Gastrointestinal epithelial innate immunity—regionalization and organoids as new model

Özge Kayisoglu¹ · Nicolas Schlegel² · Sina Bartfeld¹

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Abstract
The human gastrointestinal tract is in constant contact with microbial stimuli. Its barriers have to ensure co-existence with the commensal bacteria, while enabling surveillance of intruding pathogens. At the centre of the interaction lies the epithelial layer, which marks the boundaries of the body. It is equipped with a multitude of different innate immune sensors, such as Toll-like receptors, to mount inflammatory responses to microbes. Dysfunction of this intricate system results in inflammation-associated pathologies, such as inflammatory bowel disease. However, the complexity of the cellular interactions, their molecular basis and their development remains poorly understood. In recent years, stem cell–derived organoids have gained increasing attention as promising models for both development and a broad range of pathologies, including infectious diseases. In addition, organoids enable the study of epithelial innate immunity in vitro. In this review, we focus on the gastrointestinal epithelial barrier and its regional organization to discuss innate immune sensing and development.

Keywords Gastrointestinal tract · Immunity · Regionalization and organoids

Introduction
The gastrointestinal (GI) tract is required for the digestion of food and spans from the oral cavity via the oesophagus, stomach, small intestine and large intestine to the anus. The GI lumen is colonized by a vast variety of commensals, symbionts and occasionally pathogens. The microbial colonization follows a gradient with less than 10³ microbes/ml in the stomach to 10³–10⁷ microbes/ml in the small intestine and 10¹¹–10¹² microbes/ml in the colon (reviewed in [1–3]). From the glandular stomach onwards, the GI tract is lined by a single layer of columnar epithelial cells. This epithelial layer comprises of different specialized cells, tightly interlocked by junctional protein complexes, enforcing the physical barrier (Fig. 1).

The interaction between the gut microbiota as well as ingested pathogens and epithelial cells is mediated by pattern recognition receptors (PRRs), including Toll-like receptors (TLRs), nucleotide-binding oligomerization domain (NOD)–like receptors (NLRs) and other cytosolic receptors (Fig. 2, reviewed in [7, 11]). These PRRs play a key role in recognizing microbe-associated molecular patterns (MAMPs) and damage-associated molecular patterns (DAMPs). Upon PRR activation in epithelial cells, downstream signalling cascades induce expression of different cytokines and chemokines via inflammatory pathways, such as the NF-κB pathway, to direct the professional immune cells (reviewed in [8, 12–16]). In addition to the classical PRRs, further sensors of bacterial activities, such as alpha-kinase 1 (ALPK1) have recently been discovered [17–19]. Inflammasomes, cytoplasmic complexes composed of NLR proteins, recognize additional molecular patterns, such as bacterial metabolites (reviewed in [7, 9]).

Gastrointestinal innate immune responses, including PRR sensing, must balance the need for providing protection from potentially harmful pathogens with being able to tolerate exposure to the diverse luminal microbiome. Hence, the expression and function of PRR signalling are expected to have a major impact not only on pathogen sensing but also on tissue homeostasis (Box 1) and inflammatory diseases including acute gastrointestinal, gastritis and inflammatory bowel diseases (IBDs)
Although each of these diseases displays completely different pathogeneses, they share a critical feature: the interface between the environment and the body is significantly disturbed. However, there is an ongoing debate whether changes of the intestinal epithelial layer, in particular those affecting its innate immune function, are a cause or a consequence of the above-mentioned diseases. Most knowledge about PRR signalling has been gathered from research on haematopoietic cells. It has been a challenge to discriminate the functions of PRR signalling in epithelial cells from those of the infiltrating immune cells. Major obstacles include difficulties in isolating pure epithelial cells and raising reliable antibodies against PRRs. With the advancement of organoids that derive from intestinal epithelial stem cells, a reductionist experimental model is now available that enables investigations into the innate immune response of the primary epithelial cells. Organoids are defined as stem cell-derived, 3-dimensional cell cultures that have self-organizing capacity and retain some of the function of the original organ (e.g. secretion, filtration, absorption, contraction). Since organoids can be grown either from tissue-resident adult stem cells (ASCs) or from pluripotent stem cells (PSCs), the cells are non-transformed. Together, the two types of organoids cover an extensive repertoire of organs that can be mimicked (reviewed in [4, 22, 23]). Both technologies have their individual advantages (reviewed in [4, 22, 23]). For example, cultures of ASC-derived organoids residing at the base of the crypt, have a lifespan of about 3–6 weeks. Similarly, gastric glandular cells, such as chief cells, residing below the isthmus region have a lifespan of several months. In contrast, cells above the amplification region move conveyor belt-like towards the surface, finally reaching the gastric pit, the small intestinal villus or the colon crypt opening, where they are shed into the lumen after a lifetime of only 3–5 days [4–6]. Both organs have absorptive enterocytes and secretory cells, such as mucus-producing goblet cells, hormone-producing enteroendocrine cells and tuft cells. In addition, the small intestinal epithelium also contains Paneth cells, which produce antibacterial peptides, as well as specialized microfold (M) cells on the Peyer’s patches, enabling crosstalk between the microbiota and the immune system.
have a tremendous expansion potential and are relatively homogeneous, and PSC-derived organoids are more complex in the sense that they combine cells of very different developmental origins (e.g. epithelial and mesenchymal cells). PSC-derived organoids allow the analysis of the developmental steps but may not reach the full level of differentiation into epithelial layer a sf o u n di nv i

