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The tumor immune response is in a dynamic balance between antitumor mechanisms, which serve to decrease cancer growth, and the protumor inflammatory response, which increases immune tolerance, cell survival, and proliferation. Hypoxia and expression of HIF-1α and HIF-2α are characteristic features of all solid tumors. HIF signaling serves as a major adaptive mechanism in tumor growth in a hypoxic microenvironment. HIFs represent a critical signaling node in the switch to protumorigenic inflammatory responses through recruitment of protumor immune cells and altered immune cell effector functions to suppress antitumor immune responses and promote tumor growth through direct growth-promoting cytokine production, angiogenesis, and ROS production. Modulating HIF function will be an important mechanism to dampen the tumor-promoting inflammatory response and inhibit cancer growth.

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Hypoxia-inducible factors: a central link between inflammation and cancer

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REVIEW SERIES: HYPOXIA AND INFLAMMATION
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Hypoxia-inducible factors (HIFs) are a family of transcription factors that are activated by low oxygen tension. HIFs play a critical role in various biological processes including stress response, angiogenesis, cell proliferation, and apoptosis. HIFs are composed of two subunits: HIF-α and HIF-β. The HIF-α subunits are oxygen-sensing proteins, whereas the HIF-β subunit is constitutively expressed. In normoxia, HIF-α subunits are degraded by the von Hippel-Lindau (VHL) tumor suppressor/E3 ubiquitin ligase complex, ubiquitin conjugation, and 26S proteasomal enzymes. In hypoxia, HIF-α subunits are stabilized, dimerize with the HIF-β subunit, and translocate to the nucleus to regulate transcription by binding to hypoxia response elements (HREs) in promoters of target genes.

Introduction

The tumor microenvironment is similar to an inflammatory focus, as it consists of a complex milieu of both innate and adaptive immune cells. Hypoxia is a characteristic feature of both tumors and inflammatory foci. Increased metabolic demand from rapid cell turnover, immune cell infiltration, and vascular disruption cause local oxygen tension to decline. The decreased oxygen tension of tumors or inflamed tissue promotes activation of HIFs. HIFs are basic helix-loop-helix-Per-ARNT-Sim–containing transcription factors consisting of a heterodimer of an oxygen-sensitive α subunit (HIF-1α, HIF-2α, and HIF-3α) and a constitutively expressed β subunit (ARNT) (2). HIF-1α is ubiquitously expressed, whereas HIF-2α and HIF-3α expression is largely tissue restricted (3–5). HIF-α subunits are regulated by O2-dependent posttranslational hydroxylation of two specific proline residues by prolyl hydroxylase domain–containing enzymes. In normoxia, HIF hydroxylation leads to association with the von Hippel-Lindau (VHL) tumor suppressor/E3 ubiquitin ligase complex, ubiquitin conjugation, and 26S proteasomal degradation. As oxygen homeostasis is disrupted and O2 concentration declines under inflammatory conditions or in tumors, HIFs are stabilized, dimerize with ARNT, and translocate to the nucleus to regulate transcription by binding to hypoxia response elements (HREs) in promoters of target genes.

In addition to O2-dependent regulation, inflammation and direct HIF regulation are intimately linked. NF-κB, a master transcription factor in the inflammatory response, is a direct transcriptional regulator of Hif1a. In response to NF-κB–activating stimuli such as bacterial lipopolysaccharide (LPS), NF-κB directly increases Hif1a mRNA in macrophages (6). LPS-induced NF-κB can also enhance HIF-1α protein stability by increasing intracellular ferritin, which sequesters the labile iron pool and leads to decreased PHD activity (7). Independently of NF-κB, several cytokines and intermediate metabolites such as succinate can lead to HIF activation (8, 9). In macrophages, IL-4 and IL-13 selectively induce Hif2a mRNA (10). Several studies have demonstrated that cytokine-induced ROS and specifically mitochondrial ROS directly activate HIF (11–13), and recently it was shown that mitochondrial membrane potential increases mitochondrial ROS to modulate HIF activation (14) (Figure 1B).

