INTRODUCTION

1.1 | In vivo and in vitro models of H. pylori infection have both advantages and limitations

Over the years, one of the major challenges in science has been the lack of suitable model systems for infections affecting humans. Host-pathogen research has mainly relied on in vitro and in vivo models in rodents. Recently, the development of model systems such as organoids and lab-on-a-chip systems have opened up more opportunities to increase our understanding of host-pathogen interactions. Helicobacter pylori is a significant cause of chronic gastric diseases, such as ulcers and gastric cancer—the third leading cause of cancer-related death globally.\(^1\) Increasing resistance to antibiotics and the risk of reinfection have prevented a comprehensive understanding of the host immune response and the development of gastric cancer.\(^{2,3}\) Thus, H. pylori infection studies would benefit from the availability of better model systems that more accurately mimic H. pylori pathogenesis in humans.

In vivo studies on H. pylori pathogenesis have largely relied on the use of mice and gerbils. Ethical and practical constraints have limited the degree to which vaccines and therapies can be tested in primates. Although a challenge model in human volunteers has been established for H. pylori infection,\(^4\) only a limited number of trials have been performed.\(^5\) An interesting outcome of the most recent trial has been that challenge of adults resulted in spontaneous clearance in some volunteers, which has implications for future vaccine development.
trials. Field trials in areas with a high prevalence of *H. pylori* infections may be the best way forward for clinical testing of vaccines and therapeutics (reviewed in Ref. [6]). Although mouse models have permitted detailed immunological studies, and some models of carcinogenesis exist, there remains the difficulty in translating results obtained in animal models to human patients. The latter has been a particular problem for vaccine development where promising candidates from animal studies have been poorly immunogenic or showed no protective effect in clinical trials.\(^5,6\) The difficulty in translating from mouse to human vaccines may be attributed to factors such as the relative genetic uniformity of experimental animals as compared to human populations, regional and population differences in *H. pylori* strains, differences in bacterial virulence in rodent models, immune response mechanisms, and also to differences in the microbiota of mice and humans.\(^7\) While mouse models have a relatively close genetic similarity to humans and are easily amenable to genetic manipulation, they sometimes produce varying results based on differences in their genetic backgrounds.\(^8,9\) There are only a few knockout and transgenic mouse models that develop cancer. Their inability to proceed to the metaplastic stage\(^10\) also limits their application in the study of gastric cancer. Other models such as gerbils (*Meriones unguiculatus*) can develop tumors; however, it occurs rapidly, they are outbred and the suite of reagents available for immunological studies in mice are not available for gerbils.\(^11-13\)

In *in vitro* studies have been chiefly focused on the use of epithelial cells derived from cancerous lesions (e.g., AGS, MKN28) or viral transformation (e.g., GES-1). These studies have been key to our understanding of the cell signaling events in early *H. pylori* infection, including the key role of the *H. pylori* type IV secretion system (TIVSS) in translocating CagA in pathogenesis (reviewed in Ref. [14]), and the role of other bacterial virulence factors using deletion mutants (reviewed in Ref. [15]) including the key discovery that the TIVSS actually binds to β1-integrin which is located only on the basolateral side in *in vivo*.\(^16\) Continuous cell lines are, however, either transformed in *in vitro*, or originally derived from cancerous cells, are non-polarized, and it becomes difficult to properly evaluate the host innate immune response at the early stage of *H. pylori* infection, and the sequence of events that leads to metaplasia.\(^11\) Efforts to use primary cells for these studies have been limited by the short lifespan in vitro, with infected cells not lasting beyond 48 h.\(^17-19\)

More recently, in infection biology, the use of stem cell-derived organoid cultures is being adopted as a suitable model largely due to its ability to closely mimic human physiology. Its potential for genetic manipulation with novel gene-editing tools such as CRISPRcas9 and siRNA at a faster rate and with less labor compared to animal models also makes it a promising model for infection studies.\(^2,20,21\) These methods have already facilitated a better understanding of some human diseases. For instance, organoid cultures have been successfully applied to understand the biology of Zika virus and the emergence of microcephaly during brain development.\(^22,23\) Organoid systems have also permitted the culturing of Noroviruses\(^24\) and have opened up the possibility of precision medicine for patients with cystic fibrosis.\(^25,26\)

For *H. pylori* infection, the use of organoids derived from primary cells for *in vitro* studies offers advantages over conventional *in vivo* and *in vitro* techniques in terms of amenability, accessibility, polarization, and longevity. This potentially eliminates the dependence on neoplastic cells in the investigation of the initiating steps in gastric carcinogenesis and could facilitate an in-depth understanding of the mechanisms of pathogenesis. There have been a number of reviews on gastric organoids, particularly in the study of gastric cancer and *H. pylori* pathogenesis,\(^20,27-29\) and the generation of human gastric organoids.\(^30\) Recent reviews have discussed advances in techniques,\(^21\) advances and discoveries in the area of cancer initiation,\(^28\) however, the use of organoid models for studies on *H. pylori* biology, and innate inflammation have not been comprehensively reviewed since 2017.\(^31\) Thus, this review aims to aggregate and discuss recent advances from studies with gastric organoids in terms of how *H. pylori* interacts with the gastric epithelium, triggers inflammation, and its role in the initiation of gastric cancer. We also propose that the use of more complex organoid models will be beneficial for the study of gastric epithelial repair, the role of gastric hormones, and mechanisms of vaccine-induced protection.

