

TITLE:

Escape from immunotherapy: possible mechanisms that influence tumor regression /
progression.

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Abstract

Tumor escape is one major obstacle that has to be addressed prior to designing and delivering successful immunotherapy. There is compelling evidence to support the notion that immunogenic tumors, in murine models and cancer patients, can be rejected by the immune system under optimum conditions for activating adaptive and non-adaptive anti-tumor immune responses. Despite this capability, a large number of tumors continue to grow and evade recognition and/or destruction by the immune system. The limited success in current immunotherapeutic strategies may be due to a variety of reasons: failure of effector cells to compete with the growing tumor burden, production of humoral factors by tumors that locally block cytotoxicity, antigen / MHC loss, T-cell dysfunction, production of suppressor T – cells to name but a few or therapy ineffectiveness for the particular malignancy being treated. To optimise immunotherapy strategies correction of immune activating signals, eradication of inhibitory factors and the evasion from newly developed immunoresistant tumor phenotypes need to be simultaneously considered.

Introduction

The immune defence mechanisms are the least effective and the final barrier in the natural mechanisms against cancer formation (carcinogenesis) (49). Both innate and adaptive immunity induce anti-tumor effects via the activity of NK cells, NKT cells, macrophages, neutrophils, eosinophils, various cytokines and specific CTLs to name but a few. However spontaneous regression is rarely observed in malignant tumors, hence the need for immunotherapeutic intervention is required in order to eradicate the tumor.

Advances in both molecular and cellular immunology have improved our understanding of tumor-host interactions, although much still remains elusive despite the progress made in identifying a large number of tumor antigens and new immunotherapeutic strategies. The majority of tumors manage to “escape recognition” and currently there are a number of mechanisms known for tumor escape which have been described – loss / downregulation of HLA Class I, downregulation, mutation or loss of tumor antigens, alterations in cell death receptor signalling, production of immunosuppressive cytokines and suppressor T-cells and the involvement of

indoleamine 2,3-dioxygenase in suppression - all of which will be discussed in detail later.

It is believed that a T-cell response does exist but is ineffective as the effector phase of the immune response is inadequate. This notion may explain why current immunotherapeutic strategies have limited success. From the immunological perspective efficient therapeutic intervention should focus on boosting existing anti-tumor responses and to sustain large numbers of effector T cells at the tumor site.

Recently, this was clearly demonstrated in melanoma patient where T cell responses to vaccines occurred primarily in the presence of pre-activated T cells prior to vaccination (111). Analyses of cell mediated immunity against defined melanoma antigens using tetramer staining and an IFN- γ secretion assay suggested that specific T cell responses often exist during tumor progression (99). Adjuvants such as cytokines have been used when immunising with peptides and proteins (48,54,100,118). Immunisation with DNA vaccines leads to the expression of tumor antigens and their processing by antigen presenting cells (APCs) (112), whereas the direct delivery of antigens as proteins by cell fusion or cDNA to dendritic cells (DCs) represents a more direct attempt to generate antitumor responses (2).

Some anti-tumor therapeutic interventions have met with limited success which has been attributed mainly to the “tumor – escape” phenomena i.e. tumor cells have the ability to evade immune recognition and/or destruction, causing major obstacles in using this modality for the treatment of cancer. This was clearly demonstrated in prostate tumor (TRAMP-C1) model where the injection of Fms-like tyrosine kinase-3 ligand (flt3-L) induced short-term tumor stabilization and regression followed by tumor relapse (21). Failure of therapy in this model revealed several important immunosuppressive characteristics of the prostate tumor microenvironment including the down regulation of MHC class II in DCs, profound deficiency in the expression of CD3 ϵ (CD3epsilon) and the beta chain of the T cell receptor (TCR) of tumor-associated T cells. Cancer cells can be detected and destroyed by cytotoxic T lymphocytes (CTLs) in many experimental models and human tumors but on diagnosis the metastasis of many human tumors can be shown to correlate with unresponsive T cells. Progressive tumor growth occurs, in part, because the cancer cell is capable of escaping immune recognition and not because the immune system is defective (16). CML patients for example possess CTLs detectable by tetramer staining and

cytotoxicity that recognise HLA-A3 restricted peptide, although the number of CTLs may be too few to mediate complete rejection of the cancer (23).

