T granules in human platelets function in TLR9 organization and signaling

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T granules in human platelets function in TLR9 organization and signaling

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Introduction

Human adults contain nearly one trillion blood PLTs in circulation. These small (2–3 µm), anuclear cells have a highly organized cytoskeleton, unique receptors, and specialized secretory granules. Among their primary functions, PLTs serve as the “Band-Aids” of the bloodstream and respond to blood vessel injury by changing shape, secreting their granule contents, and aggregating to form a PLT clot. PLTs also play secondary roles in lymphatic development and inflammation (Smyth et al., 2011). Among their most recognized and least understood immunomodulatory functions, PLTs express multiple members of a family of pattern recognition receptors called toll-like receptors (TLRs). TLRs have been best characterized in neutrophils, macrophages, and dendritic cells and promote immune activation in response to conserved molecular motifs expressed by pathogens (Janeway and Medzhitov, 2002). Although human and mouse PLTs express TLRs 1–9, our current understanding of TLR function in PLTs is limited to TLRs 1–6, which are typically expressed on the PLT surface and thought to trap bacteria for elimination by professional phagocytes (Andonegui et al., 2005; Aslam et al., 2006; Clark et al., 2007). Less understood is the role of TLR9, which is expressed in the endosomes of monocytes, macrophages, and plasmacytoid dendritic cells and is thought to act as a receptor for unmethylated CpG islands found in bacterial and viral DNA (Henissy et al., 2010).

Although reported to localize internally in PLTs, neither the site nor function of subcellular TLR9 is currently known (Aslam et al., 2006). In macrophages and dendritic cells,
Figure 1. MKs up-regulate Tlr9 transcript expression during pro-PLT production and express TLR9 in distinct granules that partially colocalize with VAMP 8 at every stage in thrombocytopoiesis. (A and B) Shown is a screenshot from Integrated Genome Browser of reads mapping to the Tlr9 (A) and Vamp8 (B) locus. The black bars represent piled up sequencing reads aligning to the genomic coordinates encoding the respective RNAs. More reads will align to more highly expressed RNA regions, and the height of the black bars correlates with RNA expression level. Below the read distributions are the RefSeq annotations: thick horizontal lines represent exons, and thin horizontal lines represent introns. Quantification was based on three round MK and four pro-PLT–MK replicates and is expressed in reads per kilobase of exon model per million mapped reads. (C) Intermediates of PLT production from murine fetal liver cell cultures were spun down onto poly-l-lysine-coated glass cover slips, permeabilized with 0.5% Triton X-100 for 5 min, and probed for TLR9 and either β1-tubulin or VAMP 8. Slides were examined by fluorescence microscopy, and image fluorescence intensity is normalized to the round MK fraction. Round MKs, pro-PLT–MKs, released pro-PLTs, and individual PLTs revealed distinct, punctate/granular localization of TLR9 similar to that observed in whole-blood PLTs. β1-Tubulin antibodies were used to delineate the cell periphery and denote the different intermediate stages in PLT production. Insets (from top to bottom) represent a PLT, pre-PLT, and barbell pro-PLT. (D) Background fluorescence was subtracted, and image brightness/contrast was adjusted linearly for each micrograph to resolve individual granules. TLR9 showed significant colocalization with VAMP 8 in round MKs, along the pro-PLT shafts of pro-PLT–MKs, and within released pro-PLTs and individual PLTs. A Manders’ coefficient of 0.64 was calculated for VAMP 8 colocalization with TLR9 throughout the entirety of MK cell culture. Insets represent the magnified region outlined by the yellow box for each image.
Platelet TLR9 localization and function

TLR9 is recruited to lysosomes only after cells are stimulated with CpG DNA (Latz et al., 2004). The fact that all intracellular TLRs identified share specificity for nucleic acids suggests that this localization may be related to the recognition of this class of ligand (Barton et al., 2006). Synthetic oligodeoxynucleotides (ODNs) that contain unmethylated CpG motifs are typically used to activate TLR9. Type C CpG motifs contain a complete phosphorothioate backbone and a CpG-containing palindromic motif that is present at a 20-fold greater frequency in bacterial DNA compared with mammalian DNA (Bauer et al., 2001). Although type C CpG ODNs induce strong IFN-α production from both plasmacytoid dendritic cells and B cells, the effect of TLR9 signaling in PLTs remains to be established.

In this paper, we use confocal and electron microscopy to track TLR9 during pro-PLT production to a new electron-dense tubular system-related compartment in PLTs. Second, we use known PLT agonists to describe a novel mechanism of TLR9 signaling whereby PLTs are primed during activation to express TLR9 on their surface before secondary activation through pathogen-associated molecular patterns (PAMPs) on invading organisms. Finally, we reveal a unique role for human PLTs as mediators of innate immunity at sites of vascular damage.

Results

TLR9 is expressed in murine cell culture megakaryocytes (MKs) and is distributed to PLTs during pro-PLT production

Because TLR9 is a transmembrane receptor, we hypothesized that TLR9 should be translated by bone marrow MKs and transported along pro-PLT extensions to nascent PLTs. To test this hypothesis, mouse fetal liver cell cultures were generated, and round MKs, pro-PLT–producing MKs, released pro-PLTs, and individual PLTs were isolated on day 5 of culture. Next generation RNA sequencing of round MKs and pro-PLT–MKs confirmed the presence of the Tlr9 and Vamp8 transcripts in murine cell cultures (Fig. 1, A and B). Comparison of Tlr9 expression profiles between the two revealed that MKs experience a burst of Tlr9 mRNA expression during pro-PLT production (0.34 ± 0.21 vs. 2.44 ± 0.75 RPKM, n ≥ 3). Pro-PLT–producing MKs did not experience a similar burst of Vamp8 mRNA expression during pro-PLT production, which was used as an internal control (70.9 ± 14.5 vs. 60.4 ± 3.2 RPKM, n ≥ 3). These results confirm that TLR9 is generated by MKs during pro-PLT production. Intermediates in PLT production were probed for TLR9 and β1-tubulin to delineate the cell periphery, and samples were visualized by immunofluorescence microscopy (Fig. 1 C). TLR9 was present in mouse MKs, pro-PLT–MKs, released pro-PLTs, and individual PLTs and showed punctate/granular expression that partly colocalized with VAMP 8 at each stage of PLT production (Fig. 1 D). SNAREs, such as VAMP 8, associate with PLT granules and may mediate TLR9 localization to nascent PLT tips during thrombocytopoiesis.

