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Dynamic NETosis is Carried Out by Live Neutrophils in Human and Mouse Bacterial Abscesses and During Severe Gram-Positive Infection

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Abstract

Neutrophil extracellular traps (NETs) are released, as neutrophils die *in vitro*, in a process requiring hours, leaving a temporal gap for invasive microbes to exploit. Functional neutrophils undergoing NETosis have not been documented. During Gram-positive skin infections, we directly visualized live PMN *in vivo* rapidly releasing NETs, which prevented bacterial

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Author Contributions

B.G.Y. and B.P. designed the overall study, performed experiments, analyzed the data and wrote the manuscript, D.S. designed, performed and analyzed the TEM experiments, C.N.J. designed, performed and analyzed the immunofluorescence microscopy experiments, B.N.V.S performed intravital imaging, L.D.Z performed intravital imaging, Xenogen and dissemination experiments and analyzed the data, K.P. performed human neutrophil experiments, M.A. performed *in vitro* NETs assays, K.W. developed the transgenic bacteria, H.C.M helped develop the imaging analysis protocols, S.E.M and A.B.C designed and performed the human cytoplasm experiments, K.Z. designed and supervised the development of the transgenic bacteria and provided expert advice on microbiology, J.C. designed the clinical experiments, supervised the acquisition of patient samples and clinical histories and provided expert advice on infectious diseases and microbiology, P.K. provided overall supervision, analyzed the data and wrote the manuscript.

dissemination. NETosis occurred during crawling thereby casting large areas of NETs. NET-releasing PMN developed diffuse decondensed nuclei ultimately becoming devoid of DNA. Cells with abnormal nuclei displayed unusual crawling behavior highlighted by erratic pseudopods and hyperpolarization consistent with the nucleus being a fulcrum for crawling. A combined requirement of Tlr2 and complement mediated opsonization tightly regulated NET release. Additionally live human PMN developed decondensed nuclei and formed NETS *in vivo* and intact anuclear neutrophils were abundant in Gram-positive human abscesses. Therefore early in infection, non-cell death NETosis occurs *in vivo* during Gram-positive infection in mice and humans.

Gram-positive bacteria such as *Staphylococcus aureus*¹⁻⁴ and *Streptococcus pyogenes*⁵ are highly invasive pathogens causing severe skin infections and sepsis in humans. Neutrophils are critical for host survival and NETs may represent a novel defense mechanism⁶⁻⁹. *In vitro* NETs are a cell-death event occurring hours after initial stimulation^{10,11}; however questions remain concerning NETosis *in vivo*. Conventional PMN killing involves cell recruitment, emigration, chemotax and phagocytosis of microbes^{12,13}. However chasing individual microbes is ineffective to deal with rapidly disseminating Gram-positive pathogens. Utilizing both phagocytosis and NETosis might be counter-productive if neutrophils engulf bacteria and then lyse releasing all their bacterial contents. Thus non-lytic NETosis may prevent bacteria from escaping following phagocytosis. To date NETosis has only been observed in immobile and incapacitated cells. *In vitro*, we described rapid NET release (< 10 min) without cell lysis, however the fate and function of these PMN could not be evaluated^{14,15}. In this study we hypothesize that rapid NET release may be carried out *in vivo* by live viable PMN.

To understand the relationship between conventional PMN-mediated and NET-mediated host defense, we directly visualize PMN behavior during Gram-positive skin infections in mice and humans. Using spinning-disk confocal intravital microscopy (SDCIM) surprising insights, not in line with the current NETosis paradigm, were revealed. Cell death and lytic release of NETs were not observed *in vivo*, nor was nuclear condensation and budding off *en bloc*, observed in red blood cell enucleation¹⁶. Instead, live functional PMN crawled, phagocytosed and rapidly NETosed simultaneously, resulting in widespread tissue NETs that limited bacterial dissemination. NETosis occurred in emigrated PMN during localized infection and this was tightly regulated through Tlr2 and complement mediated opsonization. NETosing PMN crawled with erratic pseudopod formation and extreme cell morphologies due to the loss of the nuclear structure acting as a fulcrum for the cytoskeleton^{17,18}. Importantly, human neutrophils released NETs in an *in vivo* model and patients with acute Gram-positive abscesses demonstrated intact NETosing PMN identical to the animal model. Here we describe an alternate fate of *in vivo* NETosing PMN that is distinct from cell-death NETosis.

Results

Gram-positive bacteria induced rapid NET formation *in vivo*

S. aureus infected mouse skin was imaged using SDCIM and the cell-impermeable DNA specific dye SYTOX Orange was used to visualize the NETs (Method 1) (Fig. 1a). We observed sheets of DNA, in contrast to the thin strands of *in vitro* NETs (15 nm to 100 nm)^{6,11,19} (Fig. 1a and Supplementary Video 1). Chemokine induced sterile inflammation recruited PMN but NETs were not released (Fig. 1a), despite undergoing the same surgical procedure. Emigrated PMN crawled through infected tissues thereby widening NET distribution (Supplementary Video 1). NETs were inducible between 1×10^6 to 1×10^8 colony-forming units (CFUs). Acute infection did not result in dead PMN as demonstrated by lack of SYTOX uptake.

NETs were not visible in blood vessels within the infected skin; therefore muscle was imaged for improved visualization of the intravascular compartment (Supplementary Fig. 1a) NETosis was not observed, consistent with the view that systemic infection is required for intravascular NETs. Emigrated PMN (peritoneal) had increased NET release, without cell lysis, compared to non-emigrated PMN (marrow) exposed to bacteria in an *in vitro* assay (Supplementary Fig. 1b). We hypothesize that emigration may enhance DNA release and prevent NET spread through the blood, thereby decreasing bystander damage.

NETs are rapidly released from the nucleus

To confirm authentic *in vivo* NETs (histones + granular proteins + DNA^{9,20}), a second assay was developed using specific fluorescently conjugated antibodies to visualize NET constituents (Method 2). Significant release of histones and neutrophil elastase occurred within minutes in response to Gram-positive human pathogens (*S. aureus* and *S. pyogenes*) (Fig. 1b,c). Negligible NETs occurred during sterile inflammation. Histone stained areas expanded in large sheets (Fig. 1c, Supplementary Fig. 2a and Supplementary Video 2) identical to extracellular DNA. Injection of dead, washed bacteria resulted in NET release (Fig. 1b), demonstrating innate recognition is sufficient independent of active toxin release.

Mitochondrial DNA NETs exist^{21,22}, however *in vivo* mitochondrial imaging showed unaltered intracellular mitochondria during infection (Supplementary Fig. 2a,b). This does not exclude mitochondrial DNA, however the nucleus seems to be the major source of NETs in our model of infection.

Using a third technique, *in vivo* PMN nuclei were pre-stained with a cell-permeable DNA specific dye (SYTO 60, Method 3) allowing nuclear imaging throughout NETosis. Without infection, SYTO 60 is restricted to normal multilobar distinct nuclei however, abnormal PMN with indistinct diffuse nuclear remnants were observed during infection (Fig. 1d). The ratio of extracellular to intracellular DNA was quantified to demonstrate the increase in NETs during infection (Fig. 1d).