### 1.2 Development of mouse and human gastric organoids

The successful development of organoids has hinged on the identification of the necessary growth factors and conditions, and pivotal lineage tracing studies that led to the identification of multiple adult stem cell populations in the gastric gland (reviewed in Ref. \(^[32]\)). The initial studies on the development of intestinal organoids from human embryonic and induced pluripotent stem cells (PSC) laid the foundation for the identification of the growth factors required for gastric organoids.\(^33,34\) The generation of gastric organoids from PSC involves the use of growth factors which mimic embryonic development and was first described by McCracken et al.\(^35\) Human ES cell lines were first differentiated into definitive endoderm by the addition of activin A. Further treatment with FGF4, Noggin, retinoic acid and WNT3A or its agonist CHIR99021 induces differentiation into posterior foregut spheroids which are then seeded in Matrigel™ to generate gastric organoids which developed a complex three-dimensional organization and differentiated antral cell types (surface mucous cells, antral gland cells, LGR5<sup>+</sup> cells, endocrine cells), but not parietal or chief cells which are characteristic of the fundus/corpus.\(^35\)

Gastric organoids generated from adult stem cells from gastric glands depend on growth factors that mimic tissue regeneration. Studies on gastric homeostasis employing both organoids and animal model studies have also identified the key factors necessary for gastric stem cell proliferation including Wnt3a/β-catenin signaling which is augmented by R-spondin\(^36\) and differentiation (Notch signaling).\(^37\) A recent study by Wölflling et al.\(^38\) showed that epidermal growth factor (EGF) is a major fate determining factor in the gastric gland, and that gradients of EGF and bone morphogenic factor and Noggin signals are responsible for differentiation and foveolar, chief and acid-producing
parietal cells to generate a gland-type or a pit-type phenotype. As a result of these studies, there are now well-established protocols for gastric organoid cultures, and *H. pylori* infection and commercially available media for organoid cultures.

The development of gastric organoids from adult gastric glands began with the discovery by Barker and co-workers that Lgr5	extsuperscript{+} stem cells are responsible for the constant renewal of the gastric epithelium and that these stem cells can differentiate into the different cell types in the gastric crypts. These studies also revealed that Wnt signaling is necessary for the expression of Lgr5 and that its agonist R-spondin further increases this expression. Organoids were generated from the Lgr5	extsuperscript{+} stem cells in mouse stomach; however, while the Lgr5	extsuperscript{+} stem cells were found in the pyloric region, they were absent in the corpus, suggesting that other stem cells may also be responsible for the renewal of the gastric epithelium. In another study, Stange et al. proposed that a subset of chief and parietal cells that express Troy and are located near the base of the corpus glands express Wnt target genes and can act as a “reserve” of stem cells that can renew the corpus epithelium. These stem cells were also shown to act as adult stem cells which could be successfully grown into organoids that differentiated into mucus neck cells and pit cells. Gene expression analysis revealed that Troy	extsuperscript{+} cells were enriched for expression of Lgr5, and CD44, a marker of epithelial stem cell marker, and a Wnt3A target gene. A later study, using murine fundic glands, generated organoids by co-culture with immortalized stomach mesenchymal cells as a source of growth factors that were able to differentiate into all fundic cell types including parietal cells.

The development of murine organoids has been followed by the successful development of human gastric organoids (Figure 1). Initial studies involved the use of the commercial extracellular matrix mixture - Matrigel™. McCracken et al. first developed human antral spheroids and organoids using induced pluripotent stem cells seeded in Matrigel, while Bartfeld et al. recorded the development of the first human corpus spheroidal organoids using gastric glands seeded in Matrigel. *H. pylori* infection in these earlier studies was performed by microinjection into the spheroid lumen because the interior formed the luminal edge of the epithelial cells. However, a later study involving the initial generation of antral and corpus spheroids in Matrigel was further developed such that spheroids could be successfully cultured in 2D monolayers on collagen-coated surface. Although these cultures barely lasted for 24 h, infection did not require microinjection. A later study by the same group described the generation of mucosoid cultures from antral gland using an air–liquid interface (ALI) system. This was made up of collagen-coated transwell system and permitted simple addition of the bacteria to the luminal side of the culture for infection. Mucosoid cultures can be maintained as ALI for up to a month and the transwell format permits investigation of soluble factors and cells added to different compartments of the ALI culture. Frequent removal of mucus and change of media in the basal compartment are required. *H. pylori* can be co-cultured in the apical compartment for several days for experiments.

An interesting feature of the various gastric organoid culture systems is their ability to replicate the characteristic events of a *H. pylori* infection which has permitted a range of detailed studies on the biology of the interaction of the bacteria with the gastric epithelium which are discussed below.

### 1.3 Application of organoids to study epithelial interactions and the initiation of inflammation by *H. pylori*

At the level of initial epithelial interactions, organoid culture models have further increased our understanding of how chemotaxis provides the persistent colonization of the host by *H. pylori*. Notably, it permitted the identification of TlpB (HP0103) as one of the chemoreceptors responsible for the attraction of *H. pylori* to the gastric epithelium. Huang et al. reported that *H. pylori* are attracted to urea secreted by the gastric epithelium and that this is how the pathogen locates the gastric epithelium for colonization. The polarized nature of gastric organoids enabled this discovery. Dulbecco's modified eagle medium (DMEM) was conditioned with epithelial secretions from uninfected human gastric organoids. Free-swimming wildtype (WT) *H. pylori* were attracted to the conditioned medium and not attracted to the unconditioned media. Mutant *H. pylori* lacking just the TlpB gene were not attracted to the conditioned medium, while mutants lacking either TlpA, TlpC, or TlpD were all attracted to the conditioned media, implying that TlpB is responsible for this chemotaxis.

This discovery enabled the researchers to further test the effect of TlpB on the ability of *H. pylori* to colonize persistently in vivo. Mice infected with WT *H. pylori* had higher colonization six weeks postinfection compared to mice infected with the mutant lacking TlpB. Additionally, the role of TlpB in sensing urea during infection was investigated in another study where the TlpB chemoreceptor was found to be responsible for the accumulation of *H. pylori* at damaged epithelial sites. Using murine-derived gastroids as a restitution model, cellular injury was generated on gastric organoids by photodamage and the accumulation of WT and *H. pylori* mutants lacking each of the four chemoreceptors TlpA, TlpB, TlpC, and TlpD at the injury sites was monitored using confocal microscopy. Only TlpB mutants were unable to colonize the injury site whereas wildtype and other *H. pylori* mutants accumulated at the damage sites and also inhibited epithelial restitution. However, unlike the initial study where the TlpB-mediated response was due to the attraction to urea, the chemotactant responsible for accumulation at the damage site remains unknown. The microinjection of urea to the lumen of the organoids prevented the accumulation of *H. pylori* at the damage sites. Thus, implying that the TlpB chemoreceptor may sense different chemicals to attract *H. pylori* to the epithelium under different conditions.

