The gastrointestinal tract is a major site of host invasion by pathogenic organisms. Indeed, even during periods of health, the gut lumen is the site of a large population of bacteria, which under normal conditions contribute to the viability of the host (1, 2). However, this balanced state of coexistence is not autonomous but actively maintained by local protective mechanisms such as defensins (3) and ongoing adaptive immune responses that help ensure that the commensal bacteria remain resident on the luminal side of the gut lining (4). Defects in these protective responses, for example, sIgA deficiency, can result in breach of this barrier, with potential consequences ranging from septicemia to inflammatory bowel disease (5).

Both T-dependent and T-independent B cell antibody production contribute to ongoing anti-commensal immunity as well as to acute responses against pathogenic organisms. Antigen associated with DCs plays a key role in the well-recognized activation of T cells that can provide help for B cells, but also in the direct presentation of antigen to B cells (5–9). Thus, the mechanisms by which gut-localized DCs acquire antigenic information are of significance in both the maintenance of homeostasis and in the development of protective immunity to pathogens.

Specialized cells (M cells) situated in the dome region of Peyer’s patches are a major route for the transfer of antigen, including intact bacteria and viruses, from the gut lumen to underlying DCs (10, 11). Recent studies suggest that M cells also exist in villi where they may contribute to the transfer of organisms from bowel lumen to lamina propria DCs.
(12, 13). Under nonpathogenic conditions, these antigen-laden DCs can interact with associated T and B lymphocytes to generate a noninflammatory immune response biased through local signals such as TGFβ (14, 15) and in humans, TSLP (16). However, during noncommensal bacterial invasion, signals provided directly or indirectly by these organisms can alter the differentiation of freshly recruited DCs and deviate the immune response to a more inflammatory character (16).

In the past few years, another mode of antigen uptake within the gastrointestinal tract has been reported. In vitro studies using a model epithelial cell line first revealed the capacity of DCs on the basolateral (ablumenal) side of the epithelium to extend processes across the tight junctions between these cells and capture bacteria from the apical (lumenal) side (17, 18). Remarkably, this extension occurred without compromising the integrity of the epithelial barrier, apparently as a result of the creation of a tight junction-like structure between the dendrites and the contiguous epithelial cells. Immunohistochemical studies with gut tissue provided initial limited evidence that such DC extensions existed in vivo, a result confirmed and extended by more recent high resolution static imaging studies (13, 19). These latter papers used fractalkine receptor (CX3CR1)-driven expression of enhanced GFP (EGFP) to visualize the gut DC population and suggested that CX3CL1 may play a role in guiding extension of this DC subset across the epithelium. These experiments also showed that the EGFP-expressing DCs acquired microorganisms from the gut lumen when fractalkine signaling was present, although the study by Vallon-Eberhard et al. (13) did not find the extensions to be required for such trans-epithelial transport.

In both of these previous analyses, exposure of the small bowel luminal surface to bacterial or fungal products was shown to increase the extent of trans-epithelial DC extension, but no information was obtained on the mechanism of this induced response. Likewise, the question of whether the steady-state level of DC trans-epithelial extension is a constitutive activity of the DCs or sustained by stimuli from commensal flora has not been addressed, although the addition of commensal organisms to the gut lumen under defined conditions has been shown to increase the number of such extensions (19). Here, we explore this issue of regulated trans-epithelial extension of processes by lamina propria DCs using dynamic explant and intravital imaging methods to visualize these events under a variety of experimental conditions. We find that these DC-derived processes, which often have an unusual spherical (“balloon body”) shape and whose frequency varies along the small bowel length, are generated in large measure in response to MyD88-dependent signaling by a subset of Toll-like receptors (TLR4) expressed by non-hematopoietic (epithelial) cells. Our findings strongly support the previously proposed model of gastrointestinal lumen sampling by DCs and provide new insight into the location and structural characteristics of the trans-epithelial extensions as well as initial characterization of the signals inciting this sampling of the gut luminal flora.

**RESULTS**

**Differences in the frequency of lamina propria DC extensions along the length of individual villi and of the small bowel in normal mice**

Two high resolution microscopic studies have documented the presence of abundant trans-epithelial DC extensions in the small bowel of mice on the C57BL/6 background (13, 19). To determine the optimal site(s) for dynamic imaging of such extensions, we first conducted a similar static imaging analysis using explants from distinct subregions of the small bowel. Previous studies examined DCs marked by EGFP expression under the control of the CX3CR1 gene regulatory region. Here, we studied small bowel segments from mice whose DCs expressed a fluorescent MHC class II molecule in which the Aβ chain is covalently linked to EGFP as a consequence of gene targeting (MHC CII-EGFP; reference 20) or from mice whose DCs expressed cytoplasmic EGFP under control of the CD11c promoter-enhancer region (CD11c-EGFP; reference 21). In the case of the tissues from the Aβ gene-targeted animals, the identity of the fluorescent cells as DCs was confirmed using RAG-2−/− animals that lack B cells and by FACS analysis of dissociated tissue (Fig. S1, available at http://www.jem.org/cgi/content/full/jem.20061884/DC1).

