INTRODUCTION

As the most abundant myeloid cell in the body, neutrophils play a crucial role in innate immune responses to infectious disease. Neutrophils account for more than half of circulating immune cells and largely outnumber other granulocytes such as eosinophils (2%–5%) and basophils (1%) in circulation. These granulocytes display a diverse set of effector functions including phagocytosis and degranulation in response to small pathogens, and the release of DNA to form extracellular traps (ETs or neutrophil extracellular traps [NETs]) to fight larger pathogens such as fungal hyphae and multicellular parasites.

The process of NET formation, NETosis, was first described after phorbol 12-myristate 13-acetate (PMA) stimulation. Brinkmann and colleagues then discovered the bactericidal activity of NETs, heralding the intensive study of the NETosis mechanism and its role in pathogen control and disease development. The molecular pathways involved in NET formation still lack consensus, although chromatin decondensation is considered the endpoint. This decondensation ultimately precipitates nuclear membrane rupture, whereupon the DNA mixes with cytoplasmic and granular proteins. This “decorated” DNA is released into the environment, trapping pathogens and creating a micro-environment rich in enzymes and other toxic molecules for pathogens, and simultaneously protecting neighboring tissue from immunopathology.

DNA decondensation can either occur due to (i) the release of neutrophil elastase (NE) from granules, which then degrades histones in the nucleus; and/or (ii) the citrullination of histones by peptidyl arginine deiminase 4 (PAD4). In the first case, termed suicidal NETosis (with a loss of cell membrane), NADPH oxidase 2 (NOX2) leads to the formation of reactive oxygen species (ROS). H$_2$O$_2$, formed by superoxide dismutase (SOD) then becomes the substrate of myeloperoxidase (MPO), resulting in the release of NE. In the second case, also termed, vital NETosis (without loss of membrane integrity), the mechanisms are less understood, but ROS are produced by mitochondria in a NOX-independent manner. Of note, PAD4 can also be required for suicidal NETosis.

While neutrophils were not conventionally considered to be protective against helminth infection, here we review recent clinical and experimental evidence challenging this dogma, and suggest these...
neutrophils might have exerted a selective pressure on co-evolved nematode parasites due to the larvicidal potential of NETs.

2 | HOOKWORMS AND NEUTROPHILS: A CLINICAL PERSPECTIVE

Hookworms are clade V nematodes infecting a vast variety of mammals such as canids, felids and humans. They are considered the most prevalent of the soil-transmitted helminths (STH) and infect about 0.5 billion people worldwide principally in low- and middle-income countries, where poverty and lack of adequate sanitation contribute to high infection rates. Necator americanus, Ancylostoma ceylanicum and Ancylostoma duodenale are the three main species responsible for human infections. Hookworms have a free-living stage in humid soil, where eggs develop into infective larvae. Depending on the species, the infectious third-stage larvae (L3) either are ingested or infect the host via skin penetration. From the skin, the larvae enter the blood circulation to reach the lungs. There, they further mature and are coughed up and swallowed, ultimately reaching the intestine. For all species, the adults reside in the intestine and reproduce, laying eggs that are passed on in the feces. Adult parasites can maintain themselves in their host for years, and individuals in endemic areas usually harbor increasing numbers of parasites with age.

To date, the hookworm research community is still unsure whether natural immunity is raised against hookworms, distinguishing these parasites from other STHs. This is attributed to the extensive immunomodulatory abilities of hookworms. Researchers now propose that hookworms could be “Old Friends”: part of a natural human microbiome with a near-symbiotic relationship. Like other STH parasites, they are typically associated with raised levels of eosinophils in circulation and a modified Type 2 immune response. However, in-depth characterization of the immune response triggered by hookworms is lacking, due in part to the absence of species that naturally infect rodents used in immunology research. Helminth infection is commonly associated with elevated granulocyte counts. Most reports focus on eosinophil levels, which can increase from 2% to 5% in a healthy individual to 40% in a helminth-infected patient. But are eosinophils the only granulocytes recruited in the context of helminth infection? Or are neutrophils also a common feature of hookworm infection?

To date, clinical evidence for neutrophil recruitment after hookworm infection is relatively poorly documented. Despite their role as “early responders,” they appear to have been overlooked because of their reputation as “professional phagocytes” unlikely to participate in the control of large pathogens.

Hookworms of cats and dogs can often cause zoonotic infections in human hosts, leading to a pathology called “cutaneous larvae migrans” in humans. As humans are not the natural host of these animal parasites, the hookworm larvae are unable to migrate from the skin where they entered the host. The impeded parasite causes local inflammation, where serpiginous tracks are often observed. Histological examination reveals eosinophil and neutrophil recruitment in close proximity to larvae.

