Differential neutrophil responses to bacterial stimuli: Streptococcal strains are potent inducers of heparin-binding protein and resistin-release

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Neutrophils are critical for the control of bacterial infections, but they may also contribute to disease pathology. Here we explore neutrophil responses, in particular the release of sepsis-associated factors heparin-binding protein (HBP) and resistin in relation to specific bacterial stimuli and sepsis of varying aetiology. Analyses of HBP and resistin in plasma of septic patients revealed elevated levels as compared to non-infected critically ill patients. HBP and resistin correlated significantly in septic patients, with the strongest association seen in group A streptococcal (GAS) cases. In vitro stimulation of human neutrophils revealed that fixed streptococcal strains induced significantly higher release of HBP and resistin, as compared to Staphylococcus aureus or Escherichia coli. Similarly, neutrophils stimulated with the streptococcal M1-protein showed a significant increase in co-localization of HBP and resistin positive granules as well as exocytosis of these factors, as compared to LPS. Using a GAS strain deficient in M1-protein expression had negligible effect on neutrophil activation, while a strain deficient in the stand-alone regulator MsmR was significantly less stimulatory as compared to its wild type strain. Taken together, the findings suggest that the streptococcal activation of neutrophils is multifactorial and involves, but is not limited to, proteins encoded by the FCT-locus.

While in bacterial infections, the recruitment and activation of neutrophils are critical for the immune defence; it is becoming increasingly evident that neutrophils may also contribute to immunopathology. On one hand, neutrophils play an important role in bacterial killing through mechanisms such as phagocytosis, formation of extracellular traps and production of antimicrobial effector molecules. On the other hand, the release of granule proteins can cause cell and tissue damage to the host, as well as dysregulated inflammatory response1.

In sepsis, several studies have shown that neutrophil responses are aberrant with respect to survival, migratory capacity and functionality2-4. Whereas many previous studies have focused on the detrimental role of dysregulated migration and impaired phagocytic killing, less is known about the contribution of neutrophils to the hyperinflammatory responses that characterize sepsis and its complications. A distinct feature of sepsis is the recruitment of immature neutrophils from the bone marrow into the circulation. A recent report found circulating neutrophils in sepsis patients to have a suppressed apoptosis, a longer life span and pro-inflammatory phenotype with increased TNFα/IL10 ratio5.

Heparin-binding protein (HBP) and resistin are two neutrophil-derived effector molecules that both have been reported to be associated with severity of sepsis6,7. HBP (also referred to as azurocidin or CAP37) is an inactive serine protease stored within azurophilic granules and secretory vesicles8. It has been associated with

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many functions including antimicrobial and immunostimulatory activity (reviewed in9). Importantly, as a potent inducer of vascular leakage, HBP has been suggested to be a central player in the pathophysiology of circulatory failure and hypotension13,14, thereby complicating severe infections. Resistin, a cysteine-rich adipocytokine, has in humans been identified as a potent pro-inflammatory molecule associated with acute and chronic inflammatory conditions12–14. We reported on resistin as a marker of severity of sepsis, with a sustained secretion profile compared to early cytokines and the highest systemic resistin levels noted in patients with septic shock6. Neutrophils were identified as a novel dominant source of resistin in sepsis patients15.

Group A streptococcus (GAS) is one pathogen that has been demonstrated to have a significant impact on neutrophils, by inducing an altered apoptotic differentiation program16 and triggering degranulation17,18. Considering the severity of fulminant invasive GAS infections, such as streptococcal toxic shock syndrome (STSS) and necrotizing fasciitis, it seems likely that neutrophils are key contributors to the multifaceted disease pathophysiology involving hyper-inflammation and pronounced vascular leakage; to perhaps a greater extent than in the case of invasive infections caused by other bacteria.

In this study we explore whether neutrophil responses, in particular the release of the sepsis-associated factors HBP and resistin, differ depending on stimuli and how this relates to sepsis of varying aetiology. To address this, neutrophil stimulation and activation was studied in vitro using primary human neutrophils exposed to clinical sepsis isolates or bacterial antigens, as well as through analyses of patients with severe acute bacterial infections.

**Results**

**HBP and resistin responses in sepsis patients of varying aetiology.** First we measured HBP and resistin levels in acute phase plasma of patients with severe sepsis/septic shock caused by different bacteria (Fig. 1). The results showed that septic patients, including both Gram-positive and Gram-negative bacterial infections, had significantly higher levels of both factors when compared to non-infected critically ill patients (Fig. 1A,B). There were no significant differences in HBP or resistin levels between patients infected with Gram-positive (n = 20) or Gram-negative (n = 28) bacteria (Fig. 1C). However, the Gram-positive cohort demonstrated a stronger correlation between HBP and resistin (r = 0.65, p = 0.003) as compared to Gram-negative cohort (r = 0.49, p < 0.001). The Gram-positive infections were predominantly caused by *Enterococcus* or *Staphylococcus aureus*, but did not include any streptococcal infections9. As GAS has been reported to have a strong impact on neutrophils15,16,18, a separate patient cohort consisting of streptococcal septic shock patients, i.e. GAS STSS patients, was also analysed. Similarly to the septic shock cohort, high levels of both HBP and resistin were detected in plasma of STSS patients, and demonstrated an even stronger correlation (r = 0.8, p = 0.016) (Fig 1D).

