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Targeted deletion of MyD88 in intestinal epithelial cells results in compromised antibacterial immunity associated with downregulation of polymeric immunoglobulin receptor, mucin-2, and antibacterial peptides

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Intestinal epithelial cells (IECs) form a physical and immunological barrier that separates the vast gut microbiota from host tissues. MyD88-dependent Toll-like receptor signaling is a key mediator of microbial–host cross-talk. We examined the role of epithelial MyD88 expression by generating mice with an IEC-targeted deletion of the *Myd88* gene (*MyD88*^{ΔIEC}). Loss of epithelial MyD88 signaling resulted in increased numbers of mucus-associated bacteria; translocation of bacteria, including the opportunistic pathogen *Klebsiella pneumoniae*, to mesenteric lymph nodes; reduced transmucosal electrical resistance; impaired mucus-associated antimicrobial activity; and downregulated expression of polymeric immunoglobulin receptor (the epithelial IgA transporter), mucin-2 (the major protein of intestinal mucus), and the antimicrobial peptides RegIIIγ and Defa-rs1. We further observed significant differences in the composition of the gut microbiota between *MyD88*^{ΔIEC} mice and wild-type littermates. These physical, immunological, and microbial defects resulted in increased susceptibility of *MyD88*^{ΔIEC} mice to experimental colitis. We conclude that MyD88 signaling in IECs is crucial for maintenance of gut homeostasis.

The mammalian intestine harbors an estimated 100 trillion microorganisms, which normally maintain a mutually beneficial relationship with the host.¹ The intestinal epithelium provides a physical barrier as well as innate immune defense, preventing this vast community of microbes from entering host tissues.² Defects in epithelial barrier and immune functions can lead to infections with opportunistic and pathogenic microbes³ and contribute to the etiology of inflammatory bowel disease (IBD).⁴ Cross-talk between the gut microbiota and intestinal epithelial cells (IECs) is mediated by pattern recognition receptors, including the Toll-like receptor (TLR) family,⁵ leading to expression of gene products that enhance epithelial barrier function and innate immunity. TLR signaling in the gut is a double-edged sword; while physiological TLR signaling promotes intestinal homeostasis in healthy individuals, excessive TLR signaling in response to invasive microbes can exacerbate intestinal inflammation.⁶ The cytoplasmic adaptor protein MyD88 transduces

signals from TLRs that recognize bacterial products.⁷ The finding that *Myd88*-deficient mice exhibited increased susceptibility to chemically induced colitis highlighted the importance of physiological TLR signaling for protection of the epithelial barrier.⁸ Studies using bone marrow chimeras of *Myd88*-deficient and wild-type mice in several mouse models of colitis demonstrated that MyD88 signaling in hematopoietic cells was required for the development of intestinal inflammation, whereas MyD88 signaling in non-hematopoietic cells was required for host survival.^{9,10} Here we used cell type-specific deletion of the *Myd88* gene to investigate the role of epithelial-specific MyD88 signaling in antibacterial immunity and protection against intestinal inflammation.

Secretory immunoglobulin A (SIgA) acts as the first line of antigen-specific immunity at the interface between the gut microbiota and the intestinal epithelium.¹¹ Polymeric IgA secreted by plasma cells in the intestinal *lamina propria* is

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transported across IECs by the polymeric immunoglobulin receptor (pIgR).¹² Importantly, we found that reduced expression of pIgR was correlated with greater disease severity in patients with IBD¹³ and in mouse models of experimental colitis.¹⁴ Our finding that expression of pIgR is regulated in IECs by TLR signaling¹⁵ suggests that microbial-epithelial cross-talk is crucial for optimal production of SIgA. Consistent with this concept, we found that MyD88-deficient mice had reduced colonic expression of pIgR and deficient transport of SIgA.¹² However, in that study, it was not determined whether pIgR was regulated directly by MyD88 signaling in IECs or indirectly by immune cells in the intestinal *lamina propria*. In the present study, we used targeted deletion of the *Myd88* gene to demonstrate an essential role for epithelial-specific MyD88 signaling in maintaining optimal expression of pIgR and transport of SIgA.

The thick layer of intestinal mucus acts to separate luminal bacteria from the epithelial surface.¹⁶ This mucus is rich in antimicrobial peptides¹⁷ and associates with SIgA via glycan-mediated interactions,¹⁸ thus forming an immunological as well as a physical barrier against microbial invasion. Here we report that targeted deletion of MyD88 in IECs causes downregulated expression of pIgR, mucin-2 (the major protein constituent of intestinal mucus), and antimicrobial peptides in IECs. We propose a model in which MyD88-dependent bacterial-epithelial cross-talk is crucial for the maintenance of physical and immunological barrier functions in the intestine.

RESULTS

Loss of MyD88 signaling in colonic epithelial cells (ECs) disturbs the segregation of microbiota and host

To investigate the role of epithelial MyD88 signaling in intestinal homeostasis, we generated mice with a targeted deletion of the *Myd88* gene in IECs (MyD88^{ΔIEC}) by crossing mice carrying loxP-flanked *Myd88* alleles (MyD88^{Flox}) with mice that express *Cre* recombinase under the control of the IEC-specific *Vil1* promoter.¹⁹ Levels of MyD88 mRNA were significantly reduced in isolated colonic ECs from MyD88^{ΔIEC} mice at 9 weeks of age (**Figure 1a**). Immunostaining revealed expression of MyD88 in colonic ECs and *lamina propria* mononuclear cells of MyD88^{Flox} mice, whereas MyD88^{ΔIEC} mice lacked MyD88 expression in ECs but retained expression in *lamina propria* mononuclear cells (**Figure 1b**). Compared with MyD88^{Flox} littermates, MyD88^{ΔIEC} mice showed no histological signs of intestinal inflammation at 9 weeks of age (**Figure 1b**), or when followed up to 12 months (see **Supplementary Figure S1** online). No pathological changes in colon morphology were observed. Although there were no differences in numbers of culturable bacteria in feces from MyD88^{Flox} and MyD88^{ΔIEC} mice, we observed increased mucus-associated colony-forming units (CFU) in MyD88^{ΔIEC} mice (**Figure 1c**). These differences were not restricted to culturable bacteria, as similar trends were observed when using quantitative reverse transcriptase-PCR of 16S rDNA to quantify bacterial numbers (see **Supplementary Figure S2** online). Furthermore, increased bacterial attachment to the epithelial surface in MyD88^{ΔIEC} mice was correlated with increased translocation of bacteria to the draining mesenteric