A representative three-dimensional reconstruction of a small bowel imaging dataset is shown in Fig. 1 A and Video S1, which is available at http://www.jem.org/cgi/content/full/jem.20061884/DC1. In agreement with previous findings, each villus in this image from the ileum 30 cm distal to the stomach had several DC extensions visible on the luminal side of the epithelium. Two features were notable in these imaging data. First, the extensions were more frequent in the apical region of the villi and rarer near the basal regions close to crypts (Fig. 1 B and Video S2), although no difference in DC numbers was detected along the length of the villi. Second, the extensions had an unusual shape, most often being nearly spherical rather than finger-like (Fig. 1, A and C), giving rise to what we call balloon bodies (BBs). This shape was seen using several distinct surgical preparations of the bowel segment and with various media bathing the tissue during imaging (unpublished data), but because the majority of the residual bowel contents and mucus must be removed for successful imaging in all cases, we cannot be certain that this shape is adopted by DC extensions under truly physiological conditions (see Discussion). Globular structures were also noted in at least a subset of the trans-epithelial extensions of CX3CR1-EGFP cells imaged by others (13, 19). Similar numbers, distribution, and shapes of DC trans-epithelial extensions were seen using CD11c-EGFP or MHC CII-EGFP mice. The common findings using both of these reporter animals make it unlikely that the extensions originate from mucus macrophages that have been reported to be CD11c+ (22).

When the frequency of such trans-epithelial DC extensions was measured along the length of the small bowel, clear regional differences were observed (Fig. 2 A). The extensions were present in substantial number from the jejunum through most of the ileum, with a slightly greater frequency in the...
proximal portion of the latter small bowel segment. However, villi in the terminal 5 cm of the ileum near the cecal region had a very low number of DC extensions in unmanipulated healthy animals. These observations differ from those reported previously using mice expressing EGFP under the control of the CX3CR1 gene regulatory region (13, 19), in which the terminal ileum was found to have the highest density of visualized extensions. Both these earlier reports and ours involve mice on the same genetic background (C57BL/6), making this an unlikely source of the difference. The major distinction between the studies is the method for detecting DCs involving EGFP expression by only the CX3CR1+ subset in the case of the previous reports and a broader analysis involving labeling of all CD11c+ or MHC class II+ DCs in the present case. The following sections provide evidence that analysis of DC subsets differentially marked by the various reporter constructs may account for these contrasting observations.

Small bowel bacterial content affects the frequency of trans-epithelial DC extension

The variation in the frequency of DC extension both along a villus and the length of the small bowel could reflect either intrinsic properties of the tissue or the impact of exogenous