NET formation is tightly regulated and requires both Tlr2 and C3

NETs may cause bystander injury, therefore we hypothesized that release is tightly regulated. Two critical systems involved in host defense against *S. aureus* are Tlr2 and complement²³. NETs were absent in *Myd88*^{-/-} animals demonstrating Toll dependence (Supplementary Fig. 3a). Furthermore, both *Tlr2*^{-/-} and *C3*^{-/-} animals were incapable of releasing histones (Fig. 1e) or nuclear DNA (Fig. 1f) despite recruiting normal numbers of PMN (Supplementary Fig. 4a). Normal mouse serum restored NETosis in *C3*^{-/-} animals indicating the importance of opsonization (Fig. 1f). As *Tlr2*^{-/-} mice have intact complement and the *C3*^{-/-} mice have intact Tlr2, it is clear that neither of these molecules is sufficient to cause NETs in isolation. Collectively these data suggest a multi-tiered regulation for which Tlr2 and complement are essential but not individually sufficient.

Tlr2 stimulation did not induce NETs (Supplementary Fig. 3c). In preliminary experiments, the C3a receptor played a greater role in NET production compared to the C5a receptor (Supplementary Fig. 3b), however, recombinant C3a alone or in addition to PAM-3-CYS (Tlr2 stimulant) had no effect on NETosis (Supplementary Fig. 3c), revealing that purified molecules could not mimic a pro-bono Gram-positive infection.

Viable NET-forming PMN Remain Functional *in vivo*

PMN did not take-up SYTOX (indicating death) during NETosis. We pre-labeled the neutrophil nucleus *in vivo* (Method 3) to understand if NETosing neutrophils continue to function. We observed a stark contrast between normal and NETosing PMN using SDCIM and 3D reconstruction (Fig. 2a). To visualize the innermost structures, we have adjusted the levels of 3D transparency. Non-NETosing PMN have intact, fully intracellular nuclei (left panel). The NETosing PMN has released a NET that surrounds the entire outer membrane (middle panel)(Fig. 2a). Here the NET is made transparent in order to see the underlying PMN. Remarkably, the NETosing PMN remains viable and continues to chemotax towards a GFP-bacterium (Supplementary Video 3). Further 3D images of PMN releasing intracellular DNA are shown (Supplementary Fig. 5a).

To directly visualize PMN releasing intracellular DNA that becomes a NET we developed a 4-color SDCIM strategy combining the cell permeable DNA dye SYTO 60 and the cell impermeable dye SYTOX Orange. Live PMN were visualized that contained diffuse nuclear staining with concurrent release of NETs. 3D imaging confirmed that the PMN with extracellular DNA (red) contained a diffuse abnormal nucleus (right panel)(Fig 2a). PMN that crawled while releasing NETs were quantified for velocity, meandering index and crawling track (Supplementary Fig. 5b–d and Video 4). Therefore NETosis is not limited to immobile and incapacitated cells.

Live NETosing PMN develop decondensed nuclei and become anuclear

Neutrophil nuclear swelling and changes in morphology predict chromatin decondensation and nuclear envelope breakdown^{6,14,19,24,25}. We observed three different nuclear morphologies of emigrated PMN exposed to *S. aureus*. Normal nuclei were distinct demarcated multilobar rings with uniform DNA-staining (Fig. 2b, Supplementary Fig. 6a and Video 5). The normal nucleus was malleable, but retained its structure during crawling.

Diffuse nuclei lacked structure and had a heterogeneous, lower intensity staining pattern consistent with breakdown of the nuclear envelope, chromatin decondensation and intracellular chromatin dispersion (Fig. 2b, Supplementary Fig. 6b and Supplementary Video 5). Anuclear PMN had absent nucleic acid staining (Fig. 2b, Supplementary Fig. 6b and Supplementary Video 5). These observations were not due to photobleaching as all three nuclear phenotypes were observed in the same fields of view, over the same duration and laser intensity. To confirm structural nuclear differences, we used confocal z-stacks and 3D reconstruction (Fig. 2c). Completely anuclear PMN by 2D imaging were completely devoid of a nucleus on 3D reconstruction and no longer contained any stainable nucleic acids suggesting these cells had completed the NETosing process (Fig. 2c).

GFP-histone transgenic mice (Tg(HIST1H2BB/EGFP)1Pa) confirmed *in vivo* anuclear PMN (Supplementary Fig. 7a). The PMN on the left has a normal nucleus, whereas the PMN on the right has no detectable nucleus. PMN with normal and diffuse nuclei continued to crawl within the tissues. Flow cytometry of untreated animals reveal all PMN are uniformly and brightly positive for a nucleus prior to infection (Supplementary Fig. 7b). Bone marrow chimera experiments (GFP-expression restricted to hematogenous cells) support a subpopulation of PMN within infected tissue capable of surviving and functioning without a nucleus (Supplementary Fig. 7c).

To understand if NETosis affects other host-defense functions we quantified phagocytosis of live bacteria. In wild type animals approximately 50% of PMN phagocytosed bacteria (Fig. 2d). Although NETosis and phagocytosis were closely linked, phagocytosis alone was not sufficient to induce NETosis as *Tlr2*^{-/-} PMN took up bacteria without nuclear changes (Fig. 2e). *C3*^{-/-} mice could not phagocytose and did not have altered nuclei, or make NETS, however this impairment was reversed with normal serum demonstrating the importance of opsonization (Fig. 2e).

Unusual cellular phenotypes of NETosing PMN

Unique patterns of crawling and cell morphologies were observed during NETosis. Normal PMN with multilobar nuclei crawl with a single lead pseudopod (Fig. 3a). In contrast PMN with diffuse nuclei demonstrate hyperpolarization and multiple pseudopods (Fig. 3a). To test if the loss of the nucleus explains the bizarre crawling phenotype, we studied a well-defined induced anuclear human PMN cytoplasm²⁶⁻²⁹ (Fig. 3a). Indeed cytoplasm exhibits crawling behaviors similar to NETosing PMN suggesting that the intact nucleus provides the structural fulcrum for normal crawling similar to other mobile cells^{17,18}. Crawling was further analyzed in *Tlr2*^{-/-} and *C3*^{-/-} PMN. As anticipated *Tlr2*^{-/-} and *C3*^{-/-} PMN crawled with a normal single lead pseudopod and maintained morphology similar to non-NETosing PMN with normal nuclei (Fig. 3a).

