitro* culture. In a similar pattern, when irradiation depleted the stem cell pool *in vivo*, Dll1 $^{+}$ transformed into active Lgr5 $^{+}$ stem cells that proliferated into multiple progenitor lineages [42]. Another secretory progenitor cell, Atoh1 $^{+}$, was confirmed to generate Lgr5 $^{+}$ stem cells using different injury models such as irradiation and DTR (Diphtheria toxin receptor)-induced stem cell loss strategies [93]. To clarify the mechanisms behind the dedifferentiation phenomenon, several studies have tried to unravel the factors associated with cell plasticity and regeneration, but this area is still under active investigation. Recently, some studies suggested potential underlying mechanisms involved in the processes [21,28]. For instance, recombinant WNT3A supplements promote Dll1 $^{+}$ to produce organoids populated by Lgr5 $^{+}$ cells [94]. Furthermore, *Ascl2*, a Wnt target gene, was found

to promote Paneth cell dedifferentiation following the DSS treatment [84]. According to Yu, et al. [95], Notch signaling target genes, particularly *Hes1* and *Notch1*, are upregulated during Paneth cell dedifferentiation in irradiation-induced damage studies. Likewise, *Yap1*, a key hippo signaling target gene, was significantly upregulated in a DSS-induced study during epithelial regeneration [96]. Altogether, the crosstalk among Wnt, Notch, Hedgehog, and Hippo signalings needs further studies which will be relevant to understand their contributions to epithelial regeneration [2,97]. Unraveling these mechanisms could provide key insights into therapeutic targets to alleviate mucosal healing.

Non-epithelial cells, such as immune and iMSCs, also contribute to intestinal cell plasticity and regeneration. These cells secrete factors, including growth factors and cytokines, to stimulate mechanistic signalings that regulate regenerative processes [43,98,99]. They sense and are recruited to respond to injury repair. Immune cells specifically induced inflammation, and also recognized damaged associated molecular patterns released by apoptotic cells [100]. Besides immune cells, iMSCs are essential sub-epithelial cell types that are underappreciated during regeneration responses. Injured epithelial cells can send signals to iMSCs through hedgehog ligands, specifically *Ihh* and *Shh*, thereby enhancing ISC regenerations to support repair processes after epithelial injury [101-103]. Elevated levels of these ligands increased HH target gene transcription in iMSCs, including *Cyclin D1*, to promote epithelial regeneration [45,72,73]. Recent studies are unraveling potential iMSC subsets that are involved in intestinal epithelial repair functions. For instance, a study demonstrated that Gli⁺-expressing mesenchymal cells may secrete *Rspo3* to support epithelial repair processes using the DSS-induced damage model [73]. Moreover, DSS-induced colitis increased *fgf10*, *Vegf*, *Wnt2b*, *Grem1*, and *Rspo1* expressions in CD34⁺ cells, corroborating iMSCs roles during epithelial repair responses [45]. In conclusion, depleted active stem cell populations are certainly replenished during epithelial injury repair and regeneration, but there is still a knowledge gap on iMSC roles in supporting stem cell regenerative capacity. Understanding intestinal epithelial and MSC interaction could be the key target to treating relapsing intestinal diseases, such as inflammatory bowel diseases and necrotizing enterocolitis. Thus, it is important to elicit the contribution of multicellular crosstalk during intestinal cell plasticity and regeneration programs. We will discuss more details on the role of iMSCs during regeneration in the next chapter.

4. Subepithelial mesenchymal stromal cell characteristics

The intestinal mesenchymal stromal cells (iMSCs) are among the cell population localized in the lamina propria. The term “iMSCs” represents a group of non-epithelial, non-endothelial, non-neuronal, and non-hematopoietic cells that contribute growth factors and chemokines to regulate the intestinal epithelial homeostasis, regeneration, and immune functions [104-106]. These cells share surface markers with mesenchymal cells found in other tissues, giving them similar characteristics [107,108]. They were previously categorized by non-expressing α -smooth muscle actin (α -SMA-) fibroblast, α -SMA⁺ expressing myofibroblast, pericytes, and mesenchymal stem cells [55,66,107]. Before recent evidence, iMSCs were largely focused on their structural support functions to the epithelial architecture, as they form the largest component of the extracellular matrix (ECM) of the intestine [106]. Contrary to these past views, recent studies using immunostaining and single-cell RNA sequencing have revealed iMSCs as heterogeneous cell populations capable of dynamic multidirectional crosstalk with epithelial, hematopoietic, and immune cells [39,43,109]. Accordingly, iMSCs are dispersedly distributed along the crypt-villi sub-epithelial region in the lamina propria, *muscularis mucosa*, and deep in the *submucosa* with distinct morphology and functional support to epithelial cells.