Deletion or suppression of activated CTLs can occur as well as their migration away from the tumor site. The precursor frequency can be too low, such that the effector cells fail to compete with the growing tumor burden. However, it is equally possible that the therapies are ineffective for the particular malignancy being treated. In order to fully appreciate the role of tumor escape in the failure of immunotherapy, an understanding of the basic principles of immunosurveillance and its role in determining tumor phenotypes is necessary.

The hypothesis first proposed by Thomas & Burnet in the 1970s implied that the immune response to tumor occurred at an early stage in tumor development – “eradicating the cancer before it became apparent” (28,87).

Many studies have evolved in support of this theory and evidence exists to suggest that in some models and under appropriate conditions, immunosurveillance can play an active role in suppressing the growth of early tumors. Hence in this paradigm, when tumors do successfully grow, they are thought to have escaped from surveillance and a number of reviews have been written to support this theory (56,78,97).

The following will focus on the multiple mechanisms that exist leading to “tumor-escape” from the immune system; these factors have to be taken into consideration when designing novel therapeutic strategies. The aim of this review is to highlight the required mechanisms of tumor escape and how they influence the failure of immunotherapy.

Activation versus Suppression during Tumor Progression.

Escape from immune-surveillance is a major mechanism for tumors to grow progressively. Tumors may grow undetected by the immune system, being seen as “normal tissue” hence benefiting from any mechanism that makes them appear “healthy” and less dangerous i.e. exhibiting no danger signals for immune activation (56,86). It is believed that even during progressive growth the tumor has the ability to activate the immune system, and that a fine balance between activation and suppression exists, which determines the fate of the tumor. Tumor vaccines may induce activation and expansion of specific CD8⁺ T cells and destruction of tumor cells in cancer patients; this was observed in approximately 5-20% of vaccinated

melanoma patients. However, this activation can be frustrated by the lack of appropriate co-stimulation or the presence of immunosuppressive cytokines such as IL-10 and TGF- β . The eventual fate of the tumor is therefore determined by the net effect of immune activation and inhibition. (56).

Killing of tumor cells *in situ* by suicide gene transfer to induce cell death through a nonapoptotic pathway was shown to be associated with enhanced immunogenicity (74). Similar observations were also reported by our group where immunisation with irradiated RENCA cells (murine renal carcinoma) infected with DISC-HSV (Disabled Infectious Single Cycle – Herpes Simplex Virus) that enhanced immunity to tumor challenge with live parental tumor cells that induced cell death by necrosis as opposed to apoptosis (5) providing the immune system with additional activation as the cells were undergoing necrotic cell lysis enhanced tumor antigen processing and presentation by APCs and hence an increase in T cell activation (59,70).

The mechanisms of tumor escape from immune recognition / destruction are likely to be multifactorial including downregulation of MHC Class I molecules (4,52,96,131), loss of tumor antigens (26,44), defective death receptor signalling (24,47,73,116,117), lack of co-stimulation (104), production of immunosuppressive cytokines (85) and suppressive cells (12,35,61,72,113) as shown in figure 1.

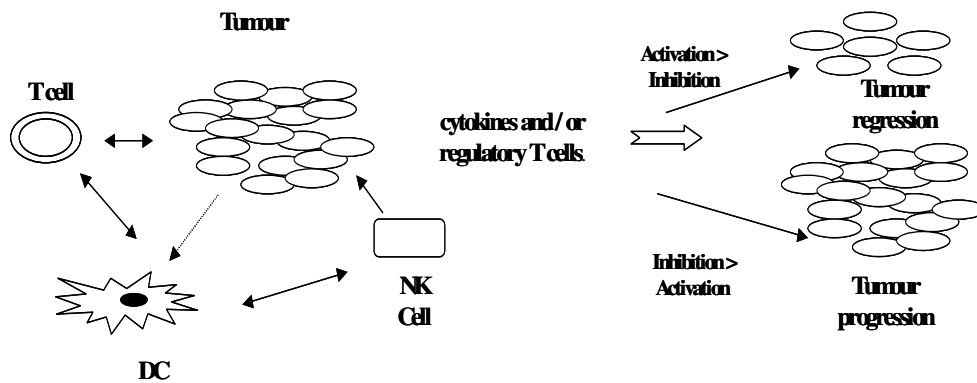


Figure 1:- A schematic diagram summarising the processes of activation and suppression during tumor progression.

