Response to Staphylococcus aureus requires CD36-mediated phagocytosis triggered by the COOH-terminal cytoplasmic domain

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Phagocyte recognition and clearance of bacteria play essential roles in the host response to infection. In an ongoing forward genetic screen, we identify the Drosophila melanogaster scavenger receptor Croquemort as a receptor for Staphylococcus aureus, implicating for the first time the CD36 family as phagocytic receptors for bacteria. In transfection assays, the mammalian Croquemort paralogue CD36 confers binding and internalization of Gram-positive and, to a lesser extent, Gram-negative bacteria. By mutational analysis, we show that internalization of S. aureus and its component lipoteichoic acid requires the COOH-terminal cytoplasmic portion of CD36, specifically Y463 and C464, which activates Toll-like receptor (TLR) 2/6 signaling. Macrophages lacking CD36 demonstrate reduced internalization of S. aureus and its component lipoteichoic acid, accompanied by a marked defect in tumor necrosis factor-α and IL-12 production. As a result, Cd36−/− mice fail to efficiently clear S. aureus in vivo resulting in profound bacteraemia. Thus, response to S. aureus requires CD36-mediated phagocytosis triggered by the COOH-terminal cytoplasmic domain, which initiates TLR2/6 signaling.

Introduction

Phagocytosis plays an important role in first line host defense. Phagocytic cells, such as macrophages and dendritic cells, must efficiently differentiate “infectious nonself” from “non-infectious self” and initiate an appropriate inflammatory response (Hoffmann et al., 1999). Central to this process is the expression of pattern recognition receptors, such as the scavenger receptors (SRs), which bind a broad array of modified and foreign ligands (Mukhopadhyay and Gordon, 2004). The B class SR CD36 is an 88-kD glycoprotein expressed by myeloid cells, platelets, endothelial cells, erythroid precursors, and adipocytes (Febbraio et al., 2001). This multifunctional receptor mediates homeostatic functions, including adhesion to thrombospondin and collagen, transport of long-chain fatty acids, and inhibition of angiogenesis (Febbraio et al., 2001). CD36 also shares with its Drosophila melanogaster parologue, Croquemort, the ability to recognize apoptotic cells (Savill et al., 1992; Franc et al., 1999). In addition to these scavenging functions, CD36 has been implicated in the innate immune response to modified host ligands such as oxidized lipoproteins and fibrillar β-amyloid, as well as foreign antigens such as the Pfemp antigen of Plasmodium falciparum (Ockenhouse and Chulay, 1988; Oquendo et al., 1989; Endemann et al., 1993; Moore et al., 2002; Medeiros et al., 2004). Intriguingly, human CD36 mutations predicted to result in the loss of the COOH-terminal transmembrane domain and cytoplasmic tail (T1264G and G1439C) are relatively common in certain African and Asian populations (occurring at high frequency between 0.5–2%), but are extremely uncommon in Caucasians (Aitman et al., 2000). This suggests that selection pressures exist to maintain or eliminate CD36 mutations within certain populations, and these may include interactions with pathogens. Two such interactions may be CD36-mediated recognition of P. falciparum in malaria and a more recently identified role as a candidate receptor for lipoteichoic acid (LTA), a cell wall component of the Gram-positive bacterium Staphylococcus aureus (Hoebe et al., 2005). However, the mechanism by which CD36 participates in the response to pathogens and their ligands and the consequences of CD36 mutations on the host defense have not been defined.
and (Mock; open bars). *, P ≤ 0.05. (c) Expression of CD36 by HEK293 cells confers a threefold increase in internalization of both FITC-labeled S. aureus and E. coli, as compared with vector-transfected cells (Mock; open bars). *, P ≤ 0.05.

Figure 1. The Drosophila protein Croquemort and its mammalian paralogue CD36 are receptors for S. aureus. (d) Croquemort and the small GTPase Rac2 are specifically required for S. aureus binding and uptake. RNAi silencing of Croquemort (Crq) and Rac2 in Drosophila S2 cells reduced phagocytosis of S. aureus (left), but not E. coli (right). *, P ≤ 0.05; **, P ≤ 0.001; significantly different from control. (b and c) Expression of CD36 confers cellular binding and internalization of S. aureus and E. coli. (b) HEK293T-CD36 expressing cells (CD36; filled bars) demonstrate a threefold increase in binding of FITC-labeled S. aureus and a twofold increase in binding of FITC-labeled E. coli over vector-transfected cells (Mock; open bars). *, P ≤ 0.05. (c) Expression of CD36 by HEK293 cells confers a threefold increase in internalization of both FITC-labeled S. aureus and E. coli, as compared with vector-transfected cells (Mock; open bars). *, P ≤ 0.05.

