Myeloid-derived suppressor cells (MDSC) expand in tumor-bearing hosts and play a central role in cancer immune evasion by inhibiting adaptive and innate immunity. They therefore represent a major obstacle for successful cancer immunotherapy. Different strategies have thus been explored to deplete and/or inactivate MDSC in vivo. Using a murine mammary cancer model, we demonstrated that doxorubicin selectively eliminates MDSC in the spleen, blood, and tumor beds. Furthermore, residual MDSC from doxorubicin-treated mice exhibited impaired suppressive function. Importantly, the frequency of CD4\(^+\) and CD8\(^+\) T lymphocytes and consequently the effector lymphocytes or natural killer (NK) to suppressive MDSC ratios were significantly increased following doxorubicin treatment of tumor-bearing mice. In addition, the proportion of NK and cytotoxic T cell (CTL) expressing perforin and granzyme B and of CTL producing IFN-γ was augmented by doxorubicin administration. Of therapeutic relevance, this drug efficiently combined with Th\(_0\) or Th\(_1\)-like lymphocytes to suppress tumor development and metastatic disease. MDSC isolated from patients with different types of cancer were also sensitive to doxorubicin-mediated cytotoxicity in vitro. These results thus indicate that doxorubicin may be used not only as a direct cytotoxic drug against tumor cells, but also as a potent immunomodulatory agent that selectively impairs MDSC-induced immunosuppression, thereby fostering the efficacy of T-cell–based immunotherapy. Cancer Res; 74(1); 1–15. ©2013 AACR.
Doxorubicin is an antineoplastic drug broadly used in the treatment of hematological malignancies, soft tissue sarcomas, and several types of carcinomas including breast cancer (21). Extensive evidence has been provided that besides its direct tumoricidal activity, doxorubicin also promotes antitumor immunity (22–26). This drug has indeed been shown to induce an “immunogenic type” of tumor cell death leading to the stimulation of dendritic cell antigen-presenting function (23). Doxorubicin administration has also been reported to enhance the proliferation of tumor-specific CD8+ T cells (27), and to increase the permeability of tumor cells to granzyme B produced by cytotoxic T lymphocytes (CTL; ref. 28). However, whether this drug may impact tumor-induced immunosuppression, specifically by negatively targeting MDSC, remains unclear. In this study, we investigated the effects of doxorubicin on MDSC in the murine breast cancer model 4T1 and explored the possibility of combining this chemotherapeutic drug with immunotherapy. We demonstrated for the first time that doxorubicin eliminated MDSC by triggering apoptosis of these cells. ROS may contribute to doxorubicin-mediated elimination of MDSC.

In addition, residual MDSC from doxorubicin-treated mice were impaired in their suppressive function. Of importance, doxorubicin administration led to improved T- and natural killer cell function. The proportion of T- and natural killer (NK) cells expressing perforin and granzyme B was indeed significantly augmented following treatment of tumor-bearing mice with doxorubicin. In addition, doxorubicin increased the proliferation status of T lymphocytes and NK cells. Substantiating these observations, doxorubicin increased the proliferation of T helper (Th) 1 or with recently identified Th17 lymphocytes impaired tumor development and metastatic spreading. These findings therefore highlight a novel property of doxorubicin as a potent selective MDSC-targeting agent, which may be used to enhance the efficacy of immunotherapeutic regimens.

Patients and Methods

Patients

A total of n = 10 patients with confirmed solid cancers (2 patients with lung, one patient with ovarian, one patient with prostate, one patient with bladder, one patient with colon, one patient with stomach, one patient with kidney, one patient with pancreatic, and one patient with breast cancer) were enrolled in the study before treatment after giving written informed consent in accordance with the Declaration of Helsinki. The study was approved by the Ethic Committee of the University Hospital of Dijon. None of the patients had received chemotherapy or any other immunosuppressive treatment during the previous 3 months.

Animals and tumor cell lines

Six- to eight-week-old Balb/c and C57BL/6 mice were purchased from the National Cancer Institute (NCI). Six- to eight-week-old gp91phox−/− (C57Bl6-Cybb−/−) were purchased from Jackson Laboratory. The mice were housed and cared for according to the University of Arizona Institutional Animal and Care Guidelines and Use Committee (IACUC). The 4T1 and EM76 murine mammary tumor cell lines (Balb/c origin) and the EL4 thymoma cell line (C57BL/6 origin) were obtained from the American Type Culture Collection. 4T1-luc was generated using luciferase reporter plasmid (PGL4-51; InvivoGen). Briefly, the transfection was performed using Fugene6 reagent (Promega) according to the manufacturer’s protocol. Stably transfected cells were clonally selected in presence of G418 (800 µg/mL) and luciferase expression was confirmed using the luciferase reporter assay system and a luminometer (Femtometer FB12; Berthold Detection System), according to the manufacturer’s recommendation (Promega).