**Figure 1.** DCs extend through the apical epithelium of villi. 

(A) Static two-photon microscopy image of the villi in the peri-cecal region of a CD11c-EGFP mouse given noninvasive Salmonella orally 3 h before imaging (top). The image is a three-dimensional reconstruction of 65 serial optical sections (1.5 μm/section) comprising a total depth of 97.5 μm. Extensions of DCs (green) into the intestinal lumen can be observed. The surface of the villi was outlined by SNARF-1 staining of the epithelial cells (red). Bar, 25 μm. Boxed region in the top panel is shown in a higher magnification in the bottom panel to better illustrate how DC extensions (green) protrude through the intestinal epithelium (red) into the luminal cavity. The arrows indicate examples of spherical DC extensions into the intestinal lumen (also see Video S1). (B) A representative static image of an ileal villus from an MHC CII-EGFP mouse stained with SNARF-1 to highlight the epithelial cells (red) shows the preferential location of DC extensions (green) near the apical region of the villus (white arrows). The villus was arbitrarily subdivided in 10 equal compartments from the bottom to the apical region (z = 0 to z = 100), and the average number of DC extensions in each compartment is displayed in the histogram. Values are expressed as the number of DC extensions per villus in the indicated section (mean ± SE). The top regions (z = 70–100; open bars) show a significant difference in the number of DC extensions as compared with the middle and lower parts (z = 40–70; striped bars) of the villi. Also see Video S 2. (C) A single optical z-slice of a villus from the ileum of an MHC CII-EGFP mouse. Epithelial cells are stained with SNARF-1 (red), and the cell nuclei are stained with Hoechst 33342 (blue). The higher magnification in the bottom panel demonstrates a DC extending an MHC CII+ (green) dendrite through the epithelium. Once reaching the intestinal lumen, the DC dendrite assumes the characteristic shape of a sphere (white arrow).
stimuli that actively induce the extension response. A likely source of such exogenous stimuli is the commensal bacteria of the gut, a possibility we examined by treating mice with broad-spectrum antibiotics for 5 d to reduce the bacterial load and then analyzing the distribution and number of DC extensions. This treatment resulted in a significant decrease in the frequency of DC extensions in the proximal regions of the small bowel where they were the most numerous in normal animals (Fig. 2 B). DC protrusions were not completely abrogated, possibly due to the residual bacterial population remaining after antibiotic treatment. In agreement with previous data implicating microbial stimuli in induction of DC extensions, an additional bacterial load consisting of live Salmonella administered by gavage led to the appearance of a substantial number of extensions in the peri-cecal region in which such DC processes were typically rare in normal mice (Fig. 2 A). The induction of BBs can be detected 3 h after bacteria gavage. The process reaches a peak a few hours later (Fig. 3), with the number of extensions then remaining constant for up to 24 h (not depicted). No significant differences in the induction of DC extensions were observed using invasive versus noninvasive strains of Salmonella in such experiments (Fig. 3). Collectively, these data suggest that many, if not most, DC trans-epithelial extensions arise as an active response to microbial products. As for the extensions seen in animals not treated with Salmonella, the extensions in infected mice were most frequent near the tip of the villi (not depicted).

MyD88-dependent signaling by diverse epithelial TLRs induces DC extension

Our observations and those reported previously suggested that pathogen-associated molecular pattern (PAMP) recognition might be directly or indirectly responsible for triggering DCs to extend processes across the epithelial cell layer. Both epithelial cells and DCs express TLRs that could mediate signaling in response to such material (16, 23, 24). Oral administration of peptidoglycan (PG), LPS, or CpGs (TLR2, TLR4, and TLR9 ligands, respectively) was sufficient to induce DC protrusion in the terminal ileum, whereas the TLR3 ligand polyinosine-polycytidylic acid (poly(I:C); reference 25) and the TLR5 ligand flagellin (26) failed to do so (Fig. 4). These findings are in accord with evidence that TLR3 is not expressed on intestinal epithelial cells and that TLR5 is expressed at a very low level in the terminal part of the small intestine and may be predominantly present in lamina propria DCs (27, 28).

To study further the role of TLR signaling in triggering DC luminal protrusions, we constructed three different bone marrow chimeras. The first was prepared using donor bone marrow from mice expressing CD11c-EGFP and recipients that were either wild-type or defective in TLR4 expression (29, 30). The second and the third chimeras were constructed using donor bone marrow from mice expressing MHC CII-EGFP (20) and recipients that were either defective in TLR2 or MyD88 expression (31, 32). After reconstitution, small bowel tissue from these mice was examined for DC extensions in the absence or presence of added TLR ligands.

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Hosts with and without epithelial cell expression of either TLR2 or TLR4 had similar numbers of DC extensions in the proximal bowel regions that normally contained such processes. However, hosts lacking epithelial TLR4 did not show the induced appearance of such extensions in the peri-cecal region after administration of the TLR4 ligand LPS, even though the DCs in these animals expressed the receptor for LPS, whereas there was a significant increase in the number of such processes in control chimeras expressing TLR4 in nonhematopoietic cells (Fig. 4). Similarly, hosts lacking epithelial TLR2 did not show DC extensions in response to the TLR2 ligand PG in this segment of the small bowel, whereas control chimeras did (Fig. 4). Trans-epithelial projections from the DCs could be induced in both of these chimeras if noninvasive Salmonella organisms were administered, consistent with the diversity of potential TLR ligands produced by these bacteria and the ability of each of several TLR ligands to induce extensions when added individually (Fig. 4). Hosts lacking epithelial expression of MyD88 did not show DC extensions in response to Salmonella and had markedly fewer extensions in the absence of TLR ligand instillation or Salmonella infection (Fig. 4). These latter results suggest a possible minor role for the MyD88-independent pathway of TLR signaling (33, 34) or a very low baseline level of protrusion unrelated to TLR stimuli (Fig. 3). Overall, these results indicate that most DC extensions across the small bowel epithelium are mediated by the epithelial cells themselves and that distinct TLRs can independently mediate this effect upon ligand exposure.