Gastric organoid cultures appear to mimic the in vivo model more accurately than gastric epithelial cell lines for studying interaction of *H. pylori* with the apical-junctional complex. Uotani et al. reported that epithelial morphology and IL-8 expression were similar to what was found in vivo model studies. The integrity of the apical-junctional complex as determined by measurement of transepithelial electrical resistance (TEER) was unaltered after 96-hour infection.
Initial studies with conventional cell lines could only observe TEER for 48 h and had therefore reported a 30%–50% decrease, whereas gastroid monolayers which permitted culturing for up to 96 hours revealed that although there was an initial decrease in TEER after 24 hours of infection, it reverted to normal after 48 hours. It is noteworthy that this study also found that contrary to reports from studies involving gastric cancer cell lines, IL-8 expression was independent of *H. pylori* virulence factors such as CagA. Wildtype *H. pylori* strain TN2GF4 induced increased expression of IL-8 in AGS and MKN28 cells compared to CagPAI deletion mutant, but no difference in IL-8 expression was observed in the gastroid monolayers infected with both strains. Clinical isolates lacking virulence factors such as CagPAI, OipA, VacA, and BabA increased IL-8 expression in gastroid monolayers, but induced no change in MKN28 cells. Additionally, inhibitors to the JNK, MEK, P38, and IKK pathways reduced IL-8 expression in gastric cancer cell lines. In the gastroid monolayers, only the inhibitors to P38 and IKK pathways reduced the expression, whereas inhibitors of the JNK and MEK pathways had no effect on IL-8 expression, implying that different pathways are involved in the production of IL-8 in gastric organoids (and presumably gastric crypts) and may be more complex than in carcinoma cell lines. While the reports of this study imply that *H. pylori* may exert different effects on cancer and non-cancer patients, a limitation of this study is that gastric organoids were derived from a single healthy patient and developed from only the antrum and not the corpus which is mainly affected by gastric cancer. Therefore, further studies with organoids from both stomach regions of cancer and non-cancer patients, and investigations into the roles of different cell types found in the antrum and corpus are required.

The organoid culture model has also helped in understanding the mechanisms of *H. pylori*-induced inflammation in both the innate and adaptive immune systems. Using murine embryo-derived organoids due to their sterility, Kayisoglu et al. reported that the expression pattern of Toll-like receptor 4 (TLR4) was independent of an interaction with gut microbiota, and was instead determined by developmental stage. Both the embryo-derived gastric organoids

**FIGURE 1** Three major types of gastric organoid culture systems that are currently in use. (A) The generation of 3D gastric organoids with *H. pylori* infection achieved by microinjection. By manipulating the growth factors, stem cells were first developed into definitive endoderm, and then into spheroids before eventually being transformed into 3D organoids. (B) 3D spheroids were generated from gastroid glands seeded in Matrigel and either used to generate organoids or cultured in a 2D cell culture system. (C) Mucosoid cultures derived either from 3D organoids or directly from gastroid glands are cultured in a transwell system. *H. pylori* infection is achieved by co-culturing for both (B) and (C).
and gastric organoids derived from adults expressed CXCL2 at a similar level when exposed to lipopolysaccharide (LPS). Morey et al. investigated interferon-gamma (IFNγ) signalling using gastric cancer cell lines, mouse, and 2D mucousoid primary cell models and reported that *H. pylori* blocks IFNγ signalling via the JAK/STAT pathway in a cholesterol-α-glucosyltransferase (cgt)-dependent manner in infected cells, while inducing inflammation in the surrounding uninfected cells of the epithelium. MKN45 gastric cancer cell lines, mouse and cells derived from antral gastric organoids infected with *H. pylori* strain P12 all blocked IFNγ compared to a mutant strain lacking cgt. Confocal microscopy observation of mucousoid cultures infected with *H. pylori* strain P12 in an air–liquid interface for 3 days and treated with IFNγ for 30 minutes revealed that *H. pylori* induced an infection foci which had lower levels of phospho-STAT1 compared to uninfected areas. This implies that *H. pylori* survives by suppressing the IFNγ signalling while inducing inflammatory responses in the surrounding epithelial cells.

The ability of organoid cultures to replicate increased expression of inflammatory cytokines in response to *H. pylori* infection has been shown in both mucousoid cultures and 3D spherical models. Sebrell et al. reported the upregulation of CXCL1, CXCL2, and CXCL8 in human 3D spheroids infected for 3 hours, and that CXCL8 secretion was higher in spheroids infected with *CagA*+ strains compared to *CagA*-deficient strains as previously reported for human gastric epithelial cell lines. Microarray analysis followed by confirmation by qPCR by Boccellato et al. also showed that, in addition to IL-8, other proinflammatory cytokines and chemokines such as CXCL1, CXCL2, and CXCL3, lymphotxin B (LTB), IL-23A and CXCL1, CXCL2, and CXCL3, lymphotoxin B (LTB), IL-23A and CXCL8 in human gastric epithelial cell lines. Microarray analysis followed by confirmation by qPCR by Boccellato et al. also showed that, in addition to IL-8, other proinflammatory cytokines and chemokines such as CXCL1, CXCL2, and CXCL3, lymphotoxin B (LTB), IL-23A and TNF-α were all upregulated, further consistent with earlier studies that reported the over-expression of these genes expressed in *H. pylori*-infected patients.

The ability to study epithelial responses to infection without interference by the host immune response in organoids has enabled the assessment of the role of *H. pylori* infection and NF-KB on the mechanism of sonic hedgehog (Shh) signalling, which is important for the initiation of gastritis. These interactions had been impossible to study in vivo due to possible interference by hematopoietic factors. After observing that *H. pylori* stimulates Shh signalling in the parietal cells of transgenic mouse expressing Shh fused to GFP within 2 days of infection, organoids were generated from both wildtype mice and parietal cell-specific Shh deletion KO mice and infected with *H. pylori* to evaluate the role of Shh signalling. Shh signalling was absent in parietal cell-specific Shh deletion KO mice, but it was observed in organoids derived from control mice. This signalling diminished following one-hour pre-treatment of control organoids with NF-κB inhibitor before infection, implying that Shh signalling in parietal cells is NF-κB-dependent.