Bioluminescence imaging

Tumor-bearing animals were injected (i.p.) with α-luciferin (4.29 mg/mouse; Xenogen). Mice were anesthetized using isoflurane (1.5 L/min oxygen, 4% isoflurane) and kept in an induction chamber. Images were captured with an AMI1000 imager (Spectral Instruments Imaging). Light emission was measured over an integration time of 1 minute, 10 minutes after injection of luciferin. Luciferase activity was analyzed using the AMI1000 Software (Spectral Instruments Imaging) to quantify tumor region flux (photons per second) and to assess tumor growth.

Generation of Th1 and Th17

Naïve CD4+CD25−CD62L− T lymphocytes were isolated from the spleen of 6- to 8-week-old Balb/c mice according to the manufacturer’s instructions (Miltenyi Biotec). Cells were subsequently cultured at a concentration of 106 cells/mL in RPMI 1640 medium supplemented with 10% FBS, 100 U/mL penicillin, 100 µg/mL streptomycin sulfate, 0.5 × MEM non-essential amino acids, and 1 mmol/L sodium pyruvate. Cells were stimulated with anti-CD3+ and anti-CD28−coated beads (Invitrogen). Th1 cells were generated in the presence of interleukin (IL)-12 (10 ng/mL), IL-2 (10 ng/mL), IL-7 (20 ng/mL), and blocking anti-IL-4 antibodies (5 µg/mL; 3-day culture). For the generation of Th17 cells, naïve T cells were cultured with IL-6 (40 ng/mL), TGF-β (0.5 ng/mL), blocking anti-IFN-γ (5 µg/mL), and anti-IL-4 (5 µg/mL) antibodies for 3 days. Th17 cultures were then prolonged in fresh medium containing IL-23 (40 ng/mL) for an additional 3 days. All cytokines were purchased from R&D Systems or Peprotech.

Chemotherapy and chemoimmunotherapy

4T1 or 4T1-Luc cells were injected (1 × 106 cells) orthotopically (mammary fat pad). In some experiments, 7 and 12 days posttumor injection, mice were administered with doxorubicin (2.5 or 5 mg/kg, i.v.), cyclophosphamide (50 mg/kg, i.p.), fludarabine (50 mg/kg, i.p.), melphalan (5 mg/kg, i.p.), vincristine (1 mg/kg, i.v.), etoposide (5 mg/kg, i.v.), or control PBS. Spleens, blood, and tumors were collected 14, 17, and 23 days posttumor cell injection. For evaluating the effects of doxorubicin plus T helper lymphocyte combination therapy, mice were injected with doxorubicin (5 mg/kg) on days 7 and 12 and...
with Th1 or Th17 cells (1 × 10⁶ i.v.; 2 × 10⁶ intratumorally) on days 9 and 14 posttumor cell injection. Mice were euthanized for ethical reasons when they exhibited severe morbidity signs because of overwhelming metastatic spreading (endpoint at 4–4.5 weeks) in compliance with IACUC regulations.

**MDSC isolation**

Spleens were harvested, dissociated, and red blood cells were lysed in lysis buffer (BD Biosciences). MDSC were purified using a mouse MDSC isolation kit according to the manufacturer’s instructions (Miltenyi Biotec). The purified cells were used in other experiments.

**Flow cytometry analysis**

Cell suspensions from blood, spleens, or tumors were filtered and red blood cells were lysed. For extracellular staining, cells were incubated with the indicated combinations of antibodies (CD11b, Ly6C, Ly6G (Gr-1), CD8, CD49b, CD4, CD25, F4/80, CD11c, CCR7, and CD206) or isotype controls (1 hour, 4°C). For intracellular staining, cells were fixed and permeabilized immediately after cell surface staining according to the manufacturer’s description (Affymetrix eBioscience). For IFN-γ staining, cells were incubated with a leukocyte stimulating kit (BD Biosciences) for 4 hours. For perforin or granzyme B staining, cells were incubated with anti-CD3– and anti-CD28–coated beads (Invitrogen) and IL-2 (4 ng/mL) for 24 hours. The proliferation status of the cells was assessed by staining with Ki67 according to the manufacturer’s protocol (Ki67 detection protocol; BD Biosciences). All antibodies and isotype controls were purchased from BD Biosciences or Affymetrix eBioscience. For the detection of apoptosis, spleen, and blood samples from tumor-bearing mice, untreated or treated with doxorubicin, were dissociated and processed as described earlier. Cells were then stained with Annexin V and propidium iodide (PI), according to the manufacturer’s protocol (Apoptosis detection kit; Affymetrix eBioscience). Fluorescence data were collected on a FACScalibur or LSRFortessa (BD Biosciences). The data were analyzed using FlowJo software (Tree star Inc.).