To further validate the finding that GAS infected patients had such strong correlation between systemic HBP and resistin, studies were undertaken to analyse these responses at the local site of infection, i.e. in infected soft tissue. Data were available from previously analysed snap-frozen tissue biopsies collected from patients with GAS necrotizing fasciitis or severe cellulitis15,20. The tissue biopsies had been analysed for expression of either HBP or resistin as well as phagocyte infiltration by intracellular immunohistochemical stainings. As previously reported, macrophages, neutrophils, HBP and resistin were detected in all infected biopsies (Fig. 1E). This can be compared to only a few resistin positive cells found in skin biopsies from healthy controls15,20. In agreement with the above-presented data from circulation, a correlation test revealed a significant correlation between HBP and resistin at the infected tissue site (r = 0.91, p = 0.0013) (Fig 1F).

**Additive effect of HBP and resistin on inflammatory response.** Next, we tested both antimicrobial and immunostimulatory properties of HBP and resistin. Antibacterial activity against GAS, *S. aureus* and *E. coli* was assessed using a Bioscreen C Microbiological Growth analyser measuring the turbidity in cultures containing a concentration range of 0.1–5 μg/ml HBP or 0.1–2 μg/ml resistin. Similar OD curves were observed for GAS and *S. aureus* in the presence or absence of resistin or HBP; thus, excluding any antimicrobial effect. In accordance with the previously reported antimicrobial activity of HBP against Gram-negative bacteria6, *E. coli* showed a slight reduction in OD during log-phase (mean % reduction: 14 ± 2), while no change in growth when resistin was added. As both HBP and resistin have been reported to exert pro-inflammatory activities9,10,12–14, it was of interest to study whether there are potential additive or synergistic effects of these factors. For this purpose, PBMCs from healthy donors were stimulated with either HBP or resistin, or the combination of the two proteins. IL-8, a classical sepsis-associated pro-inflammatory cytokine, was measured in culture supernatants. As shown in Fig. 2, IL-8 levels were significantly higher in supernatants from cells stimulated with the combination of HBP and resistin, as compared to each protein alone (p ≥ 0.03). Thus, the results indicate an additive effect of HBP and resistin in induction of inflammatory responses.

We also assessed IL-8 in the sepsis cohorts, and, as expected, elevated levels were detected in all cohorts (median ng/ml (range); STSS: 626 (306–1985); Gram-positive sepsis: 225 (28–1611); Gram-negative sepsis: 118 (51–2118)). It was noteworthy, that in STSS patients a strong correlation between IL-8 levels and HBP (r = 0.84, p = 0.009), as well as IL-8 and resistin (r = 0.86, p = 0.006) was observed, whereas no such correlation was evident in the other sepsis cohorts.

**Release of high amounts of both HBP and resistin follows neutrophil activation induced by GAS, but not *S. aureus* or *E. coli*.** As the clinical data indicated that the neutrophil response varied depending on the infectious agent, we compared the ability of different bacterial stimuli to trigger neutrophil activation and degranulation. To this end, primary human neutrophils isolated from healthy blood donors were exposed to different clinical septic shock isolates, namely *E. coli*, *S. aureus* and GAS. Filtered bacterial supernatants as well as fixed bacteria prepared from overnight cultures were used to stimulate neutrophils for 2 hours, after which HBP and resistin were measured in cell culture supernatants. Visualization of neutrophils exposed to fixed bacteria revealed starkly different responses with GAS resulting in an almost complete aggregation of the...
cells, while *S. aureus* some, and *E. coli* only marginal aggregation (Fig. 3A). Similarly, high levels of HBP and resistin were detected after stimulation with fixed GAS strains (n = 4), whereas markedly lower levels were triggered by fixed *S. aureus* (n = 2) or *E. coli* strains (n = 2) (Fig. 3B,C). Stimulation with bacterial supernatants (1:50 and 1:500 dilutions) failed to induce either HBP or resistin release. Neutrophils were also stimulated with purified bacterial proteins, including the streptococcal M1-protein and the endotoxin LPS. In agreement with the bacterial stimulation experiments, the streptococcal M1-protein resulted in release of significantly higher amounts of both HBP and resistin, as compared to LPS that was a weak trigger of release of either mediator (p < 0.008) (Fig. 3D).

