Abstract
In 2018, the Nobel Prize in medicine was awarded to James P. Allison and Tasuku Honjo for their work on the description of immune checkpoint inhibitors which contributed to the development of new anti-cancer immunotherapies. However, although these new therapeutic strategies, which are designed to limit immune escape of cancer cells, have been used or tested successfully in many different cancers, a large proportion of patients have been described to resist and not respond to these new treatments. The new incoming challenge is now therefore to overcome these resistance and new recent data presented epigenetic modifications as promising targets to restore anti-tumor immunity. Indeed, both DNA methylation and post-translational histone modifications have been described to regulate immune checkpoint inhibitor expression, tumor-associated antigen presentation or cancer cell editing by the immune system and therefore establishing epigenetic drugs as a potential complement to immunotherapies to improve their efficiency.

Keywords: Epigenetics, DNA methylation, drug resistance, immunotherapy, programmed cell death 1, programmed cell death ligand 1, cancer

INTRODUCTION
Antigen presentation by the professional antigen-presenting cells (APCs) are normally recognized by immune cells via a surface T-cell receptor (TCR), leading to the induction of naive CD8+ T-cell clonal expansion and differentiation and finally resulting in the cell death of APCs. Specific tumor-associated antigens (TAAs) expressed by cancer cells can be identified by the immune system and results in the
activation of immune system effectors and the consequent elimination of the cancer cells. Indeed, an increase in the recruitment of cytotoxic CD8+ T-lymphocytes into the tumor is mediated by chemokines such as C-C motif chemokine ligand 5, C-X-C motif chemokine ligand 9 (CXCL9) and CXCL10 and is correlated to lower metastasis risk and better outcome for patients[1-3] [Figure 1]. Three sequential phases of immune edition of cancers have been described and summarized under the rule of the “3E”: (1) Elimination; (2) Equilibrium; and (3) Escape[4]. During the Elimination phase, some newly transformed cells can be quickly eliminated by immune effector cells, such as natural killers (NKs), but this phase can also favor the development of heterogeneous tumor cancer cells resulting in the selection of new variants resistant to immune edition. Further growth of cancer cells selected during the Equilibrium phase may then lead to the acquisition of aggressive phenotypes and resistance to immune system elimination inducing clinical lesions. So, immune escape has now been included as one of the new hallmarks of cancer development described in the revised version of Hanahan and Weinberg[5] in 2011, together with resistance to cell death, replicative immortality or induced angiogenesis, for example.

RESISTANCE TO IMMUNOTHERAPY IN CANCERS

Immunotherapy protocols are designed to annihilate immune escape in cancer cells and restore elimination of cancer cells. In 2018, Nobel Prize in medicine awarded James P. Allison and Tasuku Honjo for their work on the knowledge of immune edition and the development of immunotherapies to fight against cancer cells[6,7]. Immunotherapies include: cancer vaccines, humanized monoclonal antibodies targeting TAAs, transfer of chimeric antigen receptor-T-cells or adoptive transfer of transgenic TCR-expressing T-cells. Nevertheless, immunotherapy resistances have been described in patients and are divided in innate and acquired resistance (for a review[8]). Intrinsic resistance is frequently observed in patients with a failure of systemic immune response such as immune-compromised HIV patients or specific failures induced for example by a lack of expression of specific tumour antigens by tumour cells. Adaptive resistance might be provoked by several processes [Figure 1]: (1) loss of antigens presentation due to mysfunctions.