**Suppression assays**

Spleens from naive mice were collected, dissociated, and cells were incubating on nylon wool columns (37°C, 45 minutes) after red blood cell lysis. More than 90% of the eluted cells were T lymphocytes based on T-cell receptor (TCR)β expression. These T cells were labeled with 5 mmol/L CellTrace Violet according to the manufacturer’s instructions (Invitrogen) and were plated in 96-well plates with anti-CD3– and anti-CD28–coated activation beads (Invitrogen). Isolated MDSC from 4T1 tumor-bearing mice, treated or not with doxorubicin, were then added to the culture (MDSC-to-T-cell ratio = 1:2). After 4 days, cells were harvested, stained with anti-CD4, anti-CD8, and anti-CD25 (Affymetrix eBioscience). Cell proliferation was determined by measuring the dilution of cell trace violet by flow cytometry after gating on the CD4⁺ or CD8⁺ cell populations. The proliferation index was determined using the ModFit software (Verity Software House) and percent proliferation was calculated as follows: proliferation (%) = (T⁺ − T) − (S − T)/(T⁺ − T) × 100, with T⁺ is the proliferation index of the control stimulated T cells without MDSC; T is the proliferation index of control nonstimulated T cells without MDSC; and S is the proliferation index of stimulated T cells in the presence of MDSC).

**Western blot analysis**

Cells were lysed in radioimmunoprecipitation assay buffer and sonicated. Lysates were cleared by centrifugation at 14,000 rpm and protein concentration was determined with the Thermo Scientific BCA protein assay using bovine serum albumin (BSA) as standard. Equal amounts of proteins (30 μg) were separated on 10% or 16% SDS-PAGE gels, transferred onto a polyvinylidene fluoride (PVDF) membrane (Millipore) and probed with primary Abs specific for P-STAT3, STAT3, Arginase-1, IDO, cleaved caspase-3 (Cell Signaling Company), S100A9, S100A8 (R&D Systems), and actin (Sigma-Aldrich) followed by secondary antibody (Jackson ImmunoResearch). Reactive bands were visualized by exposure to film using Super Signal Chemiluminescent Substrate (Thermo Scientific).

**ROS detection**

Spleenocytes from untreated or doxorubicin-treated mice were incubated for 30 minutes with the oxidation-sensitive dye dichlorodihydrofluorescein diacetate (DCFDA; 5 μmol/L; Molecular Probes/Invitrogen). Samples were then labeled with anti-Gr-1 and anti-CD11b antibodies. The level of ROS was detected using flow cytometry as described (29).

**Real-time PCR**

Cells were collected and total RNA was isolated using TRIzol (Invitrogen). PCR reactions were set up in 96-well plates containing 10 μL 2× IQ Supermix (Bio-Rad), 1 μL TaqMan primer/probe set (ABI), 2 μL of the cDNA synthesis reaction [10% of room temperature (RT) reaction] and 7 μL of nuclease-free water. Reactions were run and analyzed on a Bio-Rad iCycler iQ Real-Time PCR Detection System. Primers for IFN-γ and IL-17 were obtained from Applied Biosystems (Invitrogen).

**ELISA**

IL-17 and IFN-γ concentration was detected in Th1- and Th17-cell culture supernatant using ELISA according to the manufacturer’s instructions (Affymetrix eBioscience).

**Immunofluorescent staining**

Tumors and spleens were harvested and frozen. Serial sections (5 μm) were performed from each tissue and mounted. Frozen slides were first fixed in 100% cold methanol for 10 minutes. After blocking for 1 hour at RT [10% normal chicken serum in Tris-buffered saline and 0.1% Tween 20 (TBS-T)/1% BSA] slides were incubated (overnight, 4°C) with anti-Gr-1 (Affymetrix eBioscience; 0.02 mg/mL in TBS-T). Slides were washed with TBS-T and incubated with a secondary chicken anti-rat antibody conjugated with Alexa Fluor 647 for 45 minutes at RT (1:200 in TBST; Invitrogen). Slides were mounted. Frozen slides were stained with anti-CD4, anti-CD8, and anti-CD25 (Affymetrix eBioscience). Cell proliferation was determined by measuring the dilution of cell trace violet by flow cytometry after gating on the CD4⁺ or CD8⁺ cell populations. The proliferation index was determined using the ModFit software (Verity Software House) and percent proliferation was calculated as follows: proliferation (%) = (T⁺ − T) − (S − T)/(T⁺ − T) × 100, with T⁺ is the proliferation index of the control stimulated T cells without MDSC; T is the proliferation index of control nonstimulated T cells without MDSC; and S is the proliferation index of stimulated T cells in the presence of MDSC).
were washed and incubated (2 hours, RT) with a rat anti-
mouse CD11b antibody conjugated with FITC (Affymetrix
eBioscience; 0.02 mg/mL in TBS-T). Nuclear counterstaining
was performed using Sytox orange according to the manu-
facturer’s instructions (Invitrogen).Slides were mounted with
fluorescence mounting medium (Dako North America) and
visualized using a confocal microscope (Zeiss LSM 510-META
NLO).

Statistical analysis
Experiments were performed with 4 to 8 mice per groups as
indicated. Mice were individually processed and analyzed
separately unless specified otherwise. All analyses were carried
out using GraphPad Prism software (GraphPad Software).
Analyses were performed by 1- or 2-way ANOVA with a
Bonferroni or Dunnett posttest, or a paired t test where
appropriate. Statistically significant P values were labeled as
follows: ***, P < 0.001; **, P < 0.01; and *, P < 0.05.