Human PLTs sequester TLR9 and protein disulfide isomerase (PDI) in previously undescribed electron-dense tubular system-related T granules

To confirm that the majority of TLR9 in resting PLTs is expressed intracellularly, washed PLTs were spun down onto poly-1-lysine–coated glass cover slides, permeabilized with 0.5% Triton X-100 for 5 min, and were probed for either TLR9 alone or TLR9 and β1-tubulin together. (A) Samples were examined by wide-field fluorescence microscopy and revealed peripheral labeling of TLR9 in distinct, punctate granules that localized mostly (69 ± 5%) along the cell periphery, adjacent to the plasma membrane. (B) TLR9 localization to the cell edge was confirmed in human PLTs by colabeling with β1-tubulin to demarcate the PLT border. Inset represents magnified region outlined by the yellow box. (inset) Bar, 5 µm.

Figure 2. Human PLTs express TLR9 in distinct granules along the periphery of the cell, adjacent to the plasma membrane. (A and B) Washed human whole-blood PLTs were spun down onto poly-1-lysine–coated glass cover slides, permeabilized with 0.5% Triton X-100 for 5 min, and were probed for either TLR9 alone or TLR9 and β1-tubulin together. (A) Samples were examined by wide-field fluorescence microscopy and revealed peripheral labeling of TLR9 in distinct, punctate granules that localized mostly (69 ± 5%) along the cell periphery, adjacent to the plasma membrane. (B) TLR9 localization to the cell edge was confirmed in human PLTs by colabeling with β1-tubulin to demarcate the PLT border. Inset represents magnified region outlined by the yellow box. (inset) Bar, 5 µm.
Representative maximum projection z series for TLR9 and PDI (Fig. 4A) confirmed that TLR9 is distributing with PDI to a separate granular compartment. To surpass the resolution limit of conventional fluorescence microscopy (200 nm) and because Gray PLT syndrome PLTs express some α-granule membrane proteins in normal amounts (Rosa et al., 1987; Maynard et al., 2010), TLR9 and PDI localization was examined by immunogold electron microscopy. PLT cryosections were probed for TLR9 (Fig. 4B) and PDI (Fig. 4C) separately and together (Fig. 4, D and E). TLR9 distributed to previously uncharacterized electron-dense granular compartments to which PDI was also shown to colocalize (Fig. 4, B–D). These were named T granules after TLR9 electron-dense tubular system-related granular compartments.

TLR9/PDI labeling was associated with the limiting membrane, and T-granule identification was based on the colocalization of three or more immunogold molecules within a single electron-dense area (Fig. 4 B–E, denoted by white arrows). T granules had with known specific markers of α granules (fibrinogen, CD62P, PDGF-B, VEGF, CD42a, and CD42b), dense granule (serotonin), lysosome (LAMP-1), or endosome (M6P and Syntaxin-13) protein markers (Fig. 3A). To validate this observation, TLR9 labeling of whole-blood PLTs from human normal controls was compared with Gray PLT and Hermansky–Pudlack syndrome patients, which are devoid of α-granule and dense-granule content, respectively (Fig. 3B). Punctate/granular expression of TLR9 in these samples was uniminished and suggests that TLR9 is not contained in PLT α or dense granules.

In professional dendritic cells, TLR9 is distributed to intracellular endosomal compartments along with TLR7 and TLR8. PDI, by comparison, is highly expressed in the endoplasmic reticulum of endothelial cells and electron-dense tubular system-related compartments in PLTs (van Nispen Tot Pannerden et al., 2009). Interestingly, although PDI colocalized with TLR9 in resting human PLTs, TLR7 and TLR8 did not (Fig. 3 C and Videos 1 and 2). Representative maximum projection z series for TLR9 and PDI (Fig. 4A) confirmed that TLR9 is distributing with PDI to a separate granular compartment.

To surpass the resolution limit of conventional fluorescence microscopy (200 nm) and because Gray PLT syndrome PLTs express some α-granule membrane proteins in normal amounts (Rosa et al., 1987; Maynard et al., 2010), TLR9 and PDI localization was examined by immunogold electron microscopy. PLT cryosections were probed for TLR9 (Fig. 4B) and PDI (Fig. 4C) separately and together (Fig. 4D and E). TLR9 distributed to previously uncharacterized electron-dense granular compartments to which PDI was also shown to colocalize (Fig. 4, B–D). These were named T granules after TLR9 electron-dense tubular system-related granular compartments. TLR9/PDI labeling was associated with the limiting membrane, and T-granule identification was based on the colocalization of three or more immunogold molecules within a single electron-dense area (Fig. 4B–E, denoted by white arrows). T granules had
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免疫荧光显微镜，并且通过算法计算关联风险比和曼德尔斯系数。代表性图像见图5。表1汇总了共定位的结果。

TLR9 没有与 VAMPs 2 和 3（未显示）共定位，且与 VAMP 5 共定位较差，该内阈值控制（图5 A）。曼德尔斯系数为 0.5 或更高被认为是显著的，且 TLR9 在静息血小板中与 VAMPs 7 和 8 部分共定位（图5, B 和 C）。血小板激活（散开）后，VAMP 8 被重新分配到血小板的颗粒，而 VAMPs 5 和 7 分布到血小板的伪足。TLR9 继续与 VAMP 8 共定位，证明血小板中有两个主要的 T 红细胞。