Figure 1. Examples of direct implication of epigenetic modifications in immune recognition of cancer cells. Epigenetic modifications: DNA methylation (white circles) histone methylation (me) and histone acetylation (Ac, +) are directly implicated in: (1) chemotaxis by the regulation of the expression of CXCL9/10; (2) in immune checkpoint control, by the regulation of PD-L1 and PD-1 in cancer cells and CD8+ cells respectively; and in (3) antigen presentation, by the control of the expression of CTAs and MHC-I. CXCL: C-X-C motif chemokine ligand; PD-1: programed cell death 1; PD-L1: PD-1 ligand; CTA: cancer testis antigen; TCR: T-cell receptor.
in antigen presentation machinery or a decrease in major histocompatibility complex (MHC) expression; (2) dysfunctions of signaling pathways such as the Wnt/β-catenin, phosphatase and tensin homolog, phosphoinositide-3-kinase (PI3K) or Janus kinase pathways; and (3) prolonged T-cell stimulation, which is frequently observed in chronic diseases such as chronic infection or cancer. This stimulation frequently leads to the exhaustion of immune cells, phenotype which is characterized by a loss of both immune and proliferative capacities. This state is usually accompanied by a loss of expression of cytokines such as interleukin-2 (IL-2), tumor necrosis factor α (TNFα) and interferon (IFN) and induction of immune checkpoint inhibitors such as programmed cell death 1 (PD-1) (PDCD1), cytotoxic T-lymphocyte associated protein 4 (CTLA-4), lymphocyte activating 3 (LAG-3) and T-cell immunoglobulin mucin 3 (TIM-3). Immune checkpoint inhibitors are proteins controlling the immune system: PD-1 is expressed at the surface of various immune cells, such as T-lymphocytes, myeloid cells or NKS, and is particularly expressed in exhausted T-cells. PD-1 ligand (PD-L1) (also referred as CD274) or PD-L2 are expressed at the cell surface of tumor cells and can bind to the PD-1 receptor. The PD-1/PD-L1/2 interaction induces a negative signaling cascade which leads to the inhibition of CD8+ T-cell proliferation, cytokine secretion and inflammation leading to a decrease in tumor cell elimination. Moreover, interaction of the other immune checkpoints with their respective ligands (CTLA-4 with the ligands CD80/CD86 expressed on regulator T-cells, LAG-3 with MHC class II, LAG-3 with galectin-9, high mobility group protein B1 or with carcinoembryonic antigen cell adhesion molecule 1 and phosphatidylserine) also decrease the anti-tumour immune response.

Although the identification of these mechanisms is recent and most of the factors involved still remain largely unknown. Anti-PD-1 and anti-PD-L1 immunotherapy protocols (such as nivolumab or pembrolizumab) have been developed and used to restore immune editing in cancers. This process is called immune checkpoint blockade (ICB) and, in the last few years, this new treatment has been tested and shown promising results in many different types of cancers such as: non-small cell lung carcinoma (NSCLC), melanoma, mesothelioma, renal cell carcinoma (RCC), bladder cancer, head and neck squamous cell carcinoma (HNSCC). Unfortunately, although some very interesting results were obtained in some patients, resistance to ICB are observed in a large percentage of cases. For example, only 20% of NSCLC or triple negative breast cancer (TNBC) patients treated with an anti-PD-1 therapy presented a significant positive response. These results suggest that immunotherapy resistance is present in a large proportion of patients before treatment and may be due to a low immune checkpoint inhibitor expression or an absence of T-cell infiltration in the solid tumors.

