Neutrophils, the most abundant type of leukocytes in blood, can form neutrophil extracellular traps (NETs). These are pathogen-trapping structures generated by expulsion of the neutrophil’s DNA with associated proteolytic enzymes. NETs produced by infection can promote cancer metastasis. We show that metastatic breast cancer cells can induce neutrophils to form metastasis-supporting NETs in the absence of infection. Using intravital imaging, we observed NET-like structures around metastatic 4T1 cancer cells that had reached the lungs of mice. We also found NETs in clinical samples of triple-negative human breast cancer. The formation of NETs stimulated the invasion and migration of breast cancer cells in vitro. Inhibiting NET formation or digesting NETs with deoxyribonuclease I (DNase I) blocked these processes. Treatment with NET-digesting, DNase I-coated nanoparticles markedly reduced lung metastases in mice. Our data suggest that induction of NETs by cancer cells is a previously unidentified metastasis-promoting tumor-host interaction and a potential therapeutic target.

INTRODUCTION

Breast cancer metastasis is associated with very high mortality rates. Cancer cells can acquire the ability to metastasize by expressing metastasis-promoting genes, such as epithelial to mesenchymal transition–promoting transcription factors or metalloproteinases (1, 2). However, cancer cells can also recruit and activate leukocytes, including macrophages, to promote metastasis (3). Neutrophils, the most abundant leukocytes in human blood, similarly promote metastasis (4–8), although they can kill disseminated cancer cells under certain conditions (9). Neutrophils and their precursors are sensitive to many chemotherapy regimens, causing dangerously low neutrophil numbers (neutropenia) during the course of treatment. Because neutropenia carries a risk of life-threatening infections, the American Society of Clinical Oncology recommends prophylactic treatment with neutrophil-stimulating factors, including granulocyte colony-stimulating factor (G-CSF), for certain chemotherapeutic regimens (10). It is therefore important to determine the conditions under which neutrophils promote metastatic spread.

Neutrophils’ normal function is to kill harmful microorganisms in three ways: (i) phagocytosis, a process whereby bacteria or fungi are engulfed and digested; (ii) degranulation of cytotoxic enzymes into the extracellular space; and (iii) neutrophil extracellular traps (NETs), which are DNA meshes with associated cytotoxic enzymes that are released into the extracellular space where they trap microorganisms (11). NETs form in tissues and have been documented in human pancreatic, liver, and gastric cancers (12–14), but whether they participate in cancer progression remains unclear. NETs can also form intra-vascularly, and they can damage vascular cells when they form (15). Recently, it was shown that NETs induced within the vasculature by experimentally induced systemic bacterial infection or surgical stress aided in metastatic seeding of cancer cells in the liver (5, 12).

We sought to observe how disseminating cancer cells interacted with neutrophils upon arrival in the lungs, a major site of metastatic colonization in breast cancer. We developed confocal intravital lung imaging (CILI), a modification of a lung imaging approach used with two-photon microscopy (16). Here, we show that NET-like structures form around disseminated cancer cells in lungs using CILI. We also show that cancer cells stimulate neutrophils to form NETs in the absence of pathogens in vitro. Finally, we show that NETs stimulate cancer cell migration and invasion and that treatment with NET-digesting deoxyribonuclease I (DNase I)–coated nanoparticles inhibits metastasis. NET formation is a mechanism by which signals from cancer cells activate host cells to enhance metastasis. Understanding the contribution of neutrophils to metastases has pressing clinical implications because many cancer patients receiving chemotherapy also receive prophylactic treatment with neutrophil-stimulating factors.