Polarization is normal for crawling cells and *in vitro* these cells become 1.5 to 2× longer than their width, the limitation of this value being the width of the nucleus. However in infected tissue PMN that had diffuse nuclei became hyperpolarized (Fig. 3b and Supplementary Fig. 4b). Both the NETosing PMN and the cytoplasm generated multiple pseudopods simultaneously, often at obtuse angles (90–180 °), a phenotype never observed in sterile inflammation. During sterile inflammation, the majority of PMN crawled with one

dominant pseudopod, which rarely split to yield 2 pseudopods at an acute angle ($< 90^\circ$). Infection increased pseudopod formation, which was more common in cells with altered nuclei (Fig. 3c). *Tlr2*^{-/-} and *C3*^{-/-} PMN did not form multiple pseudopods. Despite having abnormal nuclei, hyperpolarization and multiple pseudopods, wild type PMN crawled with normal velocity and moved with purpose (i.e. a high meandering index)(Fig. 3d,e). We tracked individual PMN and found that NETosing PMN with diffuse nuclei crawled with high velocity and meandering index, while fully anuclear PMN crawled slowly (Supplementary Fig. 8a–c). Together these data suggest that structural nuclear breakdown is related to aberrant, but functional chemotaxis, similar to anuclear human PMN cytoplasts. By contrast, despite normal nuclei, *Tlr2*^{-/-} and *C3*^{-/-} had reduced crawling velocities and moved without purpose in random directions (Fig. 3d,e).

NETs are critical to contain an acute invasive infection *in vivo*

We followed staphylococcus skin infections over time using minimally invasive non-surgical luminescent imaging. Currently no specific inhibitor of NETosis exists, so exogenous DNase treatment was used to breakdown NETs. Animals treated with or without DNase had comparable *S. aureus* levels at 1 h, but DNase treatment decreased luminescence within 4 h (Fig. 4a–b). DNase did not alter bacterial growth in culture (data not shown). Disruption of NETs resulted in significant increase in bacteremia at 4 h and a decrease of skin CFUs at 24 h, indicating that bacteria can rapidly escape the portal of entry in the absence of NET trapping (Fig. 4c,d).

We examined dissemination in mutant mice that lacked NETosis. Wild type, *Tlr2*^{-/-} and *C3*^{-/-} animals received intraperitoneal injections of live *S. aureus* and blood CFUs were determined at 5 h. Compared to wild type mice, both *Tlr2*^{-/-} and *C3*^{-/-} mice had impaired ability to contain *S. aureus* dissemination (Supplementary Fig. 4c). Notably *C3*^{-/-} which were deficient in both phagocytosis and NETosis had more severe bacteremia than did *Tlr2*^{-/-} which could phagocytose but not NETose.

Human PMN release NETs *in vivo* and become anuclear

We investigated PMN from human subjects diagnosed with Gram-positive abscesses (Supplementary Table 1). Fresh abscess aspirates were imaged using transmission electron microscopy. In all samples, we visualized normal PMN, PMN with degraded nuclear envelopes, decondensed and dispersed chromatin (diffuse nuclei) and PMN without any intracellular DNA or chromatin (anuclear)(Fig. 5a–e). Normal PMN had abundant heterochromatin and multilobar nuclei, while NETosing cells were at various stages of nuclear envelope breakdown and NET release. Granulated cells with diffuse nuclear staining contained nuclear envelope remnants and diffuse chromatin throughout the cytoplasm however structurally intact nuclei were not present (Fig. 5b). PMN without a nucleus but still containing limited dispersed DNA and chromatin were a common finding on EM supporting our *in vivo* observation that PMN with diffuse nuclear staining have dissolved the nucleus and contain some free chromatin resulting in a diffuse DNA pattern of fluorescence (Fig. 5c). Of note, PMN with diffuse or completely absent nuclear material still contain granules, suggesting that they retain the necessary components to kill bacteria through conventional mechanisms.

NETs were identified by their typical beads-on-a-string EM appearance. To understand how NETs are released without cell lysis we imaged diffuse and anuclear PMN at higher magnification. We observed intracellular vesicles containing DNA fusing with the outer membrane resulting in delivery of NETs to the extracellular space (Fig. 5d). Electron microscopy of the abscesses also identified the late stages of NETosis (Fig. 5e).

The abscess aspirates were subjected to immunofluorescence microscopy, which demonstrated normal and anuclear neutrophils and NETs (Fig. 6a). Anuclear PMN stain positively for CD45 and CD66b but do not stain for intracellular DNA and NETs are visualized surrounding these cells (Fig. 6a). Live abscess PMN were injected directly into mouse skin to visualize NETosis in the identical manner used for mouse intravital microscopy. NETs were visualized before and after exogenous DNase treatment (Fig. 6b). A large percentage of human abscess neutrophils had decondensed nuclei with approximately 10% being anuclear (Fig. 6d), just like in mouse PMN.

Normal neutrophils isolated from healthy volunteers were injected into the mouse skin with or without live GFP-staphylococcus. Normal multilobar nuclei were observed in greater than 90% of human PMN in the absence of infection and these cells did not release NETs (Fig. 6c,d). However in the presence of infection, a large proportion of human PMN developed decondensed nuclei, released NETs and became anuclear (Fig. 6c,d).

Discussion

Despite ample *in vitro* NET research, fundamental questions about *in vivo* NETosis have remained unanswered, resulting in skepticism about the physiological relevance. For example, it has been suggested that NETs may just be an explosive necrosis, apoptosis or a process of dying^{10,11,30,31}. Moreover it has been demonstrated that NET-formation is slow, requiring hours and that NETosing PMN are non-functional thereby nullifying other critical roles that PMN play in host defense. As a result, highly invasive Gram-positive pathogens, like *S. aureus* and *S. pyogenes*, could disseminate prior to cell death NETosis, easily slipping by the incapacitated PMN. As multiple bacteria upregulate DNase and escape NETs³²⁻³⁵ it is critical that the enucleating PMN maintain conventional killing capacity. We found that viable PMN rapidly form NETs while crawling and that phagocytosis could occur at any point during NETosis. Anuclear PMN contained bacteria implying that phagolysosome maturation and NET release can be separately compartmentalized, such that bacteria cannot escape from inside the cell during NET release.

Another critique of NETs has been the inability to observe the remnants of NETosing PMN *in vivo*. Here we directly observed PMN releasing DNA and altering their nuclear structures. Interestingly only a subgroup of neutrophils became anuclear while a larger population of NETosing PMN degraded the nucleus but continued to have diffuse intracellular DNA. Our data demonstrates the existence of viable anuclear PMN consistent with the common histological finding of suppurative or necrotizing inflammation. Death of the NETosing PMN is inevitable and likely results in clearance by resident macrophages; therefore they may not be observed at later time points in tissue histology.

The nucleus is important for *de novo* protein synthesis and it would seem reasonable that loss of this organelle would lead to rapid cell death. However, anuclear platelets and red blood cells survive for long periods and red blood cells survive enucleation^{16,36}. Clearly, a terminally differentiated cell with a short half-life (8 h) should be able to survive a brief period without a nucleus. Indeed, physical removal of the nucleus from human PMN does not affect their conventional killing ability as these cytokineplasts crawl, transmigrate, phagocytose staphylococci and kill them via NO^{27–29,37} thus providing an interesting analogy to NETosing PMN. Clearly, temporary anuclear survival may be essential in this innate immune cell.