Figure 1. Systemic and local HBP and resistin responses in septic patients. HBP and resistin levels in plasma collected from patients on the day of inclusion were measured with ELISA, for details see Experimental procedures. (A,B) HBP and resistin levels in plasma from patients with severe sepsis/septic shock (n = 88) or non-infected critically ill patients (n = 31). (C) Plasma HBP and resistin levels in severe sepsis/septic shock patients with confirmed Gram-positive (G+) or Gram-negative (G-) infections (n = 48). (D) Systemic HBP and resistin in STSS patients (n = 8). The correlation was determined by Pearson's correlation test, indicated by p and r-values. (E) HBP, resistin, macrophages and neutrophils were analysed by immunohistochemical staining of cryosections of snap-frozen tissue biopsies (n = 9) from patients with severe soft tissue infections caused by GAS. A control staining where the primary antibody was omitted was also performed (control). A representative tissue biopsy is shown. The scale bar indicates 50 μm. (F) The stainings were evaluated by microscope and acquired computerized image analysis (ACIA), for details see Experimental procedures. Significant correlation was determined by Spearman test, indicated by p- and r-values.
A kinetic experiment revealed a similar profile for HBP and resistin release in response to M1-protein and fixed GAS with both factors starting to appear after 15–30 minutes depending on stimuli (Fig. 3E). Thus, the data demonstrates specific neutrophil responses to different stimuli and a seemingly synchronized release of HBP and resistin after streptococcal induced neutrophil activation. It was therefore of interest to analyse granule mobilization. For this purpose, we compared unstimulated and stimulated neutrophils for intracellular HBP, resistin and the azurophilic marker MPO through triple immunofluorescent stainings combined with confocal microscopy (Fig. 4A). In unstimulated neutrophils, numerous positive granules for each factor were detected. A large portion of HBP and resistin positive granules also contained MPO, confirming their presence in azurophilic granules. However, only few granules were double positive for both HBP and resistin (Fig. 4A, unstimulated control). In contrast, neutrophils activated with different stimuli displayed starkly different granule staining patterns depending on stimuli (Fig. 4). Streptococcal M1-protein stimulated cells displayed a greater co-localization of resistin and MPO as well as HBP and resistin, when compared to unstimulated cells (Fig. 4). Quantification of the degree of co-localization between HBP and resistin showed a significantly higher increase in cells stimulated with M1-protein as compared to LPS stimulated cells (Fig. 4D,E). Thus, the data concurs with the noted differences in in vitro responses elicited by various bacterial stimuli.

Streptococcal activation of neutrophils involves multiple surface associated proteins. To further explore whether this neutrophil-stimulatory effect is exclusive to GAS or shared by other streptococcal species, we included group B, C and G streptococcus (GBS, GCS, and GGS), as well as viridans group streptococcal strains collected from septic shock patients in addition to five GAS strains including strains of serotypes T1, T4 and T28, as well as one strain of unknown type. All strains were found to induce a strong resistin response in the two donors tested, whereas they only elicited HBP release in one of the donors (Fig. 5A,B). This inter-individual variation with low or high responders was seen particularly for streptococcal triggered HBP release (Fig. 5A,B). Donor variation in streptococcal M1-protein triggered HBP release has previously been reported, and found to be linked to the presence of high anti-M1 protein specific antibody titres in the high responder 21. In line with this report, dot blot analyses of anti-M1 protein antibodies in plasma of the high and low responder revealed a stronger signal when plasma of the higher responder (donor 2) was used as compared to the low responder (donor 1) (Fig. 5C,D).

Furthermore, the presence of other plasma proteins, in particular fibrinogen, has also been implicated as an important factor for streptococcal M1-protein triggered HBP release22,21. In accordance with these reports, stimulation experiments carried out in the presence or absence of 10% autologous plasma showed a plasma-dependency for fixed GAS triggered HBP release (Fig. 5E). Notably, this dependency was only noted for HBP and resistin release was only marginally reduced when plasma was absent (Fig. 5F), suggesting different signal requirements for the release of these two factors.

The finding that GBS, GCS, GGS, and viridans group streptococci activated neutrophils to a similar extent as GAS is of interest as they share several surface attached factors such as M- or M-like proteins, pili, as well as fibronectin- and collagen-binding proteins (reviewed in23). In an attempt to identify the stimulatory factors, we first explored the role of surface-attached M1 protein by use of an M1-protein deficient mutant of 5448. However, the results showed only a marginal, non-significant, reduction in HBP and resistin release when neutrophils were stimulated with the mutant strain (Fig. 5G,H); thus implicating also other surface-attached factors and possible involvement of multiple factors in neutrophil activation. To explore this, we exploited a set of GAS strains deficient in the gene regulatory systems including Mga which is positive regulator of emm- and emm-like
gene expression within the Mga-locus\(^2\), as well as MsmR and Ralp3, both of which influence the FCT region encoding for fibronectin- and collagen-binding proteins as well as the T-pilus\(^2\)\(^4\)\(^5\). Similarly to the 5448\(\Delta\)emm\(^1\), the 591\(\Delta\)mga mutant induced equal response as the wild type strain (Fig. 5I). In contrast, 591\(\Delta\)msm\(^R\) triggered resistin release, but at a significantly reduced level as compared to the wild type strain, while 591\(\Delta\)ralp\(^3\) triggered an increased response (Fig. 5I). Taken together, the results implicate a multifactorial activation of neutrophils involving surface attached proteins encoded by the FCT-locus, rather than M- and M-like proteins, as triggers for neutrophil activation.