This review will highlight insights from studies using organoids and discuss the potential of this technology for the study of epithelial innate immunity. We focus on the regional organization in the gastrointestinal tract.

Regional identity and innate immune signalling in the GI tract

When looking at gastrointestinal diseases it is important to note that some of them are confined to a specific section of the gastrointestinal tract. IBD includes Crohn’s disease (CD) and ulcerative colitis (UC), which show a differential and disease-specific pattern of inflammation: while UC begins in the rectum and is found in the colon, CD can affect all parts of the GI tract from the oral cavity to the anus. Furthermore, CD is characterized by segmental, discontinuous inflammation within the GI tract, while UC is usually described as a continuous inflammation of the colon (reviewed in [26, 27]). In the oesophagus, stomach and colon, cancer incidence is high, and infection and inflammation can promote development and progression of these cancers (reviewed in [28, 29]). In contrast, malignant transformation in the small intestine is very rare (reviewed in [30]). The apparent segment-specificity of these diseases in the GI tract remains enigmatic, but it is tempting to speculate that their region-specific origin is rooted in region-specific disturbances of the tightly balanced system of epithelial barrier function, innate immunity and mucosal regeneration. Therefore, it is interesting to highlight differences in the segments within the GI tract.

The GI tract comprises several anatomically defined segments with vastly distinct physical functions (reviewed in [31]). The main function of the stomach is the digestion of food and the elimination of incoming pathogens by gastric acid. Nutrients do

**Fig. 2** Simplified overview of PRR signalling pathways in gastrointestinal epithelial cells. Pattern recognition receptors (PRRs) including TLRs, NLRs or newly discovered sensors such as ALPK1 recognize the microbe-associated molecular patterns (MAMPs). Upon PRR activation in epithelial cells, downstream signalling cascades induce expression of different cytokines and chemokines via inflammatory pathways, such as the NF-κB pathway. TLR1, 2, 3, 4, 5 and 9 recognize lipoproteins, double-stranded (ds-) RNA, lipopolysaccharides (LPS), flagellin and dsDNA, respectively. ALPK1 recognizes the LPS metabolite ADP-heptose and its stimulation leads to phosphorylation of TIFA proteins, mediating the formation of TIFAsomes as a response to gram-negative bacteria. Ligands bind to a receptor which leads to the recruitment of adaptor proteins (e.g. Myd88, TRAF6 and RIP2). These adaptors drive the phosphorylation of the IκB which leads to its ubiquitination and degradation. NF-κB subunits p65 and p50 can then enter the nucleus to facilitate the expression of target genes which are proinflammatory cytokines such as IL-8 in humans and its analogue Cxcl2 in mice. Inflammasomes, which are cytoplasmic complexes composed of NLR proteins, recognize additional molecular patterns, microbial metabolites or nucleic acids. They activate caspase-1, which cleaves and thereby activates proinflammatory cytokines like pro-IL-1β and pro-IL-18, driving the downstream inflammatory pathways. In turn, proinflammatory cytokines will recruit professional immune cells of the innate and adaptive immune system, which are equipped to resolve the infection (reviewed in [7–10]).
not have to reach the epithelial layer of the stomach; thus, the body heavily invests in a protective mucus barrier, shielding the epithelial cells not only from its own acid but also from luminal content (reviewed in [32, 33]). By contrast, the main function of the small intestine is not only digestion but also the uptake of nutrients. In keeping with this, the small intestine has a significantly enlarged surface area due to the villi protruding into the gut lumen, where cell surfaces come in close contact with the nutrients. Most of the digestion takes place at the proximal small intestine, duodenum and jejunum, where the villi are long and thin. The jejunum has the highest ratio of Paneth cells secreting antimicrobial peptides [34], which decorate the rather loose mucus, thereby guarding the epithelial layer and keeping the crypts sterile (reviewed in [35]). The villi become progressively shorter and broader towards the ileum (reviewed in [36]), where again, the mucus takes over an important part of the protection: with the highest ratio of goblet cells, the ileum has a thicker mucus layer and a lower rate of digestion and absorption than the jejunum [34] (reviewed in [31]). Lastly, the colon reabsorbs water and invests into an extensive, thick and bi-layered mucus cover to be able to safely harbour trillions of commensal bacteria. The colon has no villi and the crypts are smaller than those of the small intestine. There are no Paneth cells, and the goblet cell ratio can be up to 25% of the epithelial layer [37]. Thus, the three gut segments have different strategies for maintaining a safe distance between the epithelial cells and the microbiota. MAMP recognition and activation of immune pathways is another layer of this interaction and therefore, it is only reasonable that they are also structured along the GI tract (reviewed in [38]). However, it remains a fascinating riddle what exactly shapes the structure of the PRR organization.