**IMMUNE CHECKPOINT INHIBITORS ARE EPIGENETICALLY CONTROLLED**

Epigenetics includes DNA methylation, histones post-translational modifications and non-coding RNA and regulates gene expression in a transmissible but reversible manner. DNA methylation is processed by DNA methyl transferases (DNMTs) which catalyze the addition of a 5methyl cytosine on the 5th position of cytosines in CpGs. This mark is generally repressive and many genes are normally controlled by DNA methylation during embryogenesis. Aberrant DNA methylation also frequently occurs in pathologies, specifically in cancers, where it contributes to tumor suppressor gene silencing or inactivation of apoptosis (for a review). Histones are organized in nucleosomes whose local compaction is regulated by histones post-translational modifications. These modifications are varied but the most studied are acetylation and methylation. Acetylation on lysines in histone tails is promoted by histone acetyl transferases and provokes a relaxed chromatin favorable to transcription factor recruitment, opposite to deacetylation which is catalyzed by histones deacetylases (HDACs). The pro- or anti-transcriptional role of histone lysine and arginine methylation is dependent of both the position of the amino acid and the level of methylation. For example, the H3K4me2/3 mark is permissive whereas the H3K9me2/3 or H3K27me2/3 are repressive. These modifications are catalyzed by histone methyl transferases (HMTs) and removed by histone demethylases. In
regard to their roles in tumorigenesis, the epigenetic enzymes are potential targets for anti-tumoral therapies used routinely in some specific cancers or tested in clinical trials (for review\cite{23}).

Although the regulation of expression of PD-1 and PD-L1 in tumors is still largely unknown, recent data proposed a major involvement of epigenetics. Wrangle and collaborators were the first to demonstrate, using genome-wide studies in NSCLC cell lines, that the passive DNA-demethylating agent azacytidine (Aza) potentiated innate and adaptive immune pathways. This effect was associated with an increase in PD-L1 expression\cite{24}. They first observed in a phase I study that combination of Aza with entinostat induced a significant objective response to classical anticancer agents in a small percentage of patients (4 out of 19 NSCLCs)\cite{25}. They then theorized that epigenetic drugs might restore sensitivity to anti-PD-1 therapy into specific patients. Epigenetic modifications are crucial for immunotherapy resistance, not only by controlling immune checkpoint inhibitor expression, but also by regulating immune cells infiltration within the tumor, antigen presentation or cytokine profile expression and secretion [Figure 1]. Indeed, increased recruitment

Figure 2. Crosstalk between epigenetics, PD-L1/PD-L1 expression and immunotherapy. PD-1: programed cell death 1; PD-L1: PD-1 ligand
of CD8+ tumor-infiltrating lymphocytes (TILs) are directly linked to clinical outcome in numerous cancers, such as breast cancer or ovarian cancer\textsuperscript{[26,27]}. It is now described that this infiltrate is regulated by epigenetic modifications. For example, the HMT enhancer of zeste homolog 2, which mediates the trimethylation of histone H3 lysine 27 (H3K27me3), and DNMT1 which repressed the production and secretion of CXCL9 and CXCL10 by Th1 therefore inhibited the recruitment of CD8+ T-cells within the tumor in ovarian cancer models\textsuperscript{[1]}. Moreover, decitabine was also found to increase CD80 expression on tumor cells, inducing T lymphocytes infiltration in mouse EL4 tumor model and tumor rejection\textsuperscript{[28]}. 