Finally, we explored host signals and putative receptors involved in the neutrophil activation. For this purpose, neutrophils were pre-incubated with inhibitors targeting Src-family kinases, phosphatidylinositol 3-kinases (PI3K), and the MAPK p38, extracellular signal-regulated kinase (ERK 1/2), and c-Jun N-terminal kinase (JNK), followed by stimulation with fixed GAS strains. A consistent reduction in the resistin release elicited by GAS strains was noted in neutrophils pre-treated with either Src-family kinase or p38 inhibitors, as compared to the untreated control (\(p > 0.05\)) (Fig. 6A). A small effect, although non-significant, was also seen with an inhibitor targeting PI3K, whereas inhibition of ERK1/2 and JNK had no effect (Fig. 6A). No further reduction in response
Figure 4. Subcellular localization of HBP and resistin in neutrophils. Neutrophils from healthy donors were isolated and stimulated with LPS (50 ng/ml), fMLP (5 μg/ml) or streptococcal M1-protein (1 μg/ml) for 2 h. The neutrophils were immunofluorescently stained for HBP (white), resistin (green) and MPO (red), the cell nuclei were visualized using DAPI (blue). Using scanning laser confocal microscopy, Z-stacks of the cells were acquired and vesicles positive for HBP, resistin or MPO identified using the Imaris image analysis software (processed 3D-Image). The degree of colocalization between the above-mentioned markers in the processed images was then quantified using Imaris analysis software. (A) Shows images of cells from a high responder. Bars indicate 5 μm. (B–D) Percentage of co-localization between factors as indicated. (E) Differences in co-localization in neutrophils unstimulated or stimulated with either M1-protein or LPS. Statistical significant differences were determined using the Mann Whitney test, **P < 0.01. Data represent results obtained from multiple field analyses of two experiments using different donors.
Figure 5. Pronounced neutrophil stimulatory effect can be seen across streptococcal species and involves multiple surface proteins. (A,B) HBP and resistin levels in supernatants from neutrophils isolated from two donors (D1 and D2) stimulated with different fixed streptococcal species (67 cfu/cell), as indicated in the figure. ND denotes serotype not determined. (C) Plasma from the high- and low responding donor (D1 and D2) was tested for anti-M1 protein antibodies by dot blot analyses. Purified M1-protein (100 and 10 ng) was spotted on to a PVDF-membrane, after which donor plasma was added and incubated overnight. Bound anti-M1 antibodies were detected using HRP coupled anti-human IgG. (D) The intensity of positive stain was quantified and plotted as arbitrary units (AU)/mm². (E,F) HBP and resistin levels in culture supernatants from neutrophils, either unstimulated (Unstim.) or stimulated with fixed GAS T1 (GAS), in the presence (filled symbols) or absence (open symbols) of 10% autologous plasma. The experiment was done using cells from three different donors. (G,H) HBP and Resistin levels in culture supernatants from neutrophils stimulated with fixed wild type or corresponding isogenic deletion mutants of the emm1 gene. Shown are the results from 4–5 experiments using different donors, indicated by specific symbols. (I) Resistin levels in culture supernatants from neutrophils stimulated with fixed wild type or corresponding isogenic deletion mutants of the indicated genes. Results were obtained from five experiments using different donors, the respective donors are indicated by specific symbols and the line indicates median. Wilcoxon matched-pairs signed rank test was used for comparison between groups, *P < 0.05.
was achieved through inhibition of PI3K + Src or PI3K + p38 (Fig. 6A). We also tested whether β2-integrin or TLR2 were involved in these responses by use of blocking antibodies. As previously reported, blocking of β2-integrin resulted in significant reduction of streptococcal M1-protein, but not fixed GAS, elicited responses (Fig. 6B). In contrast, the reverse was true when TLR2 was blocked, with a reduction in response to fixed GAS (Fig. 6B). Responses elicited by PMA were not affected by either antibody. It should be noted that the inhibition was far from complete, which is in line with a multifactorial activation involving several receptors and signals.

**Discussion**

In this study we demonstrate that there is a marked variation in neutrophil responses towards different bacterial pathogens. Human primary neutrophils were exposed to different clinical strains isolated from severe sepsis/septic shock cases and degranulation was assessed by measurement of the two sepsis-associated factors HBP and resistin. The stimulation experiments revealed that streptococcal strains, including not only GAS, but also GBS, GCS, GGS and viridans group streptococci as well as streptococcal M1-protein were potent triggers of both HBP and resistin release, whereas *S. aureus*, *E. coli* and LPS were all poor inducers. Although LPS is a poor inducer in comparison to the streptococcal strains, it should be noted that the magnitude of resistin release is equal to that reported in other studies. Furthermore, this relatively poor effect of the TLR-agonist LPS is in agreement with previous reports showing that, although LPS has a priming effect, it is a weak agonist of neutrophil exocytosis.