T 红细胞粒化/细胞膜融合伴随 VAMP 7 和 8 再组织

因为 SNAREs 在血小板中介导 α-粒化，我们假设 T 红细胞粒化在血小板激活和细胞膜融合中可能共定位。SNAREs 分为 t-SNAREs 和 v-SNAREs，其中包括 VAMPs 2, 3, 5, 7, 和 8 (Graham et al., 2009)。v-SNARE 与 TLR9 在静息和凝血酶活化的人血小板中共定位已被通过共聚焦免疫荧光显微镜证实，且两个风险比和曼德尔斯系数分别计算每个关联。代表性图像是图5，并且结果共定位总结在表1。TLR9 没有与 VAMPs 2 和 3（未显示）共定位，且与 VAMP 5，包含一个内阈值控制（图5 A）。曼德尔斯系数为 0.5 或更高被认为是显著的，且 TLR9 在静息和凝血酶活化的人血小板中与 VAMPs 7 和 8 部分共定位（图5, B 和 C）。血小板激活（散开）后，VAMP 8 被重新分配到血小板的颗粒，而 VAMPs 5 和 7 分布到血小板的伪足。TLR9 继续与 VAMP 8 共定位，证明血小板中有两个主要的 T 红细胞群体。

Figure 4. TLR9 colocalizes with PDI to electron-dense membrane-encapsulated regions adjacent the plasma membrane along the periphery of human PLTs. (A) Representative maximum projection z series for TLR9 and PDI by confocal immunofluorescence microscopy demonstrating colocalization. (B–E) Washed human whole-blood PLTs were fixed, frozen, and sectioned before mounting on Formvar carbon-coated copper grids. Ultrathin PLT sections were probed for TLR9, and bound antibody was labeled with immunogold. Samples were examined by electron microscopy and reveal the distribution of TLR9 (B), PDI (C), and colocalization of both (D and E) along the periphery of resting human PLTs. White arrows denote localization of TLR9 and PDI to limiting membrane of electron-dense regions adjacent the plasma membrane. (D and E) For colabeling, immunogold particles are 15 nm for PDI and 10 nm for TLR9.

a mean cross-sectional area of 4,148 nm², and there were a mean of eight T granules per PLT cross-sectional area (1,750,019 nm²) in 25 PLTs analyzed.

T-granule localization/plasma membrane fusion accompanies VAMP 7 and 8 reorganization on PLT activation

Because SNAREs mediate α-granule localization in PLTs, we hypothesized that these might also colocalize with T-granule reorganization during PLT activation and plasma membrane fusion. SNAREs are divided into t-SNAREs and v-SNAREs, which in PLTs include VAMPs 2, 3, 5, 7, and 8 (Graham et al., 2009). v-SNARE colocalization with TLR9 in resting and thrombin-activated human PLTs was determined by confocal immunofluorescence microscopy, and both risk ratio and Manders’ coefficients were calculated for each association. Representative figures are shown in Fig. 5, and results for colocalizations are summarized in Table 1. TLR9 did not associate with VAMPs 2 and 3 (not depicted) and shows poor association with VAMP 5, which was included as an internal threshold control (Fig. 5 A). Manders’ coefficients of 0.5 or greater were considered significant, and TLR9 showed partial colocalization with both VAMPs 7 and 8 in resting PLTs (Fig. 5, B and C). After PLT activation (spreading) on glass, VAMP 8 was redistributed to the PLT granulomere, whereas VAMPs 5 and 7 distributed to the PLT lamellipodium. TLR9 continued to show colocalization with VAMP 8 after spreading, which demonstrates that there are two major subpopulations of T granules in PLTs that can
Figure 5. Human PLT TLR9 localization with VAMPs 5, 7, and 8 under resting conditions and when activated (spread) on glass surface. Samples were examined by confocal fluorescence microscopy. Image analysis was completed by using the JACoP (Just Another Colocalization Plugin) plugin for ImageJ as described in Table 1. Manders' coefficients were used to compare the TLR9 relationship to the associated VAMP.

(A) Micrographs demonstrating the relationship of VAMP 5 and TLR9. As demonstrated in the micrographs, in the resting PLT, ~41% of TLR9 signal overlaps with that of VAMP 5. The portion of overlapping signal slightly increased in the adhered PLTs to 49%. (B) Additional micrographs demonstrating the relationship between VAMP 7 and TLR9. Image analysis confirms 82% of TLR9 overlapped with that of VAMP 7 in the resting PLT; however, upon spreading, the signal decreased to only 56%. (C) Micrographs of resting and spread PLTs probed with specific antiserum for VAMP 8 and TLR9. In the resting PLT, 77% of TLR9 signal overlapped with VAMP 8. Upon spreading, the signal decreased to 67% signal overlap.
be differentially distributed on PLT activation. Localization of VAMPs to the PLT granulomere or lamellipodium on contact activation has been associated with granule membrane fusion with the plasma membrane or the open canalicular system, respectively (Flaumenhaft, 2003), and accounts for TLR9 surface expression.