Involvement of Indoleamine 2,3 -dioxygenase in immunosuppression.

The mechanisms that mediate immune tolerance to cancer are not well understood, but recent findings have implicated the tryptophan/IDO (indoleamine 2,3-dioxygenase) metabolic pathway as one of many mechanisms involved. IDO is an enzyme ubiquitously distributed in mammalian tissues and cells converting tryptophan to N-formylkynurenine (34). This cytosolic enzyme catalyses the initial and rate limiting step in the catabolism of tryptophan in the kynurenine pathway (76). Low levels of tryptophan at the tumor site causes T cells to arrest in the G1 phase of the cell cycle (125). Initially the enzyme was recognised for its anti-microbial activity by allowing cells to deplete tryptophan from intracellular pool or local microenvironment (114), however recently, IDO was reported to be constitutively expressed by most human tumors and its expression by immunogenic mouse tumor cells prevents their rejection in pre-immunised mice, correlating with the lack of specific T cell accumulation at the tumor site (125).

IDO expression is an inducible feature of splenic DC subsets, and provides a potential explanation for their ability to regulate T cells. Induction of IDO completely blocked the clonal expansion of T cells from TCR transgenic mice following the administration of the immunomodulatory reagent CTLA4-Ig, whereas the same treatment did not block T cell clonal expansion in IDO-deficient recipients. Suppression of allogeneic T cell responses in vitro and in vivo by interstitial APC was also shown to be IDO-dependent and blockade of IDO activity with 1-MT (1-methyl-

DL-tryptophan (3)) or addition of exogenous tryptophan allowed the recovery of T cell proliferation in MLR assays (115). In humans, only a discrete subset of APCs that co-express the cell-surface markers CD123 and CCR6 was shown to express IDO and inhibit T cell proliferation *in vitro* (79). These results clearly demonstrated that the IDO pathway might represent a potential mechanism for DCs to regulate the immune response to tumor antigens.

Loss / Downregulation of HLA Class I.

Altered MHC class I antigen expression in tumors is a well known phenomena. The selective loss of MHC class I alleles was subsequently described in a variety of mouse models including the TL leukaemic cell line, MCA-induced tumors and murine leukaemia virus induced tumors. The altered phenotypes of MHC Class I antigen expression permits tumor cells to avoid recognition or survive attack by CD8⁺ T-cells capable of mediating cytotoxicity. This phenomenon has been studied in a series of tumor samples by immunohistochemical techniques and has been shown to be widespread (58).

HLA Class I molecule downregulation occurs frequently in many cancers and this abnormality might adversely affect the clinical course of disease and the outcome of T cell based immunotherapies. Over the past few years HLA Class I expression has been characterised in human tumors. Changes in HLA Class I expression can occur through various mechanisms – namely, mutations in genes and abnormalities in the regulation and/or defects in HLA Class I dependent antigen processing. These mutations modulate the susceptibility of tumor cells to *in vitro* lysis by CTLs and natural killer (NK) cells. Immune selection of tumor resistant CTLs and/or NK cells may explain the rapid progression and poor prognosis of cancers exhibiting HLA Class I antigen downregulation (43).

To date a number of investigators have classified the types of HLA Class I loss into different phenotypes, and Table 1 summarises the HLA phenotypes that are recognised.

Table 1: Major HLA phenotypes in tumors and the associated molecular mechanisms. (Adapted from Cabrera et al 2003 (16)).