While such reports suggest neutrophils might be recruited in the context of clinical hookworm infection, interestingly, a recent analysis of a cohort of 300 individuals living in a STH endemic area (Trichuris, hookworms and Ascaris) observed no increase in circulating neutrophils due to these infections. The authors further investigated whether the activation phenotype of the neutrophils might have been altered by infection, but once again no association with any of the STHs was found. This absence of neutrophil response has previously been reported in smaller cohorts of hookworm-infected individuals. For example, volunteers infected with N. americanus had increased levels of circulating eosinophils, but not neutrophils, 3 weeks post-infection.

So why is there an apparent discrepancy between these reports? Could it be that only the skin/infective stages of the parasite are likely to cause neutrophil recruitment? The neutrophilia in cutaneous larvae migrans results from an acute encounter with the infective stage of hookworm. In contrast, in the STH studies, individuals had chronic, established infections with adults at the time of analysis. Hookworms, particularly the adult stage, have evolved a large arsenal of immunomodulation and evasion mechanisms, which we are just beginning to harness to treat systemic anti-inflammatory diseases. Where albendazole treatment reduces neutrophilia in cutaneous larvae migrans from zoonotic sources, treatment of chronic infections results in an increase of circulating neutrophils, suggestive of active anti-neutrophil modulation.

While we could not find clinical studies reporting neutrophil migration to the site of hookworm infection, there is indirect evidence that this recruitment might occur. Indeed, total products or secreted products of several STH have been shown to cause neutrophil chemotaxis. One such protein is N. americanus Ancylostoma secreted protein 2 (Na-ASP-2), a protein involved in L3 invasion and specifically secreted at this infective stage. In an air-pouch mouse model of localized in vivo inflammation (where the dermal skin is inflated with sterile air creating a bubble in which stimuli are introduced), recombinant Na-ASP-2 was shown to cause an influx of lymphocytes to the site of injection, predominantly neutrophils. The crystal structure of Na-ASP-2 revealed a similar structure to CC-chemokines, which are generally chemotactic for neutrophils and monocytes. Beyond hookworms, infective stages of similar nematodes induce neutrophil responses, for example, Strongyloides spp. Like hookworm, this Clade IV parasite enters its host by skin penetration. Rajamanickam et al. reported that Strongyloides stercoralis infected people from South India (presenting no other STH or filariae infection) had an increased number of circulating neutrophils. Interestingly, the authors also found an increase in granular protein concentration, such as NE and MPO, in plasma suggesting that neutrophils were activated. Both neutrophils and their circulating proteins were reduced by anthelmintic treatment (ivermectin + albendazole, 6 months follow-up), as with larvae migrans.

Similarly, the veterinary-relevant hematophagous parasite Haemonchus contortus causes neutrophil recruitment to the sheep abomasum for the first 7 days of infection. Interestingly, the differential neutrophil response to STH observed between stages might not only occur due to immune evasion but also be intrinsic to the parasite antigen. Indeed, H. contortus antigens from L3 but not from the adult stage were shown to induce IL-4 release from neutrophils in vitro.
Neutrophil recruitment to STH could thus be associated with the infective stage, however, the literature is too limited to reach any significant conclusion. Nevertheless, laboratory investigations have uncovered more detailed associations between neutrophils and hookworm.

_Nippostrongylus brasiliensis_ is a rodent nematode model with a life cycle that closely resembles that of the hookworm _N. americanus_. This nematode has been shown to cause neutrophil recruitment early on after infection in both the skin and the lungs. In laboratory mice, secondary infection confers sterilizing immunity unlike other nematodes. As with clinical studies, injection of _Heligmosmoides polygyrus_ has been shown to cause neutrophil recruitment in granuloma-like formations around the larvae of both _N. brasiliensis_ and the strictly intestinal parasite _Heligmosmoides polygyrus_. As with clinical studies, injection of _Strongyloides_ spp. using a diffusion chamber model in mice caused neutrophil accumulation 1 day post-infection.

More nuanced results were observed for hookworm canine laboratory infections. _Ancylostoma caninum_, but not _Ancylostoma brasiliense_, was shown to cause an increase in circulating neutrophils between the early days of infection to the patent phase. These results point out that not all hookworms are made equal regarding neutrophil recruitment and activation, and it would certainly be interesting to characterize these differences. Altogether, such studies illustrate that the infective stages of STH parasites often cause neutrophil recruitment to the site of infection.

### 4 | NEUTROPHILS AND HOOKWORM’S EXCRETORY/SECRETORY PRODUCTS: A SELECTIVE PRESSURE?

Given the long evolutionary history hookworms share with their specific host, parasitic helminths are experts in modulating the immune system to prevent their expulsion. In the last 30 years, research has turned its attention towards studying how the excretory/secretory (ES) products of hookworms contribute to immunomodulation, with proteins, lipids, microvesicles, miRNA, and various metabolites found at this host–parasite interface. To date, only proteins have been well-studied, and these are the subject of a recent and extensive review. Hookworms express a plethora of putative or fully demonstrated neutrophil evasion mechanisms. Here, we discuss key examples (Table 1).