**TLR9 is expressed on the PLT surface during activation**

Unlike TLRs 1–6, TLR9 is exclusively expressed intracellularly in the professional dendritic cells (Barton et al., 2006). To determine whether PLTs express TLR9 on the plasma membrane under resting conditions and whether activation with a strong PLT agonist will cause TLR9 to relocate to the PLT surface, human PLTs were collected from whole blood and incubated with thrombin. Thrombin is a key regulator of the coagulation cascade and activates PLTs through the protease-activated receptors 1 and 4 in humans. Flow cytometry and immunofluorescence microscopy were used to measure the surface expression of TLR9 in human PLTs under resting conditions and after PLT activation (Fig. S3). Surprisingly, resting PLTs expressed basal levels of surface TLR9 that became significantly increased after thrombin activation, suggesting that although the majority of TLR9 is expressed intracellularly, some of it relocates to the plasma membrane on agonist exposure. Although it is unknown what local concentrations of bacterial DNA are achieved in vivo, increased TLR9 surface expression in thrombin-activated PLTs corresponded with increased surface binding of the synthetic TLR9 ligand, unmethylated CpG ODN, when ≥5 µM, and was therefore used at this concentration for all subsequent experiments (Fig. S3 A and Fig. S5). TLR9 surface expression was also regulated by PLT activation through other common PLT agonists that stimulate PLT activation and granule release through separate signaling pathways. These included ADP and type IV collagen, which were chosen because they represent well-characterized, physiologically relevant agonists, PA, which is a potent secretagogue used in both PLTs and leukocytes to evaluate granule exocytosis, and the GPVI agonist collagen-related peptide (CRP), given the importance of the GPVI receptor in PLT function. These data demonstrate that human PLTs may need to be primed to respond to bacterial/viral PAMPs in whole blood (Fig. 6). Moreover, although the direct addition of type IV collagen to resting PLTs resulted in much higher levels of TLR9 surface expression than did thrombin, these results were not matched with a significant increase in CD62P surface expression and revealed lower levels of CD61 surface expression relative to CRP controls, confirming that TLR9 and CD62P/CD61 localize to separate organelles. Conversely, CRP-activated PLT activation resulted in significantly higher levels of both CD62P and CD61 surface expression than did type IV collagen but were met with relatively lower levels of TLR9 surface expression, as confirmed by quantitative immunofluorescence microscopy (Fig. S4).

**TLR9 regulates DNA sequestration and CD62P surface expression in PLTs**

Unmethylated ODNs, including a CpG motif, can mimic the effects of bacterial DNA, inducing B cell proliferation and activating cells of the myeloid lineage. We hypothesized that the addition of type C CpG to washed human PLTs may induce PLT activation and possibly result in increased surface expression of TLR9 and CD62P and ODN sequestration. Although incubation of resting human PLTs with synthetic unmethylated type C CpG did not cause PLT shape change (Fig. S2), it did result in a 37 ± 13% increase in TLR9 and 34 ± 15% increase in CD62P surface expression followed by a 31 ± 15% increase in FITC-conjugated type C CpG sequestration over 20 min (Fig. 7, A and C). Surprisingly, when TLR9 surface expression was specifically up-regulated by preincubating PLTs with type IV collagen, subsequent incubation with type C CpG resulted in 100% of PLTs demonstrating increased type C CpG sequestration, CD62P surface expression, and PLT clumping within 30 s of addition (Fig. 7, B and C; and Videos 3 and 4). This was inhibited by preincubating the PLT culture with 20 μM IRAK-1/4 (interleukin-1 receptor-associated kinase-1/4) inhibitor for 1 h before collagen IV incubation and type C CpG addition (Videos 3 and 4). IRAK-1 and IRAK-4 are protein kinases that mediate signaling by TIR (Toll/IL1/Plant R) domain–containing receptors, including the IL-1, IL-8, and TLRs (Song et al., 2009). Pharmacological inhibition of both kinases blocked PLT clumping by type C CpG in the presence of type IV collagen and reveals downstream signaling effectors of ODN-mediated PLT activation.

Unlike in human PLTs that contain ~0.5 RPKM TLR9 RNA, TLR9 surface expression was not changed in mice after type C CpG incubation, and deep sequencing of mouse PLTs has shown that they do not contain detectable levels of TLR9 RNA (Rowley et al., 2011). Interestingly, although preincubation with type IV collagen still resulted in immediate PLT clumping when type C CpG was added (unpublished data), TLR9 knockout (KO) mice show reduced levels of ODN

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**Table 1.** Calculated risk ratio and Manders’ coefficients for each association of TLR9 with VAMPs 5, 7, and 8

<table>
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<th>Colocalization with TLR9</th>
<th>VAMP 5</th>
<th>VAMP 7</th>
<th>VAMP 8</th>
<th>Resting</th>
<th>M1 TLR9/VAMP</th>
<th>M2 VAMP/TLR9</th>
<th>Spread (glass)</th>
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<td>Rr</td>
<td>M1</td>
<td>M2</td>
<td>Rr</td>
<td>M1</td>
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<td>Rr</td>
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<tr>
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<td>0.7 ± 0.1</td>
<td>0.4 ± 0.2</td>
<td>0.4 ± 0.2</td>
<td>0.5 ± 0.1</td>
<td>0.5 ± 0.2</td>
<td>0.14 ± 0.04</td>
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<tr>
<td>VAMP 7</td>
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<td>0.82 ± 0.08</td>
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Calculated risk ratio (Rr) and Manders’ coefficients for each association of TLR9 with VAMPs 5, 7, and 8 in human whole-blood PLTs under resting conditions and when activated (spread) on glass surface. Manders’ coefficients of 0.5 or greater were considered significant and are highlighted in bold.
regard to TLR9 binding (specificity control), resting/thrombin-activated human PLTs were incubated with 5 µM FITC-conjugated type C CpG and decreasing concentrations of unlabeled sequestration and CD62P surface expression relative to isotype-matched wild-type controls. To confirm that FITC-conjugated type C CpG behaves like nonconjugated type C CpG with regard to TLR9 binding (specificity control), resting/thrombin-activated human PLTs were incubated with 5 µM FITC-conjugated type C CpG and decreasing concentrations of unlabeled