Moreover, supernatants prepared from the bacterial cultures failed to induce any HBP or resistin release. Thus, in the case of streptococcal bacterial activation, our data suggest that surface-associated factors, rather than secreted factors, are involved in triggering neutrophil degranulation. However it should be noted that in a previous report, HBP-release was observed following stimulation with GAS overnight culture supernatants, but only by supernatants containing high levels of streptolysin O (SLO). Thus we cannot exclude that the lack of a response induced by supernatants might be due to a low production of SLO by the strains used in this study. This also raises the question as to how the expression of virulence factors in vitro compares to that in patients. Humoral responses in patients show that superantigens and streptolysins are produced, but there are no data pertaining to their concentrations. Here we are using supernatants containing a mixture of the secreted virulence factors, as this is as close to the clinical setting as we can technically reproduce. Along the same lines, live infection would be associated with confounders such as varying degree of phagocytosis, intracellular replication, as well as toxin-mediated cytotoxicity, which would complicate interpretation.

Previous reports on neutrophil activation by streptococcal strains have focused on GAS and particularly on the streptococcal M-protein, which in a soluble form was identified as a key trigger of neutrophil activation and degranulation. Herwald et al. delineated the underlying mechanism and showed that it entailed complex formation with fibrinogen and subsequent β2-integrin binding. In this study, strains of different types, including T1 (coupled to emm1/M1 serotype) as well as T4 and T28 types were tested. The latter two types both express the M-related protein MRP4 that, similarly to M1-protein, has two fibrinogen binding sites. Streptococcal strains

**Figure 6. Signalling molecules and receptors involved in the release of resistin.** Resistin levels in cell culture supernatants of neutrophils from healthy donors, pre-treated for 30 min with inhibitors targeting the signalling molecules Src (PP1), PI3K (LY294002), MAPK p38 (SB203580), ERK (PD98059), and JNK (SP600125) (A) or antibodies against β2-integrins (anti-human CD18) and TLR-2 (anti-Human CD282) (B), and stimulated with fixed GAS for 2 h. In A, open and filled symbols show 5448 and the T1 isolate, respectively. Mean ± SD of results from 3–9 experiments using different donors presented as per cent relative to untreated cells, the dashed line indicates 100 per cent. The inhibitors did not influence the cell response, as unstimulated cells had similar background levels in the presence or absence of the inhibitors. Wilcoxon matched-pairs signed rank test was used for comparison between groups, *P > 0.05,**P < 0.01.
commonly express fibrinogen-binding proteins, which in GAS, GBS, GCS and GGS predominantly consist of members of the M/M-like protein family. In viridans group streptococci, a phage lysin with fibrinogen-binding capacity has been identified. Hence, streptococcal surface associated M-protein and other fibrinogen-binding proteins represent putative candidates that might contribute to the noted neutrophil activation. However, experiments using an M1-deficient GAS strain yielded only a marginal reduction in the response; thus suggesting that surface attached M-protein is not the main trigger or that there are multiple streptococcal factors involved. Streptococcal surface attached factors including M- and M-like proteins, pili structures and extracellular matrix binding proteins, such as Fibronectin and Collagen binding proteins are under the control of gene regulatory systems. We therefore tested bacterial mutants deficient in gene regulatory systems including Mga, which is a regulator influencing emm and emm-like genes, as well as MsmR and Ralp3 influencing the FCT-locus encoding for Fibronectin- and Collagen-binding proteins as well as T-pilus. Similar to the M1-mutant strain, deletion of Mga did not impair the stimulatory capacity. Furthermore, the decreased Mga-operon transcript abundance and increased FCT-locus transcription in Ralp3 knockout strain, underlines rather the involvement of FCT-region proteins than emm and emm-like genes in neutrophil activation. In addition, the MsmR-deficient strain triggered a reduced response. In msrR- and ralp3-mutants several genes are differentially expressed including the genes encoding Fibronectin binding protein F2 (prtF2) and Collagen binding protein (cpa). prtF2 and cpa were both down regulated in the msrR-mutant while upregulated in the ralp3-mutant, although the stimulatory capacity was reduced with the msrR-mutant, suggesting involvement of the FCT-region proteins as neutrophil activators, it still triggered a substantial activation. Taken together, this implies the involvement of multiple factors in streptococcal triggered neutrophil activation. Also attempts to block signal-transduction pathways using pharmacological inhibitors and blocking antibodies revealed a role for Src family kinases and TLR2 in streptococcal neutrophil activation and degranulation; however, the blocking effect was far from complete, which is expected if several bacterial ligands are involved and interact with multiple receptors.