4.1. Roles of intestinal mesenchymal stromal cells during prenatal intestinal morphogenesis

Intestinal epithelium develops from endodermal cells in the embryo [68]. During the early fetal stage, these cells in tube-like sheet layers invaginate to form pseudostratified epithelium (E9.5). This means that the proliferating epithelial cells are non-compartmentalized at the early embryonic stage [62,69,110]. Interestingly, cellular proliferation during early embryonic development survives in the

absence of Wnt/ β -catenin signaling despite the crucial role this mechanism plays in adult stem cell homeostasis [111]. Wnt target gene transcription factor (TCF) was found to be redundant during the pseudostratified epithelial development stage. Rather, mice lacking the *Tcf7l2* gene had significant epithelial shortening only after villi had been formed completely [112]. In another study, conditional β -catenin deletion did not affect pseudostratified epithelium proliferation during the early fetal stage [113]. Together, the studies indicate that there might be distinct signaling mechanisms between early-phase and late-phase embryogenesis.

The pseudostratified intestinal epithelium and mesenchyme rapidly increase from E9.5 to E14.5. By E14.5, the pseudostratified epithelium undergoes extensive remodeling into columnar epithelium, which leads to the emergence of villus structure. To gain insights into the underlying factors that potentially shape fetal gut compartments at this stage, Maimets et al. reported that CD29 and PDGFR α ⁺ cells are crucial for fetal gut vilification [20,46,69,111,114]. The CD29⁺ cells expressed high *Acta2*, *Myl4*, *Des*, and LRIG1 levels, which could be the progenitor for fetal gut *muscularis mucosa* cells. More importantly, they revealed how PDGFR α ⁺ cells guide the villigenesis of the pseudostratified epithelium [20,62]. This suggests that PDGFR α ⁺ cells could be a progenitor for stromal lineages because of their indispensable contributions to villi formation during morphogenesis, which aligns with previous studies that demonstrated iMSCs drive intestinal epithelial fate [20,66,115,116]. They hypothesized that PDGFR α ⁺ cells are crucial for villi emergence because they form clusters close to the expanding epithelium. This view is supported by their *in vitro* model study, which showed that PDGFR α ⁺ cells isolated from the fetal gut can promote organoid growth independent of essential growth factor supplements. In contrast to the early fetal stage, they also found that inhibiting *Porcn* in pregnant dams (a Wnt ligand upstream regulator) between E12.5 and E16.5 had no effects on the fetal colon, but abrogated SI villi formation. Based on these data we could deduce that Wnt gradients become relevant from mid-stage morphogenesis onward for maintaining the crypt-villus compartment. To corroborate this hypothesis, other studies have also highlighted that the underlying mechanisms of early embryonic morphogenesis are biologically different from those that regulate postnatal intestinal epithelium [113,117].

Embryonic mice at E14.5 exhibit rapidly expanding epithelial cells. By E16.5, these proliferating cells are restricted to the intervillous domain between neighboring villi and differentiated absorptive and secretory cells in villi [68]. A previous report suggests that embryonic iMSCs expressing *Dlk1* support rapid expansions of fetal gut morphogenesis, which differs from their role in adult mice [117]. To confirm this, PDGFR α ⁺ cells, which are the dominant iMSC cluster in the late fetal phase, reportedly expressed DLK1, but this expression decreased toward late gestation [117]. For these reasons, the current challenge is to better understand the PDGFR α ⁺ subset functional diversities as recent studies classified them as PDGFR α ^{hi}, PDGFR α ^{medium}, and PDGFR α ^{lo} cells during morphogenesis [20,44]. These subsets showed distinct characteristics and potentially have unique signaling properties.

Studies have previously predicted the indispensable roles of hedgehog signaling, involving SHH upregulation during villi formation [20,62,68]. Consistently, Hedgehog target genes, especially *Ptch1*, were detected in the absence of Wnt signaling. *Ptch1* is a key receptor, expressed by mesenchyme, for the SHH ligand, indicating that mesenchymal cells are indispensable during fetal gut formation. While the roles of non-Wnt pathways in fetal intestinal patterning during embryogenesis remain unclear, the available data suggest multiple signaling crosstalk such as Wnt, hedgehog, and Notch, among other factors produced by epithelial-mesenchymal cells regulate fetal gut morphogenesis [20,113].