Increased expression of PD-1 in breast cancers and colorectal carcinoma was related to both decreased DNA methylation and decreased repressive H3K9me3 and or H3K27me3 histone marks on the PD-1 promoter\textsuperscript{[29-31]}. However, no changes in epigenetic modifications were observed on the PD-L1 promoter in these cancers, probably due to its unmethylated status in normal tissues. Similarly, the expression of the immune checkpoint inhibitors CTLA-4, TIM-3 and LAG-3 were also up-regulated in breast cancers compared to normal adjacent tissues and their promoters presented lower DNA methylation and lower repressive histone marks\textsuperscript{[39]}. Only one exception was described since the decreased expression of the immune checkpoint inhibitor LAG-3 appeared to be specifically associated to variations of histone modifications (H3K9me3 and H3K27me3)\textsuperscript{[29,32]}. Similar results showed an inverse correlation between promoter DNA methylation of PD-1, PD-L1, CTLA4 and expression of these proteins in NSCLCs compared to normal tissues\textsuperscript{[33,34]}. Moreover, methylation of the CTLA4 promoter was also inversely correlated to its expression in a large cohort (470) of malignant melanoma patients treated with anti-PD-1/PD-L1 antibodies. Furthermore, a strong significant correlation was observed between low promoter methylation of CTLA4 and both positive response to immunotherapy and survival\textsuperscript{[35]}. In prostate cancers, methylation of PD-1 promoter was more frequently observed than in normal tissues and its expression was inversely correlated to its promoter methylation\textsuperscript{[36]}. Moreover, patients presenting a high level of methylated PD-1 promoter presented a higher risk of recurrence in prostate cancer. In leukemia, PD-L1 expression has also been associated with low promoter methylation and to poor prognosis\textsuperscript{[37]}. The demethylation of an enhancer region of PD-1 and the protein re-expression was reported in CD8+ T-cells in chronic lymphocytic leukemia\textsuperscript{[38]}. In diffuse low grade glioma (LGG), methylation of PD-L1, PD-L2 and CTLA-4 promoters was inversely correlated to their expression. We also recently reported in epithelial to mesenchymal transition-induced lung cancer cell models (A549) that transforming growth factor \(\beta\) (TGF\(\beta\)), a cytokine frequently expressed in the tumor microenvironment, was responsible for the demethylation of the PD-L1 promoter by inhibiting both global and local DNA methylation\textsuperscript{[39]}. In addition, the combined treatment of these cells with TNF\(\alpha\) favored NF\(\kappa\)B signaling and PD-L1 transcription by inducing the recruitment of NF\(\kappa\)B/p65 on the PD-L1 promoter. Besides the tumor microenvironment which could severely affect PD-1/PD-L1 expression, cancer drugs might also modify their expression, thus inducing drug resistance. For example, 30% of melanoma patients treated with ICB who presented a positive response towards the protocol, developed secondary tumors (for a review\textsuperscript{[40]}). Exhausted T-cells which appeared during ICB are supposed to present an epigenetic-linked genome wide regulation mediated by the de novo DNMT3A leading to the inhibition of proliferation and metabolic activity and thus restricting clonal CD8+ T-cell diversity. Indeed, DNA methylation inhibitors improved T-cell response in anti-PD-1 therapy\textsuperscript{[41]}. On the opposite, an acquired resistance was frequently found in myelodysplastic syndrome (MDS) in which the use of DNA hypomethylating agents is the standard treatment. Inhibition of DNA methylation induces an increase in PD-1 in T-cells of these patients\textsuperscript{[42]}. Although mechanisms of resistance to demethylating agents are still unclear, it has been suggested that the activation of the immune checkpoint inhibitor was more likely involved. Moreover, treatment of NSCLC patients with conventional chemotherapy or epidermal growth factor receptor inhibitors increased PD-L1 expression unlike anti-PD-L1 treatments which significantly decreased PD-L1 expression\textsuperscript{[34]}. In addition, an anti-PD-L1 treatment of primary tumors which expressed low levels of PD-L1, induced secondary tumours presenting an increase in the methylation of PD-L1 promoter.
A recent multi-country (France, Spain and Italy) cohort study of 142 high grade NSCLC samples highlighted a DNA methylation specific signature which can significantly predict which patients could positively benefit from nivolumab or pembrolizumad treatments. DNA repair pathways have been associated with efficiency of ICB treatments. Indeed, a study analyzing the methylation status of 179 genes involved in DNA repair in HNSCC, cervical carcinoma and laryngeal squamous cell carcinoma identified 15, 3 and 2 genes, respectively, whose methylation was associated with increased PD-L1 expression. Amongst these genes, RAD51B was found in the three different cancer models suggesting that alteration of homologous DNA reparation strongly favored immune checkpoint inhibitor expression.