NETosis may lead to bystander tissue damage. Certainly vascular NETs can cause small vessel vasculitis³⁸, hepatitis¹⁵, thrombosis³⁹, or lupus nephritis⁴⁰ and pulmonary NETs contribute to the pathophysiology of cystic fibrosis^{41,42}. We observed a tight regulation of NET production, potentially required to limit collateral damage.

Finally it has been questioned how NETs are catapulted²² or forcibly expelled in solid tissues. Our imaging reveals subtle release of DNA, analogous to a novel method of nuclear envelope budding and vesicular release of NETs *in vitro*¹⁴. We found that vesicular release of NETs permits the PMN to maintain membrane integrity that allows for continued function.

Together our data reveal that the physiological fate of NETosing PMN is not immediate cell death and that the release of NETs covered in elastase and histones is an important immune response to infection *in vivo*.

Methods and Materials

SYTOX Orange, SYTOX Green, SYTO 60, MitoTracker Deep Red, and Xenon Alexa Fluor 594 Goat IgG Labeling kit were from Invitrogen. Other reagents used include DNase (Sigma), MIP-2 (R&D Systems) and DRAQ5 (Biostatus Ltd). Antibody to histone H2A.X (M-20), antibody to neutrophil elastase (M-18) and goat polyclonal IgG were from Santa Cruz Biotechnology. Antibody to GR-1 (clone RB6-8C5, Ablab) was conjugated to Alexa Fluor 488 (Molecular Probes, Invitrogen) or Alexa Fluor 750 (Ablab) and PE-conjugated antibody to GR-1 was obtained from BD Biosciences. PE and FITC-conjugated antibody to Ly6G (clone 1A8) was obtained from Biolegend. PE-conjugated antibody to human CD16 (clone 3G8), PE-conjugated antibody to human CD66b (clone G10F5) and PerCP-conjugated antibody to human CD45 (clone HI30) were purchased from Biolegend.

Green fluorescence protein (GFP) reporter vector construction and GFP labelling of *S. aureus* (MRSA, USA300)

The GFP reporter vector was constructed by gene fusion of the promoterless GFP⁺ with a truncated *S. aureus* protein. Briefly, the GFP⁺ (a GFP variant with increased fluorescence) coding sequence with an in-frame polylinker was cleaved with *KpnI/SpeI* from the plasmid pMUTIN-GFP⁺⁴³ and subcloned into an *E. coli-S. aureus* shuttle vector pBT2⁴⁴ at the restriction site created by *XbaI/KpnI*. The recombinant plasmid contained the promoterless *gfp*⁺ reporter gene, designated pBT2-*gfp*⁺, was replicated in *E. coli* and verified with PCR,

sequencing and restriction analysis. The genomic DNA of a CA-MRSA USA300 local clinical strain (USA300-2406) was digested with *KpnI/EcoRI*. The resultant *KpnI-EcoRI* fragments were inserted into pBT2-*gfp*⁺ plasmid at *KpnI/EcoRI* restriction site and transformed into *E. coli* DH5 α . The clone with the brightest green fluorescence was selected and the plasmid was purified. The gene sequencing revealed that a 1.8kb *KpnI-EcoRI* fragment containing a truncated SAUSA300_1226 gene which encodes a homoserine dehydrogenase⁴⁵ was fused with the promoterless GFP⁺ in the subsequent plasmid, designated pBT2-*gfp*⁺ fused. This plasmid was further transformed into the laboratory chemical mutant strain *S. aureus* RN4220 by electroporation using tryptic soy agar (TSA) plates containing chloramphenicol. The pBT2-*gfp*⁺ fused plasmids purified from RN4220 were then used for labelling of the clinical CA-MRSA (USA300) *S. aureus* isolate by electroporation and under chloramphenicol selection⁴⁶.

Animals

C57BL/6J, Complement-3 deficient (*C3*^{-/-}) and transgenic GFP fusion to histone H2B (Histone-GFP) (B6.Cg-Tg (HIST1H2BB/eGFP) 1Pa/J) mice were purchased from The Jackson Laboratory. *Myd88*^{-/-} and *Tlr2*^{-/-} mice were provided by Prof. Shizuo Akira (Osaka University, Japan). Dr. T.Graf (Barcelona, Spain) provided the Tg(LysMeGFP). Mice (20–35 g, 6–10 weeks old) were maintained in a pathogen-free environment. The mice had access to food and water *ad libitum*. All procedures performed were approved by the University of Calgary Animal Care Committee and were in accordance with the Canadian Guidelines for Animal Research.

Bacteria Strains

Three *S. aureus* strains were used in this study. *S. aureus* (Xen8.1 and Xen29), known pathogenic strains that have been used in soft tissue, biofilm and sepsis models, were purchased from Caliper Life Sciences. The GFP-transgenic CA-MRSA USA300-2406 was developed as previously described above. Staphylococcus Xen29 and Xen8.1 were grown in LB broth (200 $\mu\text{g ml}^{-1}$ kanamycin) and CA-MRSA was grown in brain heart infusion BHI (chloramphenicol 20 $\mu\text{g ml}^{-1}$) at 37 °C overnight. Bacteria were subcultured in fresh LB or BHI and incubated for 2 h in a 37 °C shaker to obtain bacteria in mid-log phase growth prior to injection. *Streptococcus pyogenes* was obtained from ATCC (ATCC19615). *S. pyogenes* was grown on a sheep blood plate overnight at 37 °C, and then a single colony from the blood agar plate was inoculated into 25 ml fresh BHI broth for 18 h at 37 °C.

Surgical procedures

Mice were anaesthetized (10 mg kg⁻¹ xylazine hydrochloride and 200 mg kg⁻¹ ketamine hydrochloride) and body temperature was maintained using a rectal probe and heating pad. The mice were pretreated with intradermal MIP-2 (0.2 μg) diluted in sterile normal saline or MIP-2 tissue superfusion (5 nM). The right jugular vein was cannulated to administer additional anesthetic and fluorescent dyes. The microcirculation of the dorsal skin was prepared for microscopy as previously described⁴⁷. Briefly, after shaving the mouse's back, a midline dorsal incision was made extending from the tail region up to the level of the occiput. The skin was separated from the underlying tissue, remaining attached laterally to

ensure the blood supply remained intact. The area of skin was then extended over a viewing pedestal and secured along the edges using 4.0 sutures. The loose connective tissue lying on top of the dermal microvasculature was carefully removed by dissection under an operating microscope. The exposed dermal microvasculature was immersed in isotonic saline and covered with a coverslip held in place with vacuum grease. Alexa Fluor 488, Alexa Fluor 649, PE or FITC conjugated GR-1 specific antibody (10 μ l per mouse i.v.) or the neutrophil specific conjugated antibody to Ly6G (clone 1A8, Biolegend) was used to visualize neutrophils.