4.2. Roles of intestinal mesenchymal stromal cells during intestinal homeostasis

The neonatal and mature intestinal epithelium is supported by heterogeneous stromal cell populations [38,39,44]. By the time of birth, the neonate gut will transition into a compartmentalized intestine driven by polarized signaling mechanisms. Specifically, the crypt and Paneth cells in the small intestine will start to emerge by the 14th postnatal day. By postnatal day 28, crypts would have rapidly expanded and matured, forming the intestinal epithelial crypt-villus structure [68,118,119].

Adult mice maintain intestinal homeostasis through several cellular programming, including ISC stemness, TA proliferation, and differentiation [9]. Each process requires unique gradient factors to keep the intestinal compartments in normal condition, in which iMSCs serve as a crucial source for all the essential ligands required for regional specificity of epithelial cell integrity. Remarkably, iMSC subtype heterogeneity increases as gut maturation progresses during postnatal development [39,44].

4.2.1. Recent classification of iMSC lineage

Although there has been substantial progress in characterizing epithelial cells [21], subepithelial iMSCs classifications, their cellular diversities, and functionalities remain challenging due to a lack of unique surface markers. Nevertheless, they are broadly categorized according to their regional specificity and functional diversity along the intestinal region using immunostaining, FACS, and transgenic mouse models. iMSCs expressing PDGFR α^{hi} , ACTA2 $^{\text{lo}}$ myofibroblast, FOXL1 $^+$, and GLI1 $^+$ but not vascular CD31 $^+$ markers are loosely regarded as telocytes due to their proximity to epithelial cells in the villus region [69,120,121]. Most cryptal iMSCs uniquely express CD34 $^+$ /GP38 $^+$ co-localizing with CD81 $^+$ and other PDGFR α^{low} cells, forming sub- and peri-cryptal stromal populations [44-46,69]. The subcryptal stromal populations produce trophic factors to support intestinal stem cell niche, and they are distinguished from *muscularis mucosa* cells due to their lack of *Myh11* $^+$ gene expression [43]. Despite being isolated and studied, those cells still display heterogeneity and remain undifferentiated in terms of their origin and possible lineages.

Recent studies using scRNA-seq assay have revealed a robust description of possible lineage, anatomic and physiologic heterogeneity of previously obscure iMSC subtypes [38,39,43]. One common proposition is that adult iMSC subsets originate from a similar embryonic precursor identified as Gli-expressing cells, which is still debatable [39]. For instance, most adult iMSCs express PDGFR α^+ at varying magnitudes which suggests PDGFR α^+ as another potential progenitor – more details on how iMSCs support embryogenesis are presented in the morphogenesis section [20,62]. Their localization probing and gene expression profiling showed a diverse population of iMSCs despite sharing the same lineage.

iMSC subsets that are sparsely localized beneath the epithelium produce gradient factors that shape the crypt/villi compartments. To appreciate the spatial distributions of these subsets, we reviewed recently characterized iMSCs and schematically illustrated them in Figure 3. These studies revealed distinct iMSC subsets that are classed under PDGFR α^{lo} (CD34 $^{\text{hi}}$ CD81 $^+$, CD34 $^{\text{hi}}$ Igfbp5 $^+$, CD34 $^{\text{lo}}$ Fgfr2 $^+$) and PDGFR α^{hi} (CD9 $^{\text{hi}}$ CD141 $^-$, CD9 $^{\text{lo}}$ CD141 $^+$, CD141 $^{\text{int}}$) [39,66,122].

4.2.1.1. Functions of pericryptal (PDGFR α^{lo} or CD34 $^+$ Gp38 $^+$) subpopulations

PDGFR α^{lo} subsets are found beneath the *muscularis mucosae* (CD81 $^+$), pericryptal (Igfbp5 $^+$), and the lamina propria (Fgfr2 $^+$). This idea is strengthened by the data reported in recent studies [38,39]. The studies suggested that gradients regulating ISC niche and terminal differentiation support the intestinal epithelial regional specificity.

