The 5'-nucleotidase S5nA is dispensable for evasion of phagocytosis and biofilm formation in *Streptococcus pyogenes*

Marcel-Lino Dangel, Johann-Christoph Dettmann, Steffi Haßelbarth, Martin Krogull, Miriam Schakat, Bernd Kreikemeyer, Tomas Fiedler*

Rostock University Medical Centre; Institute of Medical Microbiology, Virology, and Hygiene, Rostock, Germany

¤ Current address: Miltenyi Biotec GmbH, Bergisch Gladbach, Germany

¤ Current address: Department of Medicine I, University Medical Center Hamburg-Eppendorf, Hamburg, Germany

* tomas.fiedler@med.uni-rostock.de

Abstract

5'-nucleotidases are widespread among all domains of life. The enzymes hydrolyze phosphate residues from nucleotides and nucleotide derivatives. In some pathobiontic bacteria, 5'-nucleotidases contribute to immune evasion by dephosphorylating adenosine mono-, di-, or tri-phosphates, thereby either decreasing the concentration of pro-inflammatory ATP or increasing the concentration of anti-inflammatory adenosine, both acting on purinergic receptors of phagocytic cells. The strict human pathogen *Streptococcus pyogenes* expresses a surface-associated 5'-nucleotidase (S5nA) under infection conditions that has previously been discussed as a potential virulence factor. Here we show that deletion of the S5nA gene does not significantly affect growth in human blood, evasion of phagocytosis by neutrophils, formation of biofilms and virulence in an infection model with larvae of the greater wax moth *Galleria mellonella* in *S. pyogenes* serotypes M6, M18 and M49. Hence, the surface-associated 5'-nucleotidase S5nA seems dispensable for evasion of phagocytosis and biofilm formation in *S. pyogenes*.

Introduction

*Streptococcus pyogenes* (Group A *Streptococcus*) is a Gram-positive human pathogen primarily causing purulent infections of the skin (impetigo) and throat (pharyngitis, tonsillitis) and frequently also severe invasive or systemic diseases such as necrotizing fasciitis, sepsis or streptococcal toxic shock-like syndrome [1]. The list of newly identified factors contributing to virulence mechanisms of *S. pyogenes* is constantly growing. To date, more than 50 virulence factors have been described in *S. pyogenes* [2]. Recently, the 5’-nucleotidase S5nA has been added to that list of potential virulence factors [3].

5’-nucleotidases (5’NT) are enzymes hydrolyzing phosphate groups at the 5’-end of ribose or deoxyribose in nucleotides or nucleotide-derivatives [4]. These enzymes are widely...
distributed among bacteria, plants and vertebrate tissues [4–6]. In bacteria cytoplasmic and periplasmic as well as membrane-associated 5'NT have been found [4]. Especially the membrane associated 5'NT are discussed to be associated with virulence in several pathogenic or pathobiontic bacteria such as *Vibrio cholerae* [7], *Pseudomonas aeruginosa* [8], *Staphylococcus aureus* [9] and *Streptococcus agalactiae* [10]. The immune-modulating effect of surface-exposed 5'NT of these bacteria is attributed to their ability to dephosphorylate adenosine phosphates. This can lead to a decreased concentration of the proinflammatory ATP and/or increased concentrations of adenosine, both mediating suppression of phagocytic activity via purinergic receptors on phagocytic cells [9–12]. Furthermore, in the non-pathogenic bacterium *Xylella fastidiosa*, an impact of a 5'NT on biofilm formation has been speculated [13].

In proteome analyses of *S. pyogenes* M49, a putative 5'NT with a leucine-proline-isoleucine-threonine-glycine (LPITG) cell wall anchoring motif has been found in the surface-associated sub-proteome [14]. *In vitro* the 5'NT gene of *S. pyogenes* M1 SF370 is expressed in the early exponential growth phase but not in the stationary phase [15]. The presence of antibodies against the *S. pyogenes* 5'NT in serum samples of patients indicates that the enzyme is produced under infection conditions [16]. The 5'NT S5nA of *S. pyogenes* has been associated with immune evasion, as the addition of a recombinant S5nA of *S. pyogenes* M1 SF370 increased survival of the non-pathogenic *Lactococcus lactis* in human blood [3]. Whether S5nA actually contributes to virulence in *S. pyogenes* has not been investigated so far. In the work presented here we therefore analyzed virulence traits of S5nA mutants of three different *S. pyogenes* strains of the serotypes M6, M18 and M49.

### Materials and methods

#### Bacterial strains and culture conditions

*S. pyogenes* serotype M6 strain K006 [17], M18 strain MGAS8232 [18], and M49 strain 591 were grown in Todd-Hewitt broth supplemented with 0.5% (w/v) yeast extract (THY medium) or Brain Heart Infusion (BHI) medium at 37 °C in 5% CO₂ enriched ambient air in standing cultures. *S. pyogenes* deletion mutants were cultivated in THY medium containing 30 mg/L kanamycin. *S. pyogenes* strains carrying pIB184 derivatives were grown in THY containing 5 mg/L erythromycin. *Escherichia coli* DH5α was grown in Lysogeny Broth (LB) at 37 °C under ambient air. *E. coli* strains harbouring pSF151 [19] or pUC18Erm1 [20] derivatives were grown in LB containing 50 mg/L kanamycin or 300 mg/L erythromycin, respectively.