The study revealed marked inter-individual variations particularly in HBP-responses between donors with some being low responders and others high responders. The HBP, but not resistin, response was fully dependent on the presence of plasma, and in accordance with a previous report, the donor variation in HBP release upon GAS stimulation seemed to be linked to presence or absence of anti-M protein antibodies. In the report by Kahn et al. it was proposed that the high response was related to presence of antibodies targeting the central region of the M-protein resulting in complexes containing M-protein, fibrinogen and IgG, which triggered HBP release through dual engagement of Fc-receptor and αI-integrin. In our study, the high responder did not only respond strongly to M1 protein and GAS strains but also to GBS, GCS, GGS and viridans group streptococci. At present, the underlying mechanism for this general high response to streptococcal strains remains to be elucidated, but as M and M-like proteins share homologies in the central regions, it is tempting to speculate that the effect might be related to cross-reactive antibodies. Notably, streptococcal strains triggered release of resistin in all donors and the donor variation was exclusively linked to HBP release; thus, indicating involvement of different receptors and/or signal-transducing pathways for exocytosis of HBP and resistin.

Visualization of the granule content through intracellular immunostainings of stimulated cells followed by confocal microscopy revealed that HBP and resistin were localized predominantly in separate granules, many of which were MPO-positive azurophilic granules. Notably, upon activation a pronounced increase in co-localization of HBP and resistin was evident in cells stimulated with streptococcal M1-protein, but not with LPS nor with fMLP. In addition, the granules appeared larger in size, which indicates a mobilization of HBP- and resistin-positive granules in response to M1-protein. Taken together with the similar kinetic response profile, the data is indicative of a synchronized release of these two factors in response to streptococcal stimulation. This was further supported by analyses of HBP and resistin responses in septic shock cohorts. All septic shock patients had elevated levels of plasma HBP and resistin as compared to critically ill non-infectious patients, and a positive correlation between plasma HBP and resistin levels was found in all patient cohorts (i.e. Gram-positive and Gram-infections, STSS caused by GAS. However, the correlation was particularly striking in the STSS cohort, both systemically in plasma as well in soft tissue biopsies. As it had not previously been explored how the co-presentation of these two effector molecules might influence inflammation, we also tested the response of PBMC to HBP and resistin alone or in combination. The results demonstrated that cells exposed to both factors in combination responded with a significantly higher inflammatory response, as determined by IL-8 release. Notably, measurement of IL-8 levels in the patients revealed a positive correlation between IL8 with HBP and resistin, respectively, in STSS patients, but not in the large sepsis cohort.

Taken together, the data support the concept that neutrophil activation differs depending on the bacterial stimuli, which may have clinical implications. Considering the greater stimulatory capacity of the streptococcal strains tested in this study, it seems plausible that neutrophil activation and degranulation may play a central role in the pathology of streptococcal infections. This has important implications for therapeutic strategies, especially considering that several trials have attempted to enhance neutrophil recruitment and function in sepsis. On this note, our data underline the importance of further studies on this topic to provide insight into patient groups most likely to benefit from neutrophil modulation.

**Experimental procedures**

**Patient cohorts.** Plasma samples from culture-positive severe sepsis and septic shock patients (n = 88), including 20 and 28 patients with confirmed Gram-positive and Gram-negative bacterial infections, respectively, and a reference group of non-infected critically ill patients (n = 31) enrolled at Karolinska University Hospital Huddinge were used. The three cohorts were well matched with respect to age, gender and severity of infection based on APACHE II score at the day of inclusion (median APACHE II (range) Gram-positive: 23 (12–37); Gram-negative: 25 (9–45); reference group: 22 (9–33)). Plasma from patients with STSS caused by GAS (n = 8)
were provided from a sepsis study conducted at Lund University Hospital\(^1\). No APACHE II score was available from the STSS-cohort. All plasma samples were collected during the acute septic episode.

Snapfrozen tissue biopsies from patients with necrotizing fasciitis or severe cellulitis caused by GAS (n = 9) were used. This material has been described previously\(^{34,35}\).

All studies were ethically approved by Karolinska University Hospital, the University of Toronto, and Lund University Hospital, and all experiments were carried out in accordance with the approved guidelines. Written informed consent was obtained from all patients or their legal guardians.

**Bacterial isolates and factors.** Blood isolates, including GAS (n = 4), *E. coli* (n = 2) and *S. aureus* (n = 2), collected from the septic shock patients in respective cohort described above were used. The GAS strains were of serotypes T1 (n = 2), T4 and T28. Also, streptococcal strains, including group B (GBS; n = 1), group C (GCS; n = 2), group G (GGS, n = 1), and viridans group streptococci (n = 2), isolated from patients with severe sepsis were used. In addition, isogenic mutants deficient in M1-protein (5448ΔemrR\(^{34}\)) or in gene regulatory systems including 591Δmsmr\(^{23}\), 591Δralp\(^{34}\) and 591Δmga\(^ {23}\) were used. The streptococcal strains were cultured overnight (16h) in Todd-Hewitt medium with 1.5% yeast, whereas LB medium was used for *E. coli* strains and CCY medium for *S. aureus*. Different media were used to obtain optimal growth conditions known to support exotoxin production for respective bacteria. Media controls were included in all stimulation experiments. Following centrifugation, the supernatants were sterile-filibrated and the bacterial pellet fixed with 1% paraformaldehyde for 45 minutes.