Figure 6. Human PLTs regulate surface expression of TLR9 on activation with select agonists. Human PLTs were collected from whole blood and examined under resting conditions or after activation with 1 mU/µl thrombin, 1 µM PMA, 3 µg/ml CRP, 20 µM ADP, or 50 µg/ml mouse type IV collagen for 5 min at 37°C. Samples were probed for TLR9 and either CD62P or CD61. PLT mean fluorescence intensity (relative surface expression of targeted receptor) was determined by flow cytometry for at least three independent samples, and data were subject to one-way ANOVA and Tukey HSD analysis. Error bars represent one standard deviation about the mean for at least three independent samples. (A) Human PLT activation with all listed agonists results in increased surface expression of TLR9 relative to resting control. Strikingly, TLR9 surface expression on PLT activation did not correlate with CD62P/CD61 expression for the agonists tested and suggests that TLR9 and CD62P/CD61 may localize to separate granules. (B, left) Representative forward scatter versus side scatter dot plots highlight characteristic changes in PLT morphology on activation with the listed agonists relative to resting control. (right) Representative histograms demonstrate a shift in mean fluorescence intensity for TLR9 and CD62P/CD61 on PLT activation. Outlines represent PLT gate used for sample thresholding by forward and side scatter. Mean fluorescence intensity for resting PLTs is represented in gray. Mean fluorescence intensity for agonist-activated PLTs is represented in red.
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Tested. To determine whether circulating ODNs also result in increased thrombus formation, washed human whole-blood PLTs and PLT-rich plasma (PRP) were pretreated with 5 µM control ODN, type C CpG ODN, or a vehicle control and perfused at a shear rate of 200 s⁻¹ (flow rate of 18.7 µl/min) over a surface coated with type IV collagen for (cold) type C CpG and analyzed by flow cytometry (Fig. S5). Thrombin-activated PLTs showed fluorescence profiles that correlated exactly with the relative proportion of FITC-conjugated type C CpG in the culture. Resting human PLTs showed fluorescence profiles that were comparable with the 20-µM unlabeled type C CpG control at all the concentrations tested. To determine whether circulating ODNs also result in increased thrombus formation, washed human whole-blood PLTs and PLT-rich plasma (PRP) were pretreated with 5 µM control ODN, type C CpG ODN, or a vehicle control and perfused at a shear rate of 200 s⁻¹ (flow rate of 18.7 µl/min) over a surface coated with type IV collagen or type I collagen for

Figure 7. TLR9 signaling results in type C CpG sequestration, increased CD62P surface expression, and PLT clumping. (A and B) Flow cytometric analysis showing representative forward versus side scatter profiles of human washed PLTs under resting conditions and after incubation with synthetic unmethylated type C CpG ODNs (characteristic of bacterial/viral DNA) before (A) and after (B) type IV collagen preincubation. Outlines represent PLT gate used for sample thresholding by forward and side scatter. (C) Quantification of the percentage of PLTs expressing TLR9, CD62P, and type C CpG was normalized to resting levels to resolve the difference on agonist exposure over time. Incubation of resting washed human PLTs with type C CpG ODNs resulted in a 40% increase in TLR9 surface expression followed by a 30% increased type C CpG sequestration and CD62P surface expression over 20 min. (D) Type IV collagen preincubation resulted in considerably increased type C CpG sequestration, CD62P surface expression above resting PLT controls, and significant PLT clumping within 30 s of type C CpG addition. (E) Mouse PLTs show a more pronounced ODN sequestration and comparable CD62P expression after type C CpG incubation. Unlike in human PLTs, TLR9 surface expression is not changed in mice. Although TLR9 KO mice show reduced levels of ODN sequestration and CD62P surface expression 20 min after type C CpG addition, preincubation with type IV collagen still resulted in immediate PLT clumping when type C CpG was added (not depicted). Statistical significance for marked pairings was established using a one-tailed Student’s t test for paired samples (*, P < 0.05; **, P < 0.01). Error bars represent one standard deviation about the mean.
A Collagen IV, washed platelets

(+ 5 µM Control ODN)

0 min 7 min

(+ 5 µM Type C CpG)

0 min 7 min

No ODN

0 min 7 min

B Collagen IV, platelet-rich plasma

(+ 5 µM Control ODN)

0 min 7 min

(+ 5 µM Type C CpG)

0 min 7 min

C Collagen I, platelet-rich plasma

(+ 5 µM Control ODN)

0 min 7 min

(+ 5 µM Type C CpG)

0 min 7 min

D hPLTs - 1 Hour Incubation

E hPLTs - 4 Hour Incubation

Figure 8. ODN-activated PLTs do not form thrombi on collagen type IV and endocytose type C CpG to distinct granules that do not colocalize with TLR9. (A–C) Washed human whole-blood PLTs (A) and PRP (B) were pretreated with 5 µM control ODN, type C CpG ODN, or a vehicle control and perfused at a shear rate of 200 s⁻¹ (flow rate of 18.7 µl/min) over a surface coated with type IV collagen or type I collagen (C) for 10 min. Although addition of ODN to PLTs resulted in PLT clumping in the presence of type IV collagen (white arrows), these did not form thrombi as compared with type I collagen (positive control, black arrows). Addition of type C CpG did not result in increased thrombus formation relative to the ODN control in PRP on the type I collagen-coated surface. (D) PLTs were incubated with FITC-conjugated type C CpG at 37°C and 5% CO₂ for a period of ≤4 h. PLTs were subsequently spun down onto poly-l-lysine–coated glass cover slides and probed for TLR9 as described in the Immunofluorescence microscopy section of Materials and methods. Fig. 8 (D and E) shows that within the first hour of incubation the majority of type C CpG was sequestered to the PLT surface. Although prolonged exposure of human PLTs to FITC–type C CpG (4 h) induced endocytosis of the ODN, these showed minimal colocalization with intracellular PLT TLR9.