#### Construction of recombinant vectors and *S. pyogenes* strains

For the construction of *S. pyogenes* 5'-nucleotidase gene knockout mutants, the upstream and downstream flanking regions of S5nA genes of *S. pyogenes* M6 K007 (M6_Spy0695 in the reference genome of *S. pyogenes* M6 MGAS10394), *S. pyogenes* M18 MGAS8232 (SpyM18_0933) and *S. pyogenes* M49 591 (Spy49_0686 in the reference genome of *S. pyogenes* M49 NZ131) were PCR-amplified and ligated into the pUC18Erm1 vector [20]. The kanamycin resistance gene *aphA* was amplified from vector pSF151 and ligated between the flanking regions. The resulting plasmid was electroporated into *S. pyogenes* M6 K006, M18 MGAS8232 and M49 591, respectively. Transformants were selected on THY agar plates with the respective antibiotics. Single crossover mutants were kanamycin- and erythromycin-resistant while double crossover deletion mutants were kanamycin-resistant but erythromycin-sensitive. No polar effects have to be considered as the S5nA gene is not part of an operon. For generation of complementation strains the M18 S5nA gene including promoter region was PCR-amplified and ligated into the shuttle vector pIB184 [21] via Apal and BamHI restriction sites. Primers used in this study are listed in Table 1.
Quantitative real-time PCR

For RNA isolation, bacteria were grown in 20 ml of THY, harvested in the exponential growth phase (optical density at 600 nm = 0.3) and quickly frozen in liquid nitrogen. As described by Pappesch et al. [22], RNA was isolated with the Direct-zol RNA MiniPrep Kit (Zymo Research), subsequent acid phenol:chloroform:isoamyl alcohol (125:24:1) extraction and TURBO DNase treatment. cDNA was generated with the SuperScript first-strand synthesis system for RT-PCR (Thermo Fisher Scientific). SYBR green (Thermo Fisher Scientific)-based quantitative real time PCR was carried out on a ViiA Real-Time PCR System (Applied Biosystems). The 5S rRNA gene was used as a housekeeping gene. Primers used for S5nA and 5S cDNA detection are listed in Table 1.

Table 1. Oligonucleotides.

| Primer name | Oligonucleotide sequence (5'→3') | Use                                      |
|-------------|---------------------------------|------------------------------------------|
| FR1_fw      | ATTTGTCGACAAATATCTCTAAAGTT      | Amplification of upstream and downstream flanking regions of S5nA genes |
| FR1_re      | TTAAAGATTTCCATCGAGCTTAT         |                                          |
| FR2_fw      | TAAATCTGCAGCTATCTCAAT          |                                          |
| FR2_re      | GGAAGTCGACCGCTTTAATGTAATCTAG   |                                          |
| apfA_fw     | GCCGCGACGCTCAAGACGAGGAGT       | Amplification of kanamycin resistance gene apfA |
| apfA_re     | GCCGCACTCCTACATCCTAG           |                                          |
| comp_fw     | GAGCGGGGCAATATAGCTGAGAGT       | Amplification of S5nA genes for complementation |
| comp_re     | CAGGATCCATTAGATCGAGTA          |                                          |
| 20Sp5'NT_001qF | CGGCCGCTTTGATAATACCG      | S5nA qPCR                                     |
| 20Sp5'NT_002qR | CAGAGTTAGCGAGCTGCG          |                                          |
| 5SqF        | ACAGGATCCCTATCCTCACCAG        | 5S rRNA qPCR                                  |
| 5SqR        | GAGATAACACCGTAACCATG          |                                          |

Underlined letters indicate restriction sites.

Measurement of 5'-nucleotidase activity

5'-nucleotidase activity was determined from bacteria growing exponentially in THY medium at an optical density at 600 nm of 0.5 (exponential growth between OD600 of 0.2 to 0.7 for all strains, see S1 Fig). For that purpose, bacteria from 20 ml culture were harvested by centrifugation. Pellets were washed in physiological NaCl solution, suspended in 450 μl of a buffer containing 50 mM Tris (pH 6.5), 2 mM MgCl₂ and 1 mM adenosine-5'-monophosphate (AMP) and incubated for 30 min at 37 °C. As controls, on the one hand bacteria were incubated in buffer without AMP and on the other hand the reaction mixture was incubated without bacteria. Subsequently, bacteria were pelleted by centrifugation and free phosphate in the supernatant was determined. For that purpose, 400 μl supernatant were mixed with 400 μl ammonium-vanadate solution (21 mM in 0.28 M HNO₃) and 400 μl ammonium-molybdiate solution (40 mM in 1.25 M H₂SO₄), incubated for 10 min at room temperature and extinction was measured at 405 nm. Phosphate concentrations were calculated based on a calibration curve. The calibration curve was generated with dipotassium phosphate solutions covering concentrations from 12.5 to 1000 μM. Values of AMP-free controls were subtracted from the other values. Controls without bacteria did not contain detectable amounts of inorganic phosphate.

Biofilm cultivation and quantification

For biofilm assays, bacterial overnight cultures in THY were suspended in fresh BHI supplemented with 0.5% glucose, adjusted to 10⁴ CFU/ml and inoculated into 96-well microtiter
plates. Wells were coated overnight at 4 °C with 2 μg/well human collagen I, collagen IV, or fibronectin (Sigma) in PBS. After incubation for 24, 48 or 72 h as standing cultures at 37 °C in a 5% CO₂ / 20% O₂ atmosphere, the biofilms were quantified in a Spectramax M2 plate reader after staining with crystal violet as described previously [17].