Mammalian pattern recognition receptors have evolved to identify highly conserved pathogen-associated molecular patterns essential to microbes’ existence or pathogenicity, including bacterial lipopolysaccharide (LPS), LTA, peptidoglycan, bacterial CpG-containing DNA, double-stranded RNA, or mannans. A family of receptors related to Toll, a Drosophila receptor involved in dorsal ventral patterning, have emerged as key regulators in the host response to invading pathogens (Akira and Takeda, 2004). The mammalian Toll-like receptor (TLR) family has 10 members, which each recognize a distinct subset of microbial molecules from bacteria, viruses, fungi, and protozoa to activate common signaling pathways that initiate the innate immune response (Akira and Takeda, 2004). The best-characterized TLR is perhaps TLR4, which cooperates with the glycosylphosphatidylinositol-anchored protein CD14 to mediate the signaling response to bacterial LPS (Akira and Takeda, 2004). Although the exact mechanism has yet to be defined, CD14 is believed to cluster LPS at the cell surface where it can deliver this ligand to TLR4. A precedent is emerging for similar cooperation between TLRs and other molecules. As an example, dectin-1, a C-type lectin phagocytic receptor, has been shown to cooperate with TLR2 in mediating response to zymosan (Gantner et al., 2003). Additionally, Rac1, a small Rho-GTPase involved in phagocytosis, has been shown to be required for TLR2-mediated activation of NFκB by S. aureus (Arbibe et al., 2000). Intriguingly, although many TLRs are found on the cell surface, members of this family also localize to endosomes, lysosomes, or the Golgi apparatus (Underhill et al., 1999; Underhill and Gantner, 2004), raising the possibility that these cooperative interactions exist to facilitate ligand delivery to the appropriate compartment after endocytosis or phagocytosis for efficient activation of the immune response.

Here, we show that CD36 and its Drosophila paralogue Croquemort have a similar role in tethering bacteria and derived ligands. This recognition plays an important role in initiating macrophage cytokine production in response to S. aureus, its cell wall component LTA, and, to a lesser extent, Gram-negative E. coli and its derived ligand, LPS. We demonstrate that the COOH-terminal cytoplasmic domain, which is lost in several of the common human mutations of CD36, is required for bacterial internalization and cooperation with TLR2, suggesting that individuals carrying these mutations may be more susceptible to S. aureus infection. Mice deficient in CD36 demonstrate a marked defect in the host response to S. aureus but not E. coli. A combined defect in cytokine response and bacterial phagocytosis in CD36 null mice results in profound bacteraemia and the formation of tissue abscesses. We suggest that like CD14, CD36 functions as an accessory receptor at the cell surface to present bacterial ligands to their cognate TLRs to initiate signaling (Lien et al., 1999; Schwandner et al., 1999; Morr et al., 2002). Furthermore, through its ability to internalize S. aureus and LTA, CD36 also delivers ligands to TLRs recruited to the phagosome/endosome (Underhill et al., 1999).

Results

The Drosophila CD36 paralogue Croquemort is a receptor for S. aureus

In an on-going high-throughput RNA interference (RNAi) screen for Drosophila genes involved in phagocytosis, we identified that the SR Croquemort showed a specific inhibition of phagocytosis of S. aureus but not E. coli (Fig. 1 a). Drosophila S2 cells treated with RNAi to target Croquemort showed a 35% decrease in their ability to internalize S. aureus, whereas targeting of an irrelevant control gene (GFP) showed no effect on bacterial internalization (unpublished data). Mammalian Rac has been shown to have an important role in responses to S. aureus (Arbibe et al., 2000). Therefore, we tested whether Rho-GTPase family members played a role in S. aureus uptake in Drosophila. Silencing of Rac2, but not Rac1, resulted in a specific defect in S. aureus internalization comparable to that seen with Croquemort inhibition, providing circumstantial evidence that Croquemort may activate this Rho-GTPase to mediate downstream phagocytosis. These results suggest that Croquemort and Rac2 play nonredundant roles in the recognition and internalization of S. aureus in Drosophila. However, lethality of Croquemort deficient flies precluded further study of the role of this molecule in innate immune defense in this invertebrate system.
CD36 transfection confers both binding and internalization of S. aureus and E. coli

The mammalian SR CD36 is paralogous to Croquemort, and these receptors share the ability to bind apoptotic cells, indicating that they have overlapping ligand specificities (Savill et al., 1992; Franc et al., 1999). Therefore, our observation that Croquemort mediated engulfment of S. aureus raised the possibility that CD36 might also function as a mammalian receptor for Gram-positive bacteria. Thus, to test the ability of CD36 to phagocytose bacteria, we measured binding and internalization of fluorescently labeled S. aureus and E. coli in human embryonic kidney (HEK) 293T cells transfected with a murine CD36 cDNA. Expression of CD36 conferred a threefold increase in binding of S. aureus and a twofold increase with E. coli over mock transfected control cells (Fig. 1 b). Furthermore, expression of CD36 resulted in a threefold increase in cellular internalization of bacteria (Fig. 1 c), indicating that CD36 acts as a phagocytic receptor for both Gram-positive and, to a lesser extent, Gram-negative bacteria.

