Antigen presentation in the gut

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ABSTRACT: The induction of T cell responses requires recognition of antigens in association with class II major histocompatibility complex (MHC) proteins and specialized antigen-presenting cells. Candidate antigen-presenting cells in the gut include dendritic cells, macrophages, B lymphocytes, mucosal epithelial cells and endothelial cells. Dendritic cells isolated from normal human colon are potent inducers of primary immune responses and express high levels of class II MHC proteins. Lamina propria macrophages display class II MHC proteins, can present antigens to sensitized T cells, may process antigen and release interleukin-1, and suppress antigen presentation by intestinal dendritic cells in a dose-dependent manner. Class II MHC molecules are normally expressed on small intestinal epithelial cells but not on normal colonic epithelial cells. Suppressor T cells and unresponsive T helper cells in the mucosa appear to mediate systemic T cell tolerance of dietary antigens. In the inflamed colon there is infiltration of the lamina propria by large numbers of monocytes which secrete interleukin-1, and the release of interferon-gamma appears to induce class II protein expression on colonic epithelial cells. Colonic epithelial cells from inflamed bowel may preferentially stimulate T helper cells so that local induction of class II MHC molecules on epithelial cells may amplify and localize secondary immune responses at the site of inflamed mucosa. Taken together, the aberrant expression of class II MHC molecules, breaches in epithelial cell integrity (resulting in exposure to luminal antigens including endotoxin) and the increased numbers of monocytes found in inflamed mucosa suggest that the resulting distortions in antigen presentation contribute to the localization and persistence of the inflammatory lesion in inflammatory bowel disease. Can J Gastroenterol 1990;4(7):267-270 (pour résumé, voir page 268)

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ANTIGEN PRESENTATION INVOLVES the denaturation of native antigen and the display of the processed antigen on the surface of specialized antigen-presenting cells in association with proteins of the class II major histocompatibility complex (MHC). When a resting T lymphocyte comes into contact for the first time with its cognate antigen presented in association with self MHC molecules, T cell activation occurs. By contrast, a secondary immune response is triggered by antigen presented in association with class II MHC molecules to T cells which have previously been exposed to that antigen. Activation requirements for resting and memory T cells differ from those of sensitized or activated T cells (1). The principal roles for accessory cells in the primary immune response comprise the presentation of antigen in association with class II MHC gene products to form a complex on the surface of the antigen-presenting cell that can be recognized by an antigen-specific clonotypic T cell receptor, and the delivery of an unidentified second signal to initiate T cell blast transformation, interleukin-2 secretion and clonal proliferation. By contrast, the
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sensitized T cell does not require the second signal and responds vigorously to the presentation of the antigen-MHC complex alone. The magnitude of the secondary response depends upon the concentration of antigen and the surface density of class II MHC molecules on the antigen-presenting cell.

INTESTINAL ANTIGEN UPTAKE

The mucosal surface of the gut is exposed to a vast array of antigens, including products of food digestion, microorganisms and drugs, yet hypersensitivity reactions to dietary antigens are uncommon. In the small intestine, the preferential site of antigen uptake appears to be Peyer’s patches where the specialized epithelium contains M (for ‘membranous’) cells that transport soluble and particulate antigens intact across the epithelium to the lymphoreticular cells of the dome region of the patch (2). Antigen can also cross the intact mucosal epithelium by entering the lamina propria between epithelial cells, by persorption (the passage of macromolecules or small particles by kneading between epithelial cells), by villous uptake through the extrusion zone when epithelial cells are shed from the villous tip, and by active uptake through the epithelial cell by endocytosis or by a receptor-mediated mechanism (3).

ANTIGEN-PRESENTING CELLS IN THE INTESTINE

Cell types that express class II MHC molecules have the potential to present antigen in some circumstances. The candidate antigen-presenting cells in the gut include dendritic cells, macrophages, B lymphocytes, mucosal epithelial cells and endothelial cells.

Dendritic cells isolated from the human colon have the characteristic morphology and properties of splenic dendritic cells. They are large irregular cells that display typical membrane folds or veils, are nonadherent and nonphagocytic and lack the enzyme markers, surface receptors and antigens that characterize macrophages, T cells and B cells. After antigen exposure, dendritic cells migrate as ‘veiled cells’ in the afferent lymphatics to regional lymph nodes where they become the interdigitating cells of the paracortex. Here they interact with T cells bearing the specific antigen receptor to initiate the immune response. Dendritic cells are specialized antigen-presenting cells which can induce primary T lymphocyte responses to both soluble and particulate antigens and exhibit a unique ability to form clusters with resting T and B cells – a phenomenon important to the induction of the primary response. Dendritic cells from Peyer’s patches, but not from the spleen, preferentially induce polyclonal IgA responses when mixed with B cells from either Peyer’s patch or the spleen. Interleukin-1 appears to amplify the proliferative response to limiting numbers of dendritic cells, which cluster more efficiently with T cells before the onset of mitogenesis (4). Interleukin-1 also matures dendritic cell precursors and promotes their differentiation into functionally competent antigen-presenting cells.