lymph nodes (MLNs; **Figure 1d, e**). Interestingly, about 25% of bacterial colonies cultured from MLNs of MyD88^{ΔIEC} mice had a large, mucoid morphology and were identified as *Klebsiella pneumoniae* by sequencing of 16S rRNA genes and biochemical tests (see **Supplementary Figure S3** online). No bacteria could be cultured from the spleens of MyD88^{ΔIEC} mice, suggesting that infection with gut-derived bacteria was confined to the mucosal immune system. These findings indicate that epithelial-specific MyD88 signaling is required for normal segregation of gut commensals from host tissues.

Loss of MyD88 signaling in colonic ECs compromises epithelial barrier function

We hypothesized that increased epithelial permeability in MyD88^{ΔIEC} mice, coupled with increased numbers of mucus-adherent bacteria, could result in leakage of bacteria into the *lamina propria* and travel via the lymphatics to MLNs. Measurement of transmucosal electrical resistance in isolated colonic tissues revealed significant decreases in MyD88^{ΔIEC} mice, indicative of decreased epithelial barrier function (**Figure 2a**). This *ex vivo* analysis was supported by the *in vivo* observation of elevated levels of serum albumin in the feces of MyD88^{ΔIEC} mice, reflecting leakage of serum proteins into the intestinal lumen (see **Supplementary Figure S4** online). To determine whether increased mucosal adherence and epithelial translocation of bacteria in MyD88^{ΔIEC} mice was selective for opportunistic pathogens such as *K. pneumoniae*, we treated mice orally for 7 days with an environmental, nonpathogenic strain of *Escherichia coli* (Nissle 1917),²⁰ which had been transformed to kanamycin resistance. We observed similar numbers of Kan^R *E. coli* in feces and luminal washes from MyD88^{Flox} and MyD88^{ΔIEC} mice, but significantly increased numbers in the colonic mucus of MyD88^{ΔIEC} mice (**Figure 2b**), consistent with the finding that mucus extracted from MyD88^{ΔIEC} mice had reduced antibacterial activity compared with MyD88^{Flox} mice (**Figure 2c**). However, these Kan^R *E. coli* did not translocate to the MLNs of MyD88^{ΔIEC} mice (data not shown). These results suggested that the increased mucosal adherence of bacteria in MyD88^{ΔIEC} mice may be non-selective, whereas translocation to the draining lymph nodes may be selective for more aggressive bacteria such as *K. pneumoniae*.

We hypothesized that loss of epithelial MyD88 signaling could result in increased bacterial invasion of IECs. To assess epithelial invasion independently of the confounding effects of local immune cells, we performed *in vitro* bacterial invasion assays in the human HT-29 colon carcinoma cell-line. We tested the effects of small hairpin RNA (shRNA)-mediated knockdown of MyD88 (see **Supplementary Figure S5a** online for documentation of MyD88 depletion) on attachment and invasion of three species of Gram-negative bacteria of the family *Enterobacteriaceae*: the non-invasive *E. coli* strain Nissle 1917, the opportunistic pathogen *K. pneumoniae* (a strain isolated from the MLN of a MyD88^{ΔIEC} mouse), and the invasive pathogen *Salmonella typhimurium* (strain SL1344) (**Figure 2d**). Knockdown of MyD88 expression in HT-29 cells resulted in significantly increased adherence of all three bacterial species to

Figure 1 Targeted deletion of MyD88 in intestinal epithelial cells (IECs) results in increased numbers of mucus-associated bacteria and bacterial translocation to mesenteric lymph nodes (MLNs). **(a)** Mice with a targeted deletion of the *Myd88* gene in IECs (MyD88^{ΔIEC}) were created by crossing mice in which both copies of the *Myd88* gene were flanked with *loxP* sites (MyD88^{Flox}) with mice expressing *Cre* recombinase under the control of the IEC-specific *Vil1* promoter. Levels of MyD88 mRNA were analyzed in colonic epithelial cells isolated from 9-week-old littermate MyD88^{Flox} and MyD88^{ΔIEC} mice by Nanostring nCounter hybridization. mRNA transcript abundance was normalized as described in the Methods section, and expressed as mean ± s.e.m. (*n* = 15). **(b)** Colon tissue sections from MyD88^{Flox} and MyD88^{ΔIEC} mice were stained with hematoxylin and eosin (upper panels) or fluorescein isothiocyanate-labeled antibodies to MyD88 (green) and the nuclear stain 4,6-diamidino-2-phenylindole (blue; lower panels). **(c)** Bacterial colony-forming units (CFU) were enumerated following anaerobic culture of fecal homogenates (CFU per g feces), luminal washes (CFU per ml phosphate-buffered saline), and surface mucus (CFU per 100 μl mucus) from colons of MyD88^{Flox} and MyD88^{ΔIEC} mice. Data are expressed as mean ± s.e.m. (*n* = 6). **(d)** Bacterial colonies cultured under anaerobic conditions from MLNs of representative MyD88^{Flox} and MyD88^{ΔIEC} mice. Large mucoid colonies from MyD88^{ΔIEC} mice were identified as *Klebsiella pneumoniae* (*Kp*) by sequence analysis of 16S rRNA genes and biochemical tests (see **Supplementary Figure S3** online). **(e)** CFU of total anaerobic bacteria and *K. pneumoniae* cultured from MLNs of MyD88^{Flox} and MyD88^{ΔIEC} mice. Data in the bar graph are expressed as CFU per MLN (mean ± s.e.m., *n* = 27). The numbers below the bar graph indicate the percentage of mice of each genotype from which bacteria (total and *K. pneumoniae*) could be cultured from MLNs. In all panels, asterisks indicate that the mean for MyD88^{ΔIEC} mice is significantly different from the mean for MyD88^{Flox} mice (*P* < 0.05).

the epithelial surface. As expected, the magnitude of epithelial invasion by *E. coli* and *K. pneumoniae* was dramatically lower than was observed for *S. typhimurium*, but invasion of all three species was significantly increased when expression of MyD88 was reduced. Taken together, our findings suggest that MyD88 signaling in ECs plays a crucial role in intestinal immunity by increasing the antibacterial activity of epithelial mucus, reducing

bacterial adherence to the epithelial surface, maintaining epithelial barrier function, and preventing bacterial invasion of ECs.

