PD-1 Inhibitor-induced Gastroenterocolitis: A Mini Review

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Introduction

Programmed cell death protein 1 (PD-1) is a protein present on the surface of T lymphocytes. PD-1 binds to programmed death ligand 1 (PD-L1) and ligand 2 (PD-L2) on the surface of antigen presenting cells, which prevents cytotoxic T lymphocytes from recognizing self-antigens and thus, serves as an important protein in immune tolerance [1]. Recently, antibodies directed at the PD-1 signaling pathway (nivolumab and pembrolizumab), so-called “immune checkpoint inhibitors”, have been approved by the Food and Drug administration for the treatment of advanced-stage malignancies [2]. Some tumors express PD-L1, which allows tumor cells to evade the T-cell mediated immune response and ultimately proliferate. PD-1 inhibitors prevent the interaction of PD-L1 with PD-1, thereby enabling cytotoxic T lymphocytes to fight tumor cells [3]. Unfortunately, these agents can also have autoimmune-related adverse events termed ‘immune related adverse events (irAEs),’ which have a propensity to affect the gastrointestinal tract, liver, lungs, and endocrine organs [4-7]. In fact, diarrhea is one of the most common side effects reported with PD-1 inhibitor therapy, occurring in up to 20% of patients [7,8]. Two series [9,10] have recently described the histopathologic features of colitis associated with PD-1 inhibitors and herein, we review the histopathologic features of PD-1 inhibitor colitis.

Histopathologic features of PD-1 inhibitor colitis

In the series from Gonzalez et al. [9], seventeen biopsies were from the colon. A majority of the colon specimens demonstrated neutrophilic cryptitis/abscesses (8/17, 47%; Figure 1) and expansion of the lamina propria by a mononuclear infiltrate (13/17, 76%). Other common features included crypt architectural distortion (9/17, 53%) and increased apoptosis (8/17, 47%; Figure 2). Some cases showed features of ischemic colitis (3/17, 18%) or crypt rupture with associated histiocytic aggregates, which occasionally formed granulomata (3/17, 18%). A thickened subepithelial collagen table suggestive of collagenous colitis was seen in only one case (Figure 3) and has been reported once previously in the literature [11].

In the series from Chen et al. [10], two patterns of injury were described: 1) an active colitis with apoptosis pattern and 2) a lymphocytic colitis pattern. The active colitis pattern was the most common (5/8, 63%), which demonstrated active inflammation, neutrophilic crypt abscesses, increased crypt apoptosis, and crypt atrophy/dropout. The lymphocytic colitis pattern was less common (3/8, 37%) and showed increased intraepithelial lymphocytes in the surface epithelium, injury of the surface epithelium, and mononuclear expansion of the lamina propria. Interestingly, changes of chronic colitis mimicking inflammatory bowel disease, including basal plasmacytosis, mononuclear expansion of the lamina propria, and crypt architectural

Figure 1. Colonic mucosa with active inflammation and crypt abscesses in a case of PD-1 inhibitor colitis (H&E, original magnification x100).

Figure 2. Colonic mucosa with active inflammation and crypt abscesses and increased crypt apoptotic bodies (active colitis with apoptosis pattern) in another case of PD-1 inhibitor colitis (H&E, original magnification x200).

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distortion, may be seen in patients who experience recurrent PD-1 inhibitor colitis after restarting therapy, but are typically not present without re-challenge.

The histopathologic differential diagnosis should include several entities, based upon the pattern of injury. The finding of numerous apoptotic bodies raises the possibility of either graft-versus-host disease or mycophenolate mofetil-induced injury [12,13]. Graft-versus-host disease rarely has abundant active inflammation, similar to that seen in PD-1 inhibitor colitis, while mycophenolate mofetil tends to have an inflammatory component that is rich in mucosal eosinophils and not neutrophils [12-14]. Ultimately, clinicopathologic correlation is required to differentiate these entities. If the pattern is active colitis with apoptosis, cytomegalovirus should be entertained, which can be excluded easily with an immunohistochemical stain for cytomegalovirus [9,10]. If the biopsies demonstrate the uncommon ischemic colitis-like pattern, then ischemic colitis from either vascular insufficiency or infection enters the differential diagnosis [9]. Rarely, the subepithelial collagen table may be thickened, mimicking collagenous colitis; however, more commonly, there is a lymphocytic colitis pattern. These findings raise the possibility of microscopic cryptitis and crypt abscesses, and apoptosis [15,16]. It is imperative that pathologists recognize colitis in this setting so that appropriate treatment, including withdrawal of the PD-1 inhibitor and the administration of steroids, may be instituted [17,18].

Conclusion

With the advent of immunotherapy for cancer with "checkpoint inhibitors," it is important for clinicians and pathologists alike to be aware that these agents may induce immune-related adverse events such as colitis. From the two series describing the histopathologic features of PD-1 inhibitor colitis [9,10] the most frequent findings are active inflammation, including crypt abscesses, expansion of the lamina propria, and increased apoptosis. Less common findings are lymphocytic colitis, collagenous colitis, ischemia-like colitis, or crypt rupture with histiocyte aggregates and/or granulomata [9,10]. As suggested by Gonzalez et al. [9], it may be best to refer to the histologic findings as "immunomodulatory gastroenterocolitis" as they are not specific and may be seen with other checkpoint inhibitors, such as the cytotoxic T lymphocyte antigen-4 (CTLA-4) inhibitor ipilimumab, which is well-known to cause colitis with active inflammation, cryptitis and crypt abscesses, and apoptosis [15,16]. It is imperative that pathologists recognize colitis in this setting so that appropriate treatment, including withdrawal of the PD-1 inhibitor and the administration of steroids, may be instituted [17,18].

References


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