HIFs are critical drivers of cancer and regulate a wide variety of cellular processes including metabolism, cell cycle progression, angiogenesis, invasion/metastasis, and chemoresistance. Hypoxia and expression of HIF-1α and HIF-2α are characteristic features of all solid tumors. HIF signaling serves as a major adaptive mechanism in tumor growth in a hypoxic microenvironment. HIFs represent a critical signaling node in the switch to protumorigenic inflammatory responses through recruitment of protumor immune cells and altered immune cell effector functions to suppress antitumor immune responses and promote tumor growth through direct growth-promoting cytokine production, angiogenesis, and ROS production. Modulating HIF function will be an important mechanism to dampen the tumor-promoting inflammatory response and inhibit cancer growth.

Inflammation and cancer

It is widely recognized that inflammation plays an essential role in tumororigenesis and tumor progression. This paradigm dates back to Rudolf Virchow, a 19th century pathologist whose observation of leukocytes in tumors led him to hypothesize that tumors originate in sites of chronic inflammation. The discovery of the first oncoprotein v-src in Rous sarcoma virus over a century ago led to decades
for development of hepatocellular carcinoma (HCC). For example, HCV infects more than 100 million people worldwide, and HCC due to chronic hepatitis induced by HCV infection occurs in approximately 1%–5% of infected individuals (24). One of the best-studied cancers from a genetic and inflammatory perspective has been colon cancer. Chronic intestinal inflammation associated with inflammatory bowel disease (IBD), including Crohn’s disease and ulcerative colitis, represents a significant risk factor for the development of colon cancer, termed colitis-associated colon cancer (CAC). More than 1 million Americans suffer from IBD and 12%–20% of patients will develop CAC within 30 years of developing IBD (25).

Chronic inflammation–associated cancers. As Virchow postulated, several chronic inflammatory diseases predispose the development of cancer. For example, Helicobacter pylori, a gram-negative bacterium, infects nearly 50% of the world’s population and is the major causative agent of chronic gastritis (23). Chronic gastritis associated with H. pylori is asymptomatic in the majority of infected individuals; however, this chronic gastritis represents a significant risk factor for the development of gastric cancer, which occurs in 1%–3% of H. pylori infections and is the third leading cancer type worldwide (23). Similarly, chronic viral infections predispose to the development of cancer. Hepatitis B virus (HBV) and hepatitis C virus (HCV) infections are key risk factors for development of hepatocellular carcinoma (HCC). For example, HCV infects more than 100 million people worldwide, and HCC due to chronic hepatitis induced by HCV infection occurs in approximately 1%–5% of infected individuals (24). One of the best-studied cancers from a genetic and inflammatory perspective has been colon cancer. Chronic intestinal inflammation associated with inflammatory bowel disease (IBD), including Crohn’s disease and ulcerative colitis, represents a significant risk factor for the development of colon cancer, termed colitis-associated colon cancer (CAC). More than 1 million Americans suffer from IBD and 12%–20% of patients will develop CAC within 30 years of developing IBD (25). The vast majority of colon cancers develop sporadically and proceed through a step-wise process from adenoma to invasive carcinoma (26). This process includes the acquisition of sequential mutations leading to the loss of adenomatous-polyposis coli (APC) tumor suppressor, activation of the oncogene KRAS, and loss of the TP53 tumor suppressor (27). Interestingly, in colon cancers preceded by chronic inflammation due to IBD, the kinetics of the genetic alterations are different, as TP53 is lost early in disease progression and APC mutations are not as frequently observed compared to sporadic colon cancer (28, 29). These data suggest that inflammation may induce novel mechanisms to drive cellular proliferation and survival, resulting in tumorigenesis.
**Tumor-derived inflammation.** All solid tumors elicit an inflammatory response that is critical in the tumor microenvironment. The tumor-derived inflammatory response is essential for the recruitment of immune cells, tumor cell proliferation, survival, and angiogenesis (30). Tumors initiate these responses through several mechanisms including transcriptional regulation of inflammatory genes by proto-oncogenes (31, 32). It was recently demonstrated that defects in epithelial permeability elicited an inflammatory response through a microbiota-mediated mechanism in colon cancer (33). Below, we illustrate the importance of hypoxia in mediating the tumor-derived inflammatory response.