Using CX3CR1-EGFP homozygous mice deficient in CX3CR1 expression, we confirmed that there is no increase in the number of DCs sending projections into the terminal ileum upon Salmonella infection. Surprisingly, in the proximal ileum of these same mice, we detected a significant increase in DC projections (Fig. S2, available at http://www.jem.org/cgi/content/full/jem.20061884/DC1). These results indicate that DCs lacking CX3CR1 are nonetheless able to respond to signals evoked by inflammatory stimuli by sending processes across the bowel epithelium.

To study the role of CX3CL1 in DC protrusions, we used quantitative PCR to measure CX3CL1 mRNA levels in the small bowel tissue of mice given noninvasive Salmonella orally. No significant differences in the level of CX3CL1 mRNA were detected in the terminal ileum of infected as compared with uninfected mice, even though exposure to this bacterium evoked a strong DC extension response in this region of the small bowel as assessed using CD11C-EGFP or MHC CII-EGFP animals (Fig. S3, available at http://www.jem.org/cgi/content/full/jem.20061884/DC1, and Figs. 3 and 4). Given our finding that CX3CL1 mRNA levels did not change upon Salmonella infection, we examined whether any other chemokines known to affect DC behavior in the gut might be produced in a MyD88-dependent fashion in response to this stimulus. CCL20 is known to be produced by the intestinal epithelium in response to TLR stimuli (35). We therefore measured the level of mRNA for this chemokine in response to noninvasive Salmonella infection. An 18-fold increase in the level of CCL20 mRNA was measured in the terminal ileum of wild-type infected mice as compared with untreated mice (Fig. S3). MyD88-deficient mice did not show this increase in chemokine mRNA, in accord with the preceding data showing that such animals largely lacked trans-epithelial DC extensions and that these animals extensions could not be induced by TLR ligands or Salmonella.

Collectively, these data suggest that CX3CL1 is not directly involved in actively induced DC protrusion across the intestinal epithelium in the terminal ileum and that DCs marked using CX3CR1-EGFP expression are either distinct...
from or a subpopulation of those marked by CD11c-EGFP or MHC CII-EGFP labeling, potentially accounting for the different distribution of trans-epithelial extensions seen in previous studies as compared with the present investigation.

**Dynamic visualization of the DC extension process**

The demonstration that DC extensions could be actively induced by TLR signaling provided a method for capturing the process using four-dimensional (space and time) imaging methods. Explants from the peri-cecal small bowel region of animals expressing MHC CII-EGFP were imaged for up to 4 h using two-photon microscopy, using the red fluorescent dye carboxylic acid acetate succinimidyl ester (SNARF)-1 to counterstain the epithelial cells. Within 3 h of exposure of these explants to LPS or bacteria, the apical regions of villi in the peri-cecal region began to show the active emergence of DC processes across the epithelial layer. Although these were occasionally finger-like in appearance (Fig. 5 A and Video S3), which is available at http://www.jem.org/cgi/content/full/jem.20061884/DC1), balloon-shaped structures dominated (Fig. 5 B and Videos S4 and S5). Once extended across the epithelium, the DC processes typically remained present for 10–40 min before being withdrawn.

In vitro data indicated that DC extension across a reconstructed epithelial cell layer occurred without disruption of the tight junction (17, 18). To examine if this was the case using normal tissue, fluorescent dextran was added to the medium bathing the explants. No accumulation of the exogenous dextran was seen in the lamina propria or in the DCs themselves (Fig. 6 and Video S5). This is consistent with the maintenance of the epithelial junctions and with limited macropinocytosis or micropinocytosis by the DC extensions.

**Bacterial interaction with DC extensions**

Previous work using in vitro models, tissue explants, and closed loop preparations demonstrated that DCs can capture...
bacteria across an epithelial barrier (13, 17, 19). We confirmed these data using the previously described experimental model (17) and static imaging (unpublished data), and then conducted four-dimensional imaging in an attempt to characterize this process dynamically. Using both explant and intravital methods, it was possible to obtain rare images of DC BBs interacting with noninvasive Salmonella (36, 37) in the luminal space (Fig. 7 A and Video S6, which is available at http://www.jem.org/cgi/content/full/jem.20061884/DC1) and BBs with a captured red Salmonella inside (Fig. 7 B and Video S7). However, the infrequent nature of these events (the total accumulation of noninvasive Salmonella after gavage was never >10–20 organisms per villus per 24 h), as well as rapid bleaching of the captured organisms, precluded direct visualization of binding and withdrawal of bacteria-containing processes into the lamina propria.