Only 15% of metastatic HNSCC presented an objective response to anti-PD-1/PD-L1 treatments. In these cancers, both PD-L1 and PD-L2 expression was increased in tumor cells compared to normal adjacent tissues and their expression was inversely correlated to methylation of their promoters. Interestingly, human papillomavirus-positive HNSCC were strongly associated with both PD-1 and PD-L1 promoter methylation and poor prognosis suggesting that these particular tumors are unlikely to respond to immunotherapy. Treatment of mesothelioma cells using the demethylating agent decitabine alone is unable to significantly restore PD-L1 expression but combination of decitabine with HDACi [valproic acid, suberoylanilide hydroxamic acid (SAHA) or NODH] strongly induced PD-L1 transcription in these cells. Expressions of PD-L1 and CTLA4 mRNA in acute myeloid leukemia (AML) patients treated with combined HDACi and DNMTi were also increased during treatment but presented very high oscillations. Interestingly, decitabine alone was sufficient to restore both mRNA and protein PD-L1, PD-1 and CTLA4 expression in a dose-dependent manner in AML cell lines.

**COMBINATION OF EPIDRUGS WITH IMMUNOTHERAPIES RESTORES TREATMENT EFFICIENCY**

**Cell models**

Although some studies showed a positive effect of 5-Aza treatment on PD-L1 expression in NSCLC cell lines, only 13% of NSCLC tumors issued from patients treated with this demethylating agent, presented a strong expression of PD-L1 protein (however, PD-L1 expression was at least detectable in 55% of cases). These data strongly support the hypothesis that different cellular mechanisms control PD-L1 expression. Indeed, the competitive inhibitor of DNMT1, procainamide or 5-Aza treatments alone failed to induce PD-L1 expression in the NSLCC cell line A549 whereas the combination of these compounds with TNFα reactivated PD-L1 expression. In NSCLC models (A549 and H838 cells), the DNA methylation inhibitor Aza, induced the demethylation of promoters, increased the expression of interferon regulating factor 1 (IRF1) and provoked an increase in PD-L1 expression in both a interferon-dependent and independent manner. Moreover, expression of CXCL9 and CXCL10, consecutive to the IRFs activation, improved the efficiency of anti-PD-L1 protocols in mice models.

**Pre-clinical studies**

Despite obtaining better therapeutic responses with combined molecules than single drug treatment, combined anti-PD-1 and anti-CTLA4 therapy could not limit the formation and tumor growth of CT26 and 4T1 cancer models in mice. However, HDACi significantly increased PD-L1 expression in the 4T1 cell model, and tri-therapies with the further addition of 5-Aza or entinostat fully eradicated the tumors. Indeed, the authors showed that the use of epigenetic drugs led to the elimination of the potent immunosuppressive myeloid-derived suppressor cells (MDSCs) at very low concentrations, doses generally well tolerated by cancer cells. Moreover, the addition of antibodies targeting MDSCs, or the use of a PI3K inhibitor known to reduce circulating MDSCs, in combination with immunotherapy led to the same efficiency. Similar results were observed in NSCLC cells treated with Mocetinostat (one inhibitor of HDAC I/IV classes). On the opposite, HDAC11 knockout mice presented very high levels of immunosuppressive
MDSCs suggesting that targeting only one epigenetic enzyme may lead to opposite results although molecular mechanisms remain unknown.

Many previous studies demonstrated the central role of epigenetic modifications on CD4+ T-cell polarization and therefore on cancer prognosis and outcome. Indeed, while Th1 polarization is associated with a good prognosis, Tregs (CD4+, FOXP3+) are immunoregulators whose presence in the tumor immune infiltrate is associated with a repression of CD8+ T-cells and NKs activity via the production of IL-10, IL-35 and TGFβ. Epigenetic drugs might also activate FOXP3 expression\(^{52}\) and immunotolerance and thus might limit the benefits of immunotherapy. However, it has been recently reported, in the breast cancer 4T1 and in NSCLC cells, that the use of large spectra HDACi or Mocetinostat increased PD-L1 expression and concomitantly down-regulate T-reg leading to increased CD8+ T-cell infiltration and survival in mice suggesting that Tregs might also be inhibited by epidrugs\(^{50,51}\).