Three different methods were used to visualize NETs *in vivo*. Method 1 used the membrane impermeable SYTOX Orange dye (diluted 1:1,000 with sterile saline, 100 μ l per mouse i.v.). MIP-2 superfusion (5 nM) was initiated 30 min prior to *S. aureus* administration (Xen29, DNase positive, 1×10^8 CFU in 100 μ l sterile saline i.d.). Method 2 used Xenon labeled histone specific antibody (M-20) or neutrophil elastase specific antibody (M-18) (20 μ g per mouse i.v.). The Xenon labeling kit was modified from manufacturer's instructions for *in vivo* experiments. Preliminary experiments were performed using various concentrations of component A and component B of the Xenon kit. Optimal visualization was found with 7.5 μ l component A and 10 μ l of antibody incubated at room temperature for 5 min, without the use of component B. Control IgG (20 μ g per mouse i.v.) was labeled using the same Xenon kit and modified procedure. MIP-2 (0.2 μ g, i.d.) was administered 60 min prior to *S. aureus* (Xen8.1, DNase negative, 1×10^8 CFU in 100 μ l of sterile saline, i.d.). For method 3 mice received an i.p injection of the cell-permeable nucleic acid dye SYTO 60 (5 μ l of stock 5 mM diluted in 200 μ l PBS, i.p.) 16 h prior to intravital microscopy or i.v. administration during surgery. MIP-2 (0.2 μ g, i.d.) was administered 60 min prior to the clinical CA-MRSA transgenic-GFP bacteria (USA300-2406, 1×10^8 CFU in 100 μ l of sterile saline, i.d.).

Following baseline visualization, all bacteria were directly administered into the field of view using a tuberculin needle (1×10^8 CFU in 100 μ l of sterile saline, i.d.). Three to six separate areas were visualized in each experiment in order to minimize selection bias. For killed bacteria experiments (Xen8.1) bacteria was heated for 30 min at 65 $^{\circ}$ C and washed twice with sterile saline ($n = 3$).

Spinning-Disk Confocal Intravital Microscopy (SDCIM)

Spinning-disk confocal intravital microscopy was performed using an Olympus BX51 (Olympus) upright microscope equipped with a 20 \times /0.95 XLUM Plan Fl water immersion objective. The microscope was equipped with a confocal light path (WaveFx, Quorum) based on a modified Yokogawa CSU-10 head (Yokogawa Electric Corporation). Laser excitation at 488, 561, 649 and 730 nm (Cobalt), was used in rapid succession and fluorescence in green, red and blue channels was visualized with the appropriate long pass filters (Semrock). Exposure time for all wavelengths was constant at 1 s. Sensitivity settings were maintained at the same level for all experiments. A 512 \times 512 pixels back-thinned EMCCD camera (C9100-13, Hamamatsu) was used for fluorescence detection. Volocity Acquisition software (Improvision Inc.) was used to drive the confocal microscope. Images captured using the spinning disk were processed and analyzed in Volocity 4.20. NET area was quantified using the Volocity software.

Bone Marrow Chimera Animals

Recipient C57BL/6 mice were irradiated with 2 doses of 5 Gy (Gammacell 40 ^{137}Cs γ -irradiation source; Nordion International). An interval of 3 h was allowed to pass between the first and second irradiations. Transgenic histone-GFP bone marrow was isolated from mice euthanized by spinal cord displacement and 8×10^6 donor bone marrow cells were injected into the tail vein of recipient irradiated mice. The mice were kept in microisolator cages for 8 weeks to allow full humoral reconstitution. This protocol previously confirmed that 99% of leukocytes in Thy1.1 and Thy1.2 congenic recipient mice were from donor bone marrow⁴⁸.

Neutrophil Behavior

Individual neutrophils were manually tracked to determine velocity, distance, displacement, pseudopod formation and cell contouring. Cells that remained within the field of view for 5 min were examined. Volocity software was used to calculate velocity, distance and displacement.

Bacterial Dissemination

S. aureus (Xen8.1) were grown to mid-log phase, washed and suspended in saline. Mice were injected i.p. with 1×10^8 CFUs of the bacterial preparation in a volume of 200 μL . At 5–6 h after inoculation, mice were anaesthetized and whole blood was collected. Serial dilutions were cultured on LB agar plates overnight at 37 °C. To directly assess the effect of NETs on bacterial dissemination mice received a dorsal subcutaneous live mid-log phase inoculation of staphylococcus (Xen8.1, 1×10^8 CFU in 100 μL injection of saline). 1 h post bacterial administration mice received an injection of either DNase (1,000 U) or saline into the infected skin. Dissemination was quantified by obtaining blood via a cardiac puncture in anaesthetized animals at 4 h post-bacterial inoculation, diluting it 1:1 with PBS and plating it on LB kanamycin for 24 h.

In vivo Bacterial Monitoring

An IVIS Imaging System 200 (Caliper Life Sciences) was used to quantify luminescent live staphylococcus (Xen8.1) over time. Live mid-log phase *Staphylococcus aureus* (Xen8.1, 1×10^8 CFU in 100 μL injection of saline) was administered subcutaneously into shaved mice. Animals were anesthetized with 1.5% isoflurane while in the imaging chamber. Photons were measured during a 30 s exposure. Total photon emissions from uniform area of each mouse were quantified by using the Living Image software (Caliper Life Sciences). Mice were imaged at 1 h, 4 h and 8 h post-bacterial inoculation. At 24 h, the mice were euthanized and a skin biopsy (3.5 mm punch) was obtained from within the injection site. The tissue was homogenized and plated on LB agar plates with kanamycin and grown overnight to quantify CFU.

Human PMN Isolation

Whole blood was collected into acid citrate dextrose (1:5 blood, v/v). Erythrocytes were removed using dextran sedimentation followed by two rounds of hypotonic lysis. Neutrophils were isolated from the resulting cell suspension using Ficoll-Histopaque density

centrifugation. PMN were stained *ex vivo* using PE-conjugated CD16 specific antibody and SYTO 60 (1:1,000 dilution) for 15 min prior to directed intradermal injection into the mouse skin prep. Ethical approval for obtaining healthy human volunteer blood was provided by The University of Calgary research ethics committee and each human subject provided informed consent.

Human Abscesses

The infectious disease specialty consultation service of the Division of Infectious Diseases in the Dept. of Medicine, Alberta Health Services – Calgary and Area, identified patients who were clinically diagnosed with cellulitis and secondary abscesses due to Gram-positive bacteria requiring wound aspiration as part of routine medical care. All patients were fully reviewed by a physician with specialty training in infectious diseases. Specimens were sent for routine Gram-stain and culture. Remaining samples were used for electron microscopy, immunofluorescence and *in vivo* spinning disk confocal experiments with consent of each patient. The University of Calgary research ethics committee approved the use of discarded human clinical samples.

Transmission Electron Microscopy

Fresh clinical aspirates were fixed with 2.5% glutaraldehyde. The cells were spun into a pellet using a microcentrifuge. After washing three times, the pellets were postfixed in 1% osmium tetroxide in cacodylate buffer for 1 h, dehydrated through a graded series of acetone, and embedded in epoxy-based resin. Ultrathin sections were cut in a Reichert-Jung Ultracut E microtome using a diamond knife and stained with 4% aqueous uranyl acetate and Reynolds' lead citrate. The sections were observed in a Hitachi H7650 transmission electron microscope at 80 kV, and images were taken with an AMT 16000 digital camera mounted on the microscope microscope¹⁴.