Discussion

Here, we show that TLR9 mRNA is expressed by mature MKs and becomes specifically up-regulated during pro-PLT production. Differential mRNA expression has previously been shown for TIMP-3 (Cecchetti et al., 2011) and suggests that TLR9 mRNA is selectively withheld or rapidly degraded in circulating PLTs. We speculate that increased TLR9 protein translation at this stage is necessary to package TLR9 protein into newly generated PLTs. VAMP 8 colocalizes with TLR9 protein throughout pro-PLT maturation and may be responsible for distributing T granules to PLT-sized swellings, where it is distributed with PDI to unique electron-dense membrane-delimited intracellular compartments we have named T granules. Localization of PDI to T granules suggests they may be derived from the endoplasmic reticulum of MKs and therefore related to the dense tubular system in PLTs (White and Gerrard, 1976). PDI facilitates the formation of disulfide bonds and proper folding of newly synthesized proteins. Secretion and targeting of PDI to the surface of activated PLTs is thought to contribute to isomerase activity, affect adhesion to CD41/61, α₂β₁, and GP Ibα, and regulate the procoagulant activity of tissue factor (Chen et al., 1992, 1995; Essex et al., 1995; Burgess et al., 2000; Lahav et al., 2003; Ahamed et al., 2006). T granules can be organized in two major subpopulations that differentially associate with either VAMP 7 or VAMP 8. Although thought to reside exclusively in the endosomes of monocytes, macrophages, and plasmacytoid dendritic cells, TLR9 surface expression is significantly increased in PLTs after incubation with thrombin, ADP, PMA, CRP, and type IV collagen, demonstrating partial activation-mediated granule release. SNAREs represent the core of the fusion machinery, and the distribution and association of v-SNAREs, such as VAMP 8, with t-SNAREs provides the basis for α-granule localization and secretion (Blair and Flaumenhaft, 2009).
Because TLR9 partly redistributes to the plasma membrane on contact activation with glass, T-granule colocalization with VAMP 8 and reposition to the PLT granulomere in spread PLTs suggest that increased TLR9 surface expression occurs through VAMP 8–mediated T-granule fusion with the open canalicular system (Flaumenhaft, 2003).

TLRs functionally modulate innate immunity by recognizing PAMPs on invading microorganisms (Hemmi et al., 2000). This is particularly true of bacterial/viral DNA, which depend on the presence of unmethylated CpG ODNs. By comparison, mammalian DNA has a lower frequency of CpG ODNs, is methylated, and does not have immune-stimulatory activity (Krieg, 1996; Lipford et al., 1998; Yamamoto et al., 2000). TLR9 has been identified as the receptor involved in the recognition of immunostimulatory CpG motifs (Hemmi et al., 2000; Bauer et al., 2001) and is uniquely expressed in peripheral granules in resting PLTs and on the human PLT surface after activation. Immunostimulatory CpG motifs have been implicated as a major contributor to the acute inflammatory response associated with nonviral vectors, most prominently seen after systemic delivery of cationic lipid–plasmid DNA complexes (Zhao et al., 2004). Unmethylated CpGs have been shown to stimulate B cell proliferation and macrophage activation and induce dendritic cell maturation (Krieg et al., 1995; Stacey et al., 1996; Sparwasser et al., 1998). Incubation of resting PLTs with synthetic unmethylated type C CpG ODNs (characteristic of bacterial/viral DNA) resulted in increased surface expression of TLR9 in humans and increased CD62P surface expression and type C CpG sequestration over 20 min in both humans and mice. Indeed, TLR9 KO mice sequester ~50% less type C CpG and have significantly reduced levels of CD62P surface expression relative to isotype-matched wild-type controls.

Upon vascular damage, PLTs are recruited to the site of injury where they adhere to exposed basement membrane, become activated, and contribute to clot formation. The basement membrane is particularly rich in type IV collagen and PLT membrane is particularly rich in type IV collagen and PLT recruitment to a site of vascular injury may serve to sensitize the PLT to bacteria/virus DNA. Interestingly, when TLR9 surface expression was specifically increased by preincubating PLTs with type IV collagen, subsequent incubation with type C CpG resulted in considerably enhanced type C CpG sequestration, CD62P surface expression, and PLT clumping within 30 s of addition but did not result in thrombi formation. These results imply that PLTs must be primed to express TLR9 on their surface before signal transduction through TLR9 and provide a mechanism for PLT regulation of the immune response to infection in human whole blood by sequestering bacterial/viral DNA and marking themselves for clearance.

Interestingly, mouse PLTs did not show increased surface expression of TLR9 on addition of the ODN. Indeed, human PLTs express several surface receptors absent in mice (e.g., PAR1). With regard to TLR expression, human myeloid dendritic cells express functional TLR8, whereas mice do not. Conversely, although both myeloid and plasmacytoid dendritic cells in mice express intracellular TLR7 and TLR9, these are not present in human myeloid dendritic cells. Nevertheless, TLR9 KO mouse PLTs still aggregated when type C CpG was added after preincubation with type IV collagen, which substantiates the previously reported prevalence of non–TLR9–mediated effects of CpG. Indeed, although the absence of CpG recognition by TLR9 significantly reduces acute inflammatory responses to systematically delivered cationic lipid–plasmid DNA complexes, it does not eliminate them, and TLR9 KO mice still exhibit a pronounced loss of lymphocytes and PLTs (Zhao et al., 2004). Collectively, these findings suggest that signaling through TLR9 does not exclusively contribute to PLT activation when exposed to bacterial/viral DNA and predict the presence of at least one other yet-uncharacterized receptor of CpG on the PLT surface. Even so, increased surface expression of TLR9 in human PLTs, and type C CpG sequestration may help regulate circulating levels of bacterial DNA in whole blood and thus control the inflammatory response after bacterial cell lysis.

The interaction of bacteria with human PLTs plays an important role in the pathogenesis of cardiovascular infection, and serious complications of bacteremia include life-threatening infective endocarditis, disseminated intravascular coagulation, immune thrombocytopenia purpura, and increased risk of myocardial infarction (Fitzgerald et al., 2006). A better understanding of the role TLRs play in regulating PLT activation will lead to targeted therapies for treating and preventing serious cardiovascular infections. Moreover, the development of small molecule inhibitors of PLT interactions with specific bacterial components have the advantage of bypassing present antibiotic resistance mechanisms and may be ideal for prophylaxis of infection in susceptible individuals. (Fitzgerald et al., 2006).