#### 4.1 | Hookworm ES products block neutrophil recruitment

Several studies report that hookworm ES reduces or abolishes neutrophil responses and immunopathology. For example, whole ES isolated from _N. brasiliensis_ abrogates Lipopolysaccharides (LPS)-induced lung neutrophilia. Similarly, _H. polygyrus_ also reduces the level of neutrophil-specific chemokines in mice in a model of contact hypersensitivity.

While hookworm ES proteins have now been studied for several decades, only a few functions are fully characterized. A platelet activating factor (PAF) hydrolase was identified in adult stages of _N. brasiliensis_ and found to inhibit PAF, a potent chemoattractant of eosinophils and neutrophils. Another notable hookworm ES product is neutrophil inhibitory factor (NIF), identified while studying _A. caninum_ extracts. This glycoprotein binds to the Mac-1 integrin (CD11b/CD18) on leukocytes, preventing the adhesion and transmigration of neutrophils. By transfecting the NIF gene into human endothelial cell cultures and murine lungs, it was demonstrated that this molecule effectively blocks the recruitment and migration of neutrophils both in vitro and in vivo. NIF has since been shown to be produced by a wide variety of hookworms and related helminths. Notably, however, despite being predicted in its genome, no protein has been found in _N. americanus_. In _H. contortus_, gp55, a NIF homologue, was further shown to reduce neutrophil effector functions by blocking H$_2$O$_2$ release by binding to Cd11b/Cd18.

As neutrophils are typically implicated in tissue damage, it has long been assumed that the anti-neutrophil activity of ES was to dampen immunopathology that would ultimately damage host and helminth alike. More recent advances in our understanding of neutrophil biology and functions have pressed the research community to focus on neutrophils as also being effector cells against hookworms. Interestingly, hookworms do also express secreted molecules that could potentially block neutrophil effector functions.

#### 4.2 | Hookworm ES products may block neutrophil effector functions

Most of the evasion molecules that could inhibit neutrophil effector functions have been studied before neutrophils were considered as potential anti-helminth effector cells. As such, the function of some ES products discussed in this section is putative or predicted (Table 1).

Indirect evidence that neutrophils might contribute to parasite control comes from approaches where hookworm ES have been blocked. For example, Ali and collaborators demonstrate that vaccination with NIF reduces _A. ceylanicum_ fecundity in hamsters, as measured by a reduction in eggs per gram of feces. This mimics clinical observations from Papua New Guinea that _N. americanus_-infected patients with IgE responses to ES products have lower levels of egg production.

Both _N. americanus_ and the ruminant nematode _H. contortus_ secrete a calreticulin-like molecule (NaCalr). _NaCalr_ was shown in vitro to block C1q-induced haemolysis via the complement pathway and therefore has a putative role in preventing opsonization and activation of leukocytes such as neutrophils. Like for NIF, immunization with _NaCalr_ conferred some protection against challenge infection, with a reduction of 43%–49% worms in the lungs of mice.46
Table 1: Immunomodulatory hookworm excretory/secretory proteins display documented or putative neutrophil-specific functions

| Neutrophil-related functions | Other immunomodulatory functions |
|-----------------------------|----------------------------------|
| **L3 hookworm immune-evasion** | | |
| DNase II | ✓ | ? |
| N.br, N.am, A.ce | Degraded DNA backbone of NETs | Could degrade ETs from other cells e.g. monocytes |
| ASP-2 | ✓ | ✓ |
| A.ce, N.am | Chemoattractant for neutrophils in vitro and in vivo | Chemoattractant for monocytes induces antibody responses |
| KI-1 | ✓ | ✓ |
| A.ce, A.ca | Inhibits neutrophil elastase (low expression in L3) | Inhibits trypsin and other elastases |
| HpARI* | ✓ | ? |
| H.po | | |
| HpBARI* | ✓ | ? |
| H.po | | |
| MIF* | ? | ? |
| A.ce | Binds the pro-inflammatory MIF receptor (CD74) which increases MPO expression | Binds the pro-inflammatory MIF receptor (CD74) on monocytes |
| MTP-2 | ✓ | ✓ |
| A.ce | | Induces TNFα and IFNγ release from macrophages |
| PAF inhibitor | ✓ | ✓ |
| N.br | Inhibits PAF, a chemoattractant of eosinophils and neutrophils | Various anti-coagulant peptides |
| **Adult hookworm immune-evasion** | | |
| TIL-1 | ✓ | ✓ |
| A.ce, A.du | Inhibits neutrophil elastase | De-activates DCs, induces T regulatory cells, and inhibits matrix metalloproteases |
| APs | ✓ | ✓ |
| A.ca | | Matrix metalloprotease inhibitor |
| NIF, Gp55 (H.co) | ✓ | ✓ |
| A.ca, A.ce | Blocks neutrophil migration via CD11b/CD18 integrin | Prevents C1q deposition (from L4 to adult stage) |
| Calreticulin-like | ? | ? |
| N.am | Prevents complement-mediated neutrophil activation | Prevents C1q deposition (from L4 to adult stage) |
| SODs | ? | ? |
| N.am, A.ce, N.br | Protects against oxidation | | |
| PRXs | ? | ? |
| A.ce | Protects against oxidation | | |
| TMP-1 | ✓ | ✓ |
| A.ca | | De-activates DCs, induces T regulatory cells, and inhibits matrix metalloproteases |
| TMP-2 | ✓ | ✓ |
| A.ca | | Matrix metalloprotease inhibitor |
| Acetylcholinesterase | ? | ✓ |
| N.am | Could prevent the release of neutrophil chemoattractant factors from epithelial cells | Various anti-neutrophil activity |