RESULTS

Metastatic cancer cells induce formation of NETs

To investigate whether neutrophils play a role in metastasis, we first compared neutrophil infiltration into tumors from orthotopically transplanted 4T1 and 4T07 murine breast cancer cells. These cells originate from the same mammary tumor of a BALB/c mouse, but only the 4T1 cells metastasize (17). We observed significantly (P = 0.0009) more neutrophils in primary 4T1 tumors than in 4T07 tumors (Fig. 1, A and B). Because the chemokine CXCL1 can recruit neutrophils (18), we
Fig. 1. NETs form during metastasis of breast cancer.  
(A and B) More neutrophils infiltrated metastatic 4T1 tumors than nonmetastatic 4T07 tumors (Ly6G immunostaining; mean ± SEM; n = 4 mice, t-test). Scale bar, 50 μm. (C) CXCL1 protein level was higher in 4T1 than in 4T07 tumors (mean ± SEM; n = 5 mice, t-test). (D) Cancer cell-derived CXCL1 promoted metastatic seeding after intravenous injection of luciferase-expressing cells [bioluminescence radiance, 19 days after cell injection; individual mice and means indicated; P = 0.0008, one-way analysis of variance (ANOVA); P < 0.01, Dunnett’s multiple comparison]. (E) CXCL1 secretion by 4T1 cells did not affect primary tumor growth in nude mice (mean ± SEM; n = 5 mice). (F) Knockdown of Cxcl1 in 4T1 cells reduced lung metastasis in nude mice (individual mice and means ± SD indicated; t-test). (G and H) NET-like structures of extracellular DNA, sensitive to intravenous DNase I, were found around 4T1 cancer cells in LysM-EGFP mice using CILI. Grayscale insert shows DNA channel (shown and quantified 30 to 60 min after cancer cell injection; values from individual mice and mean ± SD are indicated; P = 0.002, one-way ANOVA; P < 0.01, Sidak’s multiple comparison). Scale bar, 50 μm, FOV, field of view. (I and J) Extracellular DNA and neutrophil elastase (NE) activity colocalized near 4T1, but not 4T07, cancer cells (shown and quantified 30 to 60 min after cancer cell injection; Fisher’s exact test). Scale bars, 50 μm. (K and L) The number of NETs in the lungs was higher after 4T1 cell injection than in controls (colocalized myeloperoxidase and citrullinated histone H3 immune staining). White arrow, NET; yellow arrow, intact neutrophil (mean ± SEM; n = 3 mice). Scale bar, 10 μm.
measured its mRNA and protein and found higher amounts in 4T1 cells than in 4T07 cells and in 4T1-derived tumors than in 4T07-derived tumors (Fig. 1C and fig. S1). Reduction of CXCL1 (and the homologous CXCL2) in 4T1 cells with short hairpin RNAs reduced neutrophil infiltration into tumors and increased primary tumor growth but had no effect on macrophage infiltration or cancer cell proliferation in vitro (fig. S1). Tumors with reduced expression of CXCL1/2 had an approximately doubled tumor burden; despite this, metastatic burden was not increased (fig. S1). Instead, metastatic burden from CXCL1/2 knockout cells was significantly (P = 0.0008) decreased when equal numbers of cancer cells were injected intravenously into mice (Fig. 1D). We speculated that increased tumor burden after CXCL1/2 knockout was caused by reduced recruitment of tumor-reactive T lymphocytes (19). Consistent with this idea, CXCL1/2 knockdown did not increase primary tumor growth in T cell–deficient, nude mice (Fig. 1E). However, CXCL1/2 knockdown significantly (P = 0.03) reduced spontaneous metastasis in nude mice (Fig. 1F). This suggests that reduced metastasis is not caused by reduced T cell recruitment and that neutrophils mediate this effect.

The results of assays with CXCL1/2 knockdown cells suggest that neutrophils support metastasis upon their recruitment by cancer cells. To investigate the functions of neutrophils at the metastatic site, we performed CILI of mCherry-expressing 4T07 and 4T1 cells injected intravenously into LysM-EGFP mice, which express EGFP (enhanced green fluorescent protein) predominantly in neutrophils (20). Greater numbers of extracellular DNA species sensitive to intravenously injected DNase I were found around 4T1 than around 4T07 cancer cells (Fig. 1, G and H, and movie S1). To explore whether these structures could be NETs, we next used CILI to determine whether the extracellular DNA structures colocalized with the NET-associated protease neutrophil elastase using a probe that produces a fluorescent signal after specific cleavage by neutrophil elastase. Neutrophil elastase activity colocalized with extracellular DNA in 14 of 17 imaged fields of view after the injection of 4T1 cancer cells but not after the injection of 4T07 cells (P = 0.0008; Fig. 1, I and J). To further explore whether NETs are formed after the injection of cancer cells, we performed immunofluorescence staining for NETs in tissue sections from lungs of mice injected with 4T1 cancer cells. This analysis confirmed that NETs formed in lungs shortly after tail vein injection of 4T1 cells and that the amounts of NETs were elevated for days after the injection (Fig. 1, K and L, and fig. S2). These data demonstrate that metastatic cancer cells stimulate NET formation at sites of dissemination in the absence of infection.

To determine whether NETs could also be found in metastatic human breast cancer, we performed immunofluorescence staining on a small panel of clinical samples of primary tumor and matched metastatic lung lesions from breast cancer patients (Fig. 2A). We detected NETs in 16 of 20 primary tumors and in 13 of 19 metastatic lung lesions (Fig. 2B). The number of NETs varied between tumors, with the highest numbers in triple-negative tumors (six of six primary tumors and seven of seven metastases had detectable NETs) and an absence or very low numbers in luminal breast cancers (Fig. 2C). Thus, we detected NETs in human breast cancer and found the presence of NETs to be associated with an aggressive subtype of breast cancer.