HLA Phenotype	Mechanisms	References
(I) HLA Class I Total Loss	β_2m mutations Alterations in antigen processing machinery	37,55,88 37,105
(II) HLA haplotype Loss	LOH in chromosome 6	31,37,91,92
(III) HLA Allelic Loss	Mutations of HLA Class I genes	13,37
(IV) HLA A, B, C locus downregulation	Alteration of transcriptional factors	37,39,110
(V) Compound phenotype	2 or more independent mechanisms	37,45,93

A number of studies have been carried out to assess HLA Class I expression in tumors. In bladder carcinomas, 72% of the tumors studied showed at least one alteration in HLA expression and these were classified into different phenotypes depending on the type of downregulation/loss (18); expression was correlated to the degree of differentiation and tumor recurrence. To determine the type of HLA loss occurring during tumor progression, tissue samples from breast, colorectal and laryngeal carcinomas were analysed (17). HLA-B44 allele loss was observed more frequently than any other HLA Class I allele – suggesting an important role for this allele in tumor escape.

Furthermore, mechanisms of MHC Class I loss were also investigated in colorectal carcinoma (15). It was shown that β_2m mutation and LMP7 / TAP2 downregulation were responsible for the total loss of MHC class I expression-which contribute to the failure of T cell recognition during the immune response.

In murine models, the B9 primary tumor clone (H-2 negative) and its metastatic colonies were studied in immunocompetent Balb/c mice (37). They showed that 83% of the metastasis obtained in different syngeneic Balb/c mice repeatedly exhibited a phenotype different from the original B9 clone. The alterations were identical in different colonies of different syngeneic Balb/c mice.

Variation in the MHC Class I expression following immunotherapy has been observed in a murine colon carcinoma model (Ahmad, Ali, and Rees, unpublished observation). The CT26 (colon carcinoma) murine tumor model was used to investigate the expression of MHC Class I antigens in mice failing to regress their

tumors following immunotherapy (progressor) and tumor bearer (no therapy) animals (6). The DISC-HSV/mGM-CSF vector used in this study is an attractive tool for the delivery of cytokines to the tumors (11,94,120). This vector was developed by genetically inactivating the HSV-2 for use as a vaccine, by the deletion of the essential glycoprotein H (gH) gene – restricting it to one cycle of replication (11). On infection, the virus would release non-infectious particles together with the transduced cytokine e.g. GM-CSF. In our model the DISC therapy was administered intratumorally to mice with established subcutaneous (s.c) tumors, and approximately 40% of the treated animals fail therapy and the tumors continue to grow progressively (3,6). MHC Class I (H-2L^d) expression was investigated in both tumor bearing and progressor mice. The results (Table 2) clearly demonstrated that a high proportion (55%) of tumors from progressor mice showed loss of class I MHC antigen, unlike the control tumors (treated with PBS) all of which expressed MHC I antigen. Interestingly, the tumor cells from the progressor animals re-expressed Class I following overnight *in vitro* culture, suggesting that the tumor micro-environment may be responsible for the transient downregulation of Class I expression as opposed to a mutation in the β 2m gene (95) or a downregulation in LMP7 / TAP 2 as reported in other studies (15).

Table 2: A total of 18 tumors were studied (11 progressors, 7 PBS controls) for the expression of H2L^d. The results have been combined for the immunohistochemistry and FACS analysis, such that 45% of the therapy failures completely lost H2L^d surface expression whereas the tumor bearer animals retained a degree of expression (low, partial or full expression) (A.Talasila – MSc. Thesis, Nottingham Trent University, UK).

Treatment	No Expression (%)	Low Expression (%)	Partial Expression (%)	Full Expression (%)
Tumor Bearer	0	29	29	42
DISC/mGM-CSF	45	28	27	0

Recently a number of studies have investigated the expression of Class II in patients. Antigen – presenting cells are crucial for the induction of an antigen – specific immune response and downregulation / loss of expression in cell surface molecules such as HLA Class II may contribute to an impaired immune response. This phenomena was observed in PBMs of melanoma patients (123). A number of

different carcinomas (breast, ovarian and prostate) have been investigated for the expression of MHC Class I and II antigens (62, 69, 107) where the absence / downregulation of these recognition molecules correlated with the tumor's ability to avoid immune surveillance and development of metastatic dissemination of the malignancy.

Down-regulation, mutation or Loss of Tumor Antigens.