Note: ✓, demonstrated function; ?, putative or predicted function; *, also expressed in adults.

The third-stage larvae (L3) and adult stages of hookworms and related STH express a number of excretory/secretory (ES) proteins, some with known or putative anti-neutrophil activity.

Of the 8 characterized proteins secreted by hookworm infective larvae or their laboratory model counterparts, 5 have confirmed or putative activity associated with neutrophils. Two are related to chemotaxis (ASP-2 as attractant and PAF inhibitor as blocker). The other three could impair NETs formation (DNase II and KI-1, and MIF). Two ES not associated with neutrophils have been discovered in Heligmosomoides polygyrus and inhibit IL-33/ST2 pathway (HpARI and HpBARI). Finally, MTP-2 is an astacin-like metalloprotease that enhances the expression of TNFα and IFNγ in classically activated (LPS-stimulated) macrophages.

In the adult stage, more proteins with immunomodulatory properties have been characterized. Of 12 notable proteins we could find described in the literature, 7 had potential anti-neutrophil activity. NIF, and its homologue gp55 in H. contortus, block neutrophil chemotaxis. Acetylcholinesterase could also decrease neutrophil recruitment indirectly by blocking epithelial cells chemokine secretions. Once again, several ES protein activities were consistent with anti-NET activity (TIL-1, MIF, SODs, PRXs). Similar to KI-1, TIL-1 has been shown to inhibit NE. Both SODs and PRXs could affect NETosis by decreasing oxidative stress. Finally, a calreticulin-like protein, identified in Necator americanus could contribute to complement evasion, and thus indirectly decrease neutrophil trapping and NETosis. Non-neutrophil-related proteins include metalloprotease inhibitors (TMP-1 and 2), which have been shown to affect dendritic cell polarization and inhibit host matrix metalloproteases (MMP-2, -7, and -13). APs have been shown to have anticoagulant activity.

Checkmark indicates function described in vivo or in vitro, interrogation mark indicates putative function from known activity of protein or predicted function of sequence.

Abbreviations: A.ca, A. caninum; A.ce, A. ceylanicum; Ac-TMP-2, tissue inhibitor of metalloprotease 2; A.du, A. duodenale; APs, anticoagulant proteins; ASP-2, angiostrongylus-secreted proteins; H.co, H. contortus; KI-1, Kunitz-type inhibitor 1 from Ancylostoma ceylanicum; MIF homologue of macrophage migration inhibitory factor; MTP-2, metalloprotease 2; N.am, N. americanus; N.br, N. brasiliensis; TIL-1 trypsin-inhibitor like serine protease inhibitor. The color is to help the reader assess if the evasion products has been linked to neutrophil quickly. Dark blue is used for when Neutro association is demonstrated, light blue, when it is hypothetical and grey for a mechanism unrelated to neutrophil.
NETosis has recently been shown to be triggered in response to various nematodes.\textsuperscript{37} We reported that extracellular traps were released early on in both natural skin penetration and intradermal infection with \textit{N. brasiliensis}.\textsuperscript{20} Previously, in mice infected with \textit{N. brasiliensis} no NETs were found surrounding lung L3, but NETs were reported in the skin around dead L3.\textsuperscript{22,24} This apparent discrepancy can be explained by an active anti-NETosis evasion mechanism (Figure 1). Indeed, we demonstrated that while NETs could not be observed at late time points around living larvae both in vivo and in vitro, they were present around dead larvae. Using live imaging, we proved that in all cases NETs were released, but that they were quickly destroyed in the presence of ES products expressed by living parasites. We further identified a DNase-II highly conserved in Clade V nematodes (including \textit{N. americanus}, \textit{A. ceylanicum}), which is secreted and able to degrade NETs in vivo and in vitro.\textsuperscript{20}

Neutrophil elastase, a key molecule in NETs formation, has been shown in vitro to be directly toxic to the flukes \textit{Fasciola hepatica} and Schistosomes by damaging their tegument.\textsuperscript{48,49} NE has also been suggested to damage the much harder nematode cuticle of \textit{Trichinella}.\textsuperscript{50} Adult worms of \textit{A. ceylanicum} and \textit{A. caninum} have been shown to produce a Kunitz-type inhibitor-1 (AceKI-1 or AceK1), which is a tight-binding inhibitor of trypsin, chymotrypsin, pancreatic elastase, and NE.\textsuperscript{51} \textit{Ancylostoma duodenale} also expresses a serine protease inhibitor with two trypsin inhibitor-like domains (AduTIL-1).\textsuperscript{52} This likely reflects the reproductive niche of hookworms in the digestive tract, but might also be an evasion strategy from neutrophil attack (Figure 1). Unfortunately, the impact of NE inhibition via AceKI1 on NETosis has not been studied so far.