Immunofluorescence Microscopy

Slides (smears) were prepared from clinical aspirates and allowed to air dry. Slides were fixed in ice-cold acetone for 10 min and dried. Slides were then blocked for 2 h at room temperature in PBS containing 2% BSA followed by staining for 1 h with a cocktail of fluorescently-labeled Ab together with SYTOX Green. Slides were washed and coverslips mounted with ProLong Gold antifade reagent (Invitrogen). Slides were imaged using a spinning-disk confocal microscope consisting of an Olympus IX81 inverted microscope (Olympus) coupled to a confocal light path (WaveFx; Quorum Technologies) based on a modified Yokogawa CSU-10 head (Yokogawa Electric Corporation). Excitation by each of 491-, 561-, and 642-nm laser wavelengths (Cobolt) were sequentially controlled and merged into a single optic cable using an LMM5 laser merge module (Spectral Applied Research) and fluorescence was visualized through band pass emission filters (Semrock) driven by a Modular Automation Controller (Ludi Electronic Products, Ltd.) and detected with a 512×512 pixels back-thinned EMCCD camera (C9100-13, Hamamatsu). Volocity Acquisition software (V5.2.1) (Improvision Inc.) was used to drive the confocal microscope

Statistics

Data were analyzed using GraphPad Prism (version 4, GraphPad Software Inc.). Reported values are expressed as mean \pm s.e.m unless otherwise described. We assessed the statistical significance of the difference between 2 sets of data using an unpaired, 2-tailed *t* test. Where the difference between more than 2 sets of data was analyzed, we used a one-way analysis of variance, followed by Bonferroni multiple-comparisons test. All *P* values of less than 0.05 were considered statistically significant.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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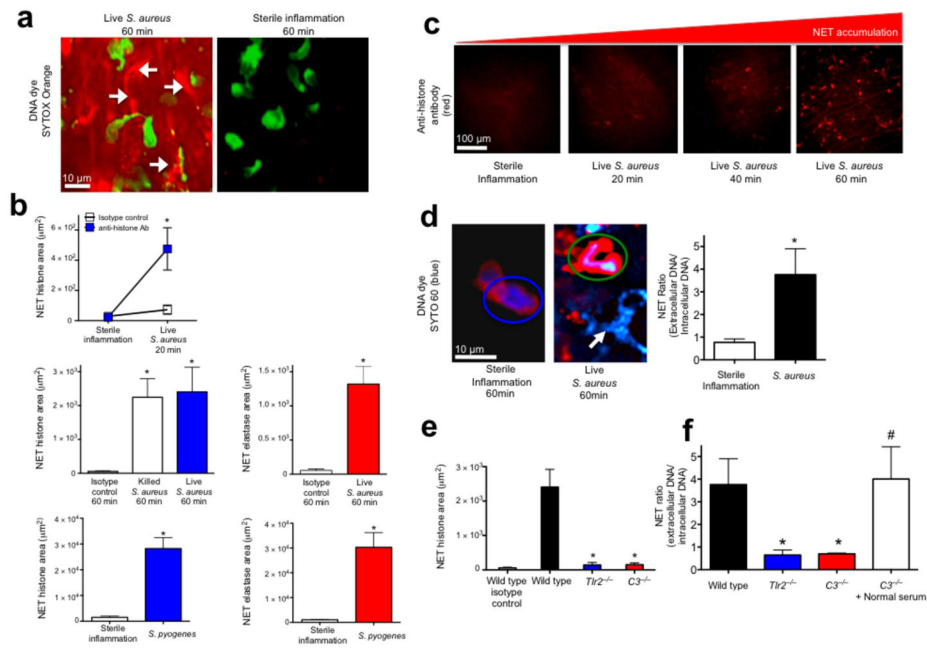
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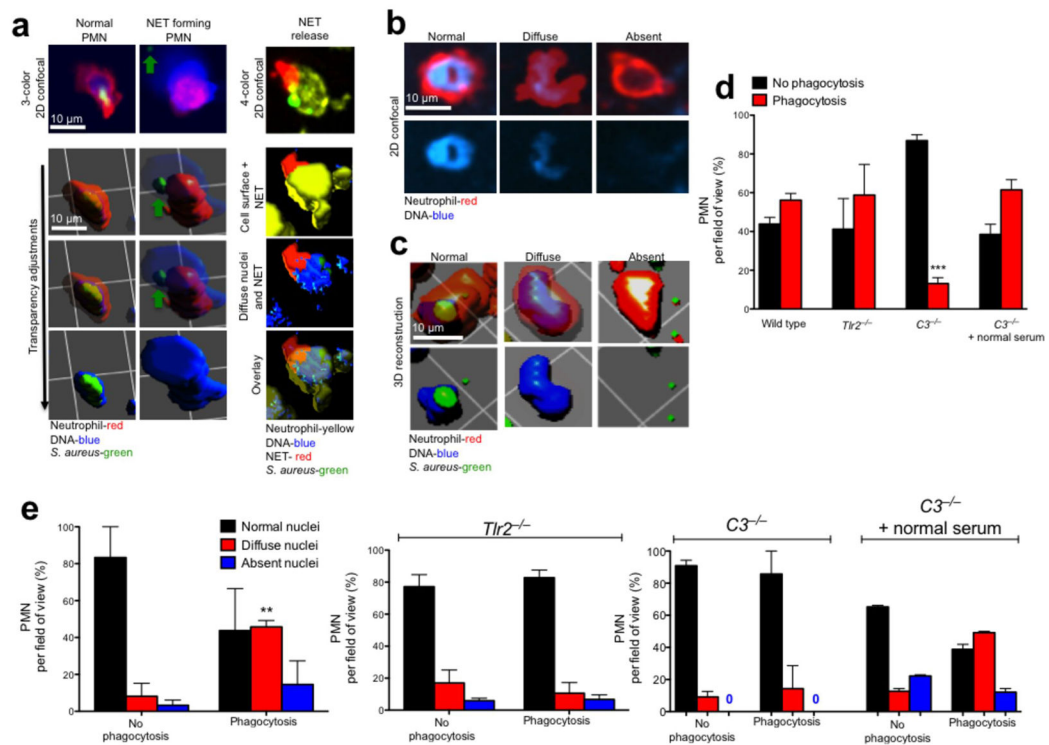
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**Figure 1.**