Results
Doxorubicin selectively eliminates tumor-induced MDSC
in mice bearing established 4T1 mammary tumors
The immunomodulatory effects of doxorubicin have been
extensively studied (23, 26). However, the possible impact of
this chemotherapeutic agent on suppressive MDSC remains
elusive. To address this question, Balb/c mice were injected
with 4T1 breast cancer cells and treated with doxorubicin
(2.5 and 5 mg/kg; nocurative doses), 7 and 12 days post-
tumor implantation (Fig. 1A). We first confirmed that 4T1
tumor growth is associated with substantial MDSC expan-
sion and determined the basal proportion of MDSC in the
absence of doxorubicin therapy at different time points
following injection of tumor cells (Supplementary Fig. S1;
refs. 30 and 31). Doxorubicin significantly reduced the
proportion and absolute number of 4T1 tumor-induced
MDSC in the spleen (Fig. 1B and C) and blood (Fig. 1D)
of treated animals. It is noteworthy that although doxorubicin-
mediated elimination of MDSC was prominent on day 14
and 17 posttumor cell injection, these cells were reconsti-
tuted by day 23. These results were further confirmed by
confocal microscopy analysis of spleen sections (Fig. 1E).
Furthermore, MDSC were also depleted within the tumor
beds in mice treated with doxorubicin (Supplementary Figs.
S2A–S2C). 4T1 tumor development is primarily associated
with the expansion of granulocytic MDSC (30, 32). Our
results indicate that doxorubicin did not significantly affect
monocytic MDSC, which were low in frequency even in
untreated animals, but significantly depleted granulocytic
MDSC (Supplementary Fig. S2D). These results thus dem-
strate that doxorubicin induced elimination of tumor-
induced MDSC.

Doxorubicin minimally affects effector immune cell
populations
To optimally promote antitumor immunity, immunomod-
ulatory drugs should negatively target immunosuppressive
cells while sparing immune effectors. We therefore sought to
investigate whether doxorubicin may affect other immune
cells, specifically antitumoral effector T lymphocytes and
natural killer cells. Interestingly, the proportion of CD4+ and
CD8+ T lymphocytes was significantly increased in the
spleen and blood of doxorubicin-treated mice (Fig. 2A and
B). NK frequency increased in the blood and was not altered
in the spleen of the treated animals (Fig. 2C). Consistent
with these results, the proliferation status of CD4+ and CD8+
T cells was augmented in doxorubicin-treated animals
(Fig. 2D). This preferential targeting of MDSC resulted in a
significant increase in the ratios of effector CD8+ T, CD4+ T,
or NK cells to suppressive MDSC (Supplementary Fig. S3A).
Importantly, the proportion of NK cells and CD8+ T lymph-
ocytes expressing perforin and granzyme B was signifi-
cantly increased in doxorubicin-treated 4T1 tumor-bearing
mice (Fig. 2E and Supplementary Fig. S3B). In line with these
results, doxorubicin administration resulted in augmented
frequency of IFN-γ–producing CD8+ T lymphocytes (Fig. 2F).
However, no change in the proportion of IFN-γ–producing
CD4+ T cells was observed (Fig. 2F). It is noteworthy that
doxorubicin did not change immunosuppressive Treg fre-
quency (Supplementary Fig. S3C). Together these results
indicate that doxorubicin-mediated MDSC depletion was
associated with augmented effector immune cell prolifera-
tion and function.

Importantly, we further determined that doxorubicin was
significantly more potent at depleting MDSC than other
chemotherapeutic agents such as cyclophosphamide, fludar-
abine, melphalan (Supplementary Fig. S4A), vincristine, or
etoposide (Supplementary Fig. S4B), which exhibited limited
effects on these cells. Of note, these drugs promoted a
decrease in tumor volume similar to that induced by doxo-
rubicin. Doxorubicin was also unique at substantially
increasing the effector lymphocytes (or NK) to suppressor
MDSC ratios, indicating that it is endowed with a higher
degree of selectivity compared with these other drugs (Sup-
plementary Fig. S4A and S4B). Furthermore, even in com-
bination with cyclophosphamide, doxorubicin significantly
triggered MDSC elimination and increased T lymphocyte
frequency (Supplementary Fig. S4C).

The observations that doxorubicin reduces MDSC fre-
quency and absolute number and increases effector lympho-
cyte proportion, resulting in an increased effector T cells to
suppressor MDSC ratios, were further confirmed in the EL4
(thymoma) and EMT6 (breast cancer) mouse models (Sup-
plementary Fig. S5A and S5B). In addition, residual MDSC
isolated from EL4 or EMT6 tumor-bearing mice treated with
doxorubicin exhibited reduced suppressive function (data
not shown).