**Materials and methods**

**Human/murine whole-blood PLTs**

Human blood was obtained by venipuncture from healthy laboratory volunteers, a Gray PLT syndrome patient, and a Hermansky–Pudlack syndrome patient as previously described (Hartwig and DeSisto, 1991). Collection was performed in accordance with ethics regulation with International Review Board approval, and informed consent was provided according to the Declaration of Helsinki. Murine blood was obtained by retro orbital bleed into 0.1 vol Aster–Jandl anticoagulant from anesthetized mice as previously described (Hoffmeister et al., 2003). Gray PLT syndrome patient whole blood was provided by A. Michelson (Harvard Medical School, Boston, MA). Hermansky–Pudlack syndrome patient whole blood was provided by R. Flaumenhaft (Harvard Medical School, Boston, MA). TLR9 KO mice were provided by D. Golenbock and B. Caetano (University of Massachusetts Medical School, Worcester, MA).

**MK suspension cultures**

Mouse fetal liver cells were collected from wild-type CD1 mice (Charles River) on embryonic day 13.5 and cultured at 37°C and 5% CO₂ in the presence of 0.1 µg/ml purified recombinant mouse c-Mpl ligand for 5 d. Round MKs, pro-PLT–MKs, and released pro-PLTs were isolated by BSA gradient sedimentation and cultured separately as previously described (Thon et al., 2010). In brief, fetal liver cell cultures were layered on a second single-step gradient (1.5–3.0% BSA) on culture day 4, and MKs were allowed to sediment for 30 min. Mouse fetal liver cell cultures were layered on a single-step gradient (1.5–3.0% BSA) on culture day 4, and MKs were allowed to sediment for 30 min. The MK pellet was then resuspended in fresh media and cultured for an additional 24 h during which pro-PLT production was readily observed and layered on a second single-step gradient (1.5–3.0% BSA) on culture day 5. MKs were allowed to sediment for 30 min, and MKs were isolated from the pelleted fraction, pro-PLT–producing MKs were isolated from the BSA fraction, and released pro-PLTs and PLTs were isolated from the upper (culture media) fraction. Preparation of polyadenylated RNA for sequencing on the Genome Analyzer IIX (Illumina) followed by alignment and analysis was performed as previously described (Rowley et al., 2011). In brief, cells were lysed in TRIzol.
Flow cytometry

Human/mouse PLTs were collected from whole blood and either (a) washed and examined under resting conditions or after activation (5 min at 37°C) with 1 µl/µl thrombin (Roche) or (b) isolated from PRP and examined under resting conditions or activation (5 min at 37°C) with 1 µM PMA (Sigma-Aldrich), 3 µg/ml CRP (Fael et al., 2010), 20 µM ADP (Bio/Data Corporation), or 50 µg/ml mouse type IV collagen (Gibco). Samples were probed with phycoerythrin (PE)-conjugated mouse anti-TLR9 antibodies (Imgenex) and one of the following: FITC-conjugated type C CpG (InvivoGen), FITC-conjugated mouse anti-CD62P antibodies, or FITC-conjugated mouse anti-CD63 antibodies (BD) and run on a flow cytometer (FACScalibur; BD). PLTs were gated by their characteristic forward and side scattering as they passed through the detector, and their total fluorescence intensity was calculated after subtraction of a PE- or FITC-conjugated IgG antibody specificity control (BD). Analyses of PLT mean fluorescence intensity were performed for at least three independent samples. Where appropriate, data were subject to one-way analysis of variance (ANOVA) and Tukey honestly significant difference (HSD) analysis. Error bars represent one standard deviation about the mean for at least three independent samples.

For examination of PLT activation, TLR9 and CD62P surface expression, and type C CpG sequestration, washed human PLTs were incubated for 0, 5, 10, 20, 40, and 60 min at 37°C with either 5 µM FITC-conjugated type C CpG (InvivoGen) or a vector control. Samples were immediately fixed in 4% formaldehyde and probed with PE-conjugated mouse anti-TLR9 antibodies (Imgenex) or PE-conjugated mouse anti-CD61P antibodies (BD) before being run on a flow cytometer (FACScalibur). PLTs were gated by their characteristic forward and side scattering as they passed through the detector, and the percentage of fluorescently conjugated PLT-sized events was calculated after subtraction of a PE- or FITC-conjugated IgG antibody specificity control (BD). Analyses of PLT mean fluorescence intensity were performed for at least three independent samples. Where appropriate, data were subject to one-way ANOVA and Tukey HSD analysis. Error bars represent one standard deviation about the mean for at least three independent samples.

Immunofluorescence microscopy

MKS, pro-PLT–MKs, released pro-PLTs, and whole-blood PLTs were isolated as described in MK suspension cultures. For type C CpG endocytosis experiments human whole blood–washed PLTs were incubated with FITC-conjugated type C CpG (InvivoGen) at 37°C and 5% CO2 for a period of ≤4 h. In brief, samples were fixed in 4% formaldehyde and centrifuged onto 1 µg/ml poly-L-lysine–coated coverslips, permeabilized with 0.5% Triton X-100, and blocked in immunofluorescence blocking buffer (0.5 g BSA, 0.25 ml of 10% sodium azide, and 5 ml FCS in 50 ml PBS) overnight before antibody labeling (Italano et al., 2003). To demarcate permeabilized samples, samples were incubated with a rabbit polyclonal primary antibody for human/mouse α-tubulin against the C-terminal peptide sequence CKAVLEEDEEVEEAMEDKPH and LEDSDEEAEAAEVEDKHKH, respectively (Genemed Synthesis, Inc.). For granule localization, samples were incubated with primary antibodies against TLR9 (Imgenex), VEGF, fibrinogen, CD62P, CD42a, PTD-BF (Santa Cruz Biotechnology, Inc.), CD42b (Dako), M6P, LAMP-1 (Abcam), Syntaxin-13 (a gift from A. Peden, University of Cambridge, Cambridge, England, UK), serotonin (EMD Millipore), or VAMPs 5, 7, and 8 (Neuromics Biotechnology; St. Louis, MO). All samples were vibrated with a secondary goat anti–rabbit or mouse antibody conjugated to an Alexa Fluor 488 or 568 nm (Invitrogen; Molecular Probes). As background controls, slides were incubated with the appropriate secondary antibody alone, and all images were adjusted to account for nonspecific binding of antibodies. For granule localization experiments with known α or dense granule, lysosomal, and endosomal markers, samples were examined with a microscope (Axiovert 200; Carl Zeiss) equipped with a 63x NA 1.4 Plan/Apochromat oil immersion objective, and images were obtained using a charged-coupled device (CCD) camera (ORCA-ER Hamamatsu Photonics). Images were analyzed using the MetaMorph image analysis software (Molecular Devices) and ImageJ (National Institutes of Health).