**Blood survival/Growth assay**

The blood survival assays were performed as described previously [23]. Briefly, overnight cultures of *S. pyogenes* were inoculated into fresh THY medium and grown to the exponential growth phase. Exponential growth was observed between OD600 of 0.2 to 0.7 for all strains (S1 Fig). Bacteria were harvested by centrifugation at an OD600 of 0.3 to 0.4, set to an optical density at 600 nm of 0.25, and further diluted 1:10,000 in PBS. The viable counts of this suspension were determined. Twenty microliters of the suspension were incubated with either 480 μl of heparinized human blood at 37 °C with rotation. After 3 h, viable counts in the blood samples were determined and correlated with the inoculum to calculate multiplication factors.

**Quantitative phagocytosis assay**

The quantitative phagocytosis assay was performed with primary neutrophils as described previously [22]. Briefly, neutrophils were isolated from human citrated blood using Polymorph-Prep (Progen Biotechnik GmbH, Heidelberg) following the manufacturer’s instructions. Neutrophils were suspended in RPMI medium. 10⁷ neutrophils and opsonized bacteria were mixed in a 1:1 ratio. After 30 min of co-incubation, the viable counts of the bacteria were determined and correlated with a mock control of bacteria mixed with an equal volume of RPMI medium instead of neutrophils.

**Ethics approval statement**

The protocol for the collection of human blood for the blood survival assays and the isolation of neutrophils was approved by the Ethikkommission an der Medizinischen Fakultät der Universität Rostock (Ethics Committee vote: A 2014–0131). The experiments were conducted in accordance with the ICH-GCP guidelines. Oral informed consent was obtained from all subjects.

**Galleria mellonella infection model**

Larvae of the greater wax moth *Galleria mellonella* were obtained from Reptilienkosmos (Niederkrüchten, Germany). Infection experiments were carried out as described elsewhere [24]. In short, *S. pyogenes* strains were grown overnight in THY, washed twice in a 0.9% NaCl solution and suspended in 0.9% NaCl to a final concentration of 1.5–2x10⁸ CFU/ml. Larvae with a weight of 150–200 mg were inoculated with 10 μl of this bacterial suspension, resulting in an infection dose of 1.5–2 x 10⁶ CFU/larva. Bacteria were injected into the hemocoel of the larvae between the last pair of legs using a microapplicator (World Precisions Instruments, Sarasota, USA) and a fine dosage syringe (Omnican-F, 0.01 ml–1 ml, 0.30x12 mm, B. Braun AG, Melsungen, Germany). As a control, larvae were mock inoculated with 10 μl of a 0.9% NaCl solution. Survival of the larvae was observed for seven days. Larvae were regarded dead when they did not move upon repeated physical stimulation with tweezers.

**Results**

5’NT have been described to contribute to virulence in several bacteria [3, 7–10]. S5nA, a recombinant 5’NT of *S. pyogenes* has been shown to increase survival of *L. lactis* in the
presence of phagocytes in vitro [3]. Therefore, we aimed to elucidate whether deletion of the S5nA gene actually affects the virulence of S. pyogenes. For this purpose the S5nA genes were replaced by a kanamycin resistance gene in three different S. pyogenes strains, i.e. the serotype M49 strain 591 (reference sequence ORF spy49_0686c), the M18 serotype strain MGAS8232 (reference sequence ORF spyM18_0933) and the M6 serotype strain K006 (reference sequence ORF M6_Spy0695). Furthermore, complementation strains expressing the S5nA gene from a pAT19-based plasmid under the control of the native promoter were constructed.

To assess whether the S5nA gene deletions led to a general growth deficiency of the bacteria, growth of all strains in complex laboratory medium THY was measured. Neither the mutants nor the complementation strains had any significant growth deficiencies as compared to the cognate wild type strains in THY medium. The respective specific growth rates in the exponential phase of growth and optical densities after overnight growth are listed in Table 2. The full growth curves are shown in S1 Fig.

The absence or presence of S5nA gene transcripts in exponentially growing wild type, mutant, and complementation strain cells was detected via qPCR with 5S rRNA as a housekeeping reference gene. As expected, no S5nA gene transcripts were detectable in the mutant strains whereas S5nA transcripts were present in all wild type and complementation strains (see S1 Table). In the complementation strains, however, the relative abundance of S5nA gene mRNA was notably higher than in the wild-type strains, likely because in these strains the gene is transcribed ectopically from a plasmid that is present in more than one copy per cell.

Since it has been proposed that S5nA of S. pyogenes contributes to virulence by degrading AMP (and to some extent ADP) and consequently increases levels of the anti-inflammatory adenosine [3], we analyzed whether the deletion of the above-mentioned S5nA gene of S. pyogenes M49 actually leads to a loss of AMP dephosphorylation activity on the surface of the bacteria. For that purpose, exponentially growing bacteria of wild type and mutant strains were harvested at an optical density of 0.5, washed, and incubated with adenosine-monophosphate (AMP) in a phosphate-free buffer. After 30 min, phosphate concentrations were measured. About 30 μM of inorganic phosphate was hydrolyzed from AMP by the S5nA deletion mutant while wild type and complementation strains released about 150 μM of inorganic phosphate from 1 mM of AMP in the same time (Fig 1).