Pro- and antitumor immune responses. In tumors, there is a mix of both innate and adaptive immune cells with anti- and pro-tumor functions (1). Precise identification of immune cells found in tumor biopsies can serve as prognostic markers for clinical outcomes. Adaptive immune responses have been correlated with positive prognosis. Several distinct T cell subsets reside within the tumor microenvironment, including cytotoxic antitumor CD8+ T cells and subsets of CD4+ T cells with both pro- and antitumor functions. In colon cancer, decreased expression of CD8 as well as the cytotoxic T cell markers granzyme B and CD45RO all correlated with disease recurrence. Furthermore, patients with high intratumoral expression of the T cell marker CD3 had improved disease-free survival compared to patients with low levels of CD3 (34). Recent meta-analysis studies have mapped the prognostic impact of 22 immune cell types on recurrence-free survival by analyzing expression signatures across more than 5,000 tumor samples (35). In general, these studies found a positive correlation between T cells and recurrence-free survival and a negative correlation for several myeloid cell types such as neutrophils and macrophages (35). In ovarian cancer, an increased CD8+/CD4+ T cell ratio was associated with improved survival, whereas increased CD4+ T cells portended poor survival (36). Expression profiles associated with antitumor T cell cytokines (IFN-γ) and effector molecules (granulysin and granzyme B) are also independent prognosticators of decreased early metastasis in colon cancer (37). These associations, however, do not highlight the complexity of the tumor immune environment. Many specific cell types have been associated with better or worse prognosis, and the plasticity of immune cells can confer both pro- and antitumor functions, which we will review here in depth.

**Hypoxia and epithelial-elicted tumoral inflammatory response**

Historically it was believed that the major function of epithelial surfaces, such as the skin or the intestinal epithelium, was to serve as a physical barrier separating the external environment from the underlying immune cells. It is now evident that epithelial surfaces play an active role in innate immunity and shape the underlying immune environment and inflammatory response (38, 39). This is also the case in epithelial-derived cancers. Colon tumor cell secretion of the C-C family chemokine, CCL2, was essential for tumorigenesis through recruitment and activation of protumorigenic myeloid cells (40). Pentraxin 3 (PTX3) is a tumor suppressor that activates complement-mediated antitumor immunity and is epigenetically silenced due to methylation in colon tumors (41). Recent studies in colon cancer have further highlighted the contribution of epithelial and stromal gene expression in patient-derived colon cancer xenograft models (42). In these studies, the epithelial and stromal gene expression was readily delineated by analyzing tumor expression of human or mouse transcripts. Analysis of genes with a greater than 50% difference in expression between the epithelium and the stroma showed that several cytokines and chemokines are directly expressed by the tumor epithelium (42). These results demonstrate that tumor epithelial cells play an active role in regulating the inflammatory response, which may impact tumorigenesis.

Several lines of evidence suggest that intratumoral hypoxia and HIFs play an essential role in sculpting the tumor immune environment in epithelial-derived tumors. For example, in a Kras-driven mouse model of non-small cell lung cancer (NSCLC), loss of HIF-2α increased tumor burden and tumor cell proliferation; however, loss of HIF-1α had no effect on tumorigenesis (43). Tumors lacking HIF-2α also displayed increased infiltration of CD45+ immune cells, specifically Gr-1+ granulocytic cells, suggesting that HIF-2α repression of granulocytic cell infiltration is in part responsible for its antitumor effects (43). Hypoxia and HIF stabilization is also a key feature of pancreatic ductal adenocarcinoma (PDAC) (16). PDAC development occurs in a step-wise manner and is preceded by precursor lesions termed pancreatic intraepithelial neoplasias (PanINs) (44). In contrast with studies showing decreased PanIN progression following pancreatic-specific disruption of HIF-2α (45), deletion of HIF-1α in a murine model of Kras-initiated PDAC significantly enhanced PanIN progression and increased tumor cell proliferation (46). Loss of HIF-1α correlated with increased pancreatic B cell infiltration and antibody-mediated depletion of B cells reversed the increased PanIN progression (46). Hypoxic inflammation is also important in colon tumorigenesis, and both HIF-1α and HIF-2α are overexpressed in colon tumors (16). Using the Apcmin/+ model of intestinal tumorigenesis, intestine-specific disruption of the tumor suppressor Vhl significantly increased colon tumorigenesis and adenoma-to-carcinoma progression (47). The increase in colon tumorigenesis was HIF-2α dependent, as double disruption of Vhl and HIF2α ameliorated the effect (47). HIF-2α-mediated inflammatory responses are essential in colon tumorigenesis. Epithelial HIF-2α regulates expression of the proinflammatory mediator TNF-α (Tnfa) (48). TNF-α has a crucial role in the progression of cancer, and inhibiting TNF-α decreases growth in several mouse models of cancer (49, 50). HIF-2α-induced inflammation was found to be critical to tumor progression, as treatment with the antiinflammatory drug nimesulide significantly reduced HIF-2α-driven colon tumorigenesis. (48). Intestine-specific overexpression of HIF-1α does not enhance tumorigenesis in colon cancer (51). These studies suggest that tumors activate HIFs have the potential to modulate tumor-associated inflammation to regulate tumor growth and progression.