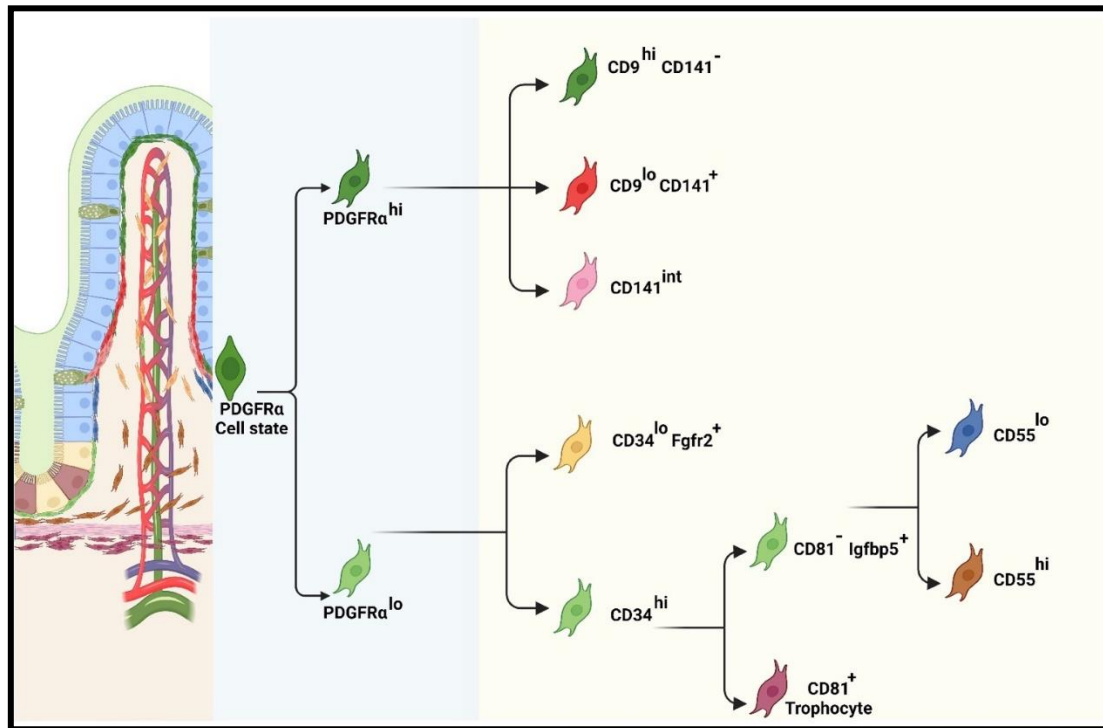


Figure 3. Schematic representation of intestinal mesenchymal stromal cell subsets and their location. The illustration here describes the subpopulations of PDGFR α expressing cells, including telocytes, CD81 $^{-}$ cells, and trophocytes. The trophocyte cluster group, CD81 $^{+}$, is confined to the subcryptal domain, beneath the *muscularis mucosa*, to secrete Wnt promoting factors that support the ISC niche. Other PDGFR α^{lo} subsets, including CD55 hi and Fgfr2 $^{+}$ cells, localize in the lamina propria and extend upward to the TA domain/villi trunk to initiate terminal differentiation. They switch the signal gradients from Wnt-promoting factors to Bmp agonists. PDGFR α^{hi} subsets form the subepithelial stromal populations that are localized in the villi core in the small intestine and the colon top [38,39].

To confirm the functional characteristics of these PDGFR α^{lo} subsets, clusters of PDGFR α^{lo} cells that expressed pericryptal CD34 $^{+}$ Gp38 $^{+}$ only emerged after birth and they are found to promote intestinal stemness [45]. The study demonstrated that CD34 $^{+}$ Gp38 $^{+}$ cells are the main Wnt ligand contributors to adult ISC homeostasis when the crypt is formed. This indicates that the intestinal epithelium requires a specific iMSC sub-population to reach maturation postnatally [45]. The idea corresponds with data showing that CD81 $^{+}$ cells located in the *muscularis mucosae*, a subset of the CD34 $^{+}$ population, are the key producer of *Grem1* that inhibits Bmp activities in the crypt [39,45]. PDGFR α^{lo} CD81 $^{+}$ *Ackr4* $^{+}$ (trophocytes) and PDGFR α^{lo} CD81 $^{-}$ CD55 hi cells are subsets of the CD34 $^{+}$ population that produce BMP antagonists for stem cell niche to promote ISC stemness near the crypt base [38]. The Shivdasani group also reported that PDGFR α^{lo} CD81 $^{+}$ cells expressed high *Wnt2b*, *R-spos*, and *Grem1* levels [55,66,104]. The depletion of *Rspo3*-secreting iMSCs that caused delayed gut maturation and reduced ISCs buttresses the functional specificity of PDGFR α^{lo} CD81 $^{+}$ cells [46,49,123].