While it is intuitive that genes important for regional functions, such as digestion and nutrient uptake, follow spatial compartmentalization along the cephalocaudal axis (reviewed in [39]), such an organization was not expected for epithelial innate immune signalling. Previous studies had reported regulation of PRR signalling in response to stimulation with MAMPs. For example, TLR4 responsiveness decreases after birth, presumably because of the exposure to LPS during delivery and subsequent colonization of the gut [40]. Also, stimulation of TLR9 with its ligand CpG-DNA leads to a decrease of Tlr4 expression and inhibits TLR4 signalling [41]. Thus, it was expected that contact with the microorganisms, their molecules and metabolites in the GI tract would lead to silencing of PRR expression towards the gut lumen (reviewed in [14]). However, contradicting results, caused by technical difficulties such as unreliable antibodies targeting TLRs, led to confusion in the field and it was unclear, whether a particular PRR was expressed or not (reviewed in [42]). Early Northern blots for mRNA of Tlr2 and Tlr4 in ex vivo isolated epithelial cells indicated that expression levels of these two Tlr molecules were segment-specific: Tlr2 was expressed mainly in the colon, while Tlr4 was mainly expressed in stomach and colon. The authors termed this “strategic compartmentalization” of these TLRs [43]. Recent studies have now discovered that this principle of segment-specific expression extends beyond these two TLRs, and have unravelled a highly complex regional organization of PRR signalling that does not always follow the microbial load [44, 45].

The first of the two studies came from the Barton group and revealed several levels of organization of TLR expression. The group generated five strains of reporter mice, enabling expression analysis of TLR2, 4, 5, 7 and 9, respectively. TLR2 and 5 were expressed in the small intestine and the proximal colon, TLR4 was expressed in colon and TLR7 and 9 were not expressed. Reporter expression in organoids from these mice closely mimicked the in vivo expression, indicating that the expression is independent of contact with the microbiota or with immune cells [45].

The second study used a biobank of freshly generated human and murine organoids covering the different segments of the GI tract: corpus, pylorus, duodenum, jejunum, ileum and colon. Transcriptional analysis of the organoids confirmed the expression patterns for the TLRs reported by the Barton group, but in addition revealed a vast extent of differential expression of TLRs, NLRs, inflammasome components and other innate immunity-related genes (Fig. 3 and [44]). For example, Nod2 was mainly expressed in the stomach, while expressions of several inflammasome pathway components like Nlrp1b, Nlrp6 and Aim2 were restricted to the intestine. Various receptors were uniformly expressed; for example, Tlr3 expression was detected in every segment of the murine GI tract in high amounts, whereas Tlr5 was expressed in every segment but in low amounts [44]. As a result, each segment appears to have its very own, specific complement of innate immune receptors and signalling components.

Comparing the murine with the human gastrointestinal organoids demonstrated that in both species the principle of complex organization of PRR signalling components was the same, but individual PRR expression profiles could differ. Some PRRs, like TLR4 and NLRP6, were similarly patterned along the GI tract of both species. TLR4 was expressed predominantly in the stomach and colon and NLRP6 expression was restricted to the segments of the intestine. On the other hand, several PRRs such as TLR1 and TLR2 showed patterns that differed between both species. While in murine organoids, expression of both increased along the GI tract and was highest in the colon, in human organoids expression was highest in the stomach and decreased along the GI tract [44]. Moreover, expression of TLR5 was higher in human organoids, with the highest levels observed in the stomach [44].

As mentioned above, organoids also allow testing of whether a particular pathway is functional by addition of the ligand and subsequent analysis of downstream target gene expression. Complementing the region-specific expression...
analysis, the study also found that human and murine organoids show region-specific function: the murine stomach responded to the TLR4 ligand LPS, but not the TLR2 ligand PAM3CSK4 or the TLR5 ligand flagellin, by upregulating the NF-κB target gene Cxcl2. Murine jejunum upregulated Cxcl2 in response to PAM3CSK4 and flagellin, but not in response to LPS. In addition, the colon expressed Cxcl2 in response to all of these 3 tested ligands [44]. In contrast, human organoids from all regions expressed the human Cxcl2 analogue IL-8 in response to flagellin, but not in response to LPS and PAM3CSK [44]. Thus, not only expression but also function of PRRs in the gut is highly organized and segment-specific. These experiments were conducted by adding the stimulus to the organoid medium, thus only stimulating the basal side, since the apical side faces the sealed-off lumen of the organoids. Other experiments demonstrated that there is an even higher level of organization, since PRR expression within a segment can be restricted to specific cell types or even subcellular locations, such as only the basal compartment. These are outlined in more detail below.