In a study that employed the use of confocal microscopy and 3D reconstruction of human and animal gastric glands, *H. pylori* was found to activate and expand the gastric stem cells and gastric progenitor cells when it colonizes the gastric gland. *H. pylori* mutants lacking ChePep which is required for chemotactic movement were incapable of colonization and did not activate the stem cells. The organoid culture model was used to study the ability of the activated stem cells to form glands ex vivo by growing antral glands from uninfected and two-month-infected mice into spherical organoids in Matrigel and measuring their size daily for 6 days. Infected glands were found to be larger in size and grew faster into organoids, suggesting the activation by *H. pylori* infection. However, due to the short time frame, it is unclear whether this effect can persist for longer periods.

Gastric organoids have also helped in the study of STAT3 signaling, known to be relevant for inflammation-associated gastric carcinogenesis. Zhu et al. reported that dopamine and cAMP-regulated phosphoprotein, Mr 32000 (DARPP-32) induces STAT3 signaling in AGS and MKN45 cells by regulating IGF1R-SRC signaling. The expression of STAT3, STAT3 target genes, and P-IGF1R was studied in mouse organoids derived from TFF-1 KO mouse, DARPP-32 KO mouse, and TFF-1/DARPP-32 double KO mouse. TFF1KO mouse showed higher expression of STAT3 target genes, STAT3 and P-IGF1R compared to DARPP-32 KO mouse and TFF-1/DARPP-32 double KO mouse. Another study reported that reduction in atrophy, metaplasia, and epithelial cell proliferation in the gastric mucosa of mouse infected with *Helicobacter felis* for 18 months, correlated with loss of STAT3 signaling. Organoids were generated from wildtype (WT) mouse and STAT3 knockout mice and stimulated with recombinant IL-6 (rIL-6) and recombinant IL-11 (rIL-11) followed by treatment with JAK inhibitor. Gene expression analysis showed that treatment with rIL-6 and rIL-11 increased the expression of intestinal metaplasia-associated genes-Intelectin1 (Itln1), and Lysosomal H⁺ transporting ATPase subunit (ATP6V0d2) but the expression was reduced upon the addition of JAK inhibitor, suggesting that these cytokines initiate intestinal metaplasia in a STAT3-dependent manner. Although these did not involve infection with *H. pylori*, dissecting these questions in the context of *H. pylori*-induced STAT3 activation may be a viable option.

### 1.4 Application of organoid cultures to understand gastric cancer initiation and progression

The emergence of organoid cultures has permitted new avenues for enquiries into the initiation of *H. pylori*-associated gastric cancer, which has previously been limited by dedifferentiation in cell culture, and limitations of animal models. While organoids may not be able to recapitulate the decades long chronic inflammatory state, a number of the mechanisms involved in cancer progression have been investigated. These studies have been reviewed in detail elsewhere, and some major findings that highlight the advantages of organoids are summarized here.

Organoids present the potential to investigate the role of IFNγ in the progression from gastritis to atrophic gastritis and metaplasia, and the development of spasmyotic polypeptide/Trefoil Factor (TFF) 2-expressing metaplasia (SPEM) in early-stage metaplasia. The latter in a cytotoxic T-cell co-culture model that showed that overall, the expression of PD-L1 on SPEM cells was found to be dependent on Shh
signaling, and the interaction of PD-L1 with PD-1 on CTLs enables the survival of SPEM cells in the presence of H. pylori infection.73

DNA damage is a major mechanism in carcinogenesis, and several studies have used organoids to address questions on H. pylori-induced damage. Bauer et al. reported that the LPS precursor β-ADP-heptose was sufficient to cause DNA damage in human-derived organoids.74 Expression of Neli-like DNA glycosylase 2 (Neil 2) which is involved in the repair of DNA damage by removing oxidase species was suppressed in H. pylori-infected mucousoid cultures.75 Consistent with this observation, mucosoids generated from Neil2 knockout mice expressed higher levels of inflammatory cytokines.76 The approach taken in this study provides a mechanistic link between H. pylori infection, inflammation, and initiation of DNA damage.

The combination of organoid culture and gene-editing techniques provides exciting potential for investigating mechanisms in chronic inflammation and cancer initiation. An example of this is the report by Nanki et al.77 on establishment of a bank of cancer or chronic inflammation and cancer initiation. An example of this is the infection are yet to be conducted with organoid systems. However, the potential for investigations on the role of gastric hormones in inflammation and repair are clear from a number of reports. Ohki et al.79 used mouse intestinal organoids, and reported a high similarity in gene expression between the enteroendocrine cells and the hormones produced by enteric organoids, when compared with that of mouse native tissues. Ghrelin plays a role in growth hormone release and food intake, and has immunomodulatory properties. Ghrelin expression is reduced in H. pylori-infected patients.80,81 An earlier study reported that mouse intestinal and colonic organoids expressed serotonin, its receptors and the serotonin uptake inhibitor in a similar manner to intact tissue.82 Gastrin secretion is important for regulation of gastric acid production, and a recent study employing mouse and organoid models showed that gastrin triggers proliferation of enterochromaffin-like cell (ECL) progenitors.83 Given the roles played by gastric hormones in H. pylori-associated disease manifestation and the potential role of other hormones such as leptin in vaccine-induced immunity to H. pylori,84,85 organoid models present great potential to investigate the contribution of hormones to gastric homeostasis, and in the inflammatory and regulatory immune responses in H. pylori pathogenesis.

1.6 | Potential applications of organoids to investigate mechanisms of vaccine-induced protection

The ability to co-culture organoids with immune cells has enabled more in-depth information to be gathered about the roles of both the innate and adaptive immune cells in H. pylori infection. This has been achieved by culturing gastric organoids with immune cells derived from the same host. Co-culture also offers opportunities to investigate the immune mechanisms underlying vaccine-induced protection, an area which has proven difficult to dissect using animal models (Figure 2). To date, there have been no specific organoid model studies on vaccination; however, several reports have provided insights into innate interactions with the epithelium that provide support for the usefulness of organoids to investigate complex interactions that involve both innate and adaptive immune responses.