Tyrosine 463 within the COOH-terminal cytoplasmic tail of CD36 is required to signal engulfment

The COOH-terminal cytoplasmic tail of CD36 is lacking in several common human CD36 mutations (Aitman et al., 2000), and this domain is postulated to interact with signaling molecules (Huang et al., 1991; Bull et al., 1994; Jimenez et al., 2000; Moore et al., 2002). Thus, we addressed the consequences of the lack of the COOH-terminal region on phagocytosis of S. aureus using structure–function analysis. To determine the minimum requirements for CD36 to signal to the actin cytoskeleton and trigger engulfment, HEK293T cells were transfected either with wild-type CD36 (CD36WT) or with a mutant form of CD36 in which the 11 COOH-terminal cytoplasmic amino acid residues had been replaced by alanines (CD36Ala; Fig. 2 a). Equivalent cell surface expression of CD36WT and CD36Ala was confirmed by flow cytometry (Fig. 2 b) and bacterial phagocytosis was measured. Importantly, mutant constructs demonstrated a ≈3% difference in cell surface expression in all experiments, indicating that the tail was not required for cell surface localization. Transfection of CD36WT and CD36Ala resulted in a twofold increase in bacterial binding; however, only CD36WT conferred S. aureus phagocytosis, suggesting that the COOH-terminal tail of CD36 is required to signal the actin cytoskeleton (Fig. 2 c). Included in these COOH-terminal amino acids is a tyrosine residue which, as a potential site of phosphorylation, could provide a docking site for SH2 domain–containing proteins, such as the Src family of tyrosine kinases and the actin regulator p130Cas.

Thus, to further define the region responsible for signaling internalization, we mutated this tyrosine at position 463 to phenylalanine (CD36ΔY463F). Cells transfected with the CD36ΔY463F mutant bound S. aureus, but, similar to CD36Ala, showed an impairment of phagocytosis, indicating that tyrosine 463 is essential for CD36-mediated bacterial internalization (Fig. 2 c). Together, these data indicate that the COOH-terminal cytoplasmic tail of CD36 regulates receptor-mediated internalization of S. aureus and implicates tyrosine 463 in signaling to the actin cytoskeleton. Thus, we would predict that the reported human CD36 mutations would also be unable to trigger internalization in response to this organism.

CD36-deficient macrophages demonstrate impaired phagocytosis of S. aureus and cytokine production

To define the consequences of lack of functional CD36 signaling, we used CD36 null mice to evaluate phagocytosis and response to S. aureus, E. coli, and their derived ligands. Cd36−/− macrophages showed a 40–50% reduction in phagocytosis of S. aureus, but comparable levels of internalization of E. coli relative to wild-type macrophages (Fig. 3, a and c). This suggests a relative specificity for Gram-positive organisms conserved between the Drosophila Croquemort protein and mammalian CD36. LTA is an abundant cell wall component of S. aureus that is released after exposure to antibiotics or leukocytic mediators. To better identify the ligand present on S. aureus that is recognized by CD36, we tested whether CD36 could bind and internalize LTA. In concordance with our findings with whole S. aureus, Cd36−/− macrophages demonstrated a 60% reduction in their ability to internalize BODIPY-LTA (Fig. 3 b), indicating that CD36 mediates phagocytic/endoctytic recognition of this important TLR ligand.

Cd36−/− macrophages treated with S. aureus for 4 h showed a profound reduction (65–75%) in TNFα and IL-12 protein expression as compared with wild-type macrophages...
CD36-deficient macrophages demonstrate impaired phagocytosis of *S. aureus* and its component LTA.

Peritoneal macrophages from wild-type and *Cd36*−/− mice were incubated with FITC-*S. aureus* or FITC-*E. coli* (10:1 bacteria/cell; panel a) or 10 μg/ml BODIPY-LTA (b), and phagocytosis was measured by flow cytometry in the presence of trypan blue. **, P < 0.005, compared with wild-type macrophages. [c] Representative photographs of wild-type and *Cd36*−/− macrophages incubated with FITC-*S. aureus* (green) and stained with DAPI (blue) to show individual nuclei. Macrophages were incubated with 10:1 bacteria/cell and allowed to internalize bacteria for 30 min at 37°C. Cells were washed 3× in PBS to remove extracellular bacteria and fixed with formaldehyde.