Every region of the GI tract expresses its specific set of innate immune genes. Although a generalization is difficult, the current data suggest that TLRs are expressed most highly in the stomach and colon, while the small intestine is characterized by expression of inflammasome components. Currently, the underlying molecular mechanism and the evolutionary benefit are unclear. We suspect that the physical and chemical barriers covering the epithelial layers in the different gut segments each require monitoring of specific threats. For example, the stomach and colon both have a bi-layered, thick mucus layer with the inner mucus layer firmly attached to the epithelial layer, while the small intestine is covered by a thin, viscous mucus layer [46]. The different constitutions of the mucus layers in turn are likely due to the different functions of the respective gut segments (digestion vs nutrient uptake vs water resorption). It is conceivable that the mucus thickness influences the type of PRRs necessary at the specific sites.

**Specialized cells with immune function in the epithelium**

A long-standing concept in epithelial innate immunity is the existence of cell type-specific innate immune recognition. Prototypes of the specialized epithelial cells with defence function are the microfold (M) cells and the Paneth cells—recently complemented by a specialized goblet cell, the sentinel goblet cell. The existence of such specialized cells highlights the intricate and redundant systems that ensure the balance of nutrient uptake, co-existence with the microbiota, and monitoring of possible invaders. Directed differentiation of organoids now also allows the study of these cells in cell culture.
M cells are located in the follicle-associated epithelium covering the Peyer’s patches. They have a unique morphology with irregular brush border and reduced microvilli structures. Their role is to transport antigens in the gut lumen across the epithelial layer to the underlying lymphoid tissue for regulation of immune responses [47]. M cells themselves as well as the follicle-associated epithelium were shown to express several TLRs [48] (reviewed in [14]). However, M cells have been difficult to study since there are not many in the epithelial tissue and they are only found near the complex structure of Peyer’s patches [47]. Nevertheless, M cells can be generated in organoid cultures using directed differentiation. For this, receptor activator of NF-κB ligand (RANKL) is added to the medium, which upregulates the transcription factor SpiB, which is characteristic for M cell differentiation [49]. Organoids generated from mice genetically deficient for the NF-κB subunit RelB are not able to produce M cells after stimulation with RANKL, indicating that NF-κB activation is essential for development of M cells [50]. In human organoid culture, in addition to RANKL, lymphotoxin and retinoic acid are crucial for inducing differentiation towards M cells. These M cells specifically take enteric viruses like rotavirus and reovirus, indicating correct phenocopying of the natural M cell function also in organoid cultures [51]. Future organoid work will have to continue to unravel the interconnection of epithelial NF-κB signalling, M cell development and communication with immune cells.

Paneth cells are intermingled with the intestinal stem cells at the base of the crypt and have long been considered to be the guardians of the stem cell compartment because they secrete antimicrobial peptides. For example, Paneth cells secrete alpha-defensins, a process shown to be regulated by microbial patterns and innate immune mechanisms [52, 53]. In addition, NOD2, which was first detected in the crypt region of the murine small intestine [54], regulates the secretion of several alpha-defensins by Paneth cells, which in turn leads to activation of adaptive immunity. Paneth cells also express TLR5 and Paneth cell–enriched organoids express especially high levels of TLR5. RNA sequencing after stimulation reveals that while normal small intestinal organoids only express moderate levels of the TLR5 downstream target genes in response to stimulation with flagellin, such as NF-κB-induced cytokines, organoids directed to contain high numbers of Paneth cells mount a much stronger response, indicating that Paneth cells are the main responders to flagellin in the small intestine [44, 45]. In contrast to target gene expression, the most dramatic response of Paneth cells (degranulation, extrusion and cell death) is not triggered by stimulation with TLR ligands rather requires stimulation with the professional immune cell–derived cytokine interferon-gamma [55]. These results from organoids are in agreement with the observations showing Paneth cell degranulation and extrusion after interferon-gamma stimulation in vivo [55]. This elegantly underlines the system of checks and balances in the epithelial layer.