Doxorubicin preferentially triggers MDSC apoptotic
program
We next investigated the mechanisms underlying doxo-
rubicin-mediated elimination of MDSC and specifically
explored whether this drug may trigger the MDSC apoptotic
program. Flow cytometry analysis after staining of spleen
cells with anti-Gr-1, anti-CD11b antibodies, Annexin V and
PI, and gating on the Gr-1+CD11b+ population indicated
Figure 1. Doxorubicin eliminates tumor-induced MDSC. A, schematic of the experimental design followed to evaluate the effects of doxorubicin on MDSC in the 4T1 breast cancer model. Mice were injected orthotopically (mammary fat pad) with 4T1 tumor cells (1 × 10⁶). Doxorubicin (2.5 or 5 mg/kg) was administered intravenously on day 7 and 12 posttumor injection. Spleen and blood samples were harvested and evaluated on days 14, 17, and 23. B, proportion of MDSC (CD11b⁺Gr-1⁺) in the spleen of 4T1 tumor-bearing mice post-doxorubicin treatment (right) and representative flow cytometry analysis 17 days posttumor injection (left). C, absolute number of MDSC in tumor-bearing mice treated or not with doxorubicin. D, proportion of MDSC in the blood of tumor-bearing mice after doxorubicin treatment. E, confocal microscopy analysis of CD11b⁺Gr-1⁺ cells in the spleens from untreated or doxorubicin-treated mice 17 days posttumor injection (5 days after the last doxorubicin treatment). CD11b (green), Gr-1 (red), and sytox orange nuclear staining (Nuc, blue). Scale bar, 20 μm. *, P ≤ 0.05; **, P ≤ 0.01; ***, P ≤ 0.001. n = 4 mice per group. Data represent one of 3 experiments performed and analyzed independently.
that doxorubicin increased the proportion of apoptotic (Annexin V⁺, PI⁻) and secondary necrotic (Annexin V⁺, PI⁺) MDSC (Fig. 3A). Consistent with these data, caspase-3 cleavage was detected in MDSC isolated from 4T1 tumor-bearing doxorubicin-treated animals (Fig. 3B). Importantly, in line with the results depicted in Fig. 2A and B, the number of dead (apoptotic and necrotic) T lymphocytes (CD4⁺ or CD8⁺) was not significantly modified by doxorubicin (Fig. 3C). These data thus demonstrate that doxorubicin preferentially induced apoptosis of MDSC with no detectable toxic effect on effector T lymphocytes.

MDSC isolated from untreated tumor-bearing mice were more sensitive to doxorubicin in vitro than the MDSC-depleted cell population or than 4T1 cells (Fig. 3D and data not shown). Therefore, to examine the effects of doxorubicin in vivo, a similar experimental design was performed. Consistent with these results, doxorubicin increased the proportion of apoptotic (Annexin V⁺, PI⁻) and secondary necrotic (Annexin V⁺, PI⁺) MDSC (Fig. 3A). Caspase-3 cleavage was detected in MDSC isolated from 4T1 tumor-bearing doxorubicin-treated animals (Fig. 3B). Importantly, the number of dead (apoptotic and necrotic) T lymphocytes (CD4⁺ or CD8⁺) was not significantly modified by doxorubicin (Fig. 3C). These data thus demonstrate that doxorubicin preferentially induced apoptosis of MDSC with no detectable toxic effect on effector T lymphocytes.

Figure 2. Doxorubicin increases the frequency, proliferation, and cytotoxic activity of effector T lymphocytes and NK. A similar experimental design as described in Fig. 1 was followed. A, frequency of CD4⁺ T cells in the spleen (left) and blood (right) of tumor-bearing mice after doxorubicin treatment. B, proportion of CD8⁺ T cells in the spleen (left) and blood (right) of tumor-bearing mice after doxorubicin treatment. C, NK cell frequency in the spleen (left) and blood (right) of doxorubicin-treated mice. D, analysis of Ki67 expression after gating on CD4⁺, CD8⁺ T lymphocytes, or NK (DX5⁺) cells as indicated (left) and related mean fluorescent intensity (MFI; right; day 17). E, percent of CD8⁺ T and NK cells expressing granzyme B or perforin in the spleen of doxorubicin-treated mice 17 days posttumor cell injection. F, percent of CD3⁺, CD4⁺, and CD8⁺ T lymphocytes expressing IFN-γ in the spleen of tumor-bearing mice treated or not with doxorubicin (day 17). *: P < 0.05; **: P < 0.01; ***: P < 0.001. n = 4 mice per group. Data represent one of 3 experiments performed and analyzed independently.
shown). Treatment of these purified MDSC with N-acetylcycteine (NAC, an ROS scavenger) prevented their killing by doxorubicin, suggesting that ROS may play a role in this process (Fig. 3D). Consistent with this result, the levels of ROS were increased in the immediate hours following doxorubicin treatment of MDSC in vitro (Fig. 3E), and MDSC isolated from EL4 tumor-bearing gp91phox−/− mice (lacking the gp91phox glycosylated subunit of the NADPH oxidase flavocytochrome b558, responsible for the production of superoxide ion O2−) were less sensitive to doxorubicin-mediated cytotoxicity in vitro (Fig. 3F). Interestingly, in vivo, the effects of doxorubicin administration on MDSC were partially impaired in EL4 tumor-bearing gp91phox−/− mice compared with wild-type mice (data not shown).
Doxorubicin impedes the suppressive activity of residual MDSC isolated from tumor-bearing mice