NETs promote cancer cell migration and invasion

To determine whether cancer cell–induced NET formation required direct interaction with neutrophils or could be induced over a distance, we performed a Transwell chamber assay. Specifically, neutrophils isolated from the mouse bone marrow were plated in the lower wells, and cancer cells were added on top of Matrigel-coated membranes in the upper wells (fig. S3). Neutrophils cocultured in this manner with metastatic 4T1 cells formed extensive NETs, whereas neutrophils similarly cocultured with nonmetastatic 4T07 cells formed few NETs (Fig. 3, A and B).

Coculturing with neutrophils increased the invasion of 4T1 cells but had little effect on the invasion of 4T07 cells (Fig. 3C). To test whether NETs promoted cancer cell invasion, we digested extracellular DNA by adding DNase I to the cultures (Fig. 3, D and E). The neutrophils’ ability to stimulate invasion of 4T1 cells was lost when the DNA of the NETs was digested, whereas the addition of DNase I had no effect on fetal calf serum (FCS)–stimulated invasion (Fig. 3F).

To test whether other metastatic cancer cells also induce NETs, we isolated primary cancer cells from C3(1)-Tag mice, a genetically engineered mouse model of metastatic basal/triple-negative breast cancer (21). Tumors in this model have high numbers of infiltrating neutrophils and high expression of CXCL1 (22). Like 4T1 cells, C3(1)-Tag primary cancer cells induced NETs when cocultured with mouse neutrophils (Fig. 3G). C3(1)-Tag cells have a limited ability to invade through Matrigel, but neutrophils significantly (P = 0.047) increased the migration of C3(1)-Tag cells across uncoated membranes. This increased migration was blocked by DNase I treatment (Fig. 3H). Extensive NETs were also formed when triple-negative human BT-549 breast cancer cells were cultured with neutrophils from healthy female volunteers (Fig. 3, I and J, and fig. S3). As observed with 4T1 cells, digesting NETs with DNase I blocked the neutrophils’ ability to stimulate BT-549 invasion through Matrigel (Fig. 3K). Thus, both human and mouse triple-negative breast cancer cells can promote the formation of promigratory/invasive NETs, whereas treatment with NET-digesting DNase I blocks migration and invasion induced by species-matched neutrophils.

Cancer cells induce lytic NETs, and blocking NET formation reduces invasion

Next, we wanted to determine how tumor-induced NETs were formed. G-CSF can prime neutrophils for NET formation and is secreted by 4T1 cells (23). Anti-G-CSF blocking antibodies significantly (P = 0.03) reduced the ability of 4T1 cells to induce NETs in the Transwell chamber assay (Fig. 4A). Conversely, human recombinant G-CSF–induced NET formation by human neutrophils (fig. S4A). Together, this suggests that G-CSF secreted by cancer cells induces NETs.

An initial step in the intracellular signaling cascade in neutrophils that results in pathogen-induced NET formation is the activation of the phagocytic NADPH (reduced form of nicotinamide adenine dinucleotide phosphate) oxidase (NOX2) enzyme complex (24). In our Transwell assay, apocynin, a pharmacologic inhibitor of NOX2, reduced NET formation and neutrophil-promoted cancer cell invasion (Fig. 4, B and C). The phox47 subunit is a critical component of NOX2 in neutrophils. The extension of NETs was greatly reduced for neutrophils isolated from p47phox–/– compared to those isolated from p47phox+/+ mice, and p47phox–/– neutrophils did not promote cancer cell invasion (fig. S4). Unfortunately, the p47phox–/– strain is on the C57BL/6 background, precluding in vivo experiments with the BALB/c–derived 4T1 cells.

Next, we determined the importance of peptidylarginine deiminase type 4 (PAD4), the enzyme responsible for histone modifications required for decondensing neutrophil DNA before expulsion (25). The
PAD4 inhibitor CI-amidine reduced NET formation (Fig. 4D) and blocked the neutrophils’ ability to promote invasion (Fig. 4E). Thus, metastatic cancer cells activate signaling pathways that include NADPH oxidase and PAD4 in neutrophils, causing the formation of invasion-promoting NETs.