**Cytosolic NADP⁺-dependent isocitrate dehydrogenase 1 and glioma**

LGGs associated with a wild type cytosolic NADP⁺-dependent isocitrate dehydrogenase 1 (IDH1) gene, present a very aggressive phenotype which is comparable to the high grade glioma glioblastoma (GBM). Moreover, the tumors with a non-mutated IDH1 locus present a more immunosuppressive context, than those carrying a mutated IDH1 gene, and expressed higher levels of PD-L1 correlated to a lower level of PD-L1 promoter methylation than the one observed in the tumors with the mutated form of the gene\(^{52,53,54}\). These data could be explained by the fact that the mutated IDH1 protein produces higher levels of 2-hydroxyglutarate, a specific sub-product which favors methylation and repression of the PD-L1 promoter. Altogether, these data suggest that some specific subtypes of LGGs may respond differently to anti-checkpoint inhibitor therapies. We could then speculate that the use of anti-PD-L1 immunotherapies may improve conventional treatment outcomes when used in patients diagnosed with a LGG carrying a non-mutated IDH1 gene but not in tumors carrying the mutated version of the gene. For these last patients, these treatments should be combined with DNA demethylating agents. In addition, expression of PD-L1 mRNA in GBM has also been associated with non-mutated IDH1 gene and poor prognosis suggesting that overall survival of these patients could be significantly improved by anti-PD-L1 therapies\(^{55}\).

**Clinical trials**

Interestingly, therapy resistance linked to demethylating agents which are used in several first line anti-cancer therapies and which induce the expression of PD-L1 could be annihilated in the second line of treatment by an anti-PD-L1 strategy. Phase I/II clinical trials including patients with different cancer origins (NSCLC, diffuse large B-cell lymphoma, metastatic or normal breast cancer, melanoma, squamous cell carcinoma of the oral cavity, colorectal cancer, HNSCC, renal carcinoma, urothelial carcinoma, lymphoma, leukemia, ovarian carcinoma, MDS) and treated with demethylating agents (5-Aza or 5-AzaCdR) or HDACi (vorinostat or entinostat) combined with ICB therapy are currently in progress and summarized in\(^{56}\).

For example, the NCT02961101 phase1/2 trial is recruiting patients to analyze the effects of a treatment combining a low dose of decitabine with anti-PD-1 in relapsed or refractory malignancies in cancer of different origins.

**PROMISING NEW COMBINATIONS OF EPIDRUGS WITH ANTI-PD-1/PD-L1**

Numerous clinical trials showed a large resistance to anti-PD-1 treatment in TNBC and these data were confirmed in mouse models since anti-PD-1 antibodies also failed to produce an objective response in TNBC xenograph models\(^{57}\). However, invalidation of the lysine demethylase LSD1 which catalyzes H3K4 and H3K9 demethylation, using specific inhibitors such as HCl-2509 or tranylcypromine, combined with anti-PD-1 antibodies significantly decreased xenograph tumor growth and metastasis and increased the number of TILs. Another study reported that contrary to breast cancer, 60% to 90% of pancreatic carcinoma expressed PD-L1\(^{54}\). Interestingly, the authors showed in pancreatic cancer cell models that this expression
was controlled by the HMT mixed lineage leukemia 1 (MML1) enzyme which catalyzes the trimethylation of histone H3 lysine 4 (H3K4me3), and that targeted inhibition of HMTs, using Verticillin A, enhanced the anti-PD-L1 response in these models.