Loss of MyD88 signaling in IECs alters the composition of the gut microbiota

We hypothesized that compromised innate immunity resulting from loss of epithelial MyD88 signaling would affect the balance

Figure 2 Loss of MyD88 expression in intestinal epithelial cells (IECs) compromises epithelial barrier integrity. **(a)** Epithelial barrier function. Transmucosal electrical resistance (TER) was analyzed *ex vivo* in isolated mouse colon tissues placed in Ussing chambers. Data are expressed as mean ± s.e.m. ($n=8$). **(b)** Association of exogenous, noninvasive bacteria with colonic mucus. MyD88^{Flox} and MyD88^{ΔIEC} mice were administered *Escherichia coli* strain Nissle 1917 that had been transformed to kanamycin resistance (Kan^R *E. coli*) at a dose of 2.5×10^7 CFU ml⁻¹ in the drinking water for 7 days. Kanamycin-resistant colonies were enumerated after anaerobic culture of fecal homogenates (colony-forming units (CFU) per g feces), luminal washes (CFU per ml phosphate-buffered saline), and epithelial-associated mucus (CFU per 100 μl mucus). Data are expressed as mean ± s.e.m. ($n=6$). **(c)** Antibacterial activity of colonic mucus. Aliquots of mucus harvested from the colons of MyD88^{Flox} and MyD88^{ΔIEC} mice were incubated for 15 min with 10^6 CFU of Kan^R *E. coli*, and then cultured in the presence of kanamycin to enumerate surviving *E. coli*. Data are expressed as mean ± s.e.m. ($n=6$). **(d)** *In vitro* bacterial invasion assays. Clones of the HT-29 human colon carcinoma cell line were stably transfected with small hairpin RNA (shRNA) specific for MyD88 or a random control sequence. Cell monolayers (2 cm²) were incubated with 10^6 CFU (MOI of 10:1) of *E. coli* strain Nissle 1917, a strain of *Klebsiella pneumoniae* isolated from mesenteric lymph nodes of MyD88^{ΔIEC} mice, or *Salmonella typhimurium* strain SL1344. Membrane-bound CFU were enumerated by anaerobic culture of cell homogenates after washing to remove unbound bacteria, and intracellular CFU were enumerated in parallel plates after 2 h of treatment of cell monolayers with gentamicin to kill all extracellular bacteria. Data are expressed as mean ± s.e.m. ($n=6$). Asterisks indicate that the mean for MyD88^{ΔIEC} mice is significantly different from the mean for MyD88^{Flox} mice (panels **a–c**) or that the mean for cells expressing MyD88 shRNA is significantly different from the mean for cells expressing control shRNA (**d**) ($P<0.05$).

of the gut bacterial community, resulting in an altered composition of the gut microbiota in MyD88^{ΔIEC} mice. The richness and diversity of fecal bacteria was analyzed by the PhyloChip assay (Second Genome, Inc., San Bruno, CA), a microarray-based method that measures the relative abundance of more than 59,000 individual microbial taxa in fecal DNA by analysis

of the entire 16S ribosomal RNA gene sequence,²¹ in MyD88^{Flox} and MyD88^{ΔIEC} mice. Although the overall richness of the fecal microbiota of MyD88^{Flox} and MyD88^{ΔIEC} mice was similar (see **Supplementary Figures S6 and S7** online), significant differences were observed in the relative abundance of operational taxonomic units (OTUs, defined as a group of highly similar

Figure 3 Differences in the composition of the fecal microbiota of MyD88^{Flox} and MyD88^{ΔIEC} mice. The relative abundance of operational taxonomic units (OTUs) was quantified by Phylochip microarray analysis of 16S rRNA gene sequences in fecal DNA from 4 MyD88^{Flox} and 5 MyD88^{ΔIEC} littermate mice (see the Methods section for details of data analysis). A representative 16S rRNA gene from each of 109 differentially expressed OTUs was aligned and used to infer the phylogenetic tree shown in this figure. The rings around the tree comprise a heatmap where the inner ring includes the samples from MyD88^{Flox} mice, and the outer ring includes the samples from MyD88^{ΔIEC} mice. Blue lines indicate that the OTU was more abundant in that sample than in the mean of MyD88^{Flox} mice samples, and red lines indicate that the OTU was less abundant. The color saturation indicates the fold difference in OTU abundance for each mouse compared with the mean for MyD88^{Flox} mice.