DISCUSSION

In this paper, we have used dynamic explant and intravital two-photon imaging to study trans-epithelial DC extension into the small bowel. Our data confirm and extend previous reports on the capacity of lamina propria DCs to send processes across the columnar epithelial layer without disruption of the tight junction between these cells and to interact with luminal microbial flora (13, 17, 19). Most of the DC extensions showed an unusual spherical shape, and their frequency differed markedly along the length of the small bowel. In the more proximal small bowel segments in which the DC processes were numerous in normal uninfected animals, such extension appeared to reflect an active response to local commensal flora and bacterial products. The vast majority of such responses to endogenous bacteria and those occurring in the presence of exogenous species such as Salmonella involved MyD88-dependent signaling evoked by interaction of any of several epithelial TLRs with cognate microbial products.

Two previous studies reported the distribution of small bowel trans-epithelial extensions of lamina propria DCs, using static imaging based on expression of EGFP under the control of the CX3CR1 regulatory region (13, 19). In these reports, the highest frequency of such extensions was observed in the terminal ileum in mice of the C57BL/6 background, which were also used here. In marked contrast, we found that with a conventional gut flora, substantial numbers of such processes are present in the proximal ileum and in the jejunum, with a concentration toward the apical region of the villi, but that only small numbers of such extensions are present in the terminal few centimeters of the ileum (the peri-cecal region) of these same mice. We obtained a similar result using two different reporter strains (MHC CII-EGFP and CD11c-EGFP). Furthermore, in contrast to what was observed by Niess et al. (19) using CX3CR1-EGFP+/+ mice that are unresponsive to CX3CL1 (38), we were able to detect the presence in these mutant animals of trans-epithelial dendrites, but only in the proximal ileum of Salmonella-treated mice. At present, we do not have a definitive explanation for the difference between our findings and the results of these two previous studies. However, it may be important that our work involved imaging based on EGFP expression by what may be a much larger and potentially distinct fraction of DCs, based on control of the reporter by the CD11c (21) or MHC class II (20) gene regulatory regions. Given the absence of a measurable change in CX3CL1 mRNA expression in response to bacterial stimuli that potently induce DC extensions in the terminal ileum, and the induction of extensions in the proximal bowel of animals lacking CX3CR1 expression, these findings suggest that induced up-regulation of CX3CL1 does not play a direct role in the DC extension process, but that the absence of CX3CR1 may change the characteristics of the DCs resident in intestinal epithelium, perhaps altering the adhesive properties of these cells (39).

A primary focus of the present work was on control of the extension process. Treatment with broad-spectrum antibiotics to reduce the load of commensal organisms reduced the number of DC extensions seen in the jejunum and proximal ileum where the number of extensions was high in untreated, uninfected animals. This result suggested that active responses to microbial products contribute to the DC protrusion process, and in accord with this notion, the introduction of viable invasive or noninvasive Salmonella organisms induced DC trans-epithelial extensions in the terminal ileum (peri-cecal region) where these processes were rare in normal animals. One possible explanation for the low frequency of extensions in the pre-cecal region of normal mice is a higher level of defensin production in this site. This suggestion is in accord with the variation in protrusion frequency we observed along the length of an individual villus. It was low.

Figure 6. Epithelial cell continuity is preserved during DC extension. Three sequential images of a villus in an ileal explant from an MHC CII-EGFP mouse given Salmonella orally and immersed in buffer containing TRITC-dextran showing the exclusion of the tracer from the basolateral surfaces of the epithelial layer (also see Video S5).
near the crypts containing Paneth cells that produce high levels of defensins (40) and greater toward the apex where the antimicrobial mediators would be less abundant. Presumably, the high dose of bacteria administered was able to overcome local defense mechanisms that may otherwise limit the local availability of microbial stimuli able to induce DC extensions.

The evidence that DC protrusion could be influenced by bacterial load suggested a possible role in this process for signaling through receptors sensitive to microbial products. TLRs were the most likely candidates, and various TLRs are expressed on the apical surface of gut epithelium (41). In agreement with this notion, individual TLR ligands such as LPS (TLR4), PG (TLR2), or CpGs (TLR9) induced a substantial increase in the frequency of such extensions in the peri-cecal region, as did Salmonella. The TLR3 ligand poly(I:C) was not able to induce a change in DC extension frequency. This is consistent with the observation that TLR3 is not expressed by the intestinal epithelium. Intriguingly, flagellin, whose receptor is TLR5, was also unable to cause this response. This finding is in accord with recent studies that show a very low level or no expression of TLR5 by the epithelium of the ileum and with the ability of flagellin to induce a humoral response only after the disruption of epithelial continuity (27, 28, 42). Collectively, these data indicate that several but not all luminal PAMPs can stimulate the intestinal epithelium to induce an extension response from lamina propria DCs.