![Figure 3](image)

CD36-deficient macrophages demonstrate impaired phagocytosis of *S. aureus* and its component LTA. Peritoneal macrophages from wild-type and *Cd36*−/− mice were incubated with FITC-*S. aureus* or FITC-*E. coli* (10:1 bacteria/cell; panel a) or 10 μg/ml BODIPY-LTA (b), and phagocytosis was measured by flow cytometry in the presence of trypan blue. **, P < 0.005, compared with wild-type macrophages. [c] Representative photographs of wild-type and *Cd36*−/− macrophages incubated with FITC-*S. aureus* (green) and stained with DAPI (blue) to show individual nuclei. Macrophages were incubated with 10:1 bacteria/cell and allowed to internalize bacteria for 30 min at 37°C. Cells were washed 3× in PBS to remove extracellular bacteria and fixed with formaldehyde.

As CD36 has the additional capacity to internalize ligands, we investigated whether CD36-mediated internalization contributes to activation of TLR2/TLR6 signaling by blocking LTA internalization with cytochalasin D in HEK293-TLR2 or -TLR6 cells transfected with CD36, TLR6, or both. Inhibition of LTA internalization abrogated CD36-initiated NFκB activation, indicating a requirement for uptake to initiate TLR2/6 signaling (Fig. 6 c). Similar to our findings with *S. aureus*, the *Cd36*−/− mutation also blocked internalization of LTA, indicating a requirement for the COOH-terminal tail of the receptor for internalization of not only whole bacteria but also of its cell wall component, LTA (Fig. 6 d). To specifically map the requirement for LTA uptake, we mutated Y463, C464, and C466 found that only mutation of Y463 and C464, but not the distal cysteine C466, inhibited LTA internalization. Having shown that internalization was necessary for signaling, we hypothesized that these mutations would also disrupt CD36 cooperation with TLR2/6. Similar to the inhibition of LTA internalization, mutation of the entire COOH-terminal tail to alanine resulted in a 70% decrease in NFκB activation com-

![Figure 4](image)

**CD36** −/− macrophages demonstrate impaired early responses to whole *S. aureus* and, to a lesser extent, *E. coli*. Wild-type and *Cd36*−/− macrophages were treated with 10:1 *S. aureus* (a) or *E. coli* (b) for 4 h, and TNFα, IL-12, and IL-6 were measured in cell culture supernatants. **, P < 0.05; ***, P < 0.005, significantly different from wild-type macrophages.
pared with wild-type CD36. This deficit could be recapitulated by mutation of either Y463 or C464, but not C466, confirming that mutations that block internalization also block NFκB induction (Fig. 6 e). This is supported by the finding that whereas cytochalasin D inhibited signaling by wild-type CD36, no inhibition of this TLR2/6 signaling was observed with CD36Ala or CD36Y463F (Fig. 6 f). Together, these data indicate that CD36-mediated internalization and cooperation with TLR2/6 is required for effective activation of the inflammatory response to LTA.

**CD36 null mice fail to clear S. aureus in vivo**

To establish the role of CD36 in the innate immune response to bacteria, CD36−/− and wild-type mice were infected intravenously with S. aureus and E. coli (Fig. 7 a). When infected with encapsulated S. aureus (Reynolds strain), CD36−/− failed to clear the initial bacteremia, resulting in a mean bacterial load of $3.8 \times 10^9$ S. aureus colony forming units per milliliter of blood on day 4 (Fig. 7 b). In contrast, wild-type mice had efficiently cleared the S. aureus and no circulating bacteria were detectable. Furthermore, 20% of CD36−/− mice died 6–8 d after inoculation, whereas all wild-type mice survived the infection (Fig. 7 a). Necropsy examination of these mice on day 22 demonstrated that the prolonged early bacteremia in CD36−/− mice resulted in abscess formation in 50% of the animals (Fig. 7 c). In contrast, only 1 of 10 wild-type mice had an apparent abscess of the kidney, indicating that these mice had successfully cleared the bacteria. Thus, loss of CD36-mediated clearance and cytokine signaling appeared to have profound biological consequences on the host response to S. aureus infection. In unencapsulated S. aureus strains, LTA is exposed on the bacterial cell surface. As CD36 directly recognizes LTA, we hypothesized...
that it may make a greater contribution to host defense against unencapsulated *S. aureus*. When injected with the unencapsulated *S. aureus* strain 3825-4 (*CP5*), 50% of *Cd36*−/− mice succumbed to the infection, whereas all wild-type mice survived (Fig. 7 a). Furthermore, 100% of the surviving *Cd36*−/− mice developed perinephric and/or pericardial abscesses. Similar experiments using *E. coli* failed to demonstrate any significant difference in the survival of wild-type and *Cd36* null mice, indicating that this was not a nonspecific immune response, but rather that *Cd36* has an important and nonredundant role in combating *S. aureus* infection, particularly those unencapsulated strains in which LTA is exposed.