S. pyogenes S5nA activity releases adenosine from AMP and ADP [3]. Adenosine triggers purinergic receptors of phagocytic cells leading to a suppression of their phagocytic activity [9, 11, 12]. Therefore, we tested the growth of the S5nA gene deletion strains in blood by inoculating the exponentially growing bacteria in fresh heparinized human blood and comparing viable counts of the bacteria before and after 3 h of incubation. As can be seen in Fig 2, the S5nA gene deletion strains had no serious disadvantage in the blood environment. Although in the M18 and M49 background the S5nA gene deletion strains had a slightly lower multiplication rate than the wild types, these differences were not statistically significant (Fig 2). Furthermore, at least in the M18 background, the complementation strain was not able to reverse this tendency, indicating that there is no S5nA specific effect. These data indicate that S5nA does not significantly contribute to innate immune evasion of S. pyogenes in blood.

To further validate these results, we investigated whether S5nA gene deletion renders S. pyogenes more prone to phagocytosis. As neutrophils account for the majority of phagocytic cells in blood, opsonized bacteria were mixed 1:1 with freshly isolated neutrophils and viable counts were determined after 30 min of incubation. While neutrophils were able to reduce the viable counts of M18 and M49 strains by about one order of magnitude within 30 min, the M6 strains were rather resistant to phagocytic clearance by the neutrophils. However, in accordance with the observations in the blood survival experiments neither of the S5nA deletion
strains was killed more efficiently than its cognate wild type and complementation strain (Fig 3).

Besides the implication of 5'NT in the inhibition of phagocytosis in several bacteria [3, 9, 10, 25, 26], for X. fastidiosa an involvement of 5'NT in biofilm formation has been proposed [13]. Therefore, we also investigated the impact of the loss of S5nA on biofilm formation of S. pyogenes. Since S. pyogenes strains differ in their ability to form biofilms on abiotic or protein-coated surfaces [27, 28], we analyzed biofilm formation on polystyrene and also on collagen I, collagen IV and fibronectin coated surfaces. Biofilm masses were assessed after 24 h, 48 h and 72 h of growth in 96-well plates. While the general biofilm-formation capacity was clearly strain-dependent, no significant differences in biofilm masses were observed between wild type, S5nA gene deletion mutant and complementation strain of the same serotype (for 72 h incubation values see Fig 4; for 24 and 48 h values see S2 Fig).

Finally, the virulence of the S5nA gene deletion strains was tested in an infection model using the larvae of the greater wax moth Galleria mellonella. For this purpose the animals were infected with defined doses of bacteria and survival of the larvae was observed for 10 d post infection.

Table 2. Growth of S. pyogenes wild type, S5nA deletion and complementation strains in THY medium.

| Strain       | Specific growth rate μ in h⁻¹ | Optical density at 600 nm after 24 h |
|--------------|-------------------------------|-------------------------------------|
|              | WT                            | ΔS5nA                              | ΔS5nA::S5nA | WT        | ΔS5nA     | ΔS5nA::S5nA |
| K006 (M6)    | 0.95 ± 0.10                   | 0.96 ± 0.07                        | 1.01 ± 0.08 | 1.7 ± 0.17 | 1.7 ± 0.17 | 1.7 ± 0.25  |
| MGAS8232 (M18)| 0.69 ± 0.09                   | 0.66 ± 0.06                        | 0.91 ± 0.09* | 1.7 ± 0.19 | 1.6 ± 0.17 | 2.0 ± 0.18* |
| 591 (M49)    | 0.80 ± 0.13                   | 0.84 ± 0.08                        | 0.84 ± 0.06 | 2.1 ± 0.35 | 2.1 ± 0.10 | 2.2 ± 0.16  |

*Data represent mean values ± standard deviations of five independent experiments;
*marginally significant difference compared to WT (p<0.05, Student’s t-test, n = 5)

https://doi.org/10.1371/journal.pone.0211074.t002

Fig 1. Surface 5'NT activity. Bacteria of wild type (wt), S5nA deletion (ΔS5nA) and complementation (comp) strains were harvested in the exponential growth phase and incubated with AMP as substrate and phosphate release was measured. Phosphate concentrations are shown as means with standard deviation (n = 3 biological replicates, *p<0.05, **p<0.01, unpaired two-tailed t-test).

Fig 1. Surface 5'NT activity. Bacteria of wild type (wt), S5nA deletion (ΔS5nA) and complementation (comp) strains were harvested in the exponential growth phase and incubated with AMP as substrate and phosphate release was measured. Phosphate concentrations are shown as means with standard deviation (n = 3 biological replicates, *p<0.05, **p<0.01, unpaired two-tailed t-test).

https://doi.org/10.1371/journal.pone.0211074.g001
infection. Again only serotype-dependent differences were observed. The M18 strain was most virulent in the larvae with about 50% of the animals dead 24 h after infection, while 50% of the larvae infected with the M6 or M49 serotype strains survived for two to three days (Fig 5). However, there were no significant differences in the survival rates of the larvae after infection with S5nA deletion strains in comparison with wild type and complementation strains of the same serotype (Fig 5). Altogether, these data indicated that S5nA is rather dispensable for virulence of \textit{S. pyogenes}.