Goblet cells are important for epithelial defence because they produce the glycosylated mucins crucial for the formation of a mucus barrier on the epithelial layer (reviewed in [56]). In mice, MUC2 deficiency results in spontaneous inflammation and increases susceptibility to infection [57, 58]. Recently, a subset of goblet cells named sentinel goblet cells was described in the mouse colon. Using tissue explants, the study identified a thickening of the mucus layer in response to exposure to TLR1/2, 4 and 5 ligands, but not to TLR9, NOD1 and NOD2 ligands [59], in congruence to a previous report of Tlr 2,4 and 5 being expressed in goblet cells [60]. The authors determined that the response also depended on the presence of the Nlrp6 inflammasome and was independent of mucosal lymphocytes using tissues from a range of knockout mice [59]. Also, a previous report had demonstrated the importance of the NLRP6 inflammasome for mucus secretion by goblet cells [61]. Imaging revealed that specific goblet cells located in the apical regions of the crypt endocyted fluorescently tagged LPS [59]. These newly termed sentinel goblet cells not only undergo rapid degranulation and epithelial expulsion after treatment with the TLR ligands but also transmit a calcium signal to neighboring cells via intercellular cytoplasmic bridges formed by gap junctions, likely stimulating other goblet cells to increase mucus secretion [59]. Both goblet cells and Paneth cells belong to the secretory lineage. Organoids allow directed differentiation towards both cell identities, yielding organoids heavily enriched in either goblet cells or Paneth cells [62]. Comparing transcriptomes of these skewed organoids helped identify key regulators of the differentiation pathway [63]. Further analysis of the omics data as well as functional analysis of these organoids will allow better understanding of the role of both cell types in innate immune defence.

Lastly, stem cells themselves have also been reported to express specific PRRs such as TLR4 [64, 65], which are not found on the murine small intestinal villi or Paneth cells [66]. In addition, the majority of Nod2 expression in the murine crypt appears to be restricted to stem cells [67]. Stimulation with NOD2 ligand increased survival of stem cells and formation of organoids, indicating that stimulation of PRRs may also regulate gut epithelial regeneration directly.

Without a doubt, the current efforts to generate atlases of gene expression covering every cell type in ever more detail will soon give a more complete picture of the cellular organization of innate immune signalling in the gut as well as in the entire body [68–71].

Cell polarity and side-specific innate immune responses

Finally, it is also relevant to consider that gastrointestinal epithelial cells are highly polarized, with a specialized apical side facing the lumen of the gut with its microbiota, and a basolateral side facing the tissue. Under homeostasis,
MAMPs reach the apical side only. However, when the epithelial barrier is breached, microorganisms can also challenge the basolateral side. It has thus been hypothesized that epithelial cells may selectively mount a proinflammatory response only when stimulated from the basolateral side, in order to match the threat posed by the signal. For example, TLR9 was demonstrated to induce distinct signalling pathways when stimulated from the apical or basolateral side in cancer cell lines [72] and TLR5 only induced the NF-κB response gene IL-8 when stimulated from the basal side [73].

While earlier studies using antibody labelling against TLRs have reported specific expression on one side only (reviewed in [14, 16]), analysis of TLR reporter mice using staining of an HA tag did not confirm this, but instead demonstrated TLR2, 4 and 5 receptors on both apical and basal sides of the proximal colon as well as some intracellular TLR4 [45]. These apparent differences are likely due to the different technical approaches.

Organoids now allow direct functional testing of side-specific immune responses, since the cellular polarization is retained in organoids. Under standard conditions when organoids are grown in an extracellular matrix, the apical side faces the lumen of the organoid and the basal side faces the extracellular matrix [74-76]. When grown outside an extracellular matrix, the polarity can reverse [77, 78]. When cells from organoids are seeded onto standard cell culture surfaces, such as culture dishes or transwells, the apical side faces the lumen of the well [44, 79-82].

Several studies used organoids to test the general function of particular PRRs without addressing specific differences between apical and basal stimulation. These studies included the stimulus in the medium of the organoids, which, under standard conditions, stimulates the basal side of the cell. Using this technique, upregulation of NF-κB downstream target genes was identified after basal stimulation with ligands of TLR4 in the stomach, TLR2 and 3 in the small intestine and TLR2, 3, 4 and 5 in the colon of mice [44, 45, 83] and to ligands of TLR2 and 5 in the stomach and TLR5 in the small intestine and colon of human [44]. Furthermore, basal stimulation of murine colon organoids with TLR4 agonists induced cellular differentiation, especially towards the secretory lineage [64], while NOD2 agonists induced increased survival of stem cells [67, 84], and taurine stimulated NLRP6-dependent upregulation of the inflammasome downstream target gene IL-18 [85]. In all these studies, apical stimulation was not tested.

Only few studies have addressed the side-specific functions of PRRs. It is noteworthy that contrary to the studies using polarized cancer cell lines, so far, none of the studies using organoids has identified a side-specific activation of a typical NF-κB-dependent proinflammatory gene. Transwell monolayers derived from human colon organoids express similar levels of the NF-κB target gene IL-6 when stimulated from the apical or basal side with ligands of TLR1/2, 3, 4, 5, 7/8 and 9 [82]. Murine gastric organoids also responded to an apical stimulation with the TLR4 ligand LPS in several assays, including in transwells and microinjection of LPS into the lumen of organoids [44]. Murine small intestinal epithelial cells did not express the NF-κB target gene icam1 in response to a range of ligands, irrespective of whether they were added to intact organoids, thus stimulating the basal side, or added to single, dissociated cells, thus stimulating all sides [55].