**Discussion**

In a forward genetic screen, we identify that Croquemort, a *Drosophila* SR, plays a specific role in the phagocytosis of *S. aureus*, demonstrating for the first time that the members of this family of CD36-related molecules act as phagocytic receptors for bacteria. We extend this initial observation to confirm that the mammalian paralogue of Croquemort, *Cd36*, also directly phagocytoses bacteria and plays a role in the host immune response to *S. aureus*. Transfection of cells with *Cd36* confers the ability to bind and internalize both *S. aureus* and *E. coli*; however, macrophages deficient in *Cd36* are impaired only in their ability to phagocytose *S. aureus*. Through structure–function analysis, we show that *Cd36*-mediated internalization of *S. aureus* and LTA is dependent on the COOH-terminal cytoplasmic portion of this receptor proposed to interact with signaling molecules. Mutations of the cytoplasmic residues, specifically tyrosine 463, abolish *Cd36*-mediated phagocytosis, but not binding, indicating that this domain is essential to signal to the actin cytoskeleton and trigger engulfment. Furthermore, in the absence of *Cd36*, macrophages show a profound defect in cytokine production in response to both *S. aureus* and its cell wall component (LTA). In vivo, *Cd36* null mice fail to efficiently clear *S. aureus*, resulting in profound bacteraemia and abscess formation. Recent data has identified a role for *Cd36* in the recognition of diacylglycerides and in the activation of signaling via TLR2 and TLR6 (Hoebe et al., 2005). However, the mechanism by which *Cd36* augments this response is unknown. We now show that *Cd36*-mediated internalization of LTA, mediated by its cytoplasmic tail, is required to fully activate TLR2/6 signaling. Our data suggest that *Cd36* functions to both clear circulating bacteria and to bind bacterial ligands where they may be presented to TLR signaling pathways either at the cell surface or the phagosome.

Microbial internalization is accompanied by inflammatory responses that are driven by TLR signaling pathways. Particles are internalized into a membrane-limited organelle, the phagosome, which matures into a highly hydrolytic environment facilitating degradation of internalized bacteria into their component parts. Although TLRs are not believed to function directly as phagocytic receptors, many TLRs are recruited to the newly formed phagosome after particle ingestion, where they are thought to “sample” the phagosome for microbial ligands (Underhill et al., 1999; Underhill and Gantner, 2004). Thus, *Cd36*-mediated delivery of *S. aureus* to this organelle might facilitate the release of ligands such as LTA from the cell wall, thereby delivering these particles to TLRs to initiate inflammatory signaling and cytokine production. On the cell surface, *Cd36* would be able to cluster LTA released from *S. aureus* after exposure to antibiotics or leukocytic mediators, coordinating engagement of TLRs at the cell surface in a manner similar to CD14. An alternative hypothesis is that *Cd36* may mediate its effects independent of TLR signaling pathways. However, *Cd36* transfection alone was insufficient to stimulate NFkB activation in response to *S. aureus* or LTA, suggesting that *Cd36* cooperates with TLRs to initiate NFkB signaling. The majority of the *Cd36* protein forms an extracellular loop containing distinct ligand binding domains, and its short NH2- and COOH-terminal cytoplasmic tails are palmitoylated, localizing this receptor to lipid rafts (Tao et al., 1996). These domains are rich in signaling molecules, and *Cd36* has been shown to interact with Src kinase family members that reside in lipid rafts to engage downstream signaling cascades, including p38 and p44/42 MAPK pathways (Huang et al., 1991; Jimenez et al., 2000; McGilvray et al., 2000; Moore et al., 2002; Bamberger et al., 2003; Medeiros et al., 2004). Recent work has highlighted the importance of MAPK pathways in regulating endocytosis,
phagocytosis, and phagosome maturation in both macrophages and dendritic cells (Blander and Medzhitov, 2004; West et al., 2004). Thus, it is also possible that CD36 regulation of MAPK pathways contributes to augment S. aureus- and LTA-induced cytokine production. However, the CD36 residues vital for augmentation of TLR2/6 signaling in response to LTA (Y463 and C464) are not required for CD36-mediated activation of MAPK by its other ligands, β-amyloid and oxidized low-density lipoprotein (unpublished data). Together, these data provide compelling evidence that CD36-mediated internalization is an important mechanism by which it augments TLR2/6 signaling.