Over 90% of the macrophages harvested from normal human colon express class II proteins, equipping them to present antigen (5). Whether lamina propria macrophages can act as antigen-presenting cells and induce a primary immune response, however, is doubtful because they have been shown to suppress induction of primary immune responses by intestinal dendritic cells, an action mediated by prostaglandin E2 (6). As discussed above, however, intestinal macrophages may be involved in antigen processing, producing interleukin-1 or presenting antigen to sensitized T cells in certain circumstances, but whether macrophages are required for antigen processing or as a source of interleukin-1 is uncertain.

Activated but not resting B cells can also take up antigen using specific surface immunoglobulin receptors. B cells process antigen which can be displayed complexed to class II MHC molecules. Although activated B cells can form clusters with T cells, they are dependent on the leukocyte function antigen-1 molecule for cell-to-cell adhesion (7). Because only activated B cells can sen-
sitize T cells, the role of B cells as antigen-presenting cells appears confined to secondary immune responses.

Class II MHC molecules are also expressed on the basolateral membrane and apical surface of mature epithelial cells of the small intestine and on the epithelium of Peyer's patches, but these molecules are not found on stomach or colon epithelial cells unless they are directly associated with lymphoid follicles (8). Class II MHC molecules may be involved in the absorption of luminal macromolecules across the epithelium in a manner analogous to that suggested for class I MHC type molecules in mediating the uptake of IgG from maternal milk (9,10). Such involvement would suggest a genetically restricted, class II-dependent translocation and presentation of luminal antigen to intra-epithelial lymphocytes or to the cells of the lamina propria.

**INDUCTION OF INTESTINAL IMMUNE RESPONSES**

While the intestinal immune system can mount effective responses to potential pathogens such as viruses, bacteria and parasites, soluble dietary antigens evoke little reaction. Secretory IgA antibody responses occur but usually a state of immunological tolerance develops locally and systemically so that hypersensitivity reactions to dietary proteins are very uncommon. A single exposure to fed antigen can suppress both antibody- (especially IgE class) and cell-mediated immune responses to that specific antigen for up to two years. This state of specific immunological unresponsiveness induced by oral feeding is called 'oral tolerance'. Although the immunoregulation of oral tolerance is complex, T cells appear to play a central role. Suppressor T cells appear to mediate tolerance of systemic cell-mediated immune responses, whereas tolerance of systemic antibody responses has been attributed to T helper cell anergy (11).

**ANTIGEN HANDLING IN IBD**

The inflammatory lesion of IBD allows massive exposure of the intestinal lamina propria to luminal antigens and to highly biologically active products of commensal bacteria such as endotoxin. The destruction of epithelial integrity results in a loss of control of antigen entry across the epithelium so that the intestinal immune system that suppresses local and systemic immune responses to soluble antigens is compromised. The heavy inflammatory cell infiltrate includes a large population of monocytes which turn over rapidly in actively inflamed areas; this has been shown by scans using radiolabelled autologous monocytes (12). These monocytes represent an active population of cells that have the potential to present antigen and produce markedly enhanced amounts of interleukin-1 (13,14), resulting in further monocyte activation and stimulation of T cell cytokine production.

Abnormal expression of class II MHC proteins occurs in regions of active IBD. There is increased expression of class II antigens by epithelial cells in the small intestine and de novo expression in colonic epithelium. Class II expression appears to be induced by interferon-gamma and its expression on small intestinal epithelial cells may be modulated by intra-epithelial lymphocytes (15). The significance of aberrant expression of class II molecules in colonic epithelial cells to antigen-presenting cell and induce abnormal immune responses that contribute to inflammation and tissue injury. Isolated rat enterocytes cultured with sensitized lymph node-derived T cells did not stimulate proliferation in the presence of antigen. Rather, the results indicated that antigen-specific suppression occurred which was blocked by antibodies to class II MHC antigens, suggesting that epithelial cell presentation of antigen to the intra-epithelial lymphocyte presented a possible means of inducing oral tolerance in the normal small intestine (16). Expression of class II MHC molecules on colon epithelial cells during inflammatory responses, however, may trigger aberrant immune responses (17). There is evidence that colonic epithelial cells from active IBD bowel may present antigen and preferentially stimulate T helper cells, providing a further mechanism for amplifying and localizing immune-mediated tissue injury (17). On the other hand, the expression of class II MHC proteins on the epithelial cell may simply reflect the presence of increased levels of interferon-gamma produced by the inflammatory T cell infiltrate. Thus, rather than initiating T cell responses, the aberrant expression of class II MHC determinants would then serve to focus the effector functions of previously sensitized T cells to the site of antigen entry.

Taken together, the breaches in epithelial integrity in IBD that lead to exposure to luminal antigens and bacterial products result in greatly increased numbers of monocytes in the IBD mucosa with resulting enhanced production of interleukin-1, interferon-gamma and aberrant expression of class II MHC molecules on colonic epithelial cells. These distortions of antigen presentation in IBD-affected mucosa may contribute significantly to the localization and persistence of mucosal inflammation.

**REFERENCES**

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