\textbf{Discussion}

5’NT have been associated with virulence of certain bacteria, as it has been shown that they hydrolyze phosphate residues of adenosine phosphates and hence can interfere with the immunologically relevant ATP/adenosine ratio in human infections. This may happen via production of the anti-inflammatory adenosine and/or degradation of the pro-inflammatory ATP, both acting on phagocytic cells via purinergic receptors [9, 11, 12, 29–33].
**S. pyogenes** encodes the surface-associated 5’NT S5nA that has its highest expression levels in the exponential growth phase and that is immunogenic, as antibodies against *S. pyogenes* S5nA can be found in the serum of patients after *S. pyogenes* infections [14–16]. Although it has been shown that S5nA of the *S. pyogenes* M1 strain SF370 allows *L. lactis* to survive in human blood when added as a recombinant enzyme [3], the implication of the enzyme in virulence of *S. pyogenes* has not yet been investigated. Since the *S. pyogenes* S5nA dephosphorylates ADP and AMP but not ATP, the effect on *L. lactis* survival is mediated by increasing the adenosine concentration rather than lowering ATP levels [3]. This mode of action has also been described for the 5’NT of other streptococci such as *S. agalactiae* and *S. suis* [10, 25]. The 5’NT of *Streptococcus sanguinis*, in contrast, directly degrades ATP into adenosine [34]. In *S. agalactiae* and *S. suis* the deletion of the 5’NT genes rendered the bacteria more prone to

---

**Fig 3. Phagocytosis by neutrophils.** Phagocytic clearance was measured by determining viable counts of wild type (wt), S5nA deletion (ΔS5nA) and complementation (comp) strains after 30 min of incubation of opsonized bacteria in RPMI medium in the absence (control) or in the presence of neutrophils (PMN) at an MOI of 1. Viable counts are shown as means with standard deviation (n = 4 biological replicates).

https://doi.org/10.1371/journal.pone.0211074.g003
phagocytosis, consequently leading to a decreased survival in blood and decreased virulence in mouse (S. agalactiae) and piglet (S. suis) infection models [10, 25]. In Streptococcus equi subsp. zooepidemicus two nucleases with 5’NT activity have been shown to contribute to survival of the bacteria in neutrophil extracellular traps (NET) by degrading the NET DNA. The respective deletion mutants were also less virulent in a mouse infection model [26]. However, none of these effects were observed in the S. pyogenes S5nA gene deletion strains investigated here. The S5nA deletion strains had no survival deficiencies in human blood nor were they more efficiently killed by neutrophils. This is also reflected in the G. mellonella infection model, where the larvae did not survive in a higher proportion when infected with the deletion strains...
in comparison to those infected with wild type strains. There was however a residual phosphate release from AMP in the S5nA deletion mutant of *S. pyogenes* M49 (Fig 1). According to sequence data in the NCBI database, none of the proteins with an LPXTG cell wall anchoring motif encoded in the *S. pyogenes* M49 genome harbors a 5’-nucleotidase domain structure. The residual AMP dephosphorylating activity might therefore result from cytosolic nucleotide phosphatases leaking from defective cells or, less likely, a so far unrecognized cell wall anchored protein with minor 5’-nucleotidase activity.

It is not clear however if S5nA does not contribute to evasion of phagocytosis in *S. pyogenes* wild type bacteria at all or if the bacteria are able to compensate for the loss of S5nA. *S. pyogenes* is equipped with numerous other virulence factors mediating enhanced resistance to phagocytosis, including complement inhibitors, leucocidal toxins, immunoglobulin binding and degrading enzymes, NET degrading nucleases, and others (for an overview see review by Walker et al., 2014 [35]) that might be able to compensate for the loss of S5nA. The residual AMP-dephosphorylation activity measured in the S5nA deletion strains—if attributable to a so far unrecognized minor 5’-nucleotidase—might also contribute to a possible compensation. Furthermore, although phagocytosis resistance and survival/growth in human blood seem not
largely dependent on S5nA in *S. pyogenes* it cannot be excluded that S5nA mediated increase in adenosine levels in *S. pyogenes* infections has an impact on other immunological mechanism, as adenosine also acts as a signaling molecule regulating cytokine release of regulatory T cells and other immune cells [36, 37]. A mammalian infection model might uncover such effects. The *G. mellonella* model—although well established for investigations on virulence of *S. pyogenes* and other bacterial pathogens [24, 38–41]—naturally comes with certain limitations as the insect immune systems only resembles the innate immune mechanisms of humans. Purinergic receptors of the adenosine-binding P1-type receptors have been described to be present on insect cells [42]. To our knowledge however the role of adenosine in regulation of phagocytosis by hemocytes of *G. mellonella* has not been investigated yet.