16S rRNA gene sequences, in most cases with similarity >99% within several bacterial families (Figure 3). A large proportion of families in the phylum *Bacteroidetes* were decreased in abundance in MyD88^{ΔIEC} mice, whereas a large proportion of families in the phylum *Proteobacteria* were increased. Within the phylum *Firmicutes*, the abundance of some families were increased while others were decreased in MyD88^{ΔIEC} mice. Principal component analysis (PCA), which was used to visualize complex relationships among multiple bacterial families in individual mice, revealed significant differences between MyD88^{Flox} and MyD88^{ΔIEC} mice (Figure 4a, left panel). Interestingly, the composition of the microbiota was much more variable among individual MyD88^{Flox} mice than among individual MyD88^{ΔIEC} mice, as indicated by the tight clustering of the five MyD88^{ΔIEC} mice in the two-dimensional PCA plot. This result suggests that loss of epithelial MyD88 signaling may cause specific and reproducible changes in the bacterial community, consistent with the finding that the abundance of 2647 OTUs differed significantly between MyD88^{Flox} and MyD88^{ΔIEC} mice (Figure 4a, right panel). The Prediction Analysis for Microarrays method²² was used to identify “distinctive” OTUs whose relative abundance characterizes the unique microbiota of MyD88^{Flox} and MyD88^{ΔIEC} mice (Figure 4b). This analysis identified four distinctive OTUs in the TM7 candidate division that were characteristic of MyD88^{ΔIEC} mice, and seven OTUs within the genus

Lactobacillus (phylum *Firmicutes*) that were characteristic of MyD88^{Flox} mice. Interestingly, the relative abundance of five bacterial species that have been associated with altered innate and adaptive immunity in the gut, including *Candidatus arthromitus* (SFB),²³ *E. coli*,²⁴ *Pseudomonas fluorescens*,²⁴ *K. pneumoniae*,²⁵ and *Proteus mirabilis*,²⁵ did not differ significantly in the fecal microbiota of MyD88^{Flox} and MyD88^{ΔIEC} mice (Figure 4c). This finding suggests that translocation of *K. pneumoniae* to the MLNs of MyD88^{ΔIEC} mice (Figure 1e) was due to a specific defect in the host response to this bacterium, and not simply due to an increase in the abundance of *K. pneumoniae* in the gut microbiota.

Loss of MyD88 signaling in IECs results in altered epithelial gene expression and reduced transcytosis of IgA

To determine whether loss of epithelial MyD88 signaling resulted in altered patterns of gene expression, we isolated colonic ECs from MyD88^{Flox} and MyD88^{ΔIEC} mice and analyzed mRNA levels for 36 genes that have been implicated in innate immune responses of IECs, maintenance of epithelial barrier function, and protection against inflammation (see **Supplementary Methods** online for a detailed description of selection criteria for these genes). Among these genes, five (including MyD88, as a positive control) were downregulated in MyD88^{ΔIEC} mice, and none were upregulated (Figure 5a). The most significantly downregulated gene was the polymeric immunoglobulin receptor

Figure 4 Loss of MyD88 expression in intestinal epithelial cells (IECs) is associated with changes in the composition of the fecal microbiota. **(a)** Principal component analysis (PCA) was used to generate two-dimensional ordination plots that visualize complex relationships between the fecal microbiota of individual MyD88^{Flox} and MyD88^{ΔIEC} mice, based on weighted Unifrac distances between operational taxonomic units (OTUs) detected in fecal samples. The graph on the left is based on 14,193 OTUs present in at least one mouse, and the graph on the right is based on 2,647 OTUs with significant differences in abundance between MyD88^{Flox} and MyD88^{ΔIEC} mice. **(b)** Significant OTUs that characterize the distinctive microbiota of MyD88^{Flox} and MyD88^{ΔIEC} mice were identified using the Prediction Analysis for Microarrays method. Data in the bar graph are expressed as fold difference ($\log_2$, mean $\pm$ s.e.m.) in abundance between MyD88^{ΔIEC} ($n=5$) and MyD88^{Flox} mice ($n=4$). Bars to the right of the zero line represent distinctive OTUs that were more abundant in MyD88^{ΔIEC} mice, all of which were classified in the candidate phylum TM7. Bars to the left represent distinctive OTUs that were more abundant in MyD88^{Flox} mice, all of which were classified in the genus *Lactobacillus* (phylum *Firmicutes*). Numbers in parentheses on the y-axis are the IDs for individual OTUs (see **Supplementary Table S1** online for complete taxonomy of the distinctive OTUs). **(c)** Relative abundance of five species of fecal bacteria in individual MyD88^{Flox} and MyD88^{ΔIEC} mice. Because each of these species (except *Candidatus arthromitus*) comprised multiple OTUs, the OTU that represented the biggest difference in abundance between MyD88^{Flox} and MyD88^{ΔIEC} mice is displayed.

(*Pigr*), the epithelial transporter for polymeric IgA antibodies. Immunostaining of colon tissues revealed that expression of pIgR and colocalization with IgA were severely downregulated in ECs of MyD88^{ΔIEC} mice (**Figure 5b**), and fecal IgA levels were significantly reduced (**Figure 5c**). Importantly, this reduction in luminal IgA was similar to the reduction we previously reported in MyD88^{-/-} mice¹¹, supporting a direct role for epithelial MyD88 signaling in regulating IgA transcytosis. We confirmed a direct role for epithelial MyD88 signaling in regulating pIgR expression by demonstrating that shRNA-mediated knockdown of MyD88 expression in the human HT-29 colon carcinoma cell line significantly reduced the upregulation of pIgR in response to bacterial stimulation (see **Supplementary Figure S5b** online). Other downregulated genes included mucin-2 (*Muc2*), the major protein constituent of intestinal mucus, and the antimicrobial

peptides regenerating islet-derived III γ (*RegIII γ*) and defensin- α related sequence 1 (*Defa-rs1*). Although there were no significant differences in goblet cell numbers in the colonic epithelia of MyD88^{Flox} and MyD88^{ΔIEC} mice (see **Supplementary Figure S8** online), immunostaining revealed that the level of mucin-2 protein per goblet cell was reduced in MyD88^{ΔIEC} mice (**Figure 5d**). Downregulation of RegIII γ and Defa-rs1 likely contributed to the reduced antimicrobial activity of colonic mucus in MyD88^{ΔIEC} mice (**Figure 2c**).