**Hypoxia and immune cell recruitment**

Tumors are highly infiltrated by cells of both the innate and adaptive immune systems. This infiltration is partially mediated by tumor-derived secretion of a host of cytokines and chemokines (Figure 2A). HIF-induced secretion of tumor-derived factors can modulate immune cell recruitment to aid in tumor growth, which we will review in detail by immune cell subtype (Table 1).
T cells. CD4+ T cells can be differentiated into several different helper T (Th) cell types. Th1 and Th2 CD4+ T cells are the classical types of Th cells that play an important role in the inflammatory response to infection and cancer (52). Th1 and Th2 cells promote antitumor immunity by cytokotic lymphocyte activation and humoral-mediated immune responses, respectively (52). In addition to the classic Th1 and Th2 T cell effector populations, other subsets of T cells have been found to have an important role in cancer. For example, Th17 cells are a recently identified subset of IL-17-expressing CD4+ T cells that are highly prevalent in tumors and have a controversial role in tumor progression. IL-6 and TGF-β collaboratively promote Th17 differentiation (53) in a HIF-1α-dependent manner (54). Th17 cells have both anti- and protumor functions. In a melanoma model, Th17 cells showed more potent tumor eradication than Th1 cells (55). However, in other models, Th17 cells promote tumor growth through angiogenic and immunosuppressive effector functions (56, 57).

Immunosuppressive regulatory T cells (Tregs) are frequently increased in cancers (58). Tregs are CD4+ and defined by expression of forkhead box transcription factor 3 (FoxP3). Tregs have been largely shown to promote tumorigenesis through suppression of antitumor CD4+ and CD8+ T cell-mediated immune responses by secreting immunosuppressive molecules such as IL-10 and TGF-β. Removal of Tregs improves antitumor immunity (59–61). Primary tumor hypoxia regulated recruitment of CCR10+CD4+FoxP3+ immunosuppressive Tregs in a model of ovarian cancer, which increased immune tolerance and angiogenesis through VEGF secretion (62). This effect was regulated by hypoxia-induced excretion of the chemokine CCL28, which was dependent upon HIF-1α and to a lesser extent HIF-2α (62).

B cells. The precise role for B cells in tumor progression is controversial, and they may function to enhance or inhibit antitumor immune responses. B cells can be directly cytotoxic to tumors and promote antitumor T cell responses (63). However, B cells have also been shown to inhibit the function of antitumor T cells, as depletion of B cells increases antitumor immunity, suggesting context-specific roles for B cells in antitumor immunity (64). In pancreatic cancer, as detailed above, HIF-1α in epithelial cells decreased tumor growth by attenuating expression of B cell–recruiting chemokines, resulting in decreased B cell infiltration into the tumors (46).