Phagocyte recognition of foreign or modified ligands, such as bacteria and apoptotic cells, is an essential component of both the innate and adaptive immune responses. Particle recognition and internalization is mediated by a variety of phagocytic receptors including the Fcγ receptor, CR3, the mannose receptor, and members of the SR family, including SR-A, LOX-1, and MARCO (Mukhopadhyay and Gordon, 2004; Underhill and Gantner, 2004). Engagement of these receptors ultimately results in activation of RhoGTPases, including Rac, to induce cytoskeletal reorganization, actin polymerization, and particle internalization (Cox et al., 1997). Although, CD36 and its Drosophila homologue Croquemort have been shown to mediate binding of apoptotic cells (Savill et al., 1992; Ren et al., 1995; Franc et al., 1999), it was unclear whether CD36 alone was sufficient to trigger engulfment. We have recently found that β-amyloid engagement of CD36 leads to assembly of focal adhesion molecules that act upstream of Rac1, including p130cas, Pyk2, and paxillin (unpublished data). These observations lead us to hypothesize that CD36, either alone or in cooperation with a coreceptor, can regulate downstream targets such as Rac and the actin cytoskeleton to mediate internalization. Our finding that expression of CD36 confers both binding and internalization of bacteria supports this role of CD36 as a phagocytic receptor. Furthermore, in Drosophila S2 cells, the absence of Croquemort and Rac2 causes a similar degree of inhibition of S. aureus, but not E. coli phagocytosis, suggesting that these molecules may function in the same pathway to signal engulfment. Importantly, we show that the COOH-terminal cytoplasmic residues of CD36 are essential to trigger phagocytic engulfment, indicating that several of the human CD36 mutations known to lack this region are unlikely to support internalization or activation in response to S. aureus or LTA. However, whether this domain of CD36 interacts directly with the actin cytoskeleton through adaptor molecules such as p130Cas, or whether it recruits coreceptors such as the integrins to perform this function, remains to be determined.

To investigate the in vivo consequences of lack of CD36-mediated phagocytosis and signaling, we used mice with a targeted deletion of CD36. We demonstrate that loss of CD36 has profound biological consequences on the early response to S. aureus infection, particularly in the ability to mount a response to an unencapsulated strain (3825-4) of this pathogen. Cd36−/− mice exhibit 5–8 log order greater bacterial loads, which ultimately lead to perinephric and pericardial abscess formation. However, although CD36 is essential for early responses to S. aureus and LTA, including macrophage phagocytosis and cytokine responses, alternative pathways appear to compensate to protect the host against S. aureus–induced lethality, indicating that the innate immune system has evolved overlapping mechanisms to combat this important pathogen. Several other molecules have been implicated in the immune response to S. aureus, including the SRs AI and II and soluble opsonins such as mannose-binding lectin, thrombospondin, and LPS-binding protein (Dunne et al., 1994; Haworth et al., 1997; Thomas et al., 2000; Schroder et al., 2003; Mukhopadhyay and Gordon, 2004; Shi et al., 2004), emphasizing the complexity of this in vivo model system.

Interestingly, CD36 did appear to play a small role in response to E. coli, as TNFα and IL-6 secretion by macrophages was decreased in its absence. However, macrophage TNFα production in response to isolated LPS did not require CD36. Importantly, neither E. coli engulfment by macrophages nor in vivo host response was compromised by the absence of CD36, indicating a specific failure to respond to S. aureus rather than a general immune depression. It is likely that in these complex and partially redundant systems, other receptors such as the LPS receptor CD14 and CD11b/CD18 are more important to mediate host cytokine responses to E. coli. Nonetheless, that CD36 binds both Gram-negative and -positive bacteria is entirely consistent with its function as a multi-ligand SR.

The data presented demonstrate that CD36 acts as a phagocytic receptor able to recognize S. aureus and its cell wall component LTA. These data demonstrate that CD36 functions in a capacity analogous to the LPS receptor CD14, clustering LTA at the cell surface where it may engage TLR2 and TLR6. Moreover, CD36 has the additional capacity to internalize whole S. aureus and LTA, concentrating them in the phagosome or endosome where they can also engage TLR signaling pathways. This internalization is vital for full activation of TLR2/6 signaling in response to LTA. Loss of this ability to internalize ligands in individuals with CD36 mutations altering the COOH-terminal cytoplasmic domain would be predicted to impair the host response, rendering these individuals more susceptible to S. aureus infection. These findings provide a novel mechanism likely to be common to many SRs that links host detection of pathogens to engagement of TLR signaling pathways, both at the cell surface and intracellularly, to activate host proinflammatory responses.