Loss of MyD88 signaling in colonic ECs results in increased susceptibility to dextran sulfate sodium (DSS)-induced colitis

Inappropriate inflammatory responses to normal components of the intestinal microbiota are thought to have a major role in

Figure 5 Loss of MyD88 expression alters epithelial gene expression and compromises polymeric immunoglobulin receptor (pIgR)-mediated immunoglobulin (IgA) transport. **(a)** Colonic epithelial cells were isolated from 8-week-old MyD88^{ΔIEC} and MyD88^{Flox} mice ($n=12$). Levels of 36 individual mRNA transcripts were analyzed by Nanostring nCounter hybridization and plotted as mean fold difference (log₂) between MyD88^{ΔIEC} and MyD88^{Flox} mice vs. P value ($-\log_{10}$). mRNA transcript levels for genes above the red dashed line were significantly different in MyD88^{ΔIEC} mice compared with MyD88^{Flox} mice ($P<0.05$). Genes to the left of the zero line were downregulated, whereas genes to the right were upregulated. **(b)** Immunofluorescence staining of colon tissues from 9-week-old MyD88^{Flox} and MyD88^{ΔIEC} mice (pIgR = red, IgA = green, 4,6-diamidino-2-phenylindole (DAPI)-stained nuclei = blue). **(c)** Feces were collected from 10-week-old MyD88^{Flox} and MyD88^{ΔIEC} mice and IgA levels were determined by enzyme-linked immunosorbent assay. Data are expressed as mean $\pm$ s.e.m. ($n=5$). The asterisk indicates that the mean for MyD88^{ΔIEC} mice is significantly different from the mean for MyD88^{Flox} mice ($P<0.05$). **(d)** Immunofluorescence staining of colon tissues from 9-week-old MyD88^{Flox} and MyD88^{ΔIEC} mice (Muc2 = red, DAPI-stained nuclei = blue). IEC, intestinal epithelial cells.

the pathogenesis of IBD.^{4,26,27} Although we observed no overt inflammatory disease in MyD88^{ΔIEC} mice, they exhibited several defects that have been reported in IBD patients, including reduced expression of pIgR,¹³ Muc2,²⁸ and antimicrobial peptides,^{17,29} as well as perturbed spatial relationships between microbiota and host.^{30,31} To test the hypothesis that MyD88^{ΔIEC} mice are more susceptible to experimental colitis, acute epithelial damage was induced by administration of 2% DSS in the drinking water for 7 days. MyD88^{ΔIEC} mice had significantly higher disease activity than did MyD88^{Flox} mice, beginning as early as 1 day after initiation of DSS treatment (**Figure 6a**). At the termination of the experiment, increased inflammatory cell infiltration and

epithelial destruction were observed in colons of MyD88^{ΔIEC} mice (**Figure 6b** and **c**). PCA of gene expression in colonic ECs following DSS treatment revealed significantly different patterns in MyD88^{ΔIEC} and MyD88^{Flox} mice (**Figure 6d**; see **Supplementary Tables S3** and **S4** online for detailed gene expression data, and **Supplementary Figure S9** online for a detailed description of PCA results). Expression of 10 genes was significantly downregulated and expression of 5 genes was upregulated in colonic ECs from MyD88^{Flox} mice following DSS treatment (**Figure 6e**, top panel). Interestingly, MyD88^{ΔIEC} mice were defective in their ability to upregulate the expression of the pro-inflammatory factors tumor necrosis factor (TNF), CXCL2, and interleukin (IL)-1 β in colonic

Figure 6 Loss of MyD88 expression in intestinal epithelial cells (IECs) increases the severity of dextran sulfate sodium (DSS)-induced colitis. (a) Disease activity in MyD88^{Flox} and MyD88^{ΔIEC} mice that were given regular drinking water or 2% DSS for 7 days. Data are expressed as mean ± s.e.m. ($n=8$). One asterisk indicates that the mean for DSS-treated mice is significantly greater than the corresponding mean for untreated mice, and two asterisks indicate that the mean for DSS-treated MyD88^{ΔIEC} mice is greater than the mean for DSS-treated MyD88^{Flox} mice. (b) Colon histology from representative mice at day 7. Images of formalin-fixed colon tissues stained with hematoxylin and eosin were captured with a $\times 20$ objective. (c) Histological inflammatory scores for MyD88^{Flox} and MyD88^{ΔIEC} mice on day 7. Data are expressed as mean ± s.e.m. ($n=8$). Asterisks indicate that the means for DSS-treated mice are significantly different from the means for untreated mice of the same genotype, and that the mean for DSS-treated MyD88^{ΔIEC} mice is significantly different from the mean for DSS-treated MyD88^{Flox} mice ($P<0.05$). (d) Principal component analysis was used to generate two-dimensional ordination plots that visualize complex relationships in expression of multiple genes in colonic EC of individual MyD88^{Flox} and MyD88^{ΔIEC} mice, with or without DSS treatment. Normalized mRNA transcript levels for 19 genes were reduced to 2 PCs (see **Supplementary Figure S9** online for details), which are plotted for individual mice. (e) Levels of 36 individual mRNA transcripts were measured by Nanostring nCounter hybridization. Data are plotted as mean fold difference ($\log_2$) between MyD88^{Flox} mice with or without DSS treatment vs. P value ($-\log_{10}$; upper panel), and mean fold difference ($\log_2$) between DSS-treated MyD88^{ΔIEC} and MyD88^{Flox} mice vs. P value ($-\log_{10}$; lower panel). Individual dots represent the mean of treatment groups ($n=8$). mRNA transcript levels for genes above the red dashed line were significantly different for each comparison ($P<0.05$). Genes above the red dashed line were significantly different in DSS-treated mice compared to untreated mice ($P<0.05$). Genes to the left of the zero line were downregulated, and genes to the right were upregulated.

ECs in response to DSS-induced damage (**Figure 6e**, bottom panel). Aberrant upregulation of the antimicrobial peptide Defa-rs1 in colonic ECs of MyD88^{ΔIEC} mice may have been caused by increased microbial stimulation secondary to increased epithelial

damage in these mice. These findings suggest that lack of epithelial MyD88 expression causes specific defects in innate immune gene expression, resulting in a compromised host response to epithelial injury and increased colonic inflammation.