Several types of NET formation can occur. One requires NADPH oxidase activity and generally results in neutrophil lysis 2 to 4 hours after activation (24), whereas another process occurs rapidly (5 to 60 min), independent of NADPH oxidase activity, and leaves the neutrophil plasma membrane intact (26). Cancer cell–induced NET formation was dependent on NADPH oxidase activity, which suggested that it occurred through a lytic process. To address this directly, we performed electron microscopy. Scanning electron microscopy revealed that extracellular meshes extruded from nonintact neutrophils 3 hours after plating and stimulation with cancer cells, whereas intact neutrophils were rarely discernible (fig. S5). These findings were largely similar to those observed after stimulation with phorbol 12-myristate 13-acetate (PMA) (fig. S5), which is known to induce NETs through a lytic mechanism (24). We next analyzed the integrity of the neutrophils by transmission electron microscopy. Three hours after stimulation with cancer cells, neutrophils typically had decondensed chromatin dispersed in the cytosol and showed plasma membrane breakdown (fig. S5). This morphology was identical to the morphology of the neutrophils induced to form NETs by PMA. Of more than 50 evaluated neutrophils, none showed evidence of DNA-containing vesicles budding from the nucleus, which is the morphology described as typical for nonlytic NET-forming neutrophils (26).

Finally, to demonstrate that the extracellular meshes observed by scanning electron microscopy contained DNA, we performed immunogold labeling with anti-DNA antibodies and detected expelled DNA in the meshes after stimulation with 4T1 cancer cells (fig. S5). Because we did not observe neutrophils with extruded nuclear DNA that were otherwise intact, we conclude that, in our in vitro setting, cancer cells induce NETs through a lytic process.

NETs are defined by the association of neutrophil proteases with the extracellular neutrophilic histone-bound DNA (27). Hence, we speculated that the proinvasive effects of NETs required NET-associated protease activity. Inhibition of cathepsin G, a protease associated with NETs, blocked the neutrophils’ ability to promote invasion with no effect on the cancer cells’ ability to invade toward FCS. Surprisingly, inhibition of cathepsin G also reduced the extension of NETs (Fig. 5, A and B). This suggests that cathepsin G is required for NET release.

Cleavage of histones by neutrophil elastase is required for chromatin decondensation and NET release (28, 29), and cathepsin G may have a similar role. The neutrophil elastase inhibitor sivelestat also reduced the extension of cancer cell–induced NETs (Fig. 5C). Although the effect of sivelestat on neutrophil-induced invasion was not significant in assays with murine 4T1 cells (Figs. 5D), inhibition of neutrophil elastase significantly (P = 0.02) blocked the neutrophils’ ability to stimulate the invasion of human BT-549 breast cancer cells, as did the inhibition of NADPH oxidase (Fig. 5E).

The proinvasive effects of NET-forming neutrophils in Transwell chamber assays suggested that a factor (or factors) generated during NET formation acted on cancer cells. To test this possibility, we generated a conditioned medium (CM) from unstimulated neutrophils, from neutrophils induced to form NETs by coculturing with cancer cells, and from neutrophils cocultured with cancer cells in the presence of the NADPH oxidase inhibitor apocynin to prevent the formation of NETs. CM from cultures induced to form NETs effectively stimulated cancer cell invasion, whereas CM from cultures where NET formation...
Figure 3. Formation of NETs by metastatic 4T1 breast cancer cells is associated with cancer cell invasion. (A and B) 4T1 but not 4T07 cells increased the formation of NETs (immunostaining for histone H3 and neutrophil elastase; mean ± SEM; n = 3, t test). Scale bar, 50 μm. (C) Neutrophils promoted invasion of 4T1 but not 4T07 cells (mean ± SEM; n = 5, t test). (D and E) DNase I (1.5 U) digested NETs (mean ± SEM; n = 3, t test). Scale bar, 50 μm. (F) DNase I treatment inhibited neutrophil-stimulated invasion of 4T1 cells (mean ± SEM; n = 5 to 7, t test). (G) Primary C3(1)-Tag cancer cells induced NETs. Scale bar, 50 μm. (H) DNase I treatment inhibited neutrophil-stimulated migration of C3(1)-Tag cancer cells (mean ± SEM; n = 3, t test). (I and J) Human BT-549 breast cancer cells promoted NET formation (immunostaining for histone H3 and myeloperoxidase; mean ± SEM; n = 3, t test). Scale bar, 50 μm. (K) DNase I (1.5 U) treatment blocked neutrophil-stimulated invasion of BT-549 breast cancer cells (mean ± SEM; BT-549 cells only or BT-549 cells with neutrophils and vehicle or DNase I, n = 5; BT-549 cells and vehicle or DNase I in 10% FCS, n = 2).

was inhibited by apocynin did not (Fig. 5F). To exclude the possibility that invasion was stimulated by factors secreted by cancer cells into the CM, we also used CM from neutrophils that formed NETs in response to PMA stimulation. CM from PMA-induced, NET-forming neutrophils also stimulated cancer cell invasion. CM from both cancer cell–induced and PMA-induced NET-forming neutrophil cultures lost the ability to stimulate invasion when neutrophils were cultured in the presence of DNase I (fig. S6). In addition, the CM lost its ability to stimulate invasion when DNase I was added after collection or upon heat denaturation (fig. S6). This suggests that the proinvasive factor is not simply extracellular DNA, but protein factors associated with the DNA mesh.