Alterations in tumor associated antigen (TAA) expression is one of the mechanisms by which tumor cells can escape CTL detection. Modifications in the TAA expression can range from down-modulation to complete loss, and loss of antigen expression can occur independently of the deregulation of HLA Class I expression. Tumor antigen expression is known to be heterogeneous, even within the same tumor (56); and a decrease in the expression of gp100 and MART-1 was associated with disease progression (26,44). In spite of the presence of TAA-specific CTL, immunogenic tumors can eventually grow and kill the host in animal models. Tumor escape in the murine mastocytoma P815 tumor was shown to be due to the emergence of stable antigen-loss variants (124).

Antigenic drift, a mechanism used by viruses to escape immune recognition, has recently been described for tumors. Transgenic mice expressing TCR for a single antigenic epitope have been used extensively in establishing antigenic mutation(s) as a mechanism for viral escape of T cell recognition (22,98). A transgenic mouse line expressing TCR specific for tumor antigen P1A35-43 presented by H-2L^d was also developed and used to study tumor escape mechanisms. The recurrence of tumors in mice that have responded favourably to transgenic T cell adoptive therapy was found to correlate with the presence of tumor variants with mutations within the P1A epitope (9). These mutations severely diminish T cell recognition of the tumor antigen by a variety of mechanisms, including modulation of MHC: peptide interaction and TCR binding to MHC: peptide complex. In another study, using a SCID mouse model and MART-1/Melan-A-specific CD8⁺ T cell clone against autologous melanoma, it was found that the *in vivo* immuno-selection of antigen-loss variants was dependent on the presence of sub optimal levels of antigen expression (66); and loss of the immunodominant T cell-defined MART-1/Melan-A antigen and downregulation of the TAP-1 gene has been identified in a recurrent metastatic melanoma. Restoring the "antigen loss" in the variant tumor cell line by simultaneously providing both the

MART-1/Melan-A gene (by retroviral transfer) and the TAP-1 gene (by a bioballistic approach) resulted in tumor cell sensitivity to MART-1/Melan-A-specific cytotoxic T lymphocytes suggesting antigen loss and downregulation of the peptide-transporter protein TAP-1 expression (68). The emergence of tumor escape variants is more likely to result after effective immunotherapies.

Alterations in cell death receptor signalling.

The expression of Fas Ligand (FasL), a well-known cell death receptor ligand, on tumor cells has been implicated in their evasion of immune-surveillance (20,24,90,116,117). Apoptosis mediated via CD95 (Fas/Apo-1) is a key regulator for the biology of normal and malignant cells. However, Fas surface expression does not necessarily render cells susceptible to FasL-induced death signals, suggesting a role for inhibitors of the apoptosis-signalling pathway. Phosphatidylinositol 3'-kinase (PI3 K) and Akt (protein kinase B) mediate the survival signal and allow cells to escape from apoptosis in various human cancers. PI3 K inhibitors, such as LY294002 inhibited cell proliferation and increase apoptosis in the human gastric carcinoma cell line by the down-regulation of Mcl-2 and phosphorylated Bad proteins, which are anti-apoptotic factors and belong to the Bcl-2 family (84). Acquisition of apoptosis resistance is a typical mechanism of chemotherapy failure in small-cell lung cancer, which was correlated, with the over-expression of Bcl-2 (32).

Defects in receptors/signalling can facilitate tumor cell survival and proliferation. The Fas/FasL complex forms and engages caspase-8 which autoactivates itself and cleaves caspase-3, -6 and -7 (101). The caspase-8 inhibitor, cellular FLICE-inhibitory protein (cFLIP) is expressed in many tumors making them resistant to death receptor mediated apoptosis (47). FLICE/caspase-8-inhibitory protein (cFLIP) is a recently identified intracellular inhibitor of caspase-8 activation that potently inhibits death signalling mediated by all known death receptors, including Fas, TNF-receptor (TNF-R), and TNF-related apoptosis-inducing ligand receptors (TRAIL-Rs), The increased expression of cFLIP in tumor cells is thought to contribute to immunoresistance to T cells *in vivo* (73) while the downregulation/loss of Fas expression resists apoptosis. In patients with B-cell chronic lymphocytic leukemia (B-CLL cells), an increase in cell-surface CD95 expression on T cells was associated with reduced progression-free probability and poorer survival (40). In TRAIL-mediated apoptosis, the loss of

expression of all TRAIL receptors can occur by many mechanisms including the loss of caspase-8 by chromosomal loss or mutation (42).