DISCUSSION

Protection against bacterial invasion and infection in the intestine is dependent on the resident microbiota, the mucus layer containing SIgA and antimicrobial peptides, and the monolayer of IECs that provide a physical and innate immune barrier.³ Here we demonstrated that loss of epithelial MyD88 signaling compromises antibacterial immunity at all three layers of defense, resulting in translocation of bacteria to MLNs. Loss of MyD88 in IECs did not cause spontaneous colitis, but rendered mice more susceptible to inflammation caused by the epithelial damaging agent DSS. These findings demonstrate the critical role of epithelial MyD88 signaling in maintenance of intestinal barrier function and protection against inflammation.

Unlike the inflammatory response generated by MyD88-dependent TLR signaling in immune cells, IECs respond by producing factors involved in EC proliferation, maintenance of barrier function, innate antimicrobial immunity, and transport of SIgA.^{12,32} Here we report that targeted deletion of the *Myd88* gene in IECs was associated with reduced expression of pIgR and epithelial transport of IgA, providing direct evidence for the first time that epithelial MyD88 signaling regulates the production of SIgA. These SIgA antibodies in turn modulate the composition of the gut microbiota and form the first line of antigen-specific immunity that prevents invasion of IECs by members of the gut microbiota.³³ This connection between epithelial MyD88 signaling, SIgA production, and host-microbial homeostasis provides a potential explanation for our previous findings that reduced pIgR expression and SIgA transport were correlated with disease severity in human IBD¹³ and in mouse models of experimental colitis.¹⁴ We previously reported³⁴ that expression of pIgR was upregulated by TNF and IL-1 β , the latter of which signals through a MyD88-dependent pathway.³⁵ We found here that loss of epithelial MyD88 signaling was associated with defective production of IL-1 β and TNF in colonic ECs in response to DSS-induced inflammation, suggesting that MyD88 may coordinate the response of IECs to autocrine and paracrine stimulation with IL-1 and TNF, as well as TLR signaling.

The intestinal epithelium is protected by a mucus layer that increases in thickness from the small to the large bowel in proportion to the density of the microbiota.³⁶ In addition to its role as a physical barrier, the mucus serves to trap protective factors such as SIgA and antimicrobial peptides.^{3,37} We found that colonic ECs from MyD88 Δ IEC mice produced significantly less Muc-2, antimicrobial peptides RegIII γ and Defa-rs1, and SIgA, consistent with reduced antimicrobial activity in colonic mucus. Vaishnavi *et al.*³¹ recently reported that mice with a targeted deletion of MyD88 in IECs had reduced expression of the antimicrobial peptide RegIII- γ in the terminal ileum, and increased bacterial colonization of small intestinal surfaces. Gong *et al.*³⁸ reported that mice transgenic for an IEC-targeted, dominant-negative mutant of MyD88 had reduced expression of antimicrobial peptides in the small intestine and increased bacterial translocation to MLNs. MyD88 signaling may also provide cues to immune cells for regulating the expression of antimicrobial and innate immune factors. Ismail *et al.*³⁹ found that expression of RegIII- γ in small intestinal $\gamma\delta$ intraepithelial lymphocytes

was dependent on epithelial-specific MyD88 signaling, but not on expression of MyD88 within the intraepithelial lymphocytes themselves. Thus, epithelial MyD88 signaling provides direct and indirect signals to augment production of antimicrobial factors that segregate the microbiota from host tissues.

In our studies, a hallmark of epithelial-specific MyD88 deficiency was translocation of bacteria to the draining MLNs, including the opportunistic pathogen *K. pneumoniae*. This Gram-negative bacterium of the family *Enterobacteriaceae* is commonly found among the resident gut bacteria in both humans and mice, but has limited invasive potential in healthy individuals.⁴⁰ Although the defect in epithelial barrier function in MyD88 Δ IEC mice may have contributed to paracellular leakage of bacteria into the *lamina propria*, we did not observe translocation of commensal *E. coli* to the MLNs of MyD88 Δ IEC mice, even though these bacteria colonized the mucus layer of the colon. We conclude that virulence factors expressed by *K. pneumoniae*, but not by commensal *E. coli*, may be required for survival during transit from the mucosal surface to the MLNs. By contrast, Fukata *et al.*²⁴ reported translocation of *E. coli* to the MLNs in mice deficient in MyD88 in all cell types. MyD88-deficient mice have been shown to be highly susceptible to systemic infections with a broad range of bacteria.⁴¹ Intratracheal infection of MyD88-deficient mice with *K. pneumoniae* resulted in severe pneumonia, attenuated neutrophil influx, and systemic dissemination of bacteria.⁴² Rare cases of autosomal recessive MyD88 deficiency in humans were found to be associated with recurrent life-threatening infections with pyogenic bacteria early in life,⁴¹ including an acute urinary tract infection caused by *K. pneumoniae* in a 4-month-old boy. However, we did not observe systemic bacterial infections in MyD88 Δ IEC mice. We conclude that epithelial MyD88 signaling is required for preventing the translocation of *K. pneumoniae* and possibly other opportunistic pathogens to the MLNs, but that MyD88-sufficient immune cells can prevent the systemic spread of resident gut bacteria that have penetrated the mucosal barrier.

Our findings have revealed for the first time a unique role for epithelial MyD88 signaling in shaping the composition of the resident gut microbiota, including increased numbers of OTUs in the phylum *Proteobacteria* and decreased numbers of OTUs within the phylum *Bacteroidetes* in MyD88 Δ IEC mice. Interestingly, similar changes in the composition of the gut microbiota have been observed in human IBD patients.⁴³ Altered host factors that are believed to impact commensal bacteria in IBD patients include increased intestinal permeability, decreased production of antimicrobial peptides, and aberrant SIgA responses,⁴³ all of which characterize the immune defects in mice with a targeted deletion of MyD88 in IECs. In addition to changes at the level of bacterial phyla and families, the Prediction Analysis for Microarrays technique identified four “signature” OTUs in the candidate division TM7, whose increased abundance was characteristic of the altered microbiota of MyD88 Δ IEC mice. Importantly, increased abundance of TM7 bacteria have been reported in humans with oral inflammation⁴⁴ and IBD.⁴⁵ Increased abundance of TM7 bacteria has been reported in the gut microbiota of mice deficient in the NLRP6 inflammasome,