**Tumor antigen and future epidrugs/immunotherapies combinations**

Most of tumor antigens are proteins normally expressed in the male germline lineage which are abnormally expressed in tumor cells. These tumor antigens, whose the most studied are melanoma-antigen family A and NY-ESO-1 (New York-esophageal squamous cell carcinoma-1), are presented by MHC-I on tumor cells and are targets and activators of immune cells. Interestingly, epigenetic modifications have been linked to the control of the expression of most of these antigens and demethylating agents or HDACi increased their expression in various models (for a review, see [59]). For example, in in vitro cancer cells models (glioma: U251 and mesothelioma: Meso96), repression of NY-ESO-1 gene expression was regulated by a complex mechanism involving different DNMTs and HDAC1-including complexes which were recruited sequentially [60]. Combination of anti-disialoganglioside (GD2), which targets the tumor antigen GD2, with vorinostat, one HDACi, also decreased neuroblastoma tumor growth in mice [61]. In these tumors, the HDACi treatment led to an increase in M2-like macrophage infiltration in the tumor but decreased immunosuppressive MDSCs.

Moreover, MHC gene expression is also epigenetically controlled. Indeed, hypermethylation of MHC promoters in melanoma was correlated with decreased antigen presentation. Immunoedition in cancers could directly contribute to immunotherapy resistance since this process drives the selection of stem cell-like immune cells associated with a high resistance to treatments. Indeed, the expression of the pluripotent transcriptional factor NANOG in these cells can bind and activate the HDAC1 promoter and, consequently, induce the deacetylation and repression of several target genes, such as NOXA, therefore leading to stem cell-like phenotype and treatment resistance [62]. Combining HDACi with immunotherapies in NANOG-positive tumors may therefore limit cancer relapse and become a new line of treatment.

Stimulator of interferon genes (STING), is a protein normally activated by the cyclic GMP-AMP synthase which detects exogenous cytosolic DNA following bacterial or viral cell infection or DNA release by tumor cells. Expression of the lysine demethylase KDM5B has been shown to be inversely correlated to STING, in ovarian cystadenocarcinoma, bladder urothelial carcinoma and breast invasive carcinoma and inversely correlated to CD8⁺ T-cell infiltration and survival, as well [63]. This recent study also revealed that KDM5B can demethylate histones and repress STING promoter and that this effect was specific since it was inhibited by KDM5Bi. This new set of data suggests that KDM5Bi could be tested in the future to restore T-cell infiltration and therefore improve anti-PD-1/PD-L1 treatments.

**Transgenic TCR and epidrugs combination for future therapies**

New protocols of engineering in immunology have recently allowed to develop novel cell-based immunotherapies. Adoptive transfer of TCR transgenic T-cells has become a promising approach in the field. For example, a protocol has been designed to target MART-1, which is a major antigen present in 95% of melanomas, and therefore to design engineered T-lymphocytes which produce a transgenic TCR specific of MART-1. However, it has been shown that these strategies had no effect on melanomas presenting a resistance to BRAF inhibitors. Indeed, these cells present important defects in the apoptotic machinery which impair TCR-induced apoptosis but it has also been shown that HDACi, such as SAHA, can regulate apoptotic gene expression and restore TCR-induced tumor cell death. Even if the mechanisms and pathways involved remain still unclear, preliminary results suggested that HDACi modulated the expression of apoptotic genes, restoring the susceptibility of melanoma cells to apoptosis.
CONCLUSION

Altogether, these recent data clearly demonstrate that, although immune checkpoints and particularly PD-L1 are frequently overexpressed in tumors compared to healthy tissues, some factors might still be able to modulate their expression via the regulation of DNA methylation and histone modifications of their promoters [Figure 2]. Since immunohistochemistry staining are still difficult to quantify and standardize, mainly because of different antibody specificities, various fixation times and partially subjective interpretations between different laboratories, we speculate that quantification of promoter methylation of immune checkpoint inhibitors could be a robust and reproducible alternative method to quickly select good responsive patients which could be directed towards anti-PD-L1 therapies and those who could be good candidates for combined therapy using DNA demethylating agents and immunotherapy protocols. But, further pre-clinical and clinical studies testing the combination of ICB with epidrugs and additional regulators of immune checkpoint expression will be necessary to evaluate their efficiency to bypass the immunotherapy resistance in cancers.

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