Myeloid-derived suppressor cells. Hypoxia also regulates tumor recruitment of immunosuppressive myeloid cells. Myeloid cells include monocytes/macrophages, neutrophils, eosinophils, basophils, mast cells, and dendritic cells (DCs). During infection or in cancers, immature myeloid cells that are closely related to neutrophils and monocytes can be detected in circulation. These cells, termed myeloid-derived suppressor cells (MDSCs), dampen immune responses to infection and promote tumor growth through suppression of both NK cell- and T cell–mediated antitumor immune responses (65). This tumor-promoting effect is mediated by multiple mechanisms including arginine metabolism via increased expression of arginase 1 (ARG1), which converts the available L-arginine pool to urea and L-ornithine. Additionally, MDSCs mediate nitration of tyrosine residues in the T cell receptor (TCR) and CD8, which decreases the function of these proteins (66–68). The function of MDSCs is well described in the azoxymethane (AOM) and dextran sulfate sodium (DSS) model of CAC. Mice with CXCR2-deficient bone marrow have significantly reduced MDSC homing to colon tumors and decreased colon tumorigenesis (69). Tumor-specific hypoxia increased recruitment of MDSCs.
Table 1. Regulation of immune cells by HIF-1α and HIF-2α

<table>
<thead>
<tr>
<th>Immune cell recruitment</th>
<th>HIF-1α</th>
<th>HIF-2α</th>
</tr>
</thead>
<tbody>
<tr>
<td>TAM</td>
<td>Promotes chemotraction via SDF-1α/CXCL12 expression (82)</td>
<td>Not known</td>
</tr>
<tr>
<td>DC</td>
<td>Not known</td>
<td>Not known</td>
</tr>
<tr>
<td>PMN</td>
<td>Not known</td>
<td>Represses GR-1+ granulocyte recruitment to lung tumors (43)</td>
</tr>
<tr>
<td>MDSC</td>
<td>Increased recruitment to HNSCC through MIF secretion (71)</td>
<td>Increased recruitment to HNSCC through MIF secretion (71)</td>
</tr>
<tr>
<td>Treg</td>
<td>Regulates CCL28 expression in ovarian cancer (62)</td>
<td>Not known</td>
</tr>
<tr>
<td>Th17</td>
<td>Not known</td>
<td>Not known</td>
</tr>
<tr>
<td>CD4</td>
<td>Not known</td>
<td>Not known</td>
</tr>
<tr>
<td>B cell</td>
<td>Repress B cell infiltration into pancreatic tumors (46)</td>
<td>Not known</td>
</tr>
</tbody>
</table>

**Immune cell effector function**

<table>
<thead>
<tr>
<th>HIF-1α</th>
<th>HIF-2α</th>
</tr>
</thead>
<tbody>
<tr>
<td>TAM</td>
<td>T-cell suppression (96)</td>
</tr>
<tr>
<td>DC</td>
<td>Not known</td>
</tr>
<tr>
<td>PMN</td>
<td>Decrease apoptosis (101)</td>
</tr>
<tr>
<td>MDSC</td>
<td>T-cell suppression (98); Pd1 expression (100)</td>
</tr>
<tr>
<td>Treg</td>
<td>Regulates Foxp3 expression (110); essential for immunosuppression</td>
</tr>
<tr>
<td>Th17</td>
<td>Essential for Th17 differentiation (54)</td>
</tr>
<tr>
<td>CD4</td>
<td>Represses cytokine secretion (112); regulates O137 to augment antitumor immunity (114)</td>
</tr>
<tr>
<td>B cell</td>
<td>Essential for B cell maturation (144)</td>
</tr>
</tbody>
</table>