Rapid *in vivo* NETosis during acute Gram-positive bacterial infections is directly visualized *in vivo*. **(a)** Method 1: NET release was visualized as extracellular DNA (SYTOX Orange, white arrows) *in vivo* during infection (*S. aureus*, Xen29), but not during sterile inflammation (MIP-2 superfusion) (neutrophils are green, while NETs are red)(6 mice). **(b)** Method 2: NET quantification using fluorochrome conjugated histone or neutrophil elastase specific antibodies following live *S. aureus*, *S. pyogenes* or killed bacteria (*S. aureus* Xen8.1). Control animals received fluorochrome conjugated IgG isotype control ($n = 4$ for each group, $n = 3$ for dead bacteria). **(c)** Temporal NET tissue accumulation with PMN images removed for clarity of NETs. **(d)** Method 3: *In vivo* PMN nuclei pre-stained with the cell-permeable DNA dye (SYTO 60, blue) during sterile inflammation (left) or during infection with GFP-*S. aureus* (GFP-USA300)(right). A PMN with a normal nucleus is circled in blue, while a PMN with a diffuse nucleus is circled in green. NETs are demonstrated during infection with a white arrow. NET release is quantified by a ratio of extracellular DNA to intracellular DNA. **(e)** Impaired histone release in *Tlr2*^{-/-} and *C3*^{-/-} animals ($n = 3$ each group). **(f)** Impaired DNA release in wild type, *Tlr2*^{-/-} and *C3*^{-/-} mice ($n = 3$ for all groups). NET area was determined using Volocity imaging software (* = $P < 0.05$ for treatment versus control, or treatment versus sterile inflammation, or knockout versus wild type, # = $P < 0.05$ for treatment versus *C3*^{-/-}).

**Figure 2.**

PMN are viable and functional during nuclear breakdown and chromatin decondensation. PMN and nuclei were evaluated *in vivo* following live *S. aureus* (GFP-USA300). **(a)** A normal PMN that has captured a live bacterium (green) is shown by 2D imaging (top left) and contrasted by a NET-forming PMN (top middle) that is chemotaxing towards a GFP-bacterium (green arrow) while releasing DNA. Beneath each 2D spinning disk image is a panel of confocal 3D reconstruction using various degrees of transparency to distinguish the DNA in relation to the PMN outer membrane. The NETosing PMN in this image (middle panels) can be observed crawling toward the live bacteria in Supplementary Video 3. Four-color spinning disk confocal and 3D reconstruction reveals a PMN (yellow, Alexa 750 conjugated GR-1 specific antibody) with a diffuse nucleus (cell permeable SYTO 60, blue) releasing an extracellular NET (cell impermeable SYTOX Orange, red) while retaining live *S. aureus* (GFP-bacteria, green). **(b)** Three nuclear phenotypes were visualized; normal, diffuse, or absent. Top panels show extracellular membrane and nuclear staining, while the lower panels show the nucleus alone. **(c)** 3D-reconstruction reveals the morphology of each nuclei group. The 3D image rendering results in an artificial white reflection on the cell surface that does not represent an authentic fluorescent stain or nuclei. **(d)** Phagocytosis of live GFP-staphylococcus by neutrophils was quantified in wildtype, *Tlr2*^{-/-} or *C3*^{-/-} mice. The ability for neutrophils with decondensed or absent nuclei to phagocytose was quantified in **(e)** wildtype, *Tlr2*^{-/-} or *C3*^{-/-} animals. ($n = 3$ experiments per group). (** = $P < 0.01$ and *** = $P < 0.001$ for phagocytosis versus no phagocytosis)

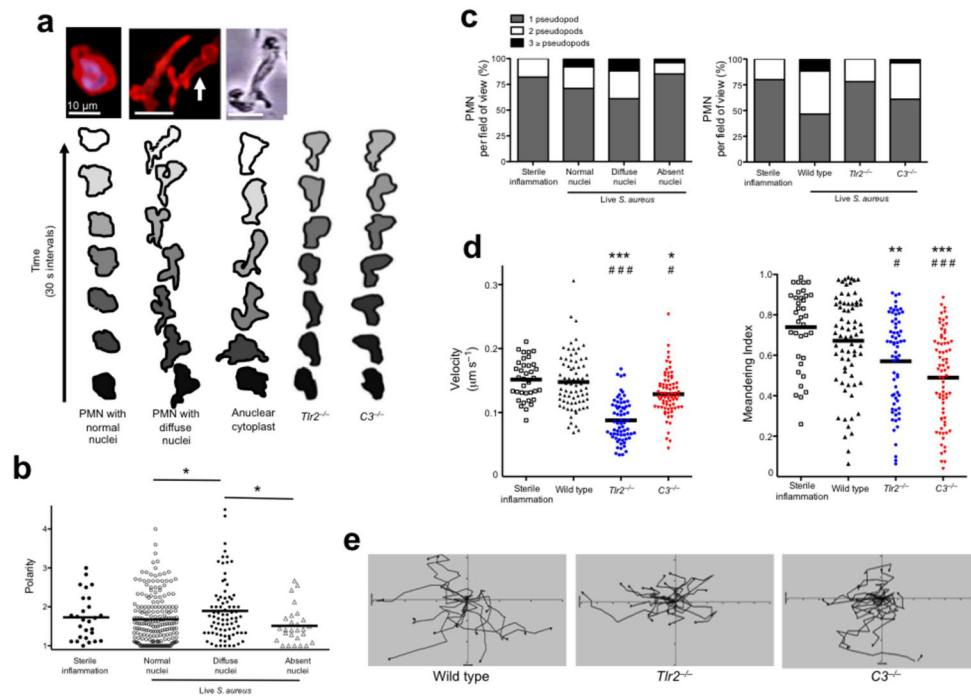
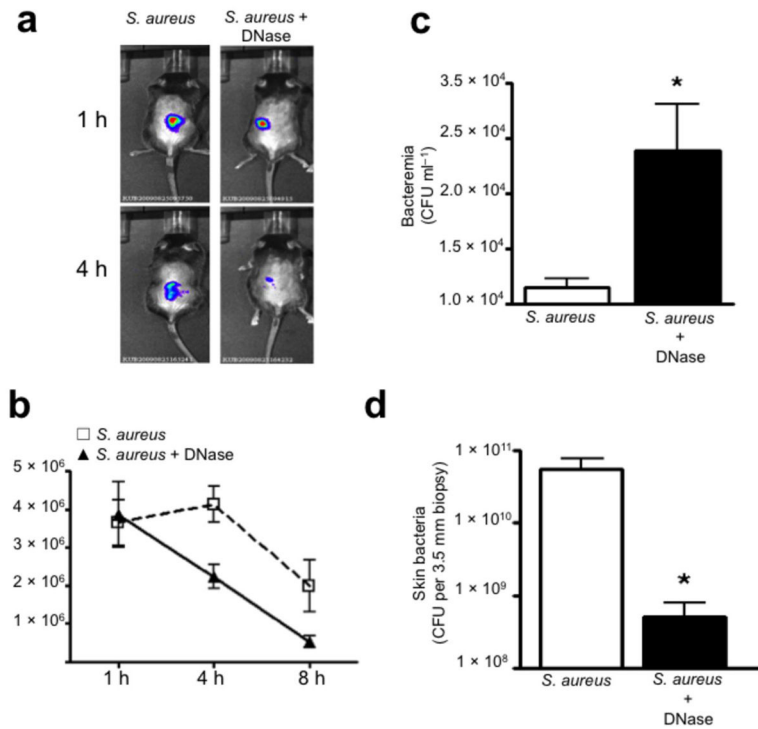


Figure 3.