The cryptal cells transitioned into differentiated cells just above the villus base. iMSCs that are found beneath the villus base and the corresponding lamina propria region produce pro-BMP factors [38,39]. For example, PDGFR α^{lo} CD81 $^{-}$ CD55 lo and CD34 lo Fgfr2 $^{+}$ cells expressed non-canonical *Wnt4* near the top of the colon crypt base, suppressed trophic factor effects, and also reinforced the BMP gradient activities, suggesting their roles in promoting terminal differentiation in the TA domain toward the villus region. Though they are PDGFR α^{lo} subsets, their transcriptomic data revealed they expressed a low level of *Wnt2b*. This is consistent with previous studies that showed BMP gradients increased apically in the SI villi or the colon crypt top [38,66]. In contrast, BMP inhibitors increased distally, especially from pericryptal PDGFR α^{lo} CD81 $^{-}$ cell to PDGFR α^{lo} CD81 $^{+}$ trophocytes beneath the

muscularis mucosae [39,44,45]. Though it is currently difficult to sort, culture, and investigate all of these distinct iMSC sub-populations, studies have shown that they may functionally overlap, express related molecular profiles to perform complementary roles, and in some cases, provide opposing cues to maintain intestinal homeostasis and development (Table 2). Collectively, the current observations depict that epithelial development relies heavily on gradients from diverse iMSCs to maintain homeostasis despite the high turnover rate of epithelial cells [104,124,125].

Table 2. Distinct intestinal stromal subsets supporting intestinal balance regulations.

Non-epithelial cells	Cell Markers	Ligands	Functions	Signaling pathways	References
PDGFRα ^{lo} Cells	CD81 ⁺ , CD55 ^{hi}	<i>Wnt2b</i> , <i>Gremlin1/2</i> , <i>Rspo3</i>	Wnt promoters	Wnt/β-catenin	[38,66]
	Fgfr2 ⁺ , CD55 ^{lo}	<i>Wnt4</i> , <i>Frzb</i> , <i>Sfrp1</i>	Wnt repressors	Non-canonical Wnt/Bmp	[38,39]
PDGFRα ^{hi} Cells	FOXL1 ⁺ , CD9 ^{hi} CD141 ⁻ , CD9 ^{lo} CD141 ⁺ , CD141 ^{int}	<i>Bmp3/4</i> , <i>Wnt5a/b</i> , <i>Dkk</i>	BMP agonists, Wnt inhibitor	Non-canonical Wnt/Bmp	[39]
PDGFRα ⁺	PDGFRα ⁺ DLK1 ⁺	<i>Dlk1</i>	Embryonic morphogenesis	Notch	[117]
LTβR ⁺	LTβR ⁺ PDGFRα ^{hi}	<i>Pdgf</i>	Stromal maturation	Bmp activation	[44]
Smooth muscle cells	<i>Tagln</i> ⁺ , <i>Acta2</i> ⁺ , <i>Myh11</i> ⁺	<i>Wnts</i>	ISC integrity and wound healing	Wnt/β-catenin	[43]
Immune cells	ILC2, ILC3	<i>Il13</i> , <i>Il22</i>	Promote regeneration	Wnt/β-catenin	[126]

4.2.1.2. Functions of PDGFRα^{hi} subpopulations

PDGFRα^{hi}Foxl1⁺ cells are reported to be a good source of BMP-promoting factors, including *Bmp4*, *Bmp5*, and *Bmp7* [38,39,45,66]. Paerregaard et al. reported that the three PDGFRα^{hi} subsets expressed *Foxl1*. 1) *Nrg1* was expressed in CD9^{hi}CD141⁻; 2) *Cxcl12*, and *Acta2* expressed in CD9^{lo}CD141⁺; and 3) *Adamdec1*, *Wnt4*, and *Acta2*⁺ were expressed in CD141^{int} cells [39]. While all these PDGFRα^{hi} subpopulations produce BMP (*Bmp5*, and *Bmp7*) gradients, CD9^{lo}CD141⁺ had the highest levels of *Wif1*, *Bmp3*, and *Bmp4*, contributing to terminal differentiation of intestinal epithelial progenitor cells. Upregulations of *Wif1*, a protein that binds and inhibits canonical and non-canonical Wnt ligands (such as *Wnt3a*, *Wnt4*, and *Wnt5a*) [127], corroborated the hypothesis that Wnt signaling activities are repressed from the villus base toward the tip. Interestingly, using RNA velocity analysis, these three PDGFRα^{hi} subsets were predicted to originate from CD34^{lo}Fgfr2⁺ found along the lamina propria of the small intestine in mice.