**Tumor-associated macrophages.** The best-characterized immune cells in the tumor microenvironment are tumor-associated macrophages (TAMs). TAMs are highly prevalent in the tumor microenvironment and can be polarized into antitumor M1 or protumor/immunosuppressive M2 phenotypes (72). M2 TAMs regulate tumor angiogenesis and are an important source of VEGF (73). TAMs directly promote tumor growth via direct secretion of cytokines such as IL-6, which induces tumor-cell STAT3 signaling to promote growth and stem cell expansion (74); these cells have been directly linked to tumor invasion and metastasis (75, 76). TAMs inhibit antitumor immune responses through secretion of immunosuppressive cytokines such as IL-10 and TGF-β (77). ARG1 expression in human monocytes and macrophages is controversial and has not been definitively shown in tumors (78). In rodents, however, there is clear evidence showing TAM expression of immunosuppressive ARG1 (79). TAMs reside in largely avascular and hypoxic regions of tumors (80). Tumor hypoxia is a potent driver of TAM recruitment and induces the secretion of chemotactants such as oncostatin M and eotaxin (81). Moreover, tumor HIF-1α directly induces recruitment of monocytes/macrophages through regulation of stromal-derived factor 1α (SDF1α; also known as CXCL12) expression (82). Expression of the SDF1α receptor CXCR4 is also regulated by hypoxia in TAMs (83). TAM polarization into a protumorigenic M2 phenotype can be directly regulated by tumor hypoxia. Tumor-derived lactic acid is induced by hypoxia in a HIF-1α-dependent manner and promotes M2 macrophage polarization and regulates expression of M2 TAM markers ARG1 and VEGF (84). Importantly, blockade of TAM recruitment to hypoxic tumor areas and trapping of TAMs in normoxic tumor microenvironments through loss of the semaphorin 3A receptor NRPI decreased tumor growth through blunted angiogenesis and increased antitumor T cell responses, showing that hypoxia-induced localization of macrophages causes a switch from anti- to protumor phenotypes (85).

**Neutrophils.** A close relationship between neutrophils (polymorphonuclear neutrophils [PMNs]) and tissue hypoxia has recently been shown. The reactive oxygen burst that is critical for neutrophil function can affect local tissue oxygenation (86). Although PMNs are highly prevalent in most solid tumor types, the specific role for PMNs is not completely understood. PMNs are highly plastic and can be differentiated into antitumor (N1) and protumor (N2) phenotypes (87). Protumor PMNs regulate tumor growth through secretion of cytokines, ROS production, generation of matrix-degrading enzymes, and angiogenesis (88). PMNs have also been shown to play an essential role in promoting metastasis in a murine breast cancer model (89). PMNs expressing the hepatocyte growth factor (HGF) receptor, c-MET, were found to be antitumorigenic in mouse models of colon cancer and liver cancer (90). PMNs have recently been demonstrated to reside in hypoxic tumor regions in epithelial uterine tumors and this effect was regulated by hypoxic tumor cell expression of PMN chemotactants such as CXCL5 (91).

**Hypoxic regulation of tumor immune cell function**

Hypoxia is a hallmark of tumors and most infiltrating immune cells function in the hypoxic tumor environment. Immune cell expression of HIF-1α and HIF-2α regulates effector function (Table 1) (92). Tumor hypoxia has an essential role in regulating tumor inflammatory cell functions in addition to regulating immune cell recruitment (Figure 2B). TAMs express both HIF-1α and HIF-2α (16, 93). TAM HIF-2α expression is highly correlated with tumor vascularity and tumor grade (94). Macrophage HIF-2α is critical in regulating macrophage inflammatory cytokine expression following LPS and IFN-γ challenge (95). Importantly, macrophage loss of HIF-2α impaired TAM infiltration of tumors and decreased tumor burden in murine models of HCC and CAC (95). TAM HIF-1α has also been shown to play an important role in TAM-mediated suppression of tumor-associated T cells (96). TAMs cultured under hypoxic conditions exhibited increased suppression of T cells in a HIF-1α-dependent manner without affecting TAM recruitment or polarization (96). Furthermore, tumor hypoxia regulates TAM expression of VEGF, suggesting a role for hypoxic TAMs in angiogenesis (97).
Hypoxia and HIF-1α have an important role in regulating tumor-associated MDSC function. MDSCs cultured with the hypoxia mimetic desferoxamine (DFO) robustly suppress T cell proliferation and loss of HIF-1α decreases MDSC-mediated T cell suppression (98). Hypoxia induces MDSC expression of miR-210 in a HIF-1α-dependent manner and miR-210 promotes MDSC-mediated T cell suppression by increasing ARG1 expression and NO synthesis (99). Notably, HIF-1α increases mRNA expression of the immune checkpoint receptor programmed death ligand-1 (PD1) in MDSCs, which is essential for their T cell immunosuppressive ability (100). MDSCs also have the capacity to differentiate into TAMs and HIF-1α is a critical mediator of this plasticity (98).