Apart from the impact of 5’NT on the phagocytosis resistance of several bacteria, there is also indirect evidence for an association of 5’NT with biofilm formation in some bacteria. In *X. fastidiosa* it has been shown that a cytosolic 5’NT is highly expressed during the initial phase of biofilm formation. The authors speculate an impact of the 5’NT on the phosphorylation status of quorum-sensing molecules that might affect biofilm formation [13]. Biofilm formation capability is remarkably heterogeneous among *S. pyogenes* strains [43]. An association between biofilm formation capacity and certain pilus types has been described [17, 44]. Pilus proteins are encoded in the “fibronectin-binding, collagen-binding, T-antigen (FCT)” region in the *S. pyogenes* genome. To date nine different FCT types have been described that differ in composition and number of genes included [45, 46]. The M18 and M49 serotype strains investigated here carry an FCT type 3 region. Biofilm formation in FCT-3 strains is described to be inhomogeneous and dependent on environmental conditions. The M6 serotype strain K007, in contrast, contains an FCT type 1 region that is described to be associated with strong biofilm formation that is less depending on environmental conditions [17, 44]. In this respect our data are in accordance with the literature, as the FCT type 1 M6 strain formed strong biofilms on uncoated and fibronectin-coated plastic surfaces. As it has been described, for the FCT type 3 strains investigated here the biofilm formation capacity was variable. While the M49 strain was able to form stable biofilms on fibronectin and uncoated plastic, the M18 was barely able to form biofilms on any of the surfaces. In any case, in neither of the strains did the deletion of the S5nA gene have a significant impact on biofilm formation.

**Conclusions**

Our study shows that deletion of the 5’NT encoding gene S5nA does not affect *S. pyogenes* with respect to phagocytosis by neutrophils, growth of the bacteria in human blood, survival rates of the *G. mellonella* larvae in the infection model or biofilm formation capacity in three different strains covering the serotypes M6, M18 and M49. Hence, although an immunomodulatory capacity of the *S. pyogenes* S5nA was demonstrated by Zheng and colleagues [3], for the evasion of phagocytosis and biofilm formation this enzyme seems to be dispensable for *S. pyogenes*.

**Supporting information**

S1 Fig. Growth curves of *S. pyogenes* M6, M18, and M49 (lower row) wild type (wt), S5nA gene deletion (ΔS5nA), and complementation (comp) strains in THY medium. The data are shown as means of n = 5 biological replicates.

(TIF)

S2 Fig. Biofilm masses of *S. pyogenes* M6 (upper row), M18 (middle row), and M49 (lower row) wild type (wt), S5nA gene deletion (ΔS5nA), and complementation (comp) strains
after 24 h (left column) and 48 h (right column) as determined via photometry after crystal violet staining. Shown are average values and standard deviations of n ≥ 3 independent experiments.

(TIF)

S1 Table. qPCR results (Cycle threshold [Ct] means, n = 3) of S5nA mRNA quantification with 5S rRNA as control.

(DOCX)

Acknowledgments

The authors would like to express their gratitude to Catur Riani, Debbie Retnoningrum, and Yvonne Humboldt for excellent technical support.

Author Contributions

Conceptualization: Bernd Kreikemeyer, Tomas Fiedler.

Investigation: Marcel-Lino Dangel, Johann-Christoph Dettmann, Steffi Haßelbarth, Martin Krogull, Miriam Schakat.

Supervision: Tomas Fiedler.

Writing – original draft: Tomas Fiedler.

Writing – review & editing: Marcel-Lino Dangel, Bernd Kreikemeyer.

References

1. Cunningham MW. Pathogenesis of group A streptococcal infections and their sequelae. Adv Exp Med Biol. 2008; 609:29–42. https://doi.org/10.1007/978-0-387-73960-1_3 PMID: 18193655

2. Fiedler T, Sugareva V, Patenge N, Kreikemeyer B. Insights into Streptococcus pyogenes pathogenesis from transcriptome studies. Future Microbiol. 2010; 5:1675–94. https://doi.org/10.2217/fmb.10.128 PMID: 21133689

3. Zheng L, Khemlani A, Lorenz N, Loh JM, Langley RJ, Proft T. Streptococcal 5'-Nucleotidase A (S5nA), a Novel Streptococcus pyogenes Virulence Factor That Facilitates Immune Evasion. J Biol Chem. 2015; 290(52):31126–37. https://doi.org/10.1074/jbc.M115.677443 PMID: 26527680

4. Zimmermann H. 5'-Nucleotidase: molecular structure and functional aspects. Biochem J. 1992; 285 (2):345–65.

5. Reis JL. Studies on 5-nucleotidase and its distribution in human tissues. Biochem J. 1950; 2:21–2.

6. Kohn J, Reis JL. Bacterial Nucleotidases. J Bacteriol. 1963; 86:713–6. PMID: 14066466

7. Punj V, Zaborina O, Dhiman N, Falzari K, Bagdasarian M, Chakrabarty AM. Phagocytic cell killing mediated by secreted cytotoxic factors of Vibrio cholerae. Infect Immun. 2000; 68(9):4930–7. PMID: 10948107

8. Zaborina O, Dhiman N, Ling CM, Kostal J, Holder IA, Chakrabarty AM. Secreted products of a nonmucoid Pseudomonas aeruginosa strain induce two modes of macrophage killing: external-ATP-dependent, P2Z-receptor-mediated necrosis and ATP-independent, caspase-mediated apoptosis. Microbiology. 2000; 146(10):2521–30.