According to a recent study, post-natal intestinal epithelium required PDGFRα⁺ cell-dependent maturation to shift from immature proliferative compartments into distinctly functional crypt-villi regions [44]. In their study, they generated reporter and inducible lineage tracing models for lymphotoxin beta receptor (LTβR) cells by crossing LTβR^{fTA} mice with Rosa26^{flloxSTOP-YFP} mice. Using the inducible lineage tracing technique, some fractions of PDGFRα⁺ subepithelial stromal cells developed from LTβR^{YFP} progenitor cell lineage before the mice reached weaning age. To confirm this hypothesis, they also generated a direct LTβR^{GFP} reporter model to show that PDGFRα^{hi} cells, found close to the villus epithelial cells, expressed GFP [44]. Additional transcriptomic analysis revealed that LTβR⁺PDGFRα⁺ expressing cells showed upregulated levels of hedgehog and pro-differentiation gene markers that are specific to PDGFRα^{hi} expressing cell population, especially *Ptch1*, *Foxf2*, *Gli1*, *Bmp4*, and *Bmp5*. In contrast, the pro-stem cell niche gene markers (i.e., *Cd34*, *Cd81*,

Rspo2/3, and *Grem1/2*) were downregulated in the transcriptome data of LT β R⁺PDGFR α ⁺ cells [39,44,66]. Inducible conditional ablation of PDGFR α in LT β R lineage (LT β R^{APDGFR α} mice) caused increased ISC markers (*Olfm4*⁺ and *Lgr5*⁺) in the gut compared with the wild type. Consistently, LT β R stromal lineage from LT β R^{APDGFR α} mice had decreased pro-differentiating factor (specifically, *Bmp2*) and increased pro-stemness factor (*Grem2* and *Chrdl1*) levels in their stromal cell transcriptome data. Collectively, these observations demonstrated in their studies suggest that the LT β R⁺PDGFR α ⁺ cells expressing pro-differentiation signals share similar characteristics with the PDGFR α ^{hi} subsets reported by Paerregaard et al., and they are essential for intestinal epithelial maturation before weaning [39,44,45]. More importantly, their results showed that the appearance of PDGFR α ^{hi} cells not only influences epithelial cell differentiations but is also essential for the transcriptomic switch of functionally distinct LT β R⁺ stromal lineage cells towards maturation and localization during early postnatal gut development [44]. Accordingly, different iMSC subsets create a functionally distinct enabling environment for polarized signaling crosstalk to regulate epithelial cell fate during development. To sum it up, iMSC populations are less diverse during intestinal morphogenesis but become heterogeneous postnatally to support mature intestinal epithelial homeostasis [38,39,43,69].

4.3. Roles of intestinal mesenchymal stromal cells during intestinal injury and repair

iMSCs are not only crucial for intestinal morphogenesis and homeostasis; recent reports have also revealed their modulatory functions using DSS, irradiation, and DTR-induced injuries in transgenic mice. For example, RBP1⁺ cells, confirmed to be a subset of GLI1⁺ cells, are suggested to sense DSS-induced colitis injury, thereby stimulating *Rspo3* production to promote the injury recovery process [73]. Likewise, conditional deletion of the *Wntless* allele in a *Villin-Wls*^{CKO}, a protein required for Wnt secretion, triggered GLI1⁺ cell expansion during injury. This observation was confirmed using recombinant Shh ligand to promote GLI1⁺ cell response to injury by compensating for epithelial Wnt loss. The results suggested that SHH signaling could be a potential regulatory target for enhancing mucosal healing by activating iMSCs. In another study, pericryptal CD34⁺ cells, in response to DSS treatment, were reported to migrate and localize under the colonic crypt-apical epithelial cells [128]. Following the induced epithelial injury, the mRNA data revealed upregulated levels of *Bmp2*, *Bmp3*, *Bmp7*, and *Wnt5a*, indicating the plasticity potential of CD34⁺ to support reepithelization during colon regeneration [128].