DNase I treatment prevents lung metastasis in mice

Digesting bacterially induced NETs with systemically administered DNase I can reduce experimental metastasis in a mouse model of
metastatic lung cancer, but the effects are relatively modest (5). Because we found that DNase I prevented NET-mediated invasion and migration in vitro with three different sources of cancer cells (Fig. 3, F, H, and K), we speculated that the modest in vivo effects of DNase I were due to its short half-life in blood (30). Immobilization of an enzyme on the surface of nanoparticles can increase enzyme stability (31); therefore, we developed DNase I-coated nanoparticles. In vitro, DNase I-coated nanoparticles digested NETs and blocked invasion as effectively as free DNase I (Fig. 6A). In vivo, a higher concentration of DNase I was found in plasma when mice were treated with DNase I–coated nanoparticles than with free DNase I (Fig. 6B). Daily intraperitoneal injection of DNase I–coated nanoparticles significantly (P = 0.002) reduced metastatic burden after intravenous injection of 4T1 cells (Fig. 6, C and D). Three of 9 mice treated with DNase I–coated nanoparticles had no detectable metastasis by histology, whereas all 10 mice treated with control nanoparticles had macroscopic or microscopic metastases. A detailed analysis of the tissue revealed that there was a significant reduction in the number of detectable, individual metastatic foci (P = 0.0003; Fig. 6E) and a reduction in the average size of the foci (P = 0.01; Fig. 6F) after treatment with DNase I–coated nanoparticles. This suggests that NETs are critical for metastatic colonization. NETs that form intravascularly in response to infection can promote extravasation of disseminated cancer cells (5). We therefore examined whether DNA digestion with DNase I–coated nanoparticles or inhibition of NET formation with the PAD4 inhibitor Cl-amidine influenced the number of cancer cells that made it out of the vasculature in our metastasis model, which does not involve infection. Despite a significant (P = 0.02) reduction in the number of NETs using either approach (fig. S7A), the number of cells that extravasated into the lung tissue 24 hours after injection was not altered (fig. S7B).

To determine whether DNase I–coated nanoparticles also reduced spontaneous metastasis, we injected 4T1 cells into both inguinal mammary glands of female mice, and after 7 days, daily treatment with DNase I–coated nanoparticles began. The DNase I–coated nanoparticles had no effect on primary tumor growth (Fig. 6G), but the lung metastatic burden was reduced: all mice treated with control nanoparticles had micrometastases, but three of six mice treated with DNase I–coated nanoparticles had no detectable metastases by histology (Fig. 6H). Together, these data suggest that therapeutic targeting of NETs can prevent lung metastasis.

**DISCUSSION**

Here, we show that cancer cells can hijack neutrophils, so that the neutrophils’ ability to eradicate pathogens through formation of NETs instead aids metastatic spread. By imaging in live mice, we...
observed that breast cancer cells can induce neutrophils to form NETs after they arrive in the lungs. We also documented the presence of NETs in the aggressive triple-negative subtype of human breast cancer. Using in vitro models, we further demonstrated that inhibiting the signaling pathways that promote NET formation or digesting NETs with DNase I blocks invasion. Our data also suggest that cathepsin G is not just an effector of NET functions but is also involved in the release of NETs, similar to what was reported for neutrophil elastase (28). Finally, we demonstrate that treatment with DNase I–coated nanoparticles effectively reduces metastasis.
Fig. 6. Targeting NETs in vivo reduces metastasis. (A) DNase I–coated nanoparticles reduced neutrophil-stimulated cancer cell invasion in vitro (mean ± SEM; n = 4, t test). (B) Injection of DNase I–coated nanoparticles results in higher plasma nuclease activity than injection of free DNase I (mice injected with 75 U of free DNase I, 75 U of nanoparticle-bound DNase I, or equivalent vehicle; mean ± SEM; n = 3 mice). (C and D) DNase I–coated nanoparticles (75 U per mouse) reduced experimental lung metastasis of 4T1 cells (mean ± SEM; n = 9 to 10 mice, t test). Arrows point to metastases. Scale bars, 4 mm. (E and F) DNase I–coated nanoparticles (75 U per mouse) reduced the number and the size of metastatic foci arising after injection of 4T1 cells [mean ± SEM; n = 9 to 10 mice, t test; data were transformed by taking the square root before performing the t test due to significantly (P = 0.0003) different variances (untransformed data graphed)]. (G) DNase I–coated nanoparticles did not affect primary tumor growth. Nanoparticle treatment was initiated 7 days after tumor cell transplantation (mean ± SEM; n = 6 mice). (H) DNase I–coated nanoparticles (75 U per mouse) reduced spontaneous metastasis of 4T1 cells [mean ± SEM; n = 6 mice, t test; data were transformed by taking the square root before performing the t test due to significantly (P = 0.007) different variances].
in vivo. Unexpectedly, cancer cell–induced NETs did not promote extravasation but did influence the number of histologically detectable metastatic foci. This suggests that NETs mediate the expansion of disseminated cells. Our findings, together with a previous report that NETs induced by infection aid in metastatic seeding of the liver (5), raise the exciting possibility of targeting NETs to prevent metastasis.

