Adenosine is an important factor that controls antitumor immunity, conditioning both the innate and adaptive immune responses.\(^1\)\(^2\) The biological effects of adenosine on the immune system, which can be stimulatory or inhibitory, are mediated by four receptor subtypes. A2a and A2b are G\(_\text{i/o}\)-coupled receptors that increase intracellular cyclic AMP (cAMP) levels, hence suppressing immune responses. A1 and A3 are G\(_\text{s}\)-coupled receptors that decrease intracellular cAMP, thereby favoring cell activation. The A2a and A2b receptors (A2aR and A2bR, respectively) play a dominant role in adenosine-induced immunosuppression in a cAMP-dependent manner (Fig. 1). The stimulation of A2aR critically impairs T-cell function during activation, by reducing the production of important cytokines and chemokines, and promotes the accumulation of regulatory T cells (Tregs).\(^1\)\(^2\) In addition, A2bR activation stimulates the expansion of myeloid-derived suppressor cells (MDSCs) and induces the release of angiogenic factors.\(^1\)\(^2\)

Extracellular adenosine results from the degradation of ATP by the sequential action of two cell-surface ectonucleotidases: CD39, which hydrolyzes ATP and ADP into AMP, and CD73, which hydrolyzes AMP into adenosine (Fig. 1). CD73 is highly expressed on the surface of several types of cancer cells and immunosuppressive cells, including Tregs and MDSCs.\(^2\)\(^3\) Therefore, ATP, which is released in high quantity from malignant cells succumbing to chemotherapy or other stressful conditions, is rapidly converted into adenosine, which accumulates in the tumor microenvironment. By activating A2aR/A2bR, intratumoral adenosine favors the escape of cancer cells from immune surveillance, hence promoting tumor progression. Hence, targeting CD73 may represent one potential strategy to increase the efficacy of anticancer therapy.\(^1\)\(^3\) Reduction of extracellular adenosine as produced from CD73 on tumor or host cells improves antitumor T-cell responses (Fig. 1).\(^1\)\(^3\)

In this regard, we have recently provided new insights into the mechanisms underlying the antitumor activity of adenosine 5’-(α,β-methylene) diphosphate (APCP), a selective inhibitor of CD73, which—in mice—promotes both a T-cell and B-cell antitumor response by lowering intratumoral adenosine levels (Fig. 1).\(^4\)

Several studies report that the majority of B cells do not express CD73, although some authors have demonstrated that CD73 is expressed on a subset of memory B cells,\(^5\) suggesting that CD73-derived adenosine might regulate memory B-cell function.\(^5\)\(^6\) Although the role of CD73 in regulating B-cell function has not yet been clearly defined, we found that CD73 inhibition in mice can indirectly affect B-cell activity in vivo, via a mechanism that relies on T cell-derived interleukin-17A (IL-17A).\(^4\) Indeed, the antitumor activity of APCP was associated with the release of Th17-like cytokines into the tumor, in turn activating B cells to produce antitumor immunoglobulins (IgGs) (Fig. 1).\(^4\) Despite the controversial role of Th17 immunity in cancer, in our model the inhibition of CD73 skewed the tumor microenvironment toward a Th17-like antitumor immune response (Fig. 1).\(^4\)

Accordingly, the neutralization of IL-17A blocked the ability of APCP to inhibit tumor growth.\(^4\) Of note, in addition to a Th17 response, Th1 and CD8\(^+\) T cell-mediated immune responses were essential for APCP to reduce tumor growth in melanoma-bearing mice (Fig. 1).\(^4\) These results provide further support to the notion that CD73-generated adenosine critically regulates the function of immune cells. However, although it has been clearly demonstrated that adenosine dampens the immune response, and hence favors tumor progression, upon ligation of A2aR/A2bR, the activation of different adenosine receptors may be associated with opposite outcomes on tumor growth. In other two reports, indeed, we have observed that NK1.1\(^+\) cells\(^7\) and CD8\(^+\) T cells\(^8\) mediate the antitumor activity of another agent, the A3R agonist Cl-IB-MECA.
achieved by targeting specific adenosine receptors, such as A3R. In conclusion, the adenosinergic system opens novel therapeutic strategies, aimed at limiting immunesuppression and potentiating protective antitumor immune responses.

Disclosure of Potential Conflicts of Interest
No potential conflicts of interest were disclosed.

conditions has been shown to operate as an endogenous stimulus for the activation of the inflammasome, which in turn promotes anticancer immune responses. Accordingly, the success of some agents currently used in anticancer chemotherapy might depend—at least in part—on their capacity to promote ATP release from cancer cells. On the other hand, an efficient antitumor immune response can also be achieved by targeting specific adenosine receptors, such as A3R. In conclusion, the adenosinergic system opens novel therapeutic strategies, aimed at limiting immunesuppression and potentiating protective antitumor immune responses.

Disclosure of Potential Conflicts of Interest
No potential conflicts of interest were disclosed.

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