Materials and methods

Mice

The Cd36−/− mice (backcrossed nine generations to C57BL/6J) were generated in our laboratory as described, and Cd36+/− mice were generated from Cd36−/− intercrosses (Moore et al., 2002). All mice were maintained in a pathogen-free facility with free access to rodent chow and water. Experimental procedures were approved by the Massachusetts General Hospital’s Subcommittee on Research Animal Care and were conducted in accordance with the US Department of Agriculture Animal Welfare Act and the Public Health Service Policy for the Humane Care and Use of Laboratory Animals.

Peritoneal macrophage culture

Elicited peritoneal macrophages were collected from mice by peritoneal lavage 4 d after i.p. injection with 3% thioglycollate as described previously (Moore et al., 2002) and cultured in DME containing 2% FBS, 100 U/ml penicillin, and 100 μg/ml streptomycin overnight at 37°C before use.
Transfection assays

COOH-terminal mutations of the murine CD36 cDNA (Fig. 2 a) were generated by PCR, verified by sequencing, and cloned into the pcDNA3.1 vector containing a CMV promoter. HEK293 EBNA T cells were maintained in DME containing 10% FBS, 100 U/ml penicillin, and 100 μg/ml streptomycin and were cultured in medium lacking penicillin and streptomycin for 24 h before transfection. Cells (6 × 10^5 cells/well; 6-well plate) were transfected with pcDNA 3.1 plasmids containing wild-type or mutant CD36 cDNAs, or no insert (mock), in OptiMEM containing Lipofectamine 2000 as previously described (Kim et al., 2004). Equivalent expression of wild-type and mutant CD36 was confirmed 48 h after transfection by flow cytometry using a monoclonal anti-murine CD36 antibody (1:650 dilution; BD Biosciences) as previously described (Medeiros et al., 2004). The anti-CD36 antibody recognizes an epitope in the extracellular portion of CD36 that is not affected by the mutations generated.

Fluorescent labeling of LTA

LTA preparations from S. aureus (Sigma-Aldrich) were labeled with 4,4-difluoro-1,3-dimethyl-5(4-methoxyphenyl)-4-bora-3a,4a-diazio-S-indacene-2-propionyl-tetramethylrhodamine (BODIPY-TMR; Molecular Probes Inc.) using the manufacturer’s protocol for oligosaccharide labeling as previously described (Levets et al., 2003).

RNAi treatment

Double stranded RNA interference was used to specifically silence genes in Drosophila S2 cells as described previously (Ramei et al., 2002). Drosophila S2 cells were cultured in Schneider’s Drosophila media (GIBCO BRL) at 26°C. S2 cells (10^5) were treated with 15 μg dsRNA for 60 h and viability was confirmed by trypan blue exclusion. The ability of RNAi-treated cells to phagocytose heat-killed FITC-labeled S. aureus and E. coli (Molecular Probes) was measured using flow cytometry as described in the following section and compared with cells treated with irrelevant GFP control dsRNA. dsRNA was generated by inverse transcription from a PCR amplicon either from cDNA, template plasmids, or PCR products using gene-specific primers flanked by a T7 polymerase site. Primer sequences were as follows: GFP forward T7, CCTACATACCTCGCTCTGC; GFP reverse T7, CCAGCCG; Crq reverse T7, CTTCATTCCACTGACAGCA; Rac1 forward T7, TTGTACGTAAC-CAAGCCG; Crq forward T7, TTGTACGTAAC-CAAGCCG; Rac1 reverse T7, CATCGCCAGTCCTTGGGGAT-TCAGCTACACGACCAATGCCTTTC; Rac1 reverse T7, GCGCTTTCTCATAGCTCAC; Crq reverse T7, CCTATCTACCTGTCCTTGC; GFP reverse T7, GGCCTTTCCTCATACTGCTAC; Cq reverse forward T7, TTGTACGTAAC-CAAGCCG; Cq reverse reverse T7, CCTATCTACCTGTCCTTGC; Rac1 forward T7; TCAGCTACAGCAATGCTTTC; Rac1 reverse T7; CAGATACTCTGACGCTGCTGATT; Rac2 forward T7; GGCATCATCAAG-TGTGTTGGTGTTG; Rac2 reverse T7; CATCGCCAGTCCTGCGGAGT-AGGT.