9. Thammanovska V, Kern JW, Missiaës DM, Schneewind O. Staphylococcus aureus synthesizes adenosine to escape host immune responses. J Exp Med. 2009; 206(11):2417–27. https://doi.org/10.1084/jem.20090097 PMID: 19808256

10. Firon A, Dinis M, Raynal B, Poyart C, Trieu-Cuot P, Kaminski PA. Extracellular nucleotide catabolism by the Group B Streptococcus ectonucleotidase NudP increases bacterial survival in blood. J Biol Chem. 2014; 289(9):5479–89. https://doi.org/10.1074/jbc.M113.456332 PMID: 24429288

11. Idzko M, Ferrari D, Eltzschig HK. Nucleotide signalling during inflammation. Nature. 2014; 509 (7500):310–7. https://doi.org/10.1038/nature13085 PMID: 24828189
12. Thiel M, Caldwell CC, Sitkovsky MV. The critical role of adenosine A2A receptors in downregulation of inflammation and immunity in the pathogenesis of infectious diseases. Microbes Infect. 2003; 5(6):515–26. PMID: 12758281

13. Santos CA, Saraiva AM, Toledo MA, Beloti LL, Crucello A, Favaro MT, et al. Initial biochemical and functional characterization of a 5'-nucleotidase from Xylella fastidiosa related to the human cytosolic 5'-nucleotidase I. Microb Pathog. 2013; 59–60:1–6. https://doi.org/10.1016/j.micpath.2013.02.007 PMID: 23474016

14. Koller T, Nelson D, Nakata M, Kreutzer M, Fischetti VA, Glicker MO, et al. PlyC, a novel bacteriophage lysin for compartment-dependent proteomics of group A streptococci. Proteomics. 2008; 8(1):140–8. https://doi.org/10.1002/pmic.200700001 PMID: 18095374

15. Severin A, Nickbarg E, Wooters J, Quazi SA, Matsuka YV, Murphy E, et al. Proteomic analysis and identification of Streptococcus pyogenes surface-associated proteins. J Bacteriol. 2007; 189(5):1514–22. https://doi.org/10.1128/JB.01132-06 PMID: 17142387

16. Reid SD, Green NM, Sylva GL, Voyich JM, Stenseth ET, Deleo FR, et al. Postgenomic analysis of four novel antigens of group A Streptococcus: growth phase-dependent gene transcription and human serologic response. J Bacteriol. 2002; 184(22):6316–24. https://doi.org/10.1128/JB.184.22.6316-6324.2002 PMID: 12399501

17. Koller T, Manetti AG, Kreikemeyer B, Lembke C, Margarit I, Grandi G, et al. Typing of the pilus-protein-encoding FCT region and biofilm formation as novel parameters in epidemiological investigations of Streptococcus pyogenes isolates from various infection sites. J Med Microbiol. 2010; 59(Pt 4):442–52. https://doi.org/10.1099/jmm.0.013581-0 PMID: 20007764

18. Smoot JC, Barbian KD, Van Gompel JJ, Smoot LM, Chaussse MS, Sylva GL, et al. Genome sequence and comparative microarray analysis of serotype M18 group A Streptococcus strains associated with acute rheumatic fever outbreaks. Proc Natl Acad Sci U S A. 2002; 99(7):4668–73. https://doi.org/10.1073/pnas.062526099 PMID: 11917108

19. Tao L, LeBlanc DJ, Ferretti JJ. Novel streptococcal-integration shuttle vectors for gene cloning and inactivation. Gene. 1992; 120(1):105–10. PMID: 13279668

20. Baev D, England R, Kuramitsu HK. Stress-induced membrane association of the Streptococcus mutans GTP-binding protein, an essential G protein, and investigation of its physiological role by utilizing an antisense RNA strategy. Infect Immun. 1999; 67(9):4510–6. PMID: 10456893

21. Biswas I, German P, McDade K, Scott JR. Generation and surface localization of intact M protein in Streptococcus pyogenes. Infect Immun. 2001; 69(11):7029–38. https://doi.org/10.1128/IAI.69.11.7029-7038.2001 PMID: 11598078

22. Pappesch R, Warnke P, Mikkat S, Normann J, Wisniewska-Kucper A, Huschka F, et al. The Regulatory Small RNA MarS Supports Virulence of Streptococcus pyogenes. Sci Rep. 2017; 7(1):12241. https://doi.org/10.1038/s41598-017-01250-7 PMID: 28947755

23. Nakata M, Koller T, Moritz K, Ribardo D, Jonas L, McIver KS, et al. Mode of expression and functional characterization of FCT-3 pilus region-encoded proteins in Streptococcus pyogenes serotype M49. Infect Immun. 2009; 77(1):32–44. https://doi.org/10.1128/IAI.00772-08 PMID: 18852238

24. Oehmcke-Hecht S, Nass LE, Wichura JB, Mikkat S, Kreikemeyer B, Fiedler T. Deletion of the L-Lactate Dehydrogenase Gene ldh in Streptococcus pyogenes Leads to a Loss of SpeB Activity and a Hypovirulent Phenotype. Frontiers in Microbiology. 2017; 8:1841. https://doi.org/10.3389/fmicb.2017.01841 PMID: 28983299

25. Liu P, Pian Y, Li X, Liu R, Xie W, Zhang C, et al. Streptococcus suis adenosine synthase functions as an effector in evasion of PMN-mediated innate immune. J Infect Dis. 2014; 210(1):35–45. https://doi.org/10.1093/infdis/jiu050 PMID: 24446521

26. Ma F, Guo X, Fan H. Extracellular Nucleases of Streptococcus equi subsp. zooepidemicus Degrade Neutrophil Extracellular Traps and Impair Macrophage Activity of the Host. Appl Environ Microbiol. 2017; 83(2).