Sebrell et al. 56 investigated the chemokine-dependent recruitment of dendritic cells to the basolateral side of the gastric epithelium during H. pylori infection. In this study, human gastric organoids were co-cultured with monocyte-derived dendritic cells and then infected with H. pylori by microinjection. A range of chemokines, including CXCL5, CXCL8, CCL20, and CXCL17, were expressed in response to infection. The recruited dendritic cells eventually ingested H. pylori by phagocytosis when co-cultured with human gastric organoids, showing the pathogen sampling activities of DC that have previously been reported in vitro.86 Further, in the study by Suarez et al. discussed above,87 the effect of H. pylori infection on cytokine production by innate immune cells was assessed by co-culturing epithelial gastric organoids from both infected WT and Nod1 KO C57BL/6 mice with macrophages from the same mice. Increased
cytokine production was observed in Nod1-deficient samples, especially when both the macrophages and gastroids lacked Nod1, implying that a functional Nod1 suppresses cytokine production.\(^87\)

The previously discussed study on the interaction between PD-L1 on epithelial cells with the PD-1 on CTLs, a CD8\(^+\) T-cell subset\(^73\) showed that both lymphocyte proliferation and epithelial cell marker interactions can be investigated in organoid models. Similar approaches could be used to investigate interactions with CD4\(^+\) T cells which are known to be the main mediators of vaccine-induced protection (reviewed in Ref. \([88]\)).

Innate and adaptive immunity also result in changes in expression of mucins and antimicrobial peptides in animal models of *H. pylori* vaccination. A study by Boccellato et al.\(^{41}\) in a two-dimensional mucosoid culture model also provided initial evidence for the bactericidal activity of the mucus and its ability to serve as a physical barrier against *H. pylori* attachment. The mucosoid organoid model co-culture offers the potential to investigate the effects of the CD4\(^+\) T-cell subsets (Th1/Th2/Th17/Treg\(^{56}\)), and innate lymphoid cells (ILC2 and ILC3) and their cytokines on the antimicrobial activity of the epithelium, and at least in principle to generate “vaccinated” organoids that recapitulate reductions in bacterial numbers seen in vivo. Such a model will be a powerful way to systematically test the roles of specific cytokines and hormones in protective responses, and to dissect the effector mechanisms.

### 1.7 Progress, challenges, and limitations

The above studies have shown how gastric organoids have been effectively used to extend and complement conventional *in vitro* and *in vivo* techniques to understand *H. pylori*-induced disease and the resulting host immune response. However, while they have revealed how certain unique features of organoids make them the ideal and perhaps the only viable medium for these studies, it also reveals that there are some limitations. Careful comparison to *in vivo* models and patient samples will still be required. Development of more complex organoid systems with co-cultured cells or factors would address some of these issues. As an emerging technique, one important question on using gastric organoids is the limits to which it is possible to mirror *in vivo* interactions.

The polarized nature of gastric epithelial organoids also provides an advantage over the non-polarized systems and many signaling
events are dependent on cell architecture. A case in point is the mislocalization of scribble, a basolateral polarization marker, by the Cag-ASPP2 binding which was not observed in AGS cells, but was observed in human antral organoids. Additionally, organoids can contain all cell types of the stomach lineage thereby enabling the study of the specific roles of each cell type in *H. pylori* pathogenesis. This has helped in the study of the role of acid-secreting parietal cells in gastric cancer development. The gastric corpus is the main region of the stomach affected by *H. pylori* infection, which is characterized by the loss of parietal cells. The absence of parietal cells in the corpus of human gastric organoids had initially limited their study to the use of murine gastric organoids. However, following a modification of the earlier protocol the detection of parietal cells in human corpus organoids and its use for the study of *H. pylori*-induced gastric cancer was reported and may permit earlier studies to be revisited.

Another unique feature of gastric organoid studies is the ability to determine the specific phenotype of generated organoids based on the selection of growth factors and source of stem cells or glands. Thus, enabling the generation of organoids specifically from the antrum/pylorus and corpus/fundus regions for the study of peptic ulcer and gastric cancer, respectively. This enables the understanding of the differential effects of *H. pylori* infection in these regions of the stomach. Manipulation of growth factors has also allowed the determination of the cell lineages to be generated. Thus, organoids with gland/basal-type or pit/foveolar-type phenotypes can be generated and a comparison between the two phenotypes can be made. Studies have reported higher expression of inflammatory cytokines in the gland-type than in the pit-type organoids, suggesting that the effect of *H. pylori* infection in the gland base is higher than in the pits. The ability to co-culture organoids with stromal cells has also been helpful in this regard. Culturing murine fundic organoids with immortalized stomach mesenchymal cells has been shown to help in the generation and maintenance of organoids expressing markers of epithelial lineages. A 2019 study revealed that the secretion of R-spondin 3 by the surrounding myofibroblasts of the gland base initiates the basal LGR5-positive cells to secrete antimicrobial factors against *H. pylori*. This finding raises interesting questions about whether it is the inflammatory cytokine milieu induced by *H. pylori* infection that drives myofibroblasts to over-express Rsps-3, and further why the antimicrobial effect is not successful in clearing *H. pylori* in the long-term in vivo?

An important advance in increased complexity of organoids was recently reported where the three primary germ layers derived from pluripotent stem cells were used in an organ assembly approach. Enteric neuroglial, mesenchymal, and epithelial precursors were used to generate human fundic and antral gastric tissue with differentiated glands, surrounding smooth muscle, and functional enteric neurons. After development in vitro, organoids expressed similar levels of gastric hormones and mucous as intact tissue. Organoids transplanted into mouse kidney capsule and grown for up to 12 weeks further developed into epithelia that were morphologically similar to Brunner’s glands with expression of mucous and gastric hormones.

While the requirement of transplantation for maturation makes this model highly complex, this represents an exciting advance in tissue engineering and for application for the study of *H. pylori* biology and immune cell interactions. Organoids replicate the genetic phenotype of the organisms they are generated from. Thus, organoids generated from knockout mice are usually devoid of the genes that were knocked out. Organoids from gastric cancer cell lines also possess the features of the cancerous tissues. The successful generation of gastric organoids from pediatric biopsies has also been reported, although further studies are required to compare the physiological characteristics with adult organoids. In the long term, this may be a means to identify aspects that differentiate childhood from adult acquisition of *H. pylori*.