A cardinal characteristic of MDSC is their ability to suppress the activation and proliferation of T cells. MDSC depletion by doxorubicin was not complete as ~15% of residual MDSC could still be detected 17 days after doxorubicin administration. Therefore, it was important to determine whether the suppressive function of these remaining MDSC was affected by doxorubicin. As expected and previously reported (32), MDSC isolated from untreated tumor-bearing mice significantly inhibited T lymphocyte (CD4+ and CD8+) proliferation (Fig. 4A) and activation (Fig. 4B). The suppressive function of residual MDSC isolated from doxorubicin-treated tumor-bearing animals was significantly impaired (Fig. 4A and B). The suppressive function of residual MDSC isolated from doxorubicin-treated tumor-bearing animals was significantly impaired (Fig. 4A and B). The suppressive function of residual MDSC isolated from doxorubicin-treated tumor-bearing animals was significantly impaired (Fig. 4A and B).

Various mechanisms have been implicated in MDSC suppressive function including the production of ROS or the expression of arginase-1 or IDO involved in the catabolism of arginine or tryptophan, respectively (1). Depletion of these amino acids from the microenvironment results in the inhibition of T-cell proliferation, notably through the downregulation of the ζ-chain of the TCR complex (1, 11). Similarly, ROS exhibit suppressive effects on T lymphocytes (1, 33).

Five days after doxorubicin treatment of 4T1 tumor-bearing mice, the production of ROS (Fig. 5A) and the expression of arginase-1 (Fig. 5B) and IDO (Fig. 5C) by residual MDSC was impaired. Additional molecules reported for their role in MDSC development and/or immunosuppressive function such as CD73, CD39, S100A8/9, or STAT-3 were not significantly altered by doxorubicin treatment (Supplementary Fig. S7A and S7B).
Previous reports have indicated that some chemotherapeutic drugs such as docetaxel can promote MDSC differentiation into macrophages (M1; ref. 20). To evaluate this possibility, MDSC from doxorubicin-treated mice were analyzed for the expression of markers expressed by M1 (CCR7) or M2 (CD206; Mannose Receptor) macrophages. The expression of these 2 cell surface markers was not detected on MDSC following doxorubicin treatment, indicating that this drug did not promote MDSC differentiation into macrophages (Supplementary Fig. S7C). Finally, further investigation indicated that the proliferation status (Ki67 expression) of residual 4T1 tumor-induced MDSC was reduced by doxorubicin therapy (not shown).

Altogether, these results indicate that 5 days after doxorubicin administration, MDSC that have not been eliminated by the drug exhibited reduced expression of arginase-1 and IDO, decreased ROS level, and curtailed immunosuppressive activity.

Figure 5. Doxorubicin decreases ROS production and arginase-1 and IDO expression by MDSC. A similar experimental design as described in Fig. 1 was followed. Spleens were harvested 5 days after the last doxorubicin treatment (day 17). A, analysis of ROS production by MDSC in tumor-free (tumor-free) or in tumor-bearing (tumor-bearing) mice treated with the indicated concentration of doxorubicin. Cells were incubated with DCFDA. Representative flow cytometry analysis of gated CD11b$^+$Gr1$^+$ cells positive for DCFDA (top). Percent of MDSC positive for DCFDA (left bottom). Mean fluorescent intensity representing ROS level in MDSC from the indicated groups (right bottom). B and C, Western blot analysis depicting expression of arginase-1 (B) or IDO (C) in MDSC isolated from doxorubicin-treated or -untreated mice. *, $P < 0.05$; **, $P < 0.01$; ***, $P < 0.001$. $n = 4$ mice per group. Data represent 1 of 3 experiments performed and analyzed independently.
Doxorubicin acts synergistically with Th1 or Th17 cell therapy

Doxorubicin depleted tumor-induced MDSC and curtailed the suppressive function of residual MDSC, thereby averting one major mechanism of cancer-mediated immunosuppression. Moreover, the proliferation status and activation of responder effector lymphocytes and NK were restored in doxorubicin-treated mice. We therefore reasoned that this drug may create a favorable environment that may allow for successful combinatory immunotherapy. To address this hypothesis, we evaluated a chemo-immunotherapy regimen consisting of doxorubicin followed by infusion of T helper lymphocytes.

Th17 lymphocytes represent a recently described subset of T helper cells with controversial effects on tumor development. Recent reports have demonstrated that Th17 generated in vitro have the potential to promote the development of CD8⁺ T lymphocyte-dependent immune response and to impair tumor growth (34). Th1 can also promote antitumor immunity (35). In our current study, mice bearing established 4T1 tumors were treated with doxorubicin and received either Th1 or Th17 lymphocytes generated in vitro from naive CD4⁺ T cells (Supplementary Fig. S8A and S8B) as depicted in Fig. 6A. The chemoimmunotherapeutic regimen significantly reduced the number of 4T1 metastatic nodules in the lungs and impaired tumor growth when compared with the monotherapies (Fig. 6B and Supplementary Fig. S8C). Importantly, MDSC depletion persisted in mice treated with doxorubicin plus Th1 or Th17 cells, whereas these cells eventually re-expanded post-treatment in mice receiving doxorubicin alone (Fig. 6C and D). Consistent with these results, the frequency of CD8⁺ and CD4⁺ T lymphocytes was significantly increased in doxorubicin plus Th1- or Th17-treated animals (Fig. 6D), but NK proportion remained unchanged (data not shown). These data thus indicate that doxorubicin administration resulted in the promotion of a favorable environment fostering the antitumoral efficacy of Th1 and Th17 lymphocytes.