The streptococcal M1-protein was purified from the AP1-derived isogenic mutant MC25 strain as previously described\(^{34}\). Commercially available lipopolysaccharide (LPS) from *E. coli* (Sigma-Aldrich, St Louis, MO) was used.

**Antibacterial activity assay.** *S. pyogenes* (T1 strain), *S. aureus* and *E. coli* were grown in liquid broth at 37°C over night. The cultures were diluted 1:50 (S. pyogenes) and 1:180 for *E. coli* and *S. aureus*. In order to determine the antibacterial effect of HBP (R&D systems, Minneapolis, MN) and resistin (PeproTech, Rocky Hill, NJ), the bacterial cultures were exposed to different concentrations of the peptides. Final concentrations of 5 μg/ml, 2 μg/ml, 1 μg/ml and 0.1 μg/ml of HBP and 2 μg/ml, 1 μg/ml, 0.1 μg/ml of resistin (Innovagyn, Lund Sweden) were tested. An optical density (OD) curve was generated based on the turbidity measurement at 420–580 nm, using a Bioscreen C over a time of 24 h. Measurements were made every 15 min and data exported to a PC. Data were analysed with GraphPad Prism version 6.0 for Mac (GraphPad software).

**Infection and stimulation of human primary cells.** Human neutrophils and peripheral blood mononuclear cells (PBMCs) were isolated from venous blood collected from healthy individuals by Polymorphophreps or Ficoll-Hypaque (Axis-Shield, Oslo, Norway) gradient centrifugation, respectively. Cells were resuspended in RPMI-1640 supplemented with 5% FCS, 10 mM L-glutamine, Penicillin (100 U/ml)/Streptomycin (100 μg/ml) and 25 mmol/L HEPES (all from Thermo Scientific, Hyclone, South Logan, UT). 5 × 10\(^{4}\) neutrophils/ml were stimulated with streptococcal M1-protein, LPS (extracted from *E. coli*), Phorbol Myristate Acetate (PMA) (Sigma-Aldrich, St Louis, MO), N-formyl-methionine-leucine-phenylalanine (fMLP) (Calbiochem, San Diego, CA), or with fixed bacteria (67 cfu/cell, unless otherwise indicated) or bacterial supernatants (1:50 and 1:500 dilutions). In case of inhibition of molecules or receptors, isolated neutrophils were pre-incubated with 10 μM of inhibitor or 10 μg blocking antibody for 30 minutes.

The dose of fixed bacteria was chosen based on dose-response experiments, which showed a stable response over a broad multiplicity of infection (MOI) range, i.e. 17–260 cfu/cell. Also the concentrations of streptococcal M1-protein and LPS were chosen based on dose-response experiments in which 0.5 or 1 μg/ml gave a peak response for M1 stimulation, whereas LPS showed an equal response over the range of 50 ng/ml–1 μg/ml. Streptococcal M1-protein, bacterial supernatants and fixed bacteria were pre-incubated for 10 minutes at 37°C with 10% autologous plasma to allow for fibrinogen-complex formation as described previously\(^{18}\). At defined time points, the cells were centrifuged and cell-free culture supernatants collected for analyses.

PBMCs (1 × 10\(^{6}\) cells/ml) were stimulated with HBP (R&D systems, Minneapolis, MN) and resistin (PeproTech, Rocky Hill, NJ), each factor alone and in combination for 2h after which culture supernatants were collected.

The following blocking antibodies and pharmacological inhibitors were used: anti-β2 integrin (anti-human CD18, clone IB4, Merk Millipore, Billerica, MA, USA), anti-TLR2 (anti-human CD282, clone TL2.1, eBioscience, San Diego, CA, USA), anti-TLR4 (clone 119, eBioscience, San Diego, CA, USA), anti-CD14 (clone M1/70, eBioscience, San Diego, CA, USA), anti-CD18, clone IB4, Merk Millipore, Billerica, MA, USA), anti-TLR2 (anti-human CD282, clone TL2.1, eBioscience, San Diego, CA, USA), anti-TLR4 (clone 119, eBioscience, San Diego, CA, USA), anti-CD14 (clone M1/70, eBioscience, San Diego, CA, USA), anti-CD3 (clone 3B5, eBioscience, San Diego, CA, USA), anti-CD2 (clone GB2, eBioscience, San Diego, CA, USA), and anti-CD247 (clone M49/3A5, eBioscience, San Diego, CA, USA). The streptococcal strains were cultured over a broad multiplicity of infection (MOI) range, i.e. 17–260 cfu/cell. Also the concentrations of streptococcal M1-protein, bacterial supernatants and fixed bacteria were pre-incubated for 10 minutes at 37°C with 10% autologous plasma to allow for fibrinogen-complex formation as described previously\(^{18}\). At defined time points, the cells were centrifuged and cell-free culture supernatants collected for analyses.