Experiments using radiation chimeras showed that microbial products required signaling via TLRs expressed by nonhematopoietic (epithelial) cells to evoke the DC response, that engagement of any of several TLRs was sufficient, and that the signaling process was largely MyD88 dependent. Indeed, in nonchimeric MyD88-deficient mice, the number of DC extensions seen in the absence of added bacteria or PAMPS was reduced almost to zero. We have not yet been able to study directly the reciprocal issue of whether TLR signaling by the lamina propria DCs is also required for the extension process. However, the fact that lamina propria DCs have been reported not to express TLR4 (27), our observations that LPS alone can induce DC extensions, and the finding that bone marrow chimeras with TLR4 expression only on hematopoietic cells do not show DC extension in response to LPS suggest that the TLR signaling by epithelial cells is necessary and sufficient to induce the extension phenomenon.

The observation of robust MyD88-dependent induction of CCL20 transcription in the ilea of animals given noninvasive Salmonella provides further evidence for an active epithelial cell response to TLR engagement. Whether this chemokine plays any role in promoting DC extension is an open question at the moment. We have not detected an increased number of DCs in the terminal ileum at early times after Salmonella administration even though CCL20 has been reported to recruit DC precursors to the small bowel (14). Therefore, it is unlikely that CCL20 contributes to an acute increase in the number of DC extensions by enlarging the pool of DCs present in the lamina propria. A possible role for CCL20-dependent cell shape changes has not yet been ruled in or out by direct experiment.

In agreement with previous in vitro and static imaging studies (13, 17, 19), we find that DC extensions are able to interact with intralumenal microbial material. Our live imaging data clearly show the interaction of DC luminal BBs with...
noninvasive labeled *Salmonella* (36, 37) for several minutes and the presence of bacteria in subepithelial Dcs. At this time, we have no evidence that DC sampling is discriminatory. Both inert beads and either invasive or noninvasive bacteria are acquired well by lamina propria Dcs based on analysis of static images, and both can be seen associating with DC extensions in live images (unpublished data). But it would be surprising if scavenger receptors, FcR, and C-type lectins did not define the preferred targets of the capture process (43). We have not yet been able to follow a DC with an internal microbial load leaving its sub-epithelial position and entering a lymphatic for migration to a local lymph node (44), migrating to a nearby Payer’s patch to initiate a response in this site (45–48), or interacting with a memory T cell within the lamina propria. New methods for maintaining animal anesthesia for more prolonged periods, together with bacteria labeled with nonbleaching probes such as quantum dots (49), may allow both the capture process and the subsequent fate of the loaded DC to be visualized.

The spherical shape of the DC extensions (BBs) was unexpected. We cannot at present rule out that this is an artifact of the imaging method, perhaps due to removal of the mucus layer that typically covers the epithelial cell apical surface and that might constrain the shape of the DC protrusion. In a buffer-filled lumen, membranes moving past the end of the tight junction and lacking support from F-actin (unpublished data) may simply take the energetically most favorable shape, that of a sphere.

A characteristic feature of the immune system is the dynamic behavior of its constituent cells. The development of methods for high resolution intravital imaging of lymphoid (50, 51) and other tissue sites (52–55) has begun to reveal how immune cells move, interact, and respond in situ, adding to our understanding of normal and pathological immune events. Our visualization here of active DC extension into the small bowel lumen and the description of the first steps in the signaling cascade controlling this process provide a clear example of the utility of this new technology in adding to the insights already generated from more conventional analyses of this important host–pathogen interface.

**MATERIALS AND METHODS**

**Animals.** CD11c-EGFP transgenic mice (21), CX3CR1-EGFP mice (38), TLR2-deficient mice (31), and TLR4-deficient mice (29, 30) were obtained from The Jackson Laboratory. MHC CII-EGFP knock-in mice (line 270) on a RAG-2−/− background; reference 20) were produced at Taconic Farms. TLR4−/− or MyD88-deficient recipient mice were sublethally irradiated with 900 rads. On the same day, 5 × 10⁴ bone marrow cells from CD11c-EGFP or MHC CII-EGFP mice were harvested and infused into a TLR4−/− or into TLR2−/− or MyD88-deficient recipient mice via the tail vein, respectively, to create CD11c-EGFP/TLR4−/−, MHC CII-EGFP/TLR2−/−, or MHC CII-EGFP/MyD88−/− bone marrow chimeras. These mice were kept in isolators and provided neomycin-containing water until use 8–10 wk later.