Neutrophils and their precursors are sensitive to chemotherapy, and cancer patients can thus develop life-threatening neutropenia. Notably, long-term survival of breast cancer patients receiving chemotherapy is higher for patients with mild chemotherapy-induced neutropenia (32). It has been proposed that this is because neutropenia represents a surrogate marker of adequate chemotherapy dosing for a given patient (33). However, our findings suggest an alternative hypothesis: mild neutropenia is associated with better survival because neutrophils play a functional role in metastasis. It could therefore be important to develop approaches to prevent or dissolve prometastatic NETs while simultaneously leaving the life-saving degranulating and phagocytic activities of neutrophils intact. Blocking antibodies against G-CSF reduced cancer cell–induced NET formation in our study and can reduce metastases in mice (34, 35). However, G-CSF is used clinically to prevent mortality associated with neutropenic infections and cannot be easily targeted. Inhibitors against NOX2 and neutrophil proteases also blocked cancer cell–induced NET formation. Unfortunately, NOX2 and the neutrophil proteases are not good targets because NOX2 is critical for bacterial killing and neutrophil proteases are required to eliminate pathogens by phagocytosis and degradation. However, the extracellular DNA mesh of NETs is an exciting and feasible target. Treatment with nuclease was reported to reduce metastasis decades ago, but with no insights as to why (36). DNase I treatment is approved by the U.S. Food and Drug Administration for the treatment of cystic fibrosis, for which it is used to decrease mucus viscosity resulting from NET accumulation triggered by persistent infections. Our results with DNase I–coated nanoparticles serve as a proof of principle that NETs are drug targets to reduce metastasis. Another strategy may be to prevent NETs from forming, for example, by targeting PAD4. Although we were able to prevent NET formation using PAD4 inhibitors in vivo in short-term assays, we could not test the effect on metastasis because long-term treatment would be needed. Commercially available PAD4 inhibitors have serum half-lives of ~15 min to 4 hours (37, 38). It was therefore impossible to compare NET digestion by DNase I–coated nanoparticles with inhibition of NET formation using PAD4 inhibition and establish which approach is the best strategy for preventing metastasis.

Results of our experimental metastasis assays suggest that NETs play a role during the establishment of metastases. They may do so through multiple mechanisms. The DNA mesh can trap circulating cancer cells at the site of dissemination (5). In addition, intravascular NETs can increase local vascular permeability (15), which would permit cancer cells to extravasate more easily. This role has been proposed for systemically formed NETs. However, our data do not support a role for cancer cell–induced NETs in enhancing extravasation. Instead, our data suggest that NETs may stimulate further invasion of the cells into the tissue and the expansion of the colonizing cells. The DNA mesh of NETs may promote invasion by acting to concentrate NET-associated proteases or protecting proteases from the endogenous inhibitors abundant in blood. We found that CM from NET-forming cultures can promote invasion but that both DNase I digestion and heat denaturation can abolish this effect. This suggests that the DNA mesh is important for mediating the effect, but because denaturation does not degrade DNA, it is not simply free DNA that stimulates invasion. Possibly, a chemotactic factor(s) is associated with NETs. HMGB1 released during NET formation may drive proliferation and migration of cancer cells (12). Because we found NETs in primary human tumors as well as metastases, another important future direction will be to elucidate whether NETs also participate in cancer cell migration and invasion in the primary tumor.