Hookworms express several antioxidant enzymes, such as SODs, which could interfere with the process of NET formation (Figure 1). Interestingly, extracts of the tapeworm \textit{Mesocestoides corti} were shown to block the formation of stress-induced NETs.\textsuperscript{53} In this study, parasite products were co-cultured with neutrophils in the presence of \textit{H}_2\textit{O}_2, and a reduction in NETs was observed. The authors then further determined that ES products block the Transient Receptor Potential Cation Channel Subfamily M Member 2 (TRPM2) as well as downstream autophagy pathways. While the authors did not define which specific parasite molecule(s) is (are) responsible for this activity, other studies have investigated the role of cestode ES in protecting against \textit{H}_2\textit{O}_2-induced cell death. One such study treated \textit{Echinococcus granulosus} with \textit{H}_2\textit{O}_2 to establish a list of parasite proteins secreted in response to reactive oxygen species.\textsuperscript{54} At the top of this list are glutathione-S-transferases, a well-characterized family of enzymes involved in detoxification. In hookworms, three GSTs have been identified, with GST-1 being a lead human vaccine candidate due to its heme detoxification function. It might thus be interesting to investigate the role of GSTs in NETosis evasion.

In this section, we have shown that hookworms have evolved a considerable defense arsenal against neutrophil activity, both blocking recruitment as well as neutrophil effector functions. We therefore argue that this reflects a selective pressure exerted by neutrophils on parasite fitness. In the next section, we discuss the proven and potential larvicidal activity of neutrophils.

5 | Neutrophil-Mediated Killing of Hookworms

We have established in the previous sections that neutrophil recruitment is a feature of hookworm infections and that hookworms have

\begin{figure}[h]
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\includegraphics[width=\textwidth]{Figure1}
\caption{Hookworms actively evade NETosis. The non-activated infectious larvae of hookworm are trapped by neutrophil extracellular traps (NETs) released by neutrophils isolated from human blood. During the transition to parasitism, heat-activation causes hookworms to secrete anti-NETs evasion molecules. Three mechanisms of evasion are illustrated: (i) a DNase-II capable of degrading NETs to evade trapping and cuticle damage, demonstrated in \textit{Necator brasiliensis} and \textit{N. americanus} and (ii) a Kunitz-type Inhibitor (Ace-KI1) identified in \textit{Ancylostoma ceylanicum} is proposed to block the formation of NETs by inhibiting NE activity, (iii) an unidentified blocker of TRPM-2 inhibits oxidative stress-induced NETs formation. This mechanism has been demonstrated in the cestode \textit{Mesocestoides corti}. Fluorescent images were obtained by co-culture of circulatory human neutrophils with \textit{N. americanus} L3 for 3 h. Activated larvae were placed at 37°C for one night before co-culture to stimulate ES release. NETs are stained using sytox green and are represented with the LUT fire in Fiji. The figure has been made using Biorender}
\end{figure}
evolved various strategies of evading the neutrophil attack. This suggests that hookworm fitness can be affected by neutrophils. But can these granulocytes kill a helminth, and by which mechanism(s), given the large size of hookworms?

Interestingly, the toxic activity of neutrophils was assessed against various nematodes in vitro in the early days of immunoparasitology. However, knowledge of neutrophil effector functions was incomplete at the time, and their role in anti-helminth immunity has only been studied in detail more recently. Two types of neutrophil-mediated killing have emerged from recent studies: (i) direct killing, associated with neutrophil trapping, NETosis and granule toxicity and (ii) indirect killing, associated with enhanced type 2 immunity.

5.1 | Direct toxicity of neutrophil to hookworm parasites: netosis killing

The web-like DNA that forms NETs is decorated with antimicrobial molecules such as histones, MPO, and NE, which have been demonstrated to kill bacterial and fungal pathogens. Here, we discuss the pathways invoking NETosis, and the mechanisms by which NETs reduce parasite infectivity and viability (Figure 2).