NET-forming PMN display a novel crawling phenotype *in vivo* related to nuclear structure. (a) 2D images of *in vivo* PMN with a normal nuclei, NET-forming cells with diffuse nuclei and an *in vitro* generated anuclear human PMN (cytoplast). Contouring analysis of each cell is shown beneath the image. The PMN outer membrane is traced every 30 s (PMN in the middle image with white arrow is traced). Typical crawling phenotypes are shown for $Tlr2^{-/-}$ and $C3^{-/-}$ animals. (b) Cell polarities in relation to nuclear morphology were quantified. (c) The relationship of pseudopod formation of crawling PMN compared to their nuclear architecture or between wild type, $Tlr2^{-/-}$ and $C3^{-/-}$ mice. (d) PMN velocity and meandering index during live staphylococcus infection. (e) Cellular tracking during live staphylococcus infection (***) = $P < 0.001$, ** = $P < 0.01$, * = $P < 0.05$ compared to sterile inflammation and ### = $P < 0.001$, # = $P < 0.05$ compared to wild type).

**Figure 4.**

NETs are essential to limit acute *S. aureus* dissemination. **(a)** Photon imaging of live luminescent *S. aureus* (Xen8.1, 1×10^8 CFU per 100 μ l saline injection) within the mouse skin at 1 h and 4 h. **(b)** Live luminescent *S. aureus* were quantified within the skin of mice pretreated with either DNase (1,000U i.p.) or saline (i.p.) at 1 h, 4 h and 8 h post-infection (6 mice). **(c)** Bacterial dissemination from the skin to the blood at 4 h post-infection (Xen8.1) (6 mice). **(d)** CFUs grown from a 3.5 mm skin biopsy of the injection site at 24 h (6 mice). (* = $P < 0.05$).

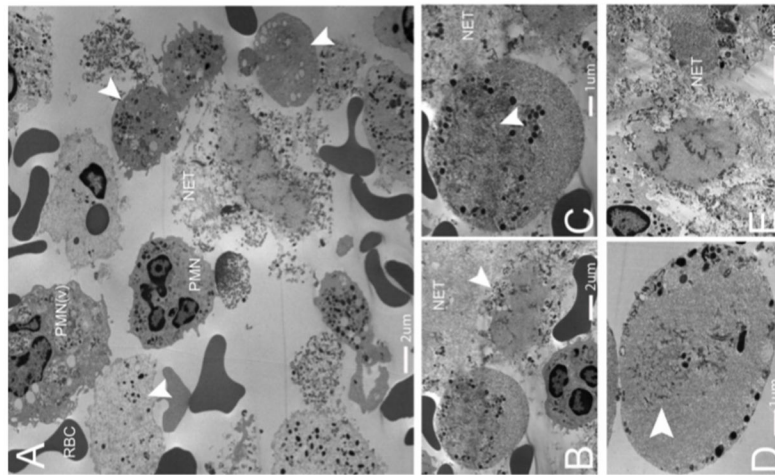


Figure 5.

NETosis occurs in human abscesses due to Gram-positive bacterial infections. Five human patients presenting with Gram-positive abscesses were evaluated. Transmission electron microscopy was performed on freshly obtained clinical samples. **(a)** The abscesses contained intact neutrophils (PMN), red blood cells (RBCs), activated neutrophils with vesicles in the cytoplasm (PMNv), as well as numerous anuclear neutrophils with cytoplasmic granules and nuclear vesicles (arrowheads). **(b)** A typical anuclear neutrophil is demonstrated undergoing NET formation (arrowhead) and a second cell is shown, enlarged in **(c)**, following nuclear envelope breakdown with dispersed chromatin and nuclear vesicles. The remnants of the nuclear envelope are highlighted (arrowhead). NETs are identified by the prototypical ‘beads on a string’ appearance on EM. **(d)** An anuclear neutrophil with nuclear envelope breakdown decondensed and dispersed chromatin (arrowhead), with vesicles and granules fusing with the outer plasma membrane (arrowhead). **(e)** Late stage neutrophils that have released chromatin and granules into the extracellular space.

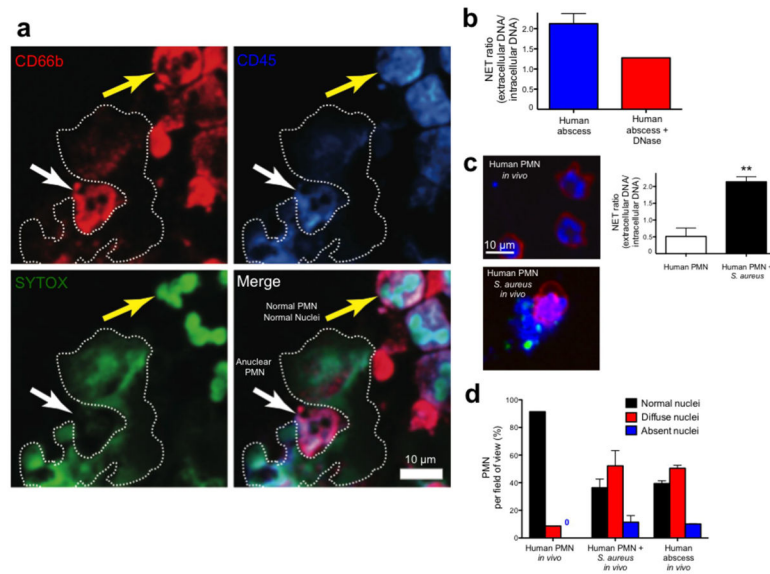


Figure 6. Immunofluorescence imaging of NETosis during human abscess formation. **(a)** Fresh abscess aspirates were stained with a PE-conjugated CD66b specific antibody, a PerCP-conjugated anti-CD45 specific antibody and SYTOX Green. The white arrow highlights an anuclear PMN adjacent to a NET that is outlined by a dotted line. The yellow arrow highlights a normal PMN with multilobar condensed nuclei. **(b)** Fresh live abscess aspirates stained with PE-conjugated CD16 specific antibody were directly injected into mouse skin to mimic our *in vivo* experiments and nuclei were pre-labeled with SYTO 60. NETs were visualized and quantified using the NET ratio. Exogenous DNase decreased visible NETs and decreased the NET ratio. **(c)** Normal PMN were stained with PE-conjugated CD16 specific antibody and SYTO 60. PMN were injected into mouse skin alone or with GFP-*S. aureus*. A NET is being released from the PMN stimulated with bacteria and the GFP-staphylococcus can be seen attached to the NET. NET release by *in vivo* human PMN is quantified as NET ratio (** $P < 0.01$ for untreated versus treated). **(d)** PMN nuclei were quantified *in vivo* from uninfected normal human PMN, human PMN infected with *S. aureus* and in human abscess PMN.