accompanied by the development of spontaneous intestinal inflammation and increased susceptibility to DSS-induced colitis.⁴⁶ Reduced production of IL-18, which transduces signals through MyD88, was characteristic of NLRP6-deficient mice. Thus, impaired MyD88 signaling in the intestine may alter the local microenvironment to favor expansion of TM7. By contrast, we did not observe changes in the gut microbiota of MyD88^{ΔIEC} mice that have been associated with other mouse models of intestinal inflammation caused by defects in immune cells, such as increased abundance of Gram-positive segmented filamentous bacteria (*C. arthromitus*), which have been implicated in the expansion of Th17 cells and inflammation of the gut and extraintestinal tissues.²³ Neither did we observe expansion of *Proteus mirabilis* nor *K. pneumoniae*, two Gram-negative species of the family *Enterobacteriaceae*, the abundance of which was increased in the gut microbiota of mice with defective immunity in dendritic cells in the intestinal *lamina propria*.²⁵ Increased colonization with *Enterobacteriaceae* and elevated titers of *Enterobacteriaceae* antibodies have been associated with IBD in humans,⁴⁷ and species of the genus *Klebsiella* are observed more frequently in the stool of ulcerative colitis patients than healthy controls.⁴⁸ Our finding of unique changes in the gut microbiota of MyD88^{ΔIEC} mice suggests that epithelial MyD88 signaling is required to protect against abnormal expansion of specific members of the gut microbial community.

Although loss of epithelial MyD88 signaling was not sufficient to cause spontaneous intestinal inflammation for at least 1 year of life, it was associated with increased severity of colitis following oral administration of the epithelial damaging agent, DSS. A potential mechanism for the increased susceptibility to DSS-induced colitis was suggested by our observation that MyD88^{ΔIEC} mice failed to upregulate expression of TNF, CXCL2, and IL-1 β in colonic ECs in response to epithelial damage. This finding was somewhat surprising, considering that these factors are generally considered to be pro-inflammatory and their increased expression has been linked to increased disease severity in experimental colitis.⁸ Our data are consistent with the notion that “physiological” inflammatory responses by IECs are a normal response to the continuous presence of the gut microbiota and are necessary for repair of damaged ECs. On the other hand, massive infiltration of gut microbes into the intestinal *lamina propria* following epithelial damage can result in “pathological” inflammatory responses and result in collateral damage to host tissues.

In conclusion, our findings shed new light on the beneficial effects of epithelial–microbial cross-talk mediated by MyD88 signaling, and demonstrate that MyD88-dependent TLR signaling in immune cells is insufficient to sustain protective innate immune responses in the intestine. Maintenance of intestinal homeostasis thus requires active communication among ECs, immune cells, and the gut microbiota.

METHODS

Mice. Mouse strains, on the C57BL/6 background, were purchased from Jackson Laboratories (Bar Harbor, ME). Mice of the B6.129P2-MyD88^{tm1Defr/J} strain were intercrossed to generate mice homozygous

for *loxP*-flanked *Myd88* (MyD88^{Flox}). The B6.SJL-Tg(Vil-cre)997Gum/J strain, which is transgenic for *Cre* recombinase under control of the intestinal epithelial-specific mouse *Vil1* gene, was intercrossed with MyD88^{Flox} mice to generate mice with a targeted deletion of the *Myd88* gene in ECs in the small and large bowel (MyD88^{ΔIEC}). Experiments were conducted with littermates generated by crossing MyD88^{ΔIEC} mice hemizygous for the *Vil1-Cre* transgene with MyD88^{Flox} mice. Acute colitis was induced by oral administration of DSS (molecular weight 36,000–50,000; MP Biomedicals, Aurora, OH), and disease activity indices were calculated as described.^{49,50} Mice were housed in microisolator cages in an AAALAC-accredited facility, and procedures were conducted in compliance with the University of Kentucky Institutional Animal Care and Use Committee.

Colonic EC isolation and analysis of mRNA levels. ECs were isolated from freshly dissected colons by dissociation with dithiothreitol and ethylenediaminetetraacetic acid and separation on a Percoll gradient, as previously described.⁵¹ Using this protocol, we typically achieve 99% purity of IECs, with contaminating CD45⁺ hematopoietic cells at levels of 0.4–1.3%.¹⁴ Abundance of mRNA transcripts was analyzed by Nanostring nCounter hybridization,⁵² and normalized using an algorithm developed by NanoString Technologies (Seattle, WA). MyD88 and pIgR mRNA in HT-29 cells were analyzed by quantitative reverse transcriptase-PCR and normalized to the level of β 2-microglobulin mRNA as described.¹⁵

Tissue histology and immunofluorescence microscopy. Dissected colons were cut lengthwise and fixed in 10% buffered formalin. Sections of formalin-fixed, paraffin-embedded colon tissues were stained with hematoxylin and eosin, or visualized by immunofluorescence using standard protocols. MyD88 was detected with a rabbit antibody (Abcam, Cambridge, MA), and pIgR was detected with a goat antibody to mouse secretory component (the extracellular domain of pIgR; R&D Systems, Minneapolis, MN), followed by horseradish peroxidase-conjugated antibody to rabbit or goat IgG (Invitrogen, Camarillo, CA) and TSA Plus-TMR Reagent (Perkin-Elmer, Waltham, MA). Co-localization with IgA was detected by fluorescein isothiocyanate-conjugated anti-mouse IgA (eBiosciences, San Diego, CA). Sections were mounted with VectaShield (Vector Laboratories, Burlingame, CA) containing 4,6-diamidino-2-phenylindole to visualize nuclei and imaged on a Zeiss Axiophot confocal microscope (Carl Zeiss Microscopy LLC, Thornwood, NY) with Axiovision image software.