5.1.1 | Neutrophil extracellular traps larvicidal activity

Neutrophils that are recruited towards larvae or their products have been shown to bind and trap larvae, which can be enhanced by antibody-dependent binding as shown during secondary infection with H. polygyrus. To bypass macrophage-mediated killing of the parasite, mice were challenged with L3 only 4 days after primary infection and passively immunized with serum from primed mice to artificially increase antibody titers. Following this transfer, worm burdens were significantly reduced. They further demonstrated in vitro that “altered” neutrophils from immunized mice swarmed and killed the larvae – although this now four-decade-old paper did not further describe this alteration, it was perhaps an early report of trained immunity. Similarly, in mice vaccinated with A. caninum, inoculation of L3 intraperitoneally caused neutrophils to bind to the larvae in an antibody-dependent manner. The authors then conducted SEM on the larvae and observed damage to the cuticle, including what they described as a swelling, focal collapse of cuticle, and deformation.

We have recently observed similar damage to the cuticle of N. brasiliensis exposed in vitro to neutrophils, as shown by increased permeability of the L3 cuticle to the impermeant DNA binding dye Sytox-Green. In this study, we further characterized the neutrophil toxicity to the larvae resulting from the release of NETs, formed around larvae after skin penetration or intradermal injection. Neutrophil depletion or NETosis blockade (PAD-4-KO, NE inhibitor, and DNase treatment) all increased parasite survival, measured by the number of adults in the intestine. We also confirmed that N. americanus-induced NETosis in vitro. As mentioned previously, N. brasiliensis and N. americanus secrete a DNase-II that cleaves extracellular traps in response to this attack. In a co-culture assay of L3 with neutrophils, neutralizing the DNase-II evasion activity with anti-serum increased the percentage of dead larvae, proving that NETs can directly impair larval survival. Altogether, this illustrates that NETs can have larvicidal activity against hookworms and that those parasites have evolved an evasion strategy against this attack (Figure 2).

NETosis appears to be initiated against various helminths, both STH4 and vector-borne. Notably it has been observed in vitro against Strongyloides stercoralis, H. contortus, Strongyloides ratti, the ruminant parasite Ostertagia ostertagia, as well as several filariae. What is most intriguing is that this response is not just limited to parasite nematodes; even if quite artificial, the free-living clade V nematode Caenorhabditis elegans is also able to trigger NETs in vitro, proving the triggers of NETosis are well-conserved amongst nematodes.

To date, no consensus has emerged regarding the importance of NETs in STH larvicidal activity. Key differences between studies are as follows: (i) So far, immuno-evasion has only been observed in hookworm, with the live larvae of other species becoming durably entrapped in NETs, (ii) Cuticle damage was only reported in N. americanus, N. brasiliensis and A. caninum; (iii) While neutrophil-induced killing was found in all models, NETs degradation was not always sufficient to reverse killing. Whether this means that NETs can damage some helminths and not others is not clear. Indeed, some of these results might be inherent to the assay, and not reflective of NETs larvicidal potential in vivo. For example, in the S. stercoralis in vitro system, incubating the larvae with a mouse or human neutrophils gave diverging results. In both cases, NETs were released and contributed to larval trapping. However, with human cells, DNase-I treatment reduced the killing of the larvae from 90% to 20%, while with murine neutrophils no reduction in the killing was observed. Further work, using in vivo studies, is required to confirm the larvicidal activity of NETs in other STH.

A potential explanation as to why some helminth larvae are killed by NETs and not others may lie in a difference in neutrophil activation, rather than in the worms themselves. Indeed, different stimuli have been shown to cause changes in the “NETome,” that is, the molecules decorating the released DNA. To date, no consensus has emerged regarding the importance of NETs in STH larvicidal activity. Key differences between studies are as follows: (i) So far, immuno-evasion has only been observed in hookworm, with the live larvae of other species becoming durably entrapped in NETs, (ii) Cuticle damage was only reported in N. americanus, N. brasiliensis and A. caninum; (iii) While neutrophil-induced killing was found in all models, NETs degradation was not always sufficient to reverse killing. Whether this means that NETs can damage some helminths and not others is not clear. Indeed, some of these results might be inherent to the assay, and not reflective of NETs larvicidal potential in vivo. For example, in the S. stercoralis in vitro system, incubating the larvae with a mouse or human neutrophils gave diverging results. In both cases, NETs were released and contributed to larval trapping. However, with human cells, DNase-I treatment reduced the killing of the larvae from 90% to 20%, while with murine neutrophils no reduction in the killing was observed. Further work, using in vivo studies, is required to confirm the larvicidal activity of NETs in other STH.

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5.1.2 | Neutrophil extracellular traps induction

Neutrophil extracellular traps can be invoked via several different pathways and signaling events, as discussed above. In all
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reported models of STH-induced NETosis, neutrophil elastase blockade was found to prevent NET formation.\textsuperscript{20,59,61} Blockade of MPO and NOX activity was also reported to block NETosis \textit{in vitro}.\textsuperscript{59,61} Interestingly, NETosis in response to \textit{Dirofilariae immitis} (a Clade III nematode) was shown to be independent of NOX activity,\textsuperscript{62} suggesting that several types of NETs could be induced by different helminths. In NOX-independent NETosis, PADs were shown to be required to decondensate DNA by hypercitrullination.\textsuperscript{57} PAD4 was shown to be required for NET killing \textit{in vivo} after \textit{N. brasilienis} infection, and NETs were heavily decorated with citrullinated Histone 3,\textsuperscript{20} though unfortunately, we did not investigate the role of NOX.