Cells expressing low levels of Fas (Fas^{low}) in the B16F10 tumor model grew more slowly in comparison to the cells expressing high levels of Fas. Fas^{low}-cells when injected showed enhanced tumor growth in mice depleted of neutrophils, suggesting that inactivation of neutrophils was an important mechanism by which tumors could escape destruction (66). The expression of FasL has also been studied in cervix adenocarcinomas, where FasL appears to play an important role in immune evasion, progression and metastases of the tumor (53). In contrast, the intracellular expression of FasL in breast cancer and normal tissue, as determined by immunohistochemistry, is unlikely to be an important marker for immune evasion (90). The expression of membrane bound FasL (mFasL) on colon cancer cells as a potential mechanism to inhibit host immune function by inducing apoptosis of host lymphocytes was investigated (109). mFasL can be cleaved to release a soluble FasL (sFasL) to spread the apoptosis induction effect within the environment. These findings have suggested sFasL as a mechanism by which tumor cells can avoid immune attack.

Immunosuppressive Cytokines.

Tumor cells produce a number of cytokines and chemokines that can have a suppressive effect on immune cells. In non-small cell lung cancer patients, the mRNA expression of IL-4, IL-10, TGF-alpha, and TGF-beta1 was significantly higher than that of IL-2, IL-12, IL-18 and INF-gamma as determined in pleural effusion and tumor tissue (64). This predominant expression of type II (immunosuppressive) cytokines mirrors an immunosuppressive state in the immunological microenvironment. Vascular endothelial growth factor (VEGF) is secreted by many tumors (121), and in the past few years VEGF has been required as a contributory factor in tumor escape. VEGF is not only important for tumor vascularisation but is also a key factor produced by solid tumors to inhibit immune recognition (82), and prevent DC differentiation and maturation by suppressing the transcription factor NF- κ B in haematopoietic stem cells (85). Blocking NF- κ B activation in haematopoietic cells by tumor-derived factors is thought to be a mechanism by which tumor cells can directly down-regulate the ability of the immune system to generate an antitumor

response (85). Elevated VEGF blood concentrations have been correlated with poor prognosis in human neoplasms (65), possibly as a result of its angiogenic properties and/or its ability to suppress DC maturation (46,65). VEGF was negatively related to DC infiltration in immunohistochemical studies on resected lung cancer samples (46) and activation status of DC and the concentration of VEGF in the peripheral blood have been shown to reflect the malignancy of NSCLC (30).

Supernatants collected from tumor cells of AML patients were shown to inhibit T cell activation, Th1 cytokine production and to prevent activated T cells from entering the cell cycle (14), however, no TGF- β , IL-10 or VEGF were detected. The T cell immunosuppression induced by AML cells provides a mechanism by which leukemic clones could evade T cell mediated lysis by inhibiting the NF- κ B, c-myc and pRb pathways (14). The production of soluble factors such as VEGF, IL-10, TNF and TGF- β is a proposed mechanism for tumor cells to avoid immune recognition, and the effects of these factors appear to be two-fold: to inhibit a) the effector function and b) the development of the immune cells by acting in the early stages of immunopoiesis (82).

IL-10 is a cytokine often quoted as being suppressive and high levels of IL-10 have been reported in patients with melanoma (33) and pancreatic cancer (10,77). IL-10 has the ability to exert its effects in many ways:- it inhibits antigen presentation, IL-12 production and the induction of Th1 responses in vivo (25,106).

High concentrations of TGF- β are also found in cancer patients (122) and are usually associated with “tumor-progression” (38) and poor responses to immunotherapy (27). The levels of TGF- β are higher in patients with disseminated melanoma when compared to those with localised disease (60). TGF- β has also been shown to induce the overproduction of IL-10 in tumors, leading to immunosuppression of antitumor responses. This suppression was reversed and Th1 responses reconstituted on the administration of anti-TGF- β antibodies in vivo (67).