The roles of PMN HIF-1α and HIF-2α in the tumor microenvironment are not well understood; however, it has been shown that PMN HIF-1α is an essential PMN survival factor that mediates its effects through an NF-κB–dependent signaling loop (101). Constitutive HIF-2α activation increases PMN inflammatory responses and loss of HIF-2α increases susceptibility to apoptosis (102).

DCs are antigen-presenting cells (APCs) that are central regulators of the adaptive immune response. DCs can sample tumor antigens and activate CD8+ T cell responses and are currently in clinical trials as a vaccination strategy to prime antitumor immune responses (103). DC function is directly regulated by hypoxia and HIFs. Activated DCs increase expression of costimulatory molecules and T cell activation in response to TLR stimulation in a HIF-1α–dependent manner when cultured under hypoxic conditions (104). Alternatively, it has been suggested that hypoxia-treated immature DCs have impaired antigen uptake and T cell activation (105, 106). Moreover, hypoxia inhibits DC maturation and T cell activation but simultaneously increases DC inflammatory cytokine secretion (107). Although the exact mechanisms are unclear, this dichotomy may be determined by the maturation state of DCs. Further work is needed to address more precisely the roles of DC HIF-1α and HIF-2α in the progression of cancer and antitumor immune responses.

Tumor hypoxia increases T cell expression of FoxP3 through hypoxia and HIF-2α signaling. Activated Tregs increase expression of FoxP3 and increase IFN-γ production (108). Tregs are more efficiently activated systemically than within the tumor microenvironment (109), and tumor secretion of cytokines such as TGF-β may be a critical source of systemic Treg activation and infiltration into tumors. CD4+ T cell HIF-1α directly targets the Foxp3 promoter and increases immunosuppressive Treg cell production and function (110, 111). Loss of HIF-1α in Tregs decreased their immunosuppressive function (110). The specific role for hypoxia signaling in antitumor CD4+ and CD8+ T cell responses is not completely clear. T cell HIF-1α represses T cell inflammatory responses, as depletion of HIF-1α significantly enhanced IFN-γ and IL-2 production (112). However, other studies have shown that hypoxia signaling has the capacity to increase CD8+ T cell function, as Vhl deletion in CD8+ T cells increased T cell effector responses and decreased tumor growth in a model of melanoma (113). Additionally, HIF-1α regulates expression of T cell CD137, which augments antitumor immunity upon antibody-mediated activation (114). More work is needed to better understand the specific roles of hypoxia and HIFs in antitumor T cell responses. In addition, hypoxia has cell-intrinsic roles in CTLs, NK, and NKT cells, which have not been completely assessed in tumor biology.

Hypoxia, stromal cells, and inflammation

In addition to immune cells, the tumor microenvironment is also composed of vascular endothelial cells, fibroblasts, and pericytes, collectively known as the stroma. In tumor endothelial cells, depletion of HIF-1α reduced tumor metastasis, whereas deletion of endothelial HIF-2α increased tumor metastasis (115). This dichotomy was due to differential regulation of NO homeostasis by HIF-1α and HIF-2α. HIF-1α regulates expression of inducible nitric oxide synthase (iNOS), which catalyzes the conversion of L-arginine to NO. iNOS was also shown to be an essential regulator of VEGF expression and promoted increased tumor vascularity (115). Myeloid cell and mesenchymal stem cell NO production has been implicated in decreasing T cell cytotoxicity; thus, it is possible that endothelial cell NO has a similar function (116–118). On the other hand, HIF-2α regulates expression of ARG1, which metabolizes L-arginine. ARG1 expression in immune cells is a potent suppressor of antitumor T cells. Future studies should determine if endothelial ARG1 also has an immunosuppressive role in the tumor microenvironment.

Fibroblasts are frequently recruited to tumors and make up a variable proportion of the tumor mass (119). Cancer-associated fibroblasts (CAFs) have a largely protumor role and promote angiogenesis through VEGF secretion and increase invasion/metastasis through secretion of extracellular matrix–degrading matrix metalloproteinase enzymes (119–121). Conditional loss of CAF HIF-1α by Fsp1-Cre expression decreased TAMs in mouse mammary tumors, suggesting that CAF HIF-1α is important for TAM tumor infiltration (122). These results demonstrate that stromal cell HIF signaling can also modulate the microenvironment of tumors; however, the specific role for HIFs in the tumor endothelial cells and CAFs and the significance of this relationship to tumor inflammatory responses have not been fully elucidated.