However, looking beyond the NF-κB response, a recent study identified a side-specific function of TLR3 and its importance in viral infection [82]. Results of experiments with human colon organoid-derived monolayers showed that the expression of the critical virus-defence genes type I and type III interferon was upregulated after basal stimulation with TLR3 agonist, but not after stimulation with other TLR agonists. When infected with a virus, similarly, the interferon response of the organoids was much stronger when infected from the basal side compared to infection from the apical side. This was visible in organoid-derived monolayers as well as in organoids microinjected with the virus. The study further identified the clathrin-sorting adapter AP-1B as the molecule responsible for polarized expression of TLR3. Correspondingly, mice deficient in Ap-1b showed exacerbated immune responses after viral infection [82]. This confirms the polarized function of PRRs and highlights the importance of this level of regulation in further modulating pathogen recognition and defence.

So far, there is no “one size fits all” answer to the polarity question and resolving the side-specific nature of PRR signalling remains a technically challenging task for the future.

**Tolerance as a response to colonization and as a default developmental program**

The mechanisms contributing to the organization of epithelial innate immunity are still unclear. The main concept in this regard has been the induction of tolerance after colonization of the sterile gut during birth, the so-called window of opportunity (reviewed in [86-88]). This concept postulates a priming period of the innate and adaptive immune system after birth, which sets the stage for immune homeostasis and subsequent host-microbial interactions.

The neonatal immune system and epithelial innate immunity are uniquely equipped to master this transition from sterility to co-existence with the microbiota. At birth, the human neonatal intestine is fully mature with intestinal villi and crypts containing Paneth cells. The murine neonatal intestinal epithelial layer is more immature and undergoes a dramatic change measurable on the transcriptome level comparing different stages of development [89] and visible in the tissue architecture and cell differentiation: The crypt-villus axis is not formed yet and cell proliferation is lower, with no cell migration or exfoliation. It does not contain mature Paneth cells; however,
enterocytes produce the cathelicidin-like antimicrobial peptide (CRAMP) [90]. Paneth cells appear when crypts form 2 weeks after birth [91]. At the time of weaning, the epithelium is fully formed with crypts and villi; enterocytes, goblet cells and enteroendocrine cells; Paneth cells have taken over the antimicrobial peptide production; and goblet cells have increased the production of mucins to form the mucus layer (reviewed in [92]).

This age-dependent transition of the epithelium goes hand in hand with a gradual decrease of TLR5 expression in the small intestinal epithelium. At the same time, expression of TLR3 increases during the neonatal period. Other PRRs, such as TLR2, 4 and 9, are expressed at similar levels in neonatal and adult mice [45, 93].

The mechanisms leading to regulation of PRR expression and function after birth remain unclear. As mentioned above, several studies have proposed a contribution of the environment, in particular microbial colonization, to the regulation of PRR expression after birth [40, 41] (reviewed in [14]). However, germ-free vs specific pathogen-free mice did not show differences in TLR expression in either small intestine or colon, indicating that neither upregulation of TLR3 nor downregulation of TLR5 in this early period depends on the microbiota [45, 93]. Also, in organoids, expression of many, but not all, PRR signalling components was already defined in organoids from tissues that had never been in contact with microbial products [44]. This indicates that a large part of the organization of the innate immune signalling pathways is defined independently of contact with the microbiota and appears to be determined by default developmental processes, such as the ones outlined above, which shape the general tissue identity along the GI tract. This does not exclude a further fine-tuning of PRR expression by environmental factors during adulthood.

The importance of the timely regulation of epithelial-microbial interactions becomes clear when the immature epithelium is prematurely confronted with microbial colonization: pre-term infants are prone to the development of necrotizing enterocolitis (NEC), characterized by intestinal necrosis, systemic sepsis and multiple organ failure. Although the pathogenesis is considered multifactorial, several studies have indicated that it develops in response to an imbalance between proinflammatory signalling and repair mechanisms in the premature gut (reviewed in [94]) and a contribution of PRRs has been suggested [41, 65, 95–97]. Several studies have used human fetal organoids [24, 98, 99], normal murine organoids exposed to bacteria and hypoxia to model NEC [100], or organoids from a murine NEC model as well as from NEC patients [101]. Future studies will use these established and new organoid models to further define the contribution of the epithelium in NEC.

Modelling epithelium-dependent aspects of IBD with organoids

Loss of intestinal epithelial barrier integrity is a defining feature of IBD, i.e. CD and UC, and seems to be caused by a multifactorial interplay of genetic predisposition, environmental factors, changes of gut microbiota and alterations of the local and systemic immune response (reviewed in [27]). Current therapeutics predominately target the (aberrant) immune responses in IBD, which are associated with high rates of non-responders and side effects (reviewed in [27]). An improved understanding of the epithelium-specific contribution to the pathophysiology of IBD is required to identify novel therapeutic targets that may also be able to directly impact intestinal epithelial barrier restoration and thus mucosal healing.