The relative longevity of organoid cultures make gene editing and other genetic manipulation approaches viable and provide a way to study the role of specific genes in *H. pylori* pathogenesis; this approach is yet to be explored with regard to *H. pylori* pathogenesis. Schlaermann et al. provided proof of principle that gastric spheroids could be transfected using a lentiviral GFP construct and the gene-editing approaches employed in cancer organoid studies both offer approaches to investigate the role of specific receptors/effectors in infection and pathogenesis. We expect that the use of gene-edited organoids will expand rapidly in the next few years.

When choosing experimental techniques, labor cost and financial commitments are important factors to consider. The growth factors for organoid cultures are relatively expensive, compared to conventional cell lines. Additionally, maintenance of organoid cultures is usually labor intensive. This is a major challenge for the long-term culture of organoids. Despite these challenges, gastric organoids are showing good potential to studying the changes that occurs in the gastric epithelium over time, such as the progression from inflammation to gastric cancer, or to investigate vaccination mechanisms. The variation in terms of longevity, labor intensity, and mode of *H. pylori* infection among the different types of organoid cultures also determines the choice of organoid culture model. Because organoids are relatively complex, in-depth investigations of protein–protein interactions for example are likely to be better suited to conventional cell line studies. Thus, the choice of techniques should be made in terms of suitability for the study and the specific questions to be addressed.

# Conclusions and Future Perspectives

The use of gastric organoids has advanced our knowledge in *H. pylori* infection studies due to their advantages over traditional in vivo and in vitro techniques in terms of polarization, longevity, amenability, and accessibility. As a result, gastric organoids have already led to novel discoveries on chemotaxis, intracellular effects of *H. pylori* virulence factors, interactions with the apical-junctional complex; innate immune activation and the initiation of inflammation by *H. pylori*, that were hitherto impossible in other models. In addition, direct
effects of *H. pylori* on gland stem cells and induction of DNA damage have greatly improved our understanding of the mechanisms of cancer initiation (reviewed in [28]) resulting in the identification of potential therapeutic targets.

The technique is not a one-size-fits-all approach, however, as certain investigations remain better suited to conventional methods. The development of long-lasting organoid cultures with reduced cost is highly desirable for the long-term monitoring of the changes that occurs after infection by the pathogen such as gastritis and the initiation of metaplasia. This would allow for the long-term monitoring of carcinogenesis, thus reducing the dependence on mice for *in vivo* carcinogenesis experimentation. Organoid culture models also have the ethical advantage of a reduction in the numbers of experimental animals that are required for host-pathogen studies because glands from a small number of animals can be cultured indeﬁnitely.

Despite numerous studies that have been carried out with gastric organoids in the context of *H. pylori* infection, the full potential of this technique is yet to be explored. As highlighted in this review, there have been impressive advances in our understanding of gastric homoeostasis. This, coupled with the capability of generating organoids from transgenic and reporter mice, and the amenability of organoids to genetic manipulation opens the door to investigate inflammatory pathways. Further, the roles of gut hormones such as ghrelin, serotonin, and leptin which are known to have both functional and cytokine activity can be addressed. The investigation of vaccination using organoids is more speculative at present, but the foundations have been laid for more complex and detailed studies on acute and chronic inflammation and gastric protective mechanisms.

**ACKNOWLEDGMENTS**

This work was supported by RMIT funds to AKW. SI is supported by an RMIT scholarship. Figures were generated using Biorender.com. Open access publishing facilitated by RMIT University, as part of the Wiley - RMIT University agreement via the Council of Australian University Librarians. [Correction added on 25 May 2022, after first online publication: CAUL funding statement has been added.]

**CONFLICT OF INTEREST**

The authors declare that they have no competing interests.

**ORCID**

Sulaimon Idowu https://orcid.org/0000-0002-8474-698X
Paul P. Bertrand https://orcid.org/0000-0003-1122-2306
Anna K. Walduck https://orcid.org/0000-0002-9624-4370