Here, we show that three different types of cancer cells can induce promigratory or proinvasive NETs, but cancer cells can acquire the ability to invade and migrate through many other means. Therefore, assays to identify patients who may benefit from NET–targeted treatments are needed. Detection of NETs in tumor biopsies may be the most accurate method of identifying patients who might benefit from NET–targeting approaches. However, because we identified G-CSF as a critical factor in the induction of NETs by cancer cells, it is possible that cancer cell expression of G-CSF could identify patients at risk of developing NET–promoted metastasis. Both 4T1 cells and C3(1)-Tag tumors express high levels of G-CSF (22, 35), and they are both models of triple-negative breast cancer. Higher amounts of G-CSF are found in human patient samples of triple-negative breast cancers than of other subtypes (34). Our analysis of clinical samples also shows that NETs are found in highest amounts in this breast cancer subtype.

In conclusion, we demonstrated that cancer cells can hijack the physiological processes of microorganism killing through NET formation to promote metastasis. This concept for how cancer cells exploit host cells represents a therapeutic opportunity to prevent metastasis, which accounts for the great majority of cancer-associated deaths.

MATERIALS AND METHODS

Study design

The objective of this research was to test the hypothesis that cancer cell–induced NETs promote invasion and metastasis in vitro and in vivo models of cancer. Animals were excluded from the study when the initial intravenous injection of the cancer cells missed the tail vein. Endpoints for animal experiments were selected before the conduction of the experiments as the time when the first animals in any experimental group had a weight loss of >10%. The number of replicates for specific experiments is listed in the figure legends. The experiments testing the effects of DNase I on invasion in vitro were repeated by three different coauthors. The investigators who assessed, measured, or quantified the results of invasion through Matrigel, NET-like structures by CILI, or metastatic burden after treatment with DNase I–coated nanoparticles were blinded to the specific intervention.

Animals

BALB/c mice were purchased from Charles River Laboratories, p47phox−/− (39) and ACTB-ECFP mice (40) were purchased from the Jackson Laboratory, and C3(1)-Tag mice (21) were purchased from the National Institutes of Health (NIH) mouse repository. LysM-EGFP mice (20) were provided by M. Looney (University of California, San Francisco). All procedures were approved by the Cold Spring Harbor Laboratory Institutional Animal Care and Use Committee and were conducted in accordance with the NIH Guide for the Care and Use of Laboratory Animals.

In vitro invasion, migration, and NET formation assays

Neutrophils (2.5 × 10⁶ cells) in 750 μl of serum-free medium were seeded on poly-L-lysine–coated coverslips (BD Biosciences) in 24-well
plates for 15 to 30 min before adding anti–G-CSF (R&D Systems), human recombinant G-CSF (R&D Systems), apocynin (Abcam), cathepsin G inhibitor I (EMD Millipore), CI-amidine (Cayman Chemical), sivelestat (Tocris Bioscience), DNase I (AM2222, Invitrogen or 10104159001, Roche), or DNase I–coated nanoparticles. Cancer cells (1 × 10^4 4T1, 4T07, or BT-549 cells) in serum-free medium were then added to rehydrated Matrigel inserts (BD Biosciences). After 22 hours at 37°C, the noninvading cancer cells from the upper surface of the membrane were wiped off, and cells on the bottom side of the membrane were fixed with 4% paraformaldehyde (PFA) and stained with hematoxylin. The number of invading cells was counted under a light microscope in six fields of view, and the average number of cells was calculated.

CM was prepared from neutrophil cultures grown alone or with cancer cells (4T1 or BT-549) present in the upper chamber of the Transwells, the latter with or without DNase I or apocynin added to the bottom well. The CM from cocultures of neutrophils and cancer cells was also heat-denatured (65°C for 15 min or 95°C for 5 min), or DNase I was added to the collected CM. The CM was added to the lower chamber, and cancer cells were seeded in serum-free medium in the upper chamber of Matrigel-coated cell culture inserts. CM was also prepared from neutrophil cultures treated with 20 nM PMA (Sigma) or without DNase I (Invitrogen). The CM was added to the lower chamber, and 4T1 cells were seeded in serum-free medium in the upper chamber of Matrigel-coated cell culture inserts. CM was also prepared from neutrophil cultures treated with 20 nM PMA (Sigma) or without DNase I (Invitrogen). The CM was added to the lower chamber, and 4T1 cells were seeded in serum-free medium in the upper chamber of Matrigel-coated FluoroBlok cell culture inserts. After 22 hours, the cells were stained with 1 μM SYTO13 (Molecular Probes), and the number of invading cells was counted in 10 random fields of view using a fluorescence microscope.