**Reagents.** RPMI 1640 medium without phenol red was purchased from Biosource International. LPS (*Escherichia coli* serotype 026:B6) and TRITC-dextran (160,000 mw) were obtained from Sigma-Aldrich. Poly(I:C) was obtained from GE Healthcare. CpGs (OCTAGACTGTTAAGT and TCACGTTGA; reference 56) were synthesized at the Facility for Biotechnology Resources at the NIH. Fluorescent beads (Fluorobrite polychromatic red; 1-μm microspheres) were obtained from Polysciences, Inc. Anti-mouse CD11c mAb labeled with APC was purchased from BD Biosciences.

**Bacterial strains and oral administration.** Unless otherwise stated, all in vivo experiments involved the administration of attenuated *Salmonella typhimurium* deficient in both the SP-1 (lnvA) and SP-2 (saa1) genes, conferring the invasive phenotype and the capacity for intraphagosomal survival, respectively (36, 37). In selected experiments, the invasive *S. typhimurium* strain SL1344 (57) was also used. A recombinant red fluorescent variant of each strain was generated by expressing the red fluorescent protein DiRed under the transcriptional control of the LacZ promoter. These bacterial strains were grown as described previously (58). Preparation and feeding of mice with bacteria was performed with 200 μl (10%, wt/vol) NaHCO₃ given 10 min before 200 μl of bacterial suspension (2 × 10⁷ bacteria in PBS) or 1 mg/ml LPS or 1 mg/ml PG. Mice were deprived of food for 4 h before and 1 h after oral treatment, which was performed using an animal feeding needle.

**Antibiotic treatment.** For experiments involving the effect of normal gut flora on luminal DC projections, we included 5 mg/ml ampicillin, 3 mg/ml kanamycin, and 5 mg/ml neomycin in the drinking water for 5 d before analysis.

**Real-time quantitative RT-PCR.** Ileal samples were collected 2 cm above the cecum, the luminal content was carefully washed out, and total RNA was extracted using the RNeasy Mini kit Protocol (QIAGEN). cDNA was synthesized from total RNA using the SuperScript III First-Strand Synthesis System for RT-PCR (Invitrogen) according to the manufacturer’s protocol. Expression levels of CCL20, CX3CL1, and GAPDH mRNA were measured by quantitative PCR based on the manufacturer’s recommended protocol with the TaqMan Universal PCR Master Mix, followed by analysis on an ABI7900 (Applied Biosystems). Triplicate reactions were performed for each cDNA preparation. Data were expressed as 2ΔΔCT relative to GAPDH levels in each sample.

**Explant preparation.** Mice were deprived of food for 4 h before and, in the case of administration of bacteria or LPS, 1 h after these procedures before being killed by CO₂ asphyxiation. The intestine was then removed surgically, and the region of interest was immediately everted and washed twice with PBS at 37°C. The epithelium was stained using 20 μM SNARF or 100 ng/ml Cell Tracker Blue (both from Invitrogen) for 3 min at 37°C in PBS. The everted intestinal segment was then immobilized on the surface of a Petri dish using Vetbond epoxy glue (3M Animal Care Products) and a mechanical support, and the entire preparation was submerged with 20 ml of PBS, and then digested in RPMI 1640 containing 0.15 mg/ml type II collagenase ( Worthington) and 0.1 mg/ml DNase (Roche Molecular Biochemicals) for 60 min at 37°C on a shaking platform. The released cells were cultured by 25% Percoll gradient (Sigma-Aldrich), stained with anti–CD11c-APC, and analyzed by flow cytometry.

**Construction of bone marrow chimeras.** 3–5-wk-old TLR4- or TLR2-deficient recipient mice were sublethally irradiated with 900 rads, whereas MyD88-deficient recipient mice were irradiated with 500 rads. On the same day, 5 × 10⁴ bone marrow cells from CD11c-EGFP or MHC CII-EGFP mice were harvested and infused into TLR4-deficient or into TLR2- or MyD88-deficient irradiated recipient mice via the tail vein, respectively, to create CD11c-EGFP/TLR4−/−, MHC CII-EGFP/TLR2−/−, or MHC CII-EGFP/MyD88−/− bone marrow chimeras. These mice were kept in isolators and provided neomycin-containing water until use 8–10 wk later.
pheno1 red–free RPM1 maintained at 37°C within a custom microscopy enclosure fitted with both an environmental heater and a thermal blanket. Experiments conducted with and without the use of medium superfused with a gas mixture of 95% O2/5% CO2 gave similar results.