Doxorubicin induces apoptosis of MDSC isolated from patients with cancer

To determine whether similar effects of doxorubicin can be observed on human MDSC, CD33⁺ cells were isolated from the blood of patients with different types of cancer (n = 10). CD33⁺ cells exhibited a phenotype consistent with that reported for human MDSC (Fig. 7A; ref. 36) and importantly were endowed with significant suppressive capabilities (Fig. 7B). Our results indicate that these immunosuppressive cells were sensitive to doxorubicin-induced cell death (Fig. 7C and E). Importantly, CD33⁺ depleted cells (CD33⁻) from the same patients were significantly less sensitive to doxorubicin (Fig. 7D and E), indicating that this chemotherapeutic molecule preferentially targets MDSC. CD3⁺ T cells were also minimally affected by doxorubicin (not shown). These results thus suggest that doxorubicin exhibit effects on human MDSC comparable to those observed on mouse MDSC, thus supporting the implementation of this drug in clinical chemoimmunotherapeutic approaches.

Discussion

The development of malignant tumors is commonly associated with the occurrence and persistence of an immunosuppressive environment. The expansion of MDSC, a main suppressive cell population (13), has been widely documented in many animal tumor models as well as in patients with different types of cancers and represents a major obstacle for efficient cancer immunotherapy (5, 9, 10). Several strategies have been explored to either eliminate or curtail the immunosuppressive function of MDSC. Depletion of Gr-1⁺ MDSC using anti-Gr-1 monoclonal antibody resulted in restored T-cell antitumor activity. However, anti-Gr-1 also led to the elimination of mature granulocytes and was associated with severe immunosuppression (13). The promotion of MDSC differentiation using ATRA, a natural metabolite of vitamin A, has also been considered. ATRA administration enhanced T lymphocyte function and fostered the efficacy of cancer vaccines (14). In recent years, evidence has been provided that chemotherapeutic drugs can not only kill tumor cells, but also enhance antitumor immunity through different mechanisms (37). Chemotherapeutic agents can promote the function of antigen presenting cells, NK, and T lymphocytes (23, 26, 37) but may also negatively target immunosuppressive cells such as Treg or MDSC (17, 19, 20, 38, 39). For instance, gemcitabine has been reported to deplete MDSC in tumor-bearing mice, resulting in enhanced antitumor immunity (17–20). Additional reports have demonstrated that 5-Fluorouracil (19), or docetaxel (20), can eliminate, impede the suppressive function, or promote the differentiation of MDSC.

Doxorubicin has been a key chemotherapeutic agent used against a variety of human cancers. More recently, this drug has been widely studied for its ability to modulate anticancer immunity (23–26, 28). Reports have demonstrated that doxorubicin induces an “immunogenic type” of tumor cell death and promotes antitumor immune responses (23). In this study, we highlight a novel property of doxorubicin: its ability to avert a major mechanism of tumor-induced immunosuppression by eliminating and inactivating tumor-induced MDSC. We established that doxorubicin administration reduced the number of MDSC in the spleen, blood, and in the tumor beds of animals bearing established 4T1 mammary tumors. Importantly, doxorubicin selectively eliminated MDSC while enhancing the proliferation status, activation, cytokine production of effector T lymphocytes and/or NK cells. This primary targeting of MDSC while effector T cells (and NK) were spared translated into a substantial increase in effector lymphocytes to MDSC ratios. Compared with the other chemotherapeutic drugs that we evaluated, doxorubicin was endowed with the highest degree of selectivity. Indeed, although associated with a reduction of tumor volume comparable to that observed after doxorubicin treatment, most of these other agents had limited effects on MDSC and some of them, such as cyclophosphamide, even led to a slight increase in MDSC as they targeted T cells.
Figure 6. The combination of doxorubicin and Th1 or Th17 impairs 4T1 tumor development. Mice were injected orthotopically (mammary fat pad) with 4T1 tumor cells ($1 \times 10^6$). Doxorubicin (6 mg/kg) was injected intravenously on day 7 and 12 posttumor cell injection. Th1 or Th17 lymphocytes were administered on day 9 and 14 posttumor cell injection, intravenously ($1 \times 10^6$) and intratumorally ($2 \times 10^6$). Tumor volume and number of metastatic nodules were evaluated on day 19 posttumor injection. A, schematic of the experimental design. B, number of metastatic nodules (left) and tumor volume (right). C, representative flow cytometry analysis of MDSC frequency in mice administered with the indicated therapies. D, proportion of MDSC, CD4+, and CD8+ T cells in mice receiving the indicated therapies; n = 8 mice per group. Data are representative of 3 independent experiments. *, P ≤ 0.05; **, P ≤ 0.01; ***, P ≤ 0.001.
Figure 7. Doxorubicin selectively kills MDSC isolated from patients with cancer. CD33\(^+\) cells were isolated from cancer patient PBMCs by magnetic cell sorting. A, phenotypic analysis of the isolated cells. Representative results of \(n = 10\) patients. B, ability of the CD33\(^+\) purified cells to impair the proliferation of cell trace violet-labeled T lymphocytes induced with anti-CD3 and anti-CD28–conjugated microbeads (at the indicated MDSC to T-cell ratios). pi, proliferation index. C and D, purified CD33\(^+\) MDSC or CD33\(^-\) cells were exposed to the indicated concentrations of doxorubicin for 24 hours and stained with Annexin V and PI (% dead cells = % of PI\(^+\) + % AnnexinV\(^+\)PI\(^+\) cells). A total of \(n = 10\) patients were analyzed. E, representative dot plots obtained with CD33\(^+\) MDSC or CD33\(^-\) cells isolated from a patient with cancer. (*, \(P \leq 0.05\); **, \(P \leq 0.01\); ***, \(P \leq 0.001\)).
Of therapeutic relevance, the addition of doxorubicin to cyclophosphamide resulted in a significant elimination of MDSC similar to that observed when doxorubicin was administered alone. This peculiar property of doxorubicin further highlights the interest and advantage of using this drug as a potent immunomodulatory agent in chemoinmunotherapeutic approaches.