To test the effect of NETs on C3(1)-Tag primary cancer cell migration, tumors (~10 to 12 mm in diameter) were resected from C3(1)-Tag mice intercrossed with ACTB-ECFP, and cancer cells were isolated as described (41). After growth to subconfluency, 5 × 10^4 cells in Dulbecco’s modified Eagle’s medium (DMEM) containing 0.5% FCS were added to a FluoroBlok cell culture insert with an 8-μm pore size (Corning). The plate was incubated for 2 hours to allow cells to adhere, and then neutrophils (2.5 × 10^5 cells) were seeded on poly-L-lysine–coated coverslips (BD Biosciences) in the lower chamber with serum-free DMEM. The cancer cell medium was then replaced by serum-free DMEM. After 22 hours, the number of migrating cells was quantified similarly to the experiments performed with neutrophil CM.

Neutrophils grown on coverslips in the lower chamber were fixed with 4% PFA, permeabilized with 0.5% Triton X-100 in phosphate-buffered saline (PBS), and incubated in 1× blocking buffer (5% goat serum and 2.5% bovine serum albumin in PBS) for 1 hour. Cells were stained with primary antibodies in blocking buffer using anti–neutrophil elastase (1:100 dilution; rabbit polyclonal antibody, ab68672, Abcam) and either anti–histone H1 (1:30 dilution; ab62884, Abcam) or anti–histone H3 (1:50 dilution; #3680, Cell Signaling Technology). After washing with PBS, the cells were stained with fluorochrome-conjugated secondary antibodies (1:150 dilution; Invitrogen) in 0.5× blocking buffer and finally stained with 4′,6-diamidino-2-phenylindole (0.05 mg/ml; Invitrogen). NET formation was determined as the percentage of the field of view positive for a histone signal.

In vivo treatment with DNase I–coated nanoparticles
To test the effect of DNase I–coated nanoparticles on experimental metastasis, female BALB/c mice were treated with DNase I–coated nanoparticles (75 U per mouse) or control, uncoated nanoparticles intraperitoneally. Two hours later, 1 × 10^5 4T1-luciferase (experiment 1) or 4T1 cells (experiment 2) were injected intravenously. Daily nanoparticle treatment was continued for 2 weeks, and the mice were euthanized when the first mouse showed >10% weight loss (33 and 26 days after cancer cell injection for experiments 1 and 2, respectively). Metastatic burden was quantified from lung sections as described above, and results from the two experiments were pooled. Notably, nanoparticles were not given intravenously because the repeated injections would damage the tail veins, making it difficult to continue treatment through the entire period. The biodistribution of intraperitoneally and intravenously administrated nanoparticles is similar (42).

To test the effect of DNase I–coated nanoparticles on spontaneous lung metastasis, 4T1 cells (2.5 × 10^5) were injected into both inguinal mammary glands of female BALB/c mice. One week later, daily intraperitoneal injections with DNase I–coated nanoparticles (75 U per mouse) or uncoated nanoparticles were initiated and continued until 19 days after cancer cell transplantation. Tumor growth and lung metastatic burden were quantified as described above.

Statistical analysis
All statistical analyses were performed using GraphPad Prism software versions 5 and 6. Data were analyzed using two-sided t tests or one-way ANOVA, as indicated in the figure legends with an α value of 0.05. For statistical analyses of the metastatic burden from orthotopically growing tumors after treatment with DNase I–coated nanoparticles and of the effect of DNase I–coated nanoparticles on the size of metastatic foci, the data were first transformed by taking the square root because the variance of the untransformed data was significantly (P = 0.007 and P = 0.0003, respectively) different between the two groups. The number of sampled units, n, is indicated in the figure legends.

SUPPLEMENTARY MATERIALS
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Materials and Methods
Fig. S1. CXCL1 promotes neutrophil recruitment to tumors.
Fig. S2. NET formation is present in lungs of mice after injection of 4T1 cancer cells.
Fig. S3. Neutrophils were harvested for in vitro assay.
Fig. S4. G-CSF and p47phox promote NET formation and cancer cell invasion.
Fig. S5. Breast cancer cells induce the formation of lytic NETs.
Fig. S6. CM from neutrophils induced to undergo NET formation promotes invasion.
Fig. S7. Cancer cell–induced NETs do not promote extravasation.
Movie S1. NET-like structures form in vivo around 4T1 cancer cells in the lung.

REFERENCES AND NOTES

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Cancer cells induce metastasis-supporting neutrophil extracellular DNA traps

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Metastasis caught in a NET

Neutrophil extracellular traps, or NETs, are DNA structures that are produced by neutrophils in response to infection and can promote the spread of cancer in the presence of infection. Park et al. discovered that even in the absence of infection, metastatic breast cancer cells can stimulate neutrophils to form NETs, which further support the spread of metastasis. The authors also demonstrated an approach to breaking this vicious cycle using nanoparticles coated with DNase I, an enzyme that breaks down DNA NETs. This treatment was effective in reducing lung metastases in mice, demonstrating the potential of NETs as a therapeutic target.