**Intravital preparation for microscopy.** Mice were anesthetized using nebulized isoflurane (2% induction, 1% maintenance) in 30% O2/68–69% air. The abdomen was shaved, and a vertical incision was performed at the caecal level to reveal the terminal ileum. The terminal 7–8 cm of the small intestine was externalized from the peritoneum while carefully preserving blood flow and lymphatic vessel continuity. The region of interest was then immobilized on the disposable stage by using both a solid support and Vetbond tissue adhesive. The lumen was surgically exposed using a battery-operated cautery (AARON Medical) and washed several times with PBS at 37°C to remove fecal material and mucus from luminal surfaces. The epithelial surface was then stained with 200 μM SNARF diluted in 20 μM PBS at 37°C for 3 min. In some cases nuclei were stained blue using Hoechst 33342. After washing with PBS at 37°C, the exposed region of the bowel was bathed in phenol red–free RPMI 1640 at 37°C, and the whole animal was placed within an enclosed microscope stage fitted with both an environmental heater and a thermal blanket.

**Two-photon microscopy.** Two-photon microscopy was conducted using a Bio-Rad Laboratories/Carl Zeiss MicroImaging, Inc. Radiance 2100MP system equipped with a Nikon 600FN upright microscope, 20× water immersion lens (N.A. 0.95; Olympus), and either a Mira 900 Sa:Ti femtosecond-pulsed laser driven by a 10-W Verdi pump laser (Coherent) or a Chameleon laser (Coherent) with similar specifications, both tuned to 880 nm. The typical pixel size of the image field was 1.09 μm, and the x-y dimensions of the scan area were 560 × 560 μm. The typical optical z-step size was 1.5 μm. Serial x-y images were collected over the entire z depth every 30–60 s, and the entire process was repeated for up to 120 min to obtain a four-dimensional dataset.

**Topographic analysis of BBs.** The terminal part of the ileum from MHC class II–EGFP mice was surgically removed, immediately everted, and stained using a Bio-Rad Laboratories/Carl Zeiss MicroImaging, Inc. Radiance 2100MP system equipped with a Nikon 600FN upright microscope, 20× water immersion lens (N.A. 0.95; Olympus), and either a Mira 900 Sa:Ti femtosecond-pulsed laser driven by a 10-W Verdi pump laser (Coherent) or a Chameleon laser (Coherent) with similar specifications, both tuned to 880 nm. The typical pixel size of the image field was 1.09 μm, and the x-y dimensions of the scan area were 560 × 560 μm. The typical optical z-step size was 1.5 μm. Serial x-y images were collected over the entire z depth every 30–60 s, and the entire process was repeated for up to 120 min to obtain a four-dimensional dataset.

**Epithelial cell continuity detection.** The explants were prepared as described previously. During image acquisition, 200 μl of 10 mg/ml TRITC Dextran in PBS was added to the medium in the Petri dish, and imaging continued.

**Image processing.** Datasets were processed using edge-preserving filters for the green channel and a median filter for the red and blue channels (Bitplane, 4.2; Imaris) to denoise the images. The same software was used to color balance the images, with all manipulations of color and intensity applied equally to an entire image stack, and the resulting files used create two-dimensional maximum intensity projections for the image stack corresponding to each time segment. These projections were then combined using Adobe After Effects to generate video sequences. The rate enhancements of these videos relative to real time are indicated in the legends.

**Statistics.** All data were expressed as average values of experimental data points, and SEs were determined using the calculated standard deviation of a dataset divided by the number of data points within each dataset. p-values for the comparison of two data samples were obtained using the two-tailed Student’s t test.

**Online supplemental material.** Fig. S1 shows that MHC class II–EGFP cells are CD11c+. Fig. S2 shows the presence of DC extensions in the lumen of a CX3CR1+/+ EGFP mouse. Fig. S3 shows CCL20 production by epithelial cells in response to Salmonella. Videos S1 and S2 show reconstructions of DC extensions across the epithelial layer of the terminal ileum. Video S3 shows different shapes of the extensions through the epithelium. Video S4 shows a dendrite reaching the luminal side and acquiring the BB shape. Video S5 shows that epithelial cell continuity is preserved during DC extension. Videos S6 and S7 show DC extension that capture fluorescent Salmonella. The online supplemental material is available at http://www.jem.org/cgi/content/full/jem.20061884/DC1.

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