Interestingly, NOX-independent NETosis has been reported to be a much faster event than NOX-dependent NETosis. Rapid induction of NETosis was reported in vitro after stimulation with \textit{S. stercoralis} and \textit{H. contortus}, respectively,\textsuperscript{57,59} which might suggest NOX-independent NETosis does occur after STH stimulation. Notably, both species induced NETs more rapidly than with highly potent artificial stimuli PMA. However, with both \textit{N. brasiliensis} and \textit{O. ostertagi}, NETs induction has been shown to be slower than with PMA.\textsuperscript{20,61} This slow NETosis has been shown to be associated with some neutrophils remaining NETosis-free several hours after culture within \textit{Candida albicans} and \textit{Group B streptococcus},\textsuperscript{66} while PMA induced NETosis in virtually all neutrophils. Similar to those slow
inducers, a large majority of neutrophils free from NETosis were reported for both *N. brasilensis* and *S. ratti*. Why such a low number of neutrophils enter NETosis in response to helminth infection is an interesting question, currently without an answer. It may be that a specific polarization of neutrophil is required (as described in Chen, 2016) or that neutrophil requires a signal above a threshold to undergo NETosis, that PMA and other “potent inducers” reliably cross, while helminth does not, maybe to prevent “NETosis storms.”

Further work is required to understand the mechanisms of NETs formation against helminths and to decipher if all NET induction mechanisms impair parasite viability.

5.2 | Indirect toxicity of neutrophils: a role for netosis in type 2 protective immunity?

The role of neutrophils in type 2 immune responses, particularly in helminth infection, is increasingly appreciated. Here, we focus on the indirect role of NETs in driving type 2 protective immunity to hookworm.

5.2.1 | Priming of type 2 immunity

As with many infections or injuries, neutrophils are quickly recruited to, and swarm, hookworm larvae after their entry into the host. Neutrophils and inflammatory monocytes have been shown to be the main cells taking up fluorescently labelled *N. brasilensis* L3 antigens in the skin. Given that NETs have been shown to interact with dendritic cells (DCs) and alter their polarization, Pellefigues and collaborators explored the role of NETs in the priming of type 2 inducing DCs after dead L3 injection. The IFN-γ signature of DCs (required for Th2 induction in helmint models) was not found to be dependent on NETosis, as this signature was not abrogated by exogenous DNase treatment or neutrophil depletion. This however does not exclude the role of NETs in shaping type 2 responses entirely. Indeed, in type 2 respiratory virus model, NETs have been recently shown to sustain the intensity and priming of the type 2 response by controlling recruitment of monocyte-derived DCs, proving that NETs can be involved in Type 2 immune priming/tuning. Further studies of NETosis in the initiation of Type 2 immunity are thus needed.

5.2.2 | Activation of effector cells

IL-4 activated macrophages are a hallmark of anti-helminth immunity, and many forms of neutrophil-macrophage communication have been reported. In *Strongyloides* infection, NETs were shown to be required for killing larvae but could do so only in the presence of macrophages. Similarly, in an *N. brasilensis* re-infection model, primed macrophages from neutrophil-depleted mice, transferred to naive animals, could no longer protect against infection, suggesting an important role for neutrophils in priming macrophage larvicidal activity. *Nippostrongylus brasiliensis* or LPS-treated neutrophils had a distinct transcriptome from one another, including prototypical M2 markers (il-13, chi3l3, retnla, arg-1). Whether this transcriptomic profile is associated with NETosis has not been investigated in this study.

Recent research, outside the field of helminthology, has begun to explore the potential for NETs to contribute to macrophage activation. Notably, NETs have been shown to activate macrophages via NLRP3 leading to increased activation of CD86+ macrophages and increased expression of IL-1β, IL-6 and TNFα (Qiongyi et al. Increased neutrophil extracellular traps activate NLRP3 and inflammatory macrophages in adult-onset Still’s disease, PMID: 30616678). Furthermore, IL-8 has been shown to cause the release of NETs and further activate macrophage activation and release of IL-8 in the context of atherosclerosis (An, 2019). Future research could thus investigate how NETs or their clearance changes macrophage polarization and function in the context of hookworm infection.

6 | HOOKWORMS AND NETOSIS: OUTSTANDING QUESTIONS REMAIN

While NETosis is an increasingly documented response against STH infection, there remain several key questions surrounding the mechanisms and requirements of this effector response. First, what are the molecular triggers of helminth-induced NETosis? Second, what other cells contribute to killing during NETosis, and could eosinophil and monocyte ETs contribute to helminth control?