**Dot-blot assay.** Streptococcal M1-protein (obtained as described above) or PBS (negative control) was spotted onto a PVDF-membrane, and left to absorb for 30 min. The M1-probed membrane was subsequently washed using a 0.1% Tween (Sigma-Aldrich, St Louis, MO) in PBS solution (PBST), and blocked for 2h using 5% milk in PBST. After blocking, the membrane was washed using PBST and incubated with donor plasma (diluted 1:10 with 2.5% milk in PBST) over night at 4°C. Following the over night incubation, the membrane was carefully washed and incubated with a horseradish peroxidase labelled secondary anti-Human IgG antibody (GE Healthcare, Piscataway, NJ) (diluted 1:10.000 with 2.5% milk in PBST). The blot was developed using an electrochemiluminescence kit (Thermo Scientific, South Logan, UT). As a positive control, the membrane was incubated with anti-M1 rabbit serum and detected using anti-rabbit IgG. Intensities were evaluated using a luminescent image...
analysed (LAS-4000 mini) and quantified using the Multi Gauge software version 3.2 (both from Fujifilm, Minato, Tokyo, Japan).

**Analysis of secreted HBP, resistin and IL-8.** HBP was analysed by ELISA as previously described. Resistin was analysed by a commercially available ELISA (BioVendor, Brno, Czech Republic), according to the manufacturer’s instructions. IL-8 levels were measured in a Luminex analysis (Invitrogen, Grand Islands, NY).

**Immunostainings of tissue biopsies and primary human cells.** Snapfrozen biopsies (n = 9) were sectioned, fixed, immunostained, and analysed as described previously. The whole section was analysed by acquired computerized image analysis (ACIA). The results are presented as ACIA values, which equals the percentage of the positively stained area x the mean intensity of positive staining.

Stimulated neutrophils were fixed with 2% paraformaldehyde for 10 min, washed with PBS and added to glass coverslips using cytoospin. Triple immunofluorescence stainings were conducted as follows; the cells were permeabilized using 0.1% saponin in PBS, and initially blocked with 2% FCS in PBS-saponin for 5 minutes at 37°C. This was followed by 30 minutes blocking with 0.1% BSA-æ (Sigma-Aldrich, St Louis, MO) in PBS-saponin, after which the primary antibodies were added for 30 minutes at 37°C. Prior to addition of the fluorophore-conjugated secondary antibodies, cells were blocked for 30 min with 1% normal goat serum in PBS-saponin. The secondary antibodies were incubated for 30 min in the dark.

The following antibodies were used: anti-neutrophil elastase (NP57); anti-CD68 (EBM11) (both from DakoCytomation, Carpinteria, CA), anti-Resistin (AF1359 and MAB1359)(both from R&D Systems, Minneapolis, MN), anti-MPO (sc-33595, Santa Cruz, Dallas, Texas), and the secondary antibodies donkey anti-mouse IgG-Alexa488 (122062, donkey anti-rabbit IgG-Alexa546 (112657) and donkey anti-goat IgG-Alexa647 (1235825) (all from Molecular Probes, Grand Islands, NY). HBP was identified with purified IgG from a polyclonal anti-HBP rabbit serum or IgG purified from a monoclonal anti-HBP mouse serum (2F23A).

For evaluation a Nikon A1R confocal microscope was used (Nikon Instruments, Amstelveen, the Netherlands). Z-stacks were acquired for 3–5 fields of view, using a 100× oil plan apochromatic objective.

**Co-localization analysis.** Z-stack images were analysed using the Imaris 3-D image analysis version 7.6.3 (Bitplane, Zürich, Switzerland). The Imaris cell function was then applied to automatically locate the vesicles inside the cells based on size and intensity thresholds defined by the user. Measurement of co-localization was carried out by masking each respective vesicle type, creating new channels for each marker. Total number of co-localized voxels was measured, and thresholds were automatically computed by using orthogonal regression analysis of the image’s scatterplots in combination with the Pearson’s coefficient approach.

**Statistical evaluation.** Data were analysed by GraphPad Prism version 4.0 for Windows (GraphPad software). Mann-Whitney U test, Wilcoxon matched-pairs signed rank test or Kruskal Wallis test was used for comparison between groups. Correlations between variables were determined by use of Pearson correlation test or, in the case of non-Gaussian distribution of the data, Spearman rank correlation coefficient. Differences were considered significant when p < 0.05.

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Author Contributions

J.S., A.L., I.J. and A.N.T. designed the study. J.S., A.L., U.I., H.H. and M.J. performed the experiments. J.S., A.L., N.S., L.J. and A.N.T. analysed the experiment data. A.L. and A.L. recruited patients, collected patient material and bacterial isolates. A.L. and I.J. analysed patient data. B.K. contributed with bacterial mutants. H.H., L.J. and A.N.T. contributed to the reagents. J.S., A.L., N.S., L.J. and A.N.T. wrote the manuscript. All authors read and approved the final version of the manuscript.

Additional Information

Competing financial interests: All authors, except HH and AL, hereby confirm that they have no conflict of interest in the publication of this research. HH and AL are listed as inventors for a patent application on the use of HBP as a diagnostic tool in sepsis, filed by Hansa Medical AB.

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