Hypoxia and intratumoral heterogeneity

Tumor growth is an evolutionary process that leads to the accumulation of genetic alterations. This process also leads to substantial spatial variation within a tumor, which can contain genetically distinct subclonal populations of cancer cells (123). A few studies have demonstrated that intratumoral heterogeneity is an independent risk factor for poor survival in several tumor types, although further studies will be required to determine the effect of such heterogeneity in additional tumor types (124). The mechanisms that drive regional heterogeneity in tumors are poorly described. Hypoxia-induced immune cell recruitment and modulation could be important processes in driving regional selective pressures in tumors. More work is needed to elucidate the mechanisms by which hypoxia and immune cells establish microenvironments leading to intratumoral heterogeneity. Enhanced imaging techniques have made it clear that distinct subpopulations of hypoxic cells are present in tumors (125–127), while histopathological and molecular analyses have shown that distinct tumor regions have different inflammatory infiltrates that can be modulated by intratumoral hypoxia (128, 129).

HIF-based therapeutics in cancer

Due to the highly complex role of HIFs in tumorigenesis, HIF-based therapies need to be assessed on an individual-tumor basis. The best-characterized approach to alter HIF signaling is through...
PHD inhibition, which leads to activation of HIF signaling (130). PHD inhibitors have been assessed in models of inflammation-induced tissue injury and were shown to be effective and safe (131–133). Through a decrease in inflammation-induced tissue injury, PHD inhibitors may have a beneficial role in several cancers. Recent work has clearly demonstrated that PHD inhibitors can selectively activate HIF-1α (134). This suggests the possibility of finding novel agents that may target each isoform. However, there are concerns about therapeutic activation of HIF signaling, as most data suggest that HIF pathway activation promotes tumor progression. Currently, there are several drugs that inhibit HIF-1α and many of them are in clinical trials. Most of the compounds that are in clinical trials were originally discovered as targeting other pathways. 2-Methoxyestradiol, a metabolite of estradiol, can decrease tumor growth through inhibition of HIF-1α but simultaneously can decrease angiogenesis and disrupt microtubes (135). Cardiac glycosides, including digoxin, can robustly reduce tumor growth through inhibition of HIF-1α (136). Furthermore, several topoisomerase inhibitors have been shown to decrease tumor growth through HIF-1α inhibition (137, 138). However, no HIF-1α–specific inhibitors have been discovered. Selective inhibition of HIF-2α can be achieved through targeting the iron response element in the 5′-UTR of HIF-2α (139). Additionally, structural analysis of HIF-2α identified a ligand-binding cavity located within the PAS-B domain, which contains a β-sheets that mediates interaction with ARNT; however, this region has not been identified on HIF-1α (140). There are no known endogenous ligands for HIF-2α, but this cavity has been targeted for drug development and several promising highly specific small-molecule inhibitors have been identified that efficaciously disrupt HIF-2α heterodimerization with ARNT and block DNA binding and transcription of target genes in cultured cells (141). These reagents coupled with recent HIF structural analysis provide a framework to specifically regulate HIF-1α and HIF-2α and will aid the development of clinical tools to alter protumoral inflammatory responses or antitumor immune responses (142).

Conclusion
Hypoxia is an important microenvironmental feature in solid tumors and is essential for tumor growth. HIF-1α and HIF-2α have been extensively studied in regulating tumor glucose metabolism, angiogenesis, cell survival, proliferation, and migration. Research in the past two decades has established an essential role for protumor inflammatory responses or antitumor immune responses in the growth of most solid tumors and with this increased focus, Hanahan and Weinberg’s “hallmarks of cancer” were recently updated to include tumor inflammation and immune evasion as a major enabling factor in cancer progression (143). It is becoming clear that hypoxia is central to regulating the inflammatory response in tumors. Future studies should be directed towards a better understanding of the precise molecular mechanisms by which hypoxia alters the balance between growth-promoting inflammation and the antitumor immune response, which may lead to better use of existing drugs that alter the HIF response as stand-alone therapies and improve the efficacy of standard chemotherapy.

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