6.1 | Triggers for nets

Size has been proposed as a trigger for NETosis. Indeed, Bransk and collaborators proposed that neutrophils selectively release NETs in response to fungal hyphae or bacterial aggregates, but not small yeasts or non-aggregated bacteria. The absence of phagosomes that form when neutrophils encounter large pathogens allows for NE to be slowly released into the cytosol. This in turn promotes the decondensation of chromatin, leading to the release of NETs. These molecular events could certainly occur when a neutrophil encounters a helminth, although it is yet to be explored. However, in this size-dependent model, NETosis is slow and can take up to 4 hours. Hookworms and helminths on the other hand have been shown to trigger NETs within 30 minutes to 1 hour post-encounter. Thus, it is likely that other triggers are involved in the induction of NETosis.
have known endosymbionts, the infective larvae are likely to be covered with soil-derived bacteria. It stands to reason that host immunity could evolve responses to this hookworm-associated microbial signature.

While hookworm NETosis induction could be bacterial dependent, it is however independent of TLR-2 and TLR-4, as NETs were still observed around dead *N. brasiliensis* L3 in TLR-2-KO and TLR-4-KO.22 Similar results were obtained with TLR-4 inhibitor treatment in *O. ostertagi* assays,61 proving that TLR-2 and TLR-4 ligands are not required for hookworm-induced NETosis. While specific helminth PAMPs have still not been characterized, several hookworm molecules could be triggering NETosis. The glycan-rich cuticle of hookworms could be recognized by neutrophils, as lipophosphoglycans of *Trypanosoma* spp. have been shown to induce NETs.74 Another potential PAMP could be aspartic protease-1 (APR-1), the lead vaccine candidate against hookworms, as fungal aspartic proteases of *C. albicans* were shown to trigger NETosis.75

Finally, NETosis could happen in response to environmental cues such as danger signals or cytokine alarmins. Indeed, cytokine (IL-8) and chemokine binding (to IFN-γ and CXCR1) has been shown to be capable of triggering NETs formation.76 To date, such a role for the cytokine milieu has not been addressed in STH. Instead, a recent study reports that IL-4, a hallmark of anti-helminth response, limits NETs formation.77

6.2 Extracellular traps from other cells

Since the discovery of NETs, DNA traps have been shown to be a general feature of leukocytes, including eosinophils (EETs), macrophages (METs), mast cells, and basophils (BETs).78,79 Literature is even more limited for these cells than neutrophils, but a few reports are pointing towards their involvement in hookworm infection.

As mentioned previously, *Strongyloides* infection has been shown to trigger egress of both neutrophils and eosinophils to the site of infection.50 Interestingly, Ehrens and collaborators, reported recently that *S. ratti* infection triggers both NETosis and EETosis with larvicidal activity. EETosis was also demonstrated against filarial parasites *Litomosoides sigmodontis* and *D. immitis*, suggesting a conserved mechanism of helminth defense.80

*Strongyloides* larvae were shown to cause the release of METs from human cells in vitro.57 The subsequent killing of larvae then required a combination of neutrophils and macrophages, and was reversed by DNase treatment. Interestingly, the authors report that in co-cultures of mouse neutrophils, macrophages and larvae, NETs but not METs are released. The killing was then still present but not reversed by DNase treatment.57 This suggests that METs might be involved in *Strongyloides* killing but that redundant mechanisms exist.

Basophils are known to be important for early priming of the immune response against *N. brasiliensis*, notably releasing IL-4 and priming macrophages in secondary responses.81,82 Nothing is yet known regarding their DNA release during helminth infection, but it would be interesting to investigate the contribution of BETs to anti-helminth immunity.

7 CONCLUSION

Hookworms appear to have evolved complex immune evasion mechanisms specifically tailored towards neutrophil effector functions, with many mechanisms still remaining to be elucidated. It also appears that neutrophils, and in particular their NETs, can have larvicidal activity directed against at least the infective stage of hookworm. Thus, we argue that neutrophils are an important player in anti-hookworm immunity and that their role in clinical infection should be explored further.

Clinical investigations of the immune response to hookworm infections are scarce, as local tissues are hard to access in humans. Quite recently, controlled challenge infections with hookworms have been designed to test the efficacy of hookworm vaccine candidates during clinical trials.83 This approach has created an interesting opportunity for researchers to study the immune response to hookworm infections in otherwise hookworm-naïve individuals, notably the early events after infection, which may be associated with neutrophil responses and potential larvicidal activity.

Attempts to develop anti-hookworm vaccines have highlighted the need to exploit early immune responses to the infective stage, rather than established adult worms. Given neutrophils role as first responders, deciphering the neutrophil/hookworm interplay might pave the way towards new vaccine targets and uncover neutrophil evasion mechanisms that could be harnessed to combat neutrophil-mediated diseases.

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CONFLICT OF INTEREST

The authors declare no conflict of interest.

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