To gain insights into the epithelial pathology, several groups have established living biobanks comprised of organoids generated from individual patients with UC or CD. Although this approach is obvious, so far, only few studies have reported results from organoids derived from this group of patients [102–104]. This may be explained by the observation that organoids from patients with IBD are more difficult to generate. Our own experience is that organoids generated from CD patients grow more slowly during the first passages and some samples were lost—which was associated with a higher rate of bacterial contamination in the cultures [105].

Characterization of organoids derived from patients with IBD revealed a phenotype with decreased size and budding capacity, increased rate of cell death, luminal debris and partially inverted polarization of epithelial cells [106]. Global comparison of organoids from UC or CD patients and healthy controls showed that transcriptional and methylation differences seen in the intestinal epithelium were maintained in vitro [102, 103, 107]. Also, an earlier study from organoids generated from patients with CD suggested permanent alterations of intestinal stem cells in IBD. This was based on the observation that organoids generated from active CD lesions maintained high expression levels of intestinal epithelial stem cell markers [108]. This was to some extent confirmed in a recent study in which colon organoids derived from paediatric IBD patients showed a prolonged-expression pattern of antigen-presenting genes [109].

Focusing on changes of intestinal epithelial barrier function, including loss of tight junctions and desmosomes, which are usually found in IBD [110], it was shown that organoids from CD patients maintain this phenotype of junctional alterations under culture conditions [105]. This was especially the case when organoids were generated from sites of severe inflammation [105]. A decrease in junctional proteins could also be induced in organoids from healthy donors by application of the proinflammatory cytokine TNF-α and/or IFN-γ [106, 111]. However, the fixed pattern of changes of junctional proteins in organoids from patients with IBD was observed only on the protein level but not on the mRNA level [105].
These observations suggest that a number of changes are fixed in organoids from IBD patients. The observation that some, but not all, permanent changes are only visible on protein, but not RNA-level, suggests that there are permanent alterations in post-transcriptional modifications or protein degradation in organoids from IBD patients. This however remains to be investigated in detail.

What exactly induces the permanent alterations in expression patterns of the inflamed epithelium is unclear. An effect of the microbiota was long suspected; however, a recent study demonstrated that the effect of the microbiota on the epithelium is lost over time [112]. The study compared several mouse facilities, because the different microbiome compositions of mouse colonies have been identified as confounding factors in vivo studies. To clarify this effect, the study compared epithelial isolates and organoid cultures from germ-free mice and two separate specific pathogen-free mouse colonies with different microbiota. While the freshly isolated epithelium showed an imprint of the microbiota exposure on RNA and protein level, this effect was lost after several weeks of culture of small intestinal organoids [112]. Also, a global comparison of gene expression of organoids generated from inflamed or non-inflamed regions of the same IBD patients demonstrated that the IBD organoids of the inflamed regions lost the inflammatory gene expression already after a few weeks in culture. The transcriptomes of the organoids were then clustered per patient; thus, the permanent differences between IBD and healthy controls remained. The inflammatory phenotype in the IBD organoids could then be re-induced by addition of a cytokine cocktail [107]. Overall, this suggests that permanent changes observed in organoids from IBD patients are independent of contact with microbiota or cytokines.

Thus, although it is reasonable to speculate that some of the permanent changes in intestinal epithelial cells from patients with IBD may be caused by changes in innate immune signaling, the evidence for this is currently scarce. In addition, not all of the epithelial changes observed in the inflamed regions in the gut are permanently conserved in the purified epithelium, indicating important contributions from the local environment. In the future, new studies with more complex organoid models, also incorporating immune cells, inflammatory cytokine stimulation and co-cultures with microorganisms, should help to address this hypothesis in more detail.

Conclusions, future perspectives and outlook

In summary, while many diseases within the GI tract remain incompletely understood, increasing evidence points to a critical role of the gastrointestinal epithelium in the pathogenesis of many of them—although its specific role remains unclear. The limited knowledge of innate immune function of the gastrointestinal epithelium has been attributed to a lack of appropriate experimental models. With the implementation of the organoid technology, a major step has been taken to overcome this problem. Organoids generated from each region of the gastrointestinal tract will add significantly to the existing knowledge. As revealed by previous studies, the