27. Lembke C, Podbielski A, Hidalgo-Grass C, Jonas L, Hanski E, Kreikemeyer B. Characterization of biofilm formation by clinically relevant serotypes of group A streptococci. Appl Environ Microbiol. 2006; 72(4):2864–75. https://doi.org/10.1128/AEM.72.4.2864-2875.2006 PMID: 16597993

28. Sugareva V, Art R, Fiedler T, Riani C, Podbielski A, Kreikemeyer B. Serotype- and strain-dependent contribution of the sensor kinase CovS of the CovRS two-component system to Streptococcus pyogenes pathogenesis. BMC Microbiol. 2010; 10:34. https://doi.org/10.1186/1471-2180-10-34 PMID: 20113532

29. Junger WG. Immune cell regulation by autocrine purinergic signalling. Nat Rev Immunol. 2011; 11(3):201–12. https://doi.org/10.1038/ni2938 PMID: 21331080

30. Piccini A, Carta S, Tassi S, Lasiglie D, Fossati G, Rubartelli A. ATP is released by monocytes stimulated with pathogen-sensing receptor ligands and induces IL-1beta and IL-18 secretion in an autocrine way.
31. Thammavongsa V, Schneewind O, Missiakas DM. Enzymatic properties of Staphylococcus aureus adenine synthase (AdsA). BMC Biochem. 2011; 12:56. https://doi.org/10.1186/1471-2091-12-56 PMID: 22035583

32. Le Moine O, Stordeur P, Schandene L, Marchant A, de Groote D, Goldman M, et al. Adenosine enhances IL-16 secretion by human monocytes. J Immunol. 1996; 156(11):4408–14. PMID: 8666814

33. Hasko G, Pacher P. A2A receptors in inflammation and injury: lessons learned from transgenic animals. J Leukoc Biol. 2008; 83(3):447–55. https://doi.org/10.1189/jl.0003059 PMID: 18160539

34. Fan J, Zhang Y, Chuang-Smith ON, Frank KL, Guenther BD, Kern M, et al. Ecto-5′-nucleotidase: a candidate virulence factor in Streptococcus sanguinis experimental endocarditis. PLoS One. 2012; 7(6): e38059. https://doi.org/10.1371/journal.pone.0038059 PMID: 22685551

35. Walker MJ, Barnett TC, McArthur JD, Cole JN, Gillen CM, Henningham A, et al. Disease manifestations and pathogenic mechanisms of group A Streptococcus. Clin Microbiol Rev. 2014; 27(2):264–301. https://doi.org/10.1128/CMR.00101-13 PMID: 24696436

36. Hasko G, Cronstein BN. Adenosine: an endogenous regulator of innate immunity. Trends Immunol. 2004; 25(1):33–9. PMID: 14698282

37. Ernst PB, Garrison JC, Thompson LF. Much ado about adenosine: adenosine synthesis and function in regulatory T cell biology. J Immunol. 2010; 185(4):1993–8. https://doi.org/10.4049/jimmunol.1000108 PMID: 20686167

38. Olsen RJ, Watkins ME, Cantu CC, Beres SB, Musser JM. Virulence of serotype M3 Group A Streptococcus strains in wax worms (Galleria mellonella larvae). Virulence. 2011; 2(2):111–9. https://doi.org/10.4161/viru.2.2.14338 PMID: 21258213

39. Cook SM, McArthur JD. Developing Galleria mellonella as a model host for human pathogens. Virulence. 2013; 4(5):350–3. https://doi.org/10.4161/viru.25240 PMID: 23799664

40. Loh JM, Adenwalla N, Wiles S, Proft T. Galleria mellonella larvae as an infection model for group A streptococcus. Virulence. 2013; 4(5):419–28. https://doi.org/10.4161/viru.24930 PMID: 23652836

41. Chalmers C, Khemlani AHJ, Sohn CR, Loh JMS, Tsai CJ, Proft T. Streptococcus pyogenes nuclease A (SpnA) mediated virulence does not exclusively depend on nuclease activity. J Microbiol Immunol Infect. 2017.

42. Burnstock G, Verkhratsky A. Evolutionary origins of the purinergic signalling system. Acta Physiol (Oxf). 2009; 195(4):415–47.

43. Fiedler T, Koller T, Kreikemeyer B. Streptococcus pyogenes biofilms-formation, biology, and clinical relevance. Front Cell Infect Microbiol. 2015; 5:15. https://doi.org/10.3389/fcimb.2015.00015 PMID: 25717441

44. Manetti AG, Koller T, Becherelli M, Buccato S, Kreikemeyer B, Podbielski A, et al. Environmental acidification drives S. pyogenes plus expression and microcolony formation on epithelial cells in a FCT-dependent manner. PLoS One. 2010; 5(11):e13864. https://doi.org/10.1371/journal.pone.0013864 PMID: 21079780

45. Kratovac Z, Manoharan A, Luo F, Lizano S, Bessen DE. Population genetics and linkage analysis of loci within the FCT region of Streptococcus pyogenes. J Bacteriol. 2007; 189(4):1299–310. https://doi.org/10.1128/JB.01301-06 PMID: 17028269

46. Falugi F, Zingaretti C, Pinto V, Mariani M, Arnodeo L, Manetti AG, et al. Sequence variation in group A Streptococcus pilis and association of pilus backbone types with Lancefield T serotypes. J Infect Dis. 2008; 198(12):1834–41. https://doi.org/10.1086/593176 PMID: 18928379