Further investigation indicated that doxorubicin selectively triggered the apoptotic program of MDSC. No change in the apoptosis rate of T or NK cells was detected post-doxorubicin treatment compared with untreated groups. This selective effect may be partly explained by the preferential targeting of highly proliferative cells by doxorubicin. Indeed, MDSC in untreated tumor-bearing mice expressed high level of Ki67, a marker of cell proliferation (data not shown), whereas T lymphocyte proliferation was low in untreated tumor-bearing mice. The triggering of apoptosis by doxorubicin through the induction of ROS production has been described in tumor and normal cells such as cardiomyocytes (40). In vitro, the ROS scavenger NAC impaired doxorubicin-mediated elimination of MDSC, and MDSC isolated from gp91phox−/− mice were less sensitive to doxorubicin. The effects of doxorubicin on MDSC were partially impaired in EL4 tumor-bearing gp91phox−/− mice. These results strongly suggest that, early after its administration, doxorubicin may induce ROS-dependent triggering of the MDSC apoptotic program, resulting in the rapid elimination of most of thesecells. Of note, MDSC that were not eliminated and that were found later (5 days) after doxorubicin treatment exhibited an overall impaired suppressive activity, which, at this time was associated with a reduction of ROS production.

A previous report has indicated that docetaxel reduced MDSC number by promoting their differentiation into M1 macrophages (20). We did not detect any changes in the expression of M1 or M2 macrophage markers on MDSC following doxorubicin treatment. In addition, no change in the frequency of macrophages or DC was detected in the treated mice. However, as mentioned earlier, the suppressive function of the limited number of residual MDSC that were not depleted by doxorubicin was significantly impaired.

Because doxorubicin administration induced MDSC depletion and was associated with restored T lymphocyte activity, we reasoned that it may create a favorable environment for efficient immunotherapy. Infusion of in vitro generated Th1 cells has been reported to promote antitumor immunity (35). Th17 lymphocytes have recently emerged as a new effector CD4+ T helper cell subset (41) exhibiting effector functions distinct from Th1 and Th2 lymphocytes (42). Th17 have been identified as major contributors to the pathogenesis of multiple autoimmune conditions in animals and humans (42). However, the role of Th17 cells in cancer remains controversial. Although some studies have documented the antitumoral efficacy of these cells (34–43), others have reported on their immunosuppressive properties (46). These conflicting results related to the protumoral versus antitumoral properties of Th17 may be explained by the high degree of plasticity of these cells (47–49). Supporting the antitumoral role of in vitro polarized Th17, Muranski and colleagues demonstrated that adoptive Th17 cell therapy has the potential to eliminate established tumors. The antitumoral efficacy of Th17 lymphocytes depended on their ability to produce both IFN-γ and IL-17 (43, 44). Consistent with these studies, our own results indicated that Th17 generated from OT-II mice exhibit antitumoral effects against B16-OVA melanoma (unpublished data). In this study, we demonstrated that the therapeutic efficacy of adoptively transferred Th1 or Th17 lymphocytes was significantly enhanced by doxorubicin administration, resulting in impaired development of the highly metastatic 4T1 mammary carcinoma.

Doxorubicin is widely used in chemotherapeutic regimen primarily for its conventional direct tumoricidal activity. Here we highlight a new application for this drug as a selective MDSC-targeting agent, which can be used to overcome a major mechanism of tumor immune evasion. These results advocate for the implementation of doxorubicin in combination strategies to enhance the efficacy of immunotherapy.

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

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