According to the reviewed studies above, mesenchymal cells support epithelial regeneration by secreting growth factors and other signaling molecules that promote the proliferation and differentiation of epithelial cells. scRNA seq analyses showed that iMSCs are intricately involved in epithelial cell response to injury recovery processes. iMSCs such as CD81⁺ and CD81⁻ subsets play significant roles during injury states, by increasing the production of Wnt-promoting factors [45,73]. For instance, *Grem1* and *Rspo3* reportedly increase in PDGFR α ^{lo} cells during colitis-induced conditions, suggesting that these iMSCs support epithelial regeneration [116]. In another study, non-coding RNAs (miR-143/145) specific to iMSCs were also involved in regulating IGF1 signaling response to repair DSS-induced epithelial damage [129]. In all, iMSCs are essential for the repair and regeneration of the intestinal epithelium following injury. However, more studies are needed to clarify the relevance of specific iMSC subsets that engage in different disease resolution processes.

5. Conclusion and Future Perspectives

The iMSC populations play a crucial role in maintaining intestinal morphogenesis, homeostasis, and regeneration by providing support and signaling to the epithelial cells. Recent studies have shown that iMSCs belong to different subsets that have distinct regulatory functions along the crypt-villi axis. However, several obstacles need to be resolved to fully understand their specific characteristics, functions, and responses during homeostasis and epithelial regeneration. To advance our understanding of iMSCs, three critical areas require immediate attention.

Firstly, researchers need to develop specific markers to further characterize homogenous iMSC subsets within clusters of notable iMSC subpopulations addressed in this paper. This approach will help to address the conflicting results on the iMSC population that have been reported to provide

opposing signaling factors they secrete at distinct locations along the crypt-villi axis. Improved iMSC sub-type sorting techniques would clarify the unique functional diversity of specific homogenous iMSCs and how each distinct subset interacts with different epithelial cells. As such, researchers will be able to define the contributions of the overlapping molecular signatures by the heterogeneous iMSCs clusters during normal and regeneration conditions. Thus, this is a call to develop novel iMSCs surface markers for cell sorting assays which will advance the current knowledge about epithelial and mesenchymal crosstalk.

Second, to unravel the roles of characterized iMSC subsets during morphogenesis and disease development, researchers should improve lineage tracing techniques that focus on examining the response of iMSCs to regeneration. For example, iMSCs are currently proposed to have multidirectional relationships with both epithelial and immune cells. While iMSCs supply Wnt and Bmp agonists during normal conditions, it remained obscure whether (1) they directly provide regenerative factors to repair epithelial cells, (2) adopt homing effects to replace damaged epithelial cells, or [41] indirectly activate anti-inflammatory immune cell responses to restore intestinal homeostasis after injury. Future research directions in this field could include investigating the mechanisms by which subepithelial stromal cells regulate stem cell behavior and the role of these cells in chronic diseases of the intestine, such as inflammatory bowel disease. Additionally, understanding the interactions between subepithelial stromal cells and other cell types may lead to the development of novel therapeutic strategies for promoting intestinal regeneration and repair.

Finally, despite the difficulty relating to investigating functional studies *in vivo* due to their delicate nature, the future approach needs to improve scRNA seq techniques, develop special iMSCs culture conditions that can support the viability of FACS cells, and improve co-culture models to better understand the complexity of mesenchymal and epithelial crosstalk.

In conclusion, this paper has provided a comprehensive review of the recent evidence about iMSCs heterogeneity, suggesting their spatial distribution which enables them to perform distinct or overlapping functions in maintaining epithelial integrity. Future studies should reconstruct 3D organoid co-culture set-ups, such as transwell and scaffold models, to elucidate the spatial organization of iMSCs in the intestine for different developmental stages and address potential communications between epithelial and sorted iMSC subsets. By addressing these challenges, we can gain a better understanding of the complexity of the iMSC niche and develop novel therapeutic strategies for promoting intestinal regeneration and repair.

Author Contributions: HA: conceptualization, image design, original writing draft, revision; EML: revision; YJ: conceptualization, image design, revision, funding.

Acknowledgments: The authors appreciate the financial support from the Maryland Stem Cell Research Fund and MAES Competitive Grant Program at the University of Maryland, College Park, USA. We used Biorender to illustrate the images presented in this paper.

Conflicts of Interest: The authors declare no conflicts of interest.

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