Alterations in the expression of signal transduction molecules

Many different mechanisms may contribute to immune evasion. In pancreatic cancer, the loss of signal transducing CD3 zeta chain (CD3 ξ) of TILs has been attributed to immune escape together with the production of immunosuppressive cytokines and

local impairment of TILs (126); CD3 zeta loss in many cases has also been correlated with elevated levels of IL-10 and TGF- β . The loss of the CD3 ξ chain has been studied extensively and has been proposed as a mechanism by which tumors are able to escape. The zeta chain is present as a large intracytoplasmic homodimer in the TCR forming part of the TCR-CD3 complex. It functions as a single transducer upon antigen binding. (50). Hence any alterations in this chain could result in changes in the signalling pathway. In pancreatic cancer, the loss of signal transducing CD3 ξ chain of tumor infiltrating lymphocytes (TILs) has also been attributed to immune escape together with the production of immunosuppressive cytokines and local inactivation of TILs (126). Alterations in the expression of CD3 ξ chain has also been correlated with elevated levels of IL-10 alone or in association with the downregulation of IFN- γ (81). A general downregulation / decreased expression the CD3 ξ chain has been reported in tumors such as cervical and colorectal carcinomas (57,80,103). In cervical cancer the reduced expression of the zeta chain was associated with a reduction in cellular functions such as the production of TNF (57). In contrast, PBMCs from breast cancer and colorectal carcinomas patients do not show a decrease in TCR zeta expression yet the former exhibit an impairment in T cell function (81).

Lack of co-stimulation

Most tumors do not regress and continue to grow in spite of the presence of spontaneous or antigen (vaccine)-induced immune responses. The existence of systemic immune responses may not by itself be sufficient to cope with the complex nature of tumor-host interactions because of factors: insufficient co-stimulation to induce T-cell response may further contribute to the lack of effective immunity. There are a number of molecules that normally perform co-stimulatory functions by interacting with their counter ligand/receptor on T cells to provide the critical second signal for T-cell and/or APC activation including: CD80 (B7-1) / CD86 (B7-2) binding to CD28 and CTLA4 respectively; CD40L on activated CD4 helper cells binding to CD40 receptors on APCs; human CD58 and mouse CD48 (LFA-3) binding to CD2; and CD54 (ICAM-1) binding to LFA-1. Members of the TNFR super-family including CD27, CD30, 4-1BB and OX40 have also been shown to transmit a co-stimulation signals to leukocytes (41,108).

It is a recognised phenomenon that T cells are rendered anergic due to the lack of co-stimulatory molecule(s) expression by tumor cells (104). Tumor cells are able to induce antigen-specific tolerance or anergy on the basis of MHC-1-restricted antigen presentation without the expression of co-stimulatory ligand(s) (1). This unresponsiveness however can be reversed when tumor cells are genetically modified to express co-stimulation molecules. Chen et al (19) showed that the insertion of genes encoding B7.1 and /or B7.2 molecules into tumors generally increases their immunogenicity.

Recently, fusogene vectors were developed to encode multiple gene products as fusion proteins from a single cistron to increase the immunogenicity of target tumor cells (36). Analysis of over 100 individual clones derived from human and murine tumor cell lines demonstrated efficient expression and biological activity of each of the proteins. Tricistronic viral vectors co-expressing IL-12 and B7.1 have been used in the immunotherapy of cancer (128); the vectors generated could be used in immunotherapy for the treatment of multiple myeloma and other cancers as they were shown to stimulate allogeneic mixed lymphocyte proliferation and provoke increases in CTL responses and IFN- γ release from normal donor lymphocytes exposed to the parental U266 cells.