For colocalization experiments with PDI, TLR7 and 8, and VAMPs 5, 7, and 8, 4.0 x 10^6 washed PLTs were spun down onto poly-L-lysine–coated cover slides in either 4% formaldehyde (resting PLTs) or in PLT resuscitation buffer (10 mM Hepes, 140 mM NaCl, 3 mM KCl, 0.5 mM MgCl2, 5 mM NaHCO3, and 10 mM glucose, pH 7.4) and allowed to spread on a glass slide for 15 min at room temperature before fixation (spread PLTs). Samples were permeabilized with 0.5% Triton X-100 and blocked in immunofluorescence blocking buffer overnight before antibody labeling and then incubated with IgG matching the primary antibody’s species to block nonspecific binding. PLTs were subsequently probed with primary antibodies against TLR9 (Imgenex) and PDI (Abcam), TLR7 (Abcam), TLR8 (Abcam), VAMP 5 (Abcam), VAMP 7 (SYBL1; Abcam), or VAMP 8 (Synaptic Systems). Fluorescence images were visualized using a confocal microscope (BX62; Olympus) equipped with a 60x NA 1.42 Plan-Apochromat oil immersion objective and captured with a cooled CCD camera (ORCA-ER). Image acquisitions were controlled by ImageJ (BioVision Technologies). Additional images were acquired by a confocal laser-scanning microscope with a TCS SP2 (for PDI, TLR7, and TLR8; Leica) or LSM 510 Meta system (for VAMPs 5, 7, and 8; Carl Zeiss). Fluorescence images of two colors were captured sequentially using laser lines at 488 nm (for Alexa Fluor 488) and 561 nm (for Alexa Fluor 568). Series of z images were collected along the z axis at 0.244-µm intervals using a Plan-Apochromat 63x, NA 1.4 oil immersion objective using an optical zoom of 2x. Image analysis and colocalization values were generated using ImageJ and Velocity (PerkinElmer) software. For PDI, TLR7, and TLR8 colocalization experiments, three individual images were averaged to generate a representative image for each sample plane. At least 10 averaged images were collected at 0.244-µm steps along the z axis of each sample for both fluorescence channels, and Pearson’s correlation coefficients above threshold were calculated using the Colocalization Threshold macro written by T. Collins and W. Rasband (National Institutes of Health, Bethesda, MD) for ImageJ.

Immunogold electron microscopy

Rapid-freeze immunogold electron microscopy of washed human PLTs was performed as previously described (Italano et al., 1999). In brief, human PLTs were fixed with 1.25% paraformaldehyde, 0.03% picric acid, and 2.5% glutaraldehyde in 0.1 M cacodylate buffer, pH 7.4. Ultrathin sections were probed for TLR9 (Imgenex) and labeled with protein A–gold (British Biocell International) before staining/embedding. Grids were examined with an electron microscope (G2 Spirit BioTwin; Tecnal) at an accelerating voltage of 80 kV. Images were recorded with an Advanced Microscopy Techniques 2k CCD camera using Advanced Microscopy Technologies digital acquisition and analysis software.

Differential interference contrast live-cell imaging of PLT clumping

Washed human PLTs were transferred onto video chambers maintained at 37°C. Cells were viewed on an inverted microscope (TE-200; Nikon) equipped with a 63x oil immersion objective, and images were obtained using a CCD camera (ORCA-ER). Videos were prepared using the MetaMorph software package and ImageJ. Pictures of PLTs cultured with either 20 µM IRAK-1/4 for 1 h or a vehicle control and 50 µg/ml type IV collagen were captured at 30 s intervals over a course of 20 min. 5 µM type C CpG was then added to the PLT culture, and images were captured for an additional 20 min. Experiments were performed for at least three different samples.

Perfusion analysis of PLT thrombus formation

Washed human whole-blood PLTs or PRP were recalcified with 5 mM CaCl2 and pretreated with 5 µM of control ODN, type C CpG ODN, or a vehicle control at 37°C for 10 min. Samples were perfused at a rate of 18.7 µl/min over a 300 µg/ml type IV collagen-coated or type I collagen-coated (Takeda Pharmaceuticals International GmbH) perfusion chamber. All samples were perfused with a secondary goat anti–rabbit or mouse antibody conjugated to an Alexa Fluor 488 or 568 nm (Invitrogen; Molecular Probes). As background controls, slides were incubated with the appropriate secondary antibody alone, and all images were adjusted to account for nonspecific binding of antibodies. For granule localization experiments with known α or dense granule, lysosomal, and endosomal markers, samples were examined with a microscope (Axiovert 200; Carl Zeiss) equipped with a 63x NA 1.4 Plan/Apochromat

Preparation of photomicrographs

The digital images produced in MetaMorph and AxioVision LE were assembled into composite images by using ImageJ and Photoshop CS3
References


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