**REFERENCES**

1. Torre LA, Bray F, Siegel RL, Ferlay J, Lortet-Tieulent J, Jemal A. Global Cancer Statistics, 2012. *CA Cancer J Clin*. 2015;65(2):87-108. doi:10.3322/caac.21262
2. Kim J, Koo BK, Knoblich JA. Human organoids: model systems for human biology and medicine. *Nat Rev Mol Cell Biol*. 2020;21:571-584. doi:10.1038/s41580-020-0259-3
3. Dutta D, Clevers H. Organoid culture systems to study host-pathogen interactions. *Curr Opin Immunol*. 2017;48:15-22. doi:10.1016/j.coi.2017.07.012
4. Graham DY, Opekun AR, Osato MS, et al. Challenge model for Helicobacter pylori infection in human volunteers. *Gut*. 2004;53:1235-1243. doi:10.1136/gut.2003.037499
5. Sutton P, Boag JM. Status of vaccine research and development for Helicobacter pylori. *Vaccine*. 2019;37(50):7295-7299. doi:10.1016/j.vaccine.2018.01.001
6. Walduck A, Raghavan S. Immunity and vaccine development against Helicobacter pylori. In: Kamiya S, Backert S, eds. *Helicobacter pylori in Human Diseases*. Advances in Experimental Medicine and Biology. Vol 1149, Springer, Cham; 2019:1-19. doi:10.1007/5584_2019_370
7. Kostic AD, Howitt MR, Garrett WS. Exploring host-microbiota interactions in animal models and humans. *Genes Dev*. 2013;27(7):701-718. doi:10.1101/gad.212522.112
8. Kaparakis M, Laurie KL, Wijburg O, et al. CD4+ CD25+ regulatory T cells modulate the T-cell and antibody responses in Helicobacter-Infected BALB/c mice. * Infect Immun*. 2006;74(6):3519-3529. doi:10.1128/IAI.01314-05
9. Nedrud JG, Czinn SJ, Ding H, et al. Lack of genetic influence on the innate inflammatory response to Helicobacter infection of the gastric mucosa. *Front Immunol*. 2012;3:181. doi:10.3389/fimmu.2012.00181
10. Toller IM, Hitzler I, Sayi A, Mueller A. Prostaglandin E2 prevents helicobacter-induced gastric preneoplasia and facilitates persistent infection in a mouse model. *Gastroenterology*. 2010;138(4):1455-1467.e4. doi:10.1053/j.gastro.2009.12.006
11. Schlaermann P, Toelle B, Berger H, et al. A novel human gastric primary cell culture system for modelling Helicobacter pylori infection in vitro. *Gut*. 2016;65(2):202-213. doi:10.1136/gutjnl-2014-307949
12. Rourke JLO, Lee A. Animal models of Helicobacter pylori infection and disease. *Microbes Infect*. 2003;5(8):741-748. doi:10.1016/S1286-4579(03)00123-0
13. Noto JM, Gaddy JA, Lee JY, et al. Iron deficiency accelerates Helicobacter pylori-induced carcinogenesis in rodents and humans. *J Clin Invest*. 2013;123:479. doi:10.1172/JCI64373
14. Backert S, Haas R, Gerhard M, Naumann M. The helicobacter pylori type IV secretion system encoded by the cag pathogenicity Island: Architecture, function, and signaling. In: Backert S, Backert E, Grohmann E, eds. Type IV Secretion in Gram-Negative and Gram-Positive Bacteria. *Current Topics in Microbiology and Immunology*. Current Topics in Microbiology and Immunology. Vol 413, Springer, Cham; 2017:187-220. doi:10.1007/978-3-319-75241-9_8
15. Sgouras D, Tegtmeier N, Wessler S. Activity and functional importance of Helicobacter pylori virulence factors. *Adv Exp Med Biol*. 2019;1149:35-56. doi:10.1007/5584_2019_358
16. Kwok T, Zabler D, Urman S, et al. Helicobacter exploits integrin for infection in a mouse model. *Helicobacter*. 2019;24(1):13-19. doi:10.1111/hel.12645
17. Boxbberger HJ, Sessler MJ, Grausam MC, Becker HD. MYD88. Isolation and culturing of highly polarized primary epithelial cells from normal human stomach (antrum) as spheroid-like vesi- cles. *Methods Cell Sci*. 1997;19(3):169-178. doi:10.1205/AM002009-1713654
18. Richter-Dahlfors A, Heuschko U, Meloche RM, Finlay BB, Buchan AMJ. Helicobacter pylori-infected human antral primary cell cultures: Effect on gastrin cell function. *Am J Physiol - Gastrointest Liver Physiol*. 1998;275(3 38-3):393-401. doi:10.1152/ajpgi.1998.275.3.g393
19. Ootani A, Toda S, Fujimoto K, Sugihara H. An air-liquid interface promotes the differentiation of gastric surface mucous cells (GSM06) in culture. *Biochem Biophys Res Commun*. 2000;271(3):741-746. doi:10.1006/BBRC.2000.2673
20. Lau HCH, Kranenburg O, Xiao H, Yu J. Organoid models of gastrointestinal cancers in basic and translational research. *Nat Rev Gastroenterol Hepatol.* 2020;17(4):203-222. doi:10.1038/s41575-019-0255-2

21. Aguilar C, Alves da Silva M, Saraiwa M, et al. Organoids as host models for infection biology – a review of methods. *Exp Mol Med.* 2021;53:1471-1482. doi:10.1038/s12276-021-00629-4

22. Dang J, Tiwari SK, Lichinchi G, et al. Zika virus depletes neural progenitors in human cerebral organoids through activation of the innate immune receptor TLR3. *Cell Stem Cell.* 2016;19(2):258-265. doi:10.1016/j.stem.2016.04.014

23. Garcez PP, Loiola EC, Madeiro da Costa R, et al. Zika virus impair growth in human neurospheres and brain organoids. *Science.* 2016;352(6287):816-818. doi:10.1126/science.aaf6116

24. McCracken KW, Howell JC, Spence JR, Wells JM. Generating of human antral and fundic gastric organoids from pluripotent stem cells. *Nat Protoc.* 2019;14(1):28-50. doi:10.1038/s41596-018-0080-z

25. Dekkers JF, Berkers G, Kruiswinkel E, et al. Characterizing responses to CFTR-modulating drugs using rectal organoids derived from subjects with cystic fibrosis. *Sci Transl Med.* 2016;8(344):344RA84. doi:10.1126/scitranslmed.aad2878

26. Berkers G, van Mourik P, Vonk AM, et al. Rectal organoids enable personalized treatment of cystic fibrosis. *Cell Rep.* 2019;26(7):1701-1708.e3. doi:10.1016/j.celrep.2019.01.048

27. Alzeeb G, Metges J-P, Corcos L, Le Jossic-Corcos C. Three-dimensional culture systems in gastric cancer research. *Cancers (Basel).* 2020;12(10):1-20. doi:10.3390/cancers12102800

28. Traulsen J, Zagan C, Daddi AA, Boccellato F. Molecular modelling of the gastric barrier response, from infection to carcinogenesis. *Best Pract Res Clin Gastroenterol.* 2021;50-51:101737. doi:10.1016/j.jbpg.2021.101737

29. Idowu S, Bertrand PP, Walduck AK. Homeostasis and cancer initiation: Organoids as models to study the initiation of gastric cancer. *Int J Mol Sci.* 2022;23(5):2790. doi:10.3390/ijms23052790

30. Eicher AK, Berns HM, Wells JM. Translating developmental principles to generate human gastric organoids. *Cell Mol Gastroenterol Hepatol.* 2018;5(3):353-363. doi:10.1016/j.jcmgh.2017.12.014

31. Pompaiah M, Bartfeld S. Gastric organoids: An emerging model for infection biology – a review of methods. *Curr Opin Gastroenterol.* 2014;30(1):25-33. doi:10.1097/MOG.0000000000000022

32. Dekkers JF, Berkers G, Kruiswinkel E, et al. Characterizing responses to CFTR-modulating drugs using rectal organoids derived from subjects with cystic fibrosis. *Sci Transl Med.* 2016;8(344):344RA84. doi:10.1126/scitranslmed.aad2878