In addition, co-stimulation through molecules like 4-1BB was also found to be critical in the expansion and differentiation of CTLs. Systemic administration of an agonistic mAb against 4-1BB enhanced the CD8 T-cell response, leading to the eradication of established AG104A sarcomas and P815 mastocytomas in vivo (75). However, resistance to this treatment modality by a number of poorly immunogenic tumors, including the TC-1 lung carcinoma and B16-F10 melanoma, was shown to be due to tumor antigen-specific CTLs “ignorance” rather than anergy or deletion; breaking CTL ignorance by peptide immunisation was necessary for the anti-4-1BB to function to enhance T-cell responses (129). Given the importance of co-stimulation in the regulation of immune responses against cancer, the manipulation of this pathway to increase immunity represents a promising therapeutic approach. Co-stimulation through OX40L for example would be advantageous, since its expression is primarily detected on recently stimulated antigen specific CD4⁺ T cells (127, our unpublished data), which is considered advantageous for generating CD8⁺ T-cell responses and anti-tumor immunity.

Immunosuppressive cells

There are at least 2 types of cells that when produced can impose a suppressive effect on the host's immune system ($CD4^+$ / $CD25^+$ T cells and $Gr1^+$ / $CD11b^+$ myeloid cells), hence providing the tumor with the opportunity to escape immune recognition. Elucidating the mechanism(s) of T cell unresponsiveness in cancer is critical for the design and application of an effective cancer immunotherapy. Inhibition of a number of T lymphocyte functions in tumor-bearing hosts has been extensively investigated (63,89), and suppressor cells have been shown to play a role in the progression of cancer (29). $CD4^+$ $CD25^+$ T cells are known to be immunoregulatory and are important in immunological tolerance to self-antigens (72) and inhibition of T cell proliferation (119). They constitute approximately 5-10% of the $CD4^+$ T cells in both humans and rodents and their removal induces autoimmune diseases in various locations in the body. Suttmuller and colleagues (113) demonstrated that the depletion of $CD4^+$ $CD25^+$ T cells followed by an injection of an antibody which blocks CTLA-4 enhanced T-cell reactivity to a known tumor associated antigen. However, it is thought that the mere depletion or blocking of T regulatory cells is not sufficient to successfully treat established tumors (8).

The existence of anergic and functionally suppressive $CD4^+$ $CD25^+$ T cells was demonstrated in patients with melanoma undergoing immunisation with known melanoma antigens. The degree of inhibition of T cell proliferation was proportional to $CD4^+$ $CD25^+$ T cells present; the addition of IL-2 reversed their hypo-responsiveness and abrogated their suppressive function (51). An increase in $CD4^+$ $CD25^+$ regulatory T cells was shown to be correlated with immunosuppression and tumor progression in patients with gastrointestinal malignancies (102). Increased numbers of $CD4^+$ $CD25^+$ T cells secreting TGF- β was detected in tumor infiltrates from patients with early and late-stage epithelial tumors (130). These observations provide clear evidence for the contribution of $CD4^+$ $CD25^+$ T cells to immune dysfunction in cancer patients. Immunotherapy aimed at decreasing the role of regulatory T-cells would be advantageous to successfully treating cancer.

In addition, immune suppression in tumor bearing mice has been attributed to the presence of cells with an immature myeloid phenotype that express the granulocyte – monocyte markers $Gr1^+$ / $CD11b^+$ (12), and accumulate in the spleens and lymph nodes (71) and blood of tumor bearing mice (Ahmad, Ali and Rees, unpublished data). They

are capable of inhibiting antibody production, CTL generation, T cell function and lymphocytic proliferation (61,71). Accumulation of Gr1⁺/CD11b⁺ cells and their ability to inhibit the T cell function has been reported in cancer patients (7). The Gr1⁺ cells are also able to decrease CD3 ξ molecule expression significantly (83), and inhibit MHC Class I dependent antigen-specific CD8⁺ T-cell (35).

Conclusion

This review has highlighted the main mechanisms by which the tumors are known to escape from immune recognition. Given the many different potential mechanisms that tumors can acquire to avoid or subvert adaptive immunity, future generation immunotherapeutic strategies will need to consider not only how to promote antigen driven T and B lymphocyte responses and their effective targeting to residual tumor, but also to understand how the mechanism(s) of tumor escape can be dealt with. Current research is exploring the application of combination therapy that utilises several treatment modalities, which may include sequential chemo-, radio- and immunotherapy protocols.

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