Bacterial culture from mouse feces and tissues. Fecal pellets were gently released from dissected colons and homogenized in phosphate-buffered saline. Luminal washes were collected by flushing colons with 5 ml of phosphate-buffered saline, and mucus was collected by gentle scraping. Aliquots of fecal homogenates, luminal washes, and mucus were cultured for 48 h on LB agar at 37°C under anaerobic conditions. For analysis of bacterial translocation, MLNs were homogenized in phosphate-buffered saline and cultured anaerobically for 48 h. *E. coli* strain Nissle 1917 (a generous gift of Dr Ulrich Sonnenborn, Ardeypharm GmbH, Herdecke, Germany) was transformed to kanamycin resistance, and Kan^R *E. coli* were delivered to the mice in the drinking water at a dose of 2.5×10^7 CFU ml⁻¹ for 7 days. Feces, luminal washes, and colonic mucus were collected as described above and cultured anaerobically in 50 μ g ml⁻¹ kanamycin to enumerate CFU.

Measurement of transmucosal electrical resistance. Segments of mouse colon were mounted in 0.3 cm² aperture Ussing chambers (Physiologic Instruments, San Diego, CA), and transmucosal electrical resistance was measured as previously described.⁵³ Apical and basolateral recording buffers contained 118 mM NaCl, 5 mM KCl, 2 mM NaH₂PO₄, 1.2 mM MgSO₄, 1 mM CaCl₂, 10 mM HEPES, 10 mM glucose, and 22 mM NaHCO₃, pH 7.4, and were bubbled with 95% O₂ and 5% CO₂ at 37°C.

Analysis of fecal IgA and albumin by enzyme-linked immunosorbent assay. Freshly collected feces were diluted in 50 mM Tris, pH 7.4,

0.14 M NaCl, 1% bovine serum albumin, 0.05% Tween 20, and analyzed by enzyme-linked immunosorbent assay as previously described¹¹ using mouse IgA (Cat. No. E90-103) and mouse albumin (Cat. No. E90-134) ELISA Quantification Sets (Bethyl Laboratories, Montgomery, TX).

Analysis of mucus-associated antimicrobial activity. Colonic mucus was isolated as described above, diluted 1:3 in phosphate-buffered saline, and concentrated by centrifugation at 2,500 r.p.m. for 10 min. In all, 10^6 CFU of Kan^R *E. coli* from a mid-log phase culture was added to the mucus, then centrifuged together at 1,500 r.p.m. for 15 min at room temperature. Serial dilutions of bacterial suspensions before and after incubation with colonic mucus were cultured anaerobically for 24 h in $50 \mu\text{g ml}^{-1}$ kanamycin to enumerate CFU.

shRNA-mediated knockdown of MyD88 in human colonic ECs and bacterial invasion assays. HT-29 cells were transfected with SureSilencing plasmids encoding shRNA specific for human MyD88 or an irrelevant control shRNA (SABiosciences, Frederick, MD, cat no. KH00911P), and puromycin-resistant subclones were isolated as described.⁵⁴ Binding and invasion of HT-29 cells was assessed for *E. coli* (strain Nissle 1917), *K. pneumoniae* (strain isolated from MLNs of MyD88^{ΔIEC} mice) and *S. typhimurium* (strain SL1344) as described. Approximately 10^6 CFU were added to monolayers of HT-29 cells at an MOI of 10:1. After 2 h at 37°C, cell monolayers were washed and lysed with 1% Triton X-100 to enumerate total cell-associated bacteria. Parallel co-cultures of bacteria and HT-29 cells were treated with gentamicin ($100 \mu\text{g ml}^{-1}$) for 90 min to kill extracellular bacteria, washed, and cultured to enumerate intracellular bacteria. Membrane-associated bacteria were calculated by subtracting internalized CFU from total cell-associated CFU.

Analysis of the fecal microbiota. Fecal samples were collected from littermate MyD88^{Flox} and MyD88^{ΔIEC} mice, and frozen samples of fecal DNA (extracted with the QIAamp DNA stool kit, Qiagen, Germantown, MD) were shipped to Second Genome (San Bruno, CA) for analysis of microbial richness and diversity by the PhyloChip microarray method²¹ (www.secondgenome.com). Bacterial 16S rRNA genes were amplified by PCR using the forward primer 5'-AGRGTTCGATCMTGGCTCAG-3' and reverse primer 5'-GGTACCTTGTTACGACTT-3', fragmented, biotin labeled, and hybridized to the PhyloChip Array, version G3. Stained arrays were scanned with a GeneArray scanner (Affymetrix, Inc., Santa Clara, CA), and analyzed by Affymetrix software (GeneChip Microarray Analysis Suite). All profiles are intercompared in a pair-wise fashion to determine a UniFrac distance metric, which utilizes the phylogenetic distance between OTUs to determine the dissimilarity between communities.⁵⁶ For weighted UniFrac (WuniFrac), both the abundance and dissimilarities among OTUs were considered. The phylogenetic tree shown in **Figure 3** was generated using the "Interactive Tree of Life" software tool.⁵⁷ Based on 14,193 OTUs present in at least one of the mice, a Welch *t*-test was performed across the two genotypes using abundance metrics. PCA was used to position points on two-dimensional ordination plots based on dissimilarity values (**Figure 4a**). Significant OTUs whose abundance characterizes each mouse genotype (**Figure 4b**) were identified by Prediction Analysis for Microarrays, which utilizes a nearest shrunken centroid method.²²

Statistical analyses. Statistical differences in mRNA abundance, CFU, and disease activity were determined by analysis of variance and Fisher's protected least significant difference test. PCA was used for multifactorial clustering of gene expression data (see **Supplementary Methods** and **Supplementary Figure S5** online for details). A one-tailed Student's *t*-test was used to compare transmucosal electrical resistance values for colon tissues. Permutational multivariate analysis of variance was utilized to compare the fecal microbiota between mouse genotypes, based on distance matrices of OTUs.

SUPPLEMENTARY MATERIAL is linked to the online version of the paper at <http://www.nature.com/mi>

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DISCLOSURE

The authors declared no conflict of interest.

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