![Fig. 4](image-url) The use of organoids for the study of epithelial innate immunity. Clockwise: Organoids have been generated from different segments of the gastrointestinal tract and have been shown to retain tissue identity. Organoids from patients can be stored in biobanks to enable research on specific pathologies such as IBD. Using culture modifications, the cells in organoids can also be directed towards specific cell identities, such as secretory cells, enterocytes and M cells. Because organoids retain the polarity of the cells, they also allow testing of apical and basal stimulation. For this, cells can be seeded in transwell systems. Organoids are a reductionist model, which is not in contact with microbial compounds under standard conditions but can be used to study infection with bacteria and viruses and to study interaction with immune cells in co-culture experiments. In the future, more complex models will also allow combinations of several organoid types in microfluidic lab-on-a-chip devices.
The generation of organoids provides the crucial advantage of being able to observe responses of primary gastrointestinal cells as opposed to transformed cell lines, which are mostly generated from gastrointestinal malignant tumours. One of the most fascinating and interesting features of organoids is that they maintain specific characteristics of the segment of the gastrointestinal tract they were generated from as part of their cell identity during adulthood. According to current experimental studies, the regional identity is fixed in the gastrointestinal stem cells. In this context, a remaining issue will be to determine exactly how and when during development this intrinsic programming occurs.

When looking at specific concepts of how the epithelium may respond to or interact with the environment, organoid technology has now enabled illumination of the differential and segment-specific expression and function of PRRs within the gastrointestinal epithelium. The overall functional consequences for the complex regulatory systems within the whole gastrointestinal tract are still unclear and will have to be addressed in the future. For this, it will also be important to co-culture organoids with immune cells (reviewed in [113]), cells of the enteric nervous system [114] and luminal factors such as bacterial co-cultures (reviewed in [115]) (Fig. 4).

A further important aspect is that organoids generated from patient tissue affected by GI diseases such as IBD maintain some of the characteristics seen in the corresponding tissue specimens they were derived from. This offers the unique possibility to further unravel the epithelium- and disease-specific contribution to the pathogenesis of GI diseases—not only those involving inflammation-induced changes but also those involving changes in malignant diseases. Both of them may turn out to involve a specific contribution of epithelium-derived innate immunity. For this, the systematic establishment of “living biobanks” would be an important step. As a vision for the future, such living biobanks could be attached to already existing biobanks, which currently provide solely “dead” biomaterials. This would represent another important step not only for research but also in facilitating individualized diagnostics and therapy for patients.

Box 1. Innate immunity and epithelial cells in the GI tract

In homeostasis, it is likely that due to the mucus layer, which poses a diffusion barrier, only low concentrations of MAMPs reach the epithelial layer [59]. At these low, MAMPs seem not to elicit a proinflammatory response but still impact the epithelial cells in several ways. Mice deficient for either TLR2, 4, 5, and 9, the TLR-pathway mediator MyD88 or the NLRP6 inflammasome are highly susceptible to experimentally induced colitis [116–119]. Mice devoid of TLR2 or MyD88 show disrupted tight junctions, more apoptotic epithelial cells and higher barrier permeability [116]. NLRP6 deficiency leads to reduced mucin secretion [61]. Mice devoid of NOD2 show higher epithelial cell death rates after treatment with doxorubicin, indicating that they are more susceptible [67]. In line with this observation, patients with mutations in the NOD2 gene are predisposed to develop IBDs especially CD [120]. While these studies clearly point to the importance of innate immune signalling for epithelial homeostasis, it is unclear whether the observed impact is due to the innate immune signalling in epithelial cells or in professional immune cells. To disentangle the interplay of professional immune cells and epithelial cells, several studies have used epithelial-specific knockouts, or more recently, epithelial organoids. In mouse models, none of the epithelial-specific deletions of PRRs leads to spontaneous inflammation. However, mice with epithelial-specific knockout of MyD88 are more susceptible to experimental colitis and show severe barrier disruption, impaired goblet and Paneth cell responses [121] and reduced production of mucin and antimicrobial peptides [121, 122]. Small intestinal organoids do not mount an inflammatory response to several purified PRR ligands [55], although this cannot be generalized and depends on the species, location and age of the tissue the organoids are generated from [44, 45].

The absence of a spontaneous inflammatory phenotype in epithelial cell-specific PRR knockout models does support the hypothesis that factors other than a general inflammatory response of the epithelium have an impact on epithelial homeostasis. For example, stimulation with the NOD2 agonist muramyl dipeptide (MDP) increased the number of organoids growing out of isolated stem cells, indicating that the innate immune signalling supported survival of the stem cells [67, 84]. Furthermore, data from mice highlight the importance of the anti-apoptotic effects of NF-κB signalling in response to other stimuli, such as TNF-α [123]. Interestingly, in humans, polymorphisms in innate immune genes including NOD2 and TLR4 are associated with an increased risk to develop IBD [124] and blockade of TNF-α is currently the most efficacious treatment for IBD in some patients (reviewed in [125]).

A picture emerges, in which a low level of innate immune stimulation is important for mucus secretion, barrier integrity and epithelial cell survival. Its impairment may allow translocation of intestinal bacteria from the lumen into the subepithelial tissue, leading to inflammation.

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Declarations

Conflict of interest The authors declare no competing interests.

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