33. Berkers G, van Mourik P, Vonk AM, et al. Rectal organoids enable personalized treatment of cystic fibrosis. *Cell Rep.* 2019;26(7):1701-1708.e3. doi:10.1016/j.celrep.2019.01.048

34. Alzeeb G, Metges J-P, Corcos L, Le Jossic-Corcos C. Three-dimensional culture systems in gastric cancer research. *Cancers (Basel).* 2020;12(10):1-20. doi:10.3390/cancers12102800

35. Traulsen J, Zagan C, Daddi AA, Boccellato F. Molecular modelling of the gastric barrier response, from infection to carcinogenesis. *Best Pract Res Clin Gastroenterol.* 2021;50-51:101737. doi:10.1016/j.jbpg.2021.101737

36. Broda TR, McCracken KW, Wells JM. Generation of human antral and fundic gastric organoids from pluripotent stem cells. *Nat Protoc.* 2019;14(1):28-50. doi:10.1038/s41596-018-0080-z

37. Choudhury D, Ashok A, Naing MW. Commercialization of organoids. *Trends Mol Med.* 2020;26(3):243-249. doi:10.1016/j.tmm.2019.12.002

38. Urbisheck M, Rannikmae H, Foets T, Ravn K, Hyvönen M, de la Roche M. Organoid culture media formulated with growth factors of defined cellular activity. *Sci Rep.* 2019;9(1):6193. doi:10.1038/s41598-019-42604-0

39. Barker N, Huch M, Kujala P, et al. Lgr5+ve stem cells drive self-renewal in the stomach and build long-lived gastric units in vitro. *Cell Stem Cell.* 2010;6(1):25-36. doi:10.1016/j.stem.2009.11.013

40. Barta-Skeirik N, Feng R, Schumacher MA, et al. CD44 plays a functional role in Helicobacter pylori-induced epithelial cell proliferation. *PLoS Pathog.* 2015;11(2):e1004663. doi:10.1371/journal.ppat.1004663

41. Huang J, Sweeney E, Sigal M, et al. Chemodetection and destruction of host urea allows Helicobacter pylori to locate the epithelium. *Cell Host Microbe.* 2015;18(2):147-156. doi:10.1016/j.chom.2015.07.002

42. Hanyu H, Engevik KA, Matthis AL, Ottemann KM, Montrose MH, Aihara E. Helicobacter pylori uses the TlpB receptor to sense sites of Helicobacter pylori infection. *Proc Natl Acad Sci U S A.* 2005;102(26):9300-9305. doi:10.1073/pnas.0409873102

43. Boccellato F, Van De Wetering M, et al. CD44 plays a functional role in Helicobacter pylori-induced epithelial cell proliferation. *Cell Host Microbe.* 2015;18(2):147-156. doi:10.1016/j.chom.2015.07.002

44. Uotani T, Murakami K, Uchida T, et al. Changes of tight junction and interleukin-8 expression using a human gastrointestinal model of Helicobacter pylori infection. *Helicobacter.* 2019;24(3):1-12. doi:10.1111/he.12583

45. Brandt S, Kwok T, Hartig R, Koo W, Backert S. NF-κB activation and potentiation of proinflammatory responses by the Helicobacter pylori CagA protein. *Proc Natl Acad Sci U S A.* 2005;102(26):9300-9305. doi:10.1073/pnas.0409873102

46. Backert S, Naumann M. What a disorder: Proinflammatory signaling pathways induced by Helicobacter pylori. *Trends Microbiol.* 2010;18(11):479-486. doi:10.1016/j.tim.2010.08.003

47. Miehlke S, Hackelsberger A, Meining A, et al. Severe expression of Helicobacter pylori CagA protein. *Proc Natl Acad Sci U S A.* 2005;102(26):9300-9305. doi:10.1073/pnas.0409873102

48. Kayisoglu Ö, Schlegel N, Bartfeld S. Gastrointestinal epithelial innate immunity—regionalization and organoids as new model of Helicobacter pylori infection. *Helicobacter.* 2019;24(3):1-12. doi:10.1111/he.12583

49. Morey P, Pfannkuch L, Peng E, et al. Helicobacter pylori depletes cholesterol in gastric glands to prevent interferon gamma signalling and escape the inflammatory response. *Gastroenterology.* 2018;154(5):1391-1404.e9. doi:10.1053/j.gastro.2017.12.008

50. Nebrell TA, Hashimi M, Sidar B, et al. Novel gastric spheroid culture model reveals chemokine-dependent recruitment of human dendritic cells to the gastric epithelium. *Cell Mol Gastroenterol Hepatol.* 2019;8(1):157-171.e3. doi:10.1016/j.jcmgh.2019.02.010
polypeptide-expressing metaplasia after acute loss of parietal cells. Gastroenterology. 2014;146(7):1727-1738.e8. doi:10.1053/j.gastro.2014.02.007

92. Keilberg D, Steele N, Fan S, Yang C, Zavros Y, Ottemann KM. Gastric metabolomics detects Helicobacter pylori correlated loss of numerous metabolites in both the corpus and antrum. Infect Immun. 2021;89(2):e00690-20. doi: 10.1128/IAI.00690-20

93. Sigal M, Reinés MDM, Müllerke S, et al. R-spondin-3 induces secretory, antimicrobial Lgr5+ cells in the stomach. Nat Cell Biol. 2019;21:812-823. doi:10.1038/s41556-019-0339-9

94. Eicher AK, Kechele DO, Sundaram N, et al. Functional human gastrointestinal organoids can be engineered from three primary germ layers derived separately from pluripotent stem cells. Cell Stem Cell. 2022;29(1):36-51.e6. doi:10.1016/j.stem.2021.10.010

95. Seidlitz T, Merker SR, Rothe A, et al. Human gastric cancer modelling using organoids. Gut. 2019;68(2):207-217. doi:10.1136/gutjnl-2017-314549

96. Jones BC, Calà G, De Coppi P, Giobbe GG. Paediatric gastric organoids as a tool for disease modelling and clinical translation. Pediatr Surg Int. 2021;37:317-324. doi:10.1007/s00383-020-04821-x