Phagocytosis assays

Quantification of binding and uptake of heat-killed FITC-labeled S. aureus and E. coli (both from Molecular Probes) was performed as described previously (Ramei et al., 2002; Shi et al., 2004). In brief, 2 × 10^5 cells in 48-well plates were cooled to 4°C for 20 min, 2 × 10^5 bacteria were added for 30 min at 4°C, and then cells were incubated at 37°C (26°C for S2 cells) for 30 min to allow internalization. Cells were placed on ice to stop internalization and were maintained on ice for the remainder of the assay. Total cell-associated fluorescence (binding and uptake) was measured by flow cytometry (10,000 events) using the CellQuest program (Becton Dickinson). Bacterial internalization was measured by quenching extracellular fluorescence with 0.2% trypan blue shortly before analysis. Binding was calculated as the difference between total cell-associated fluorescence and intracellular fluorescence. The amount of phagocytosis (phagocytic index) was expressed as the percentage of fluorescence-positive cells multiplied by the mean fluorescence of these cells. All treatments were performed in triplicate and data are presented as the mean ± SD.

Microscopy

Peritoneal macrophages were allowed to adhere to glass coverslips for 1 h before the addition of bacteria at a ratio of 10:1. FITC-bacteria were spun onto the cells, and phagocytosis was synchronized by first binding on ice for 30 min. Cells were then warmed to 37°C for 30 min. Internalization was measured by previously described, and cells were fixed using formaldehyde before mounting with Vectashield fluorescent mounting medium containing DAPI stain (Vector Laboratories). Slides were imaged on an inverted fluorescence microscope (model TE 2000-U; Nikon) equipped with a camera (Nikon) using Openlab software (version 4.0.2), original magnification ×40, NA 0.60, at RT. After acquisition, manipulation of brightness, color balance, and contrast was applied uniformly across images using Adobe Photoshop software.

Measurement of cytokine expression

TNFα, IL-12, and IL-6 mRNA levels were measured by real-time quantitative RT-PCR using an iCycler (Bio-Rad Laboratories) as described previously (Medeiros et al., 2004). Total RNA was extracted using Trizol reagent and reverse transcribed using Moloney Murine Leukemia Virus reverse transcriptase. Each reaction contained 3 μl cDNA, 0.3 μM TNFα, IL-12, IL-6, or GAPDH primers, SYBR green (QIAGEN), and HotStarTaq polymerase. PCR conditions were 15 min at 95°C, followed by 30 cycles of 30 s at 95°C, 30 s at 55°C, and 30 s at 72°C. Each sample was analyzed in triplicate and the amount of mRNA in each sample was calculated from a standard curve of known template and normalized to GAPDH. The amount of TNFα, IL-12, and IL-6 protein in cell culture supernatants was measured by ELISA (R&D Systems) as described previously (Shi et al., 2004). Data are presented as the mean of triplicate samples ± SD.

Nf-B luciferase reporter assays

Dual luciferase reporter assays for NFκB activation were performed in HEK293 cells that stably express TLR2 or TLR9 (provided by D. Golenbock, University of Massachusetts Medical School, Worcester, MA) as previously described (Latz et al., 2002). Cells were transfected with a NfκB-luciferase reporter construct and/or pcDNA3.1-CD36 and/or Trk6 cDNAs using lipofectamine 2000 as per the manufacturer’s instructions. Before assays, equivalent expression of CD36 and TIR6 was confirmed in all samples. The cells were stimulated with 1 μg/ml LTA or S. aureus (10:1) for 5 h and lysed, and reporter gene activity was measured using the Dual Luciferase Assay Reporter System (Promega) using a plate-reader luminometer. Data were normalized to cellular protein. In some assays, cells were treated with 1 μg/ml cytochalasin D (Sigma-Aldrich) for the duration of the assay. The data shown represent one of at least four separate experiments and are presented as the mean values ± SD of triplicate samples.

Bacterial infection in vivo

S. aureus was grown overnight in Columbia medium with 2% NaCl and resuspended in saline. Wild-type and Cd36−/− mice (n = 10; 6–8 wk old) were inoculated with S. aureus Reynolds capsular serotype 5 (10^8), unencapsulated S. aureus 3825-4 strain (2 × 10^9), or E. coli 018K (5 × 10^5) by tail vein injection and monitored for survival and complications from infection as described previously (Shi et al., 2004). Moribund animals were killed by CO2 inhalation. In mice injected with S. aureus, blood was collected from the tail vein after 4 and 15 d and mixed with heparin, and serial dilutions of the blood were cultured on tryptic soy agar plates supplemented with 5% sheep’s blood to determine the number of S. aureus colony forming units. Necropsies were performed on surviving mice at the end of the study and abscess formation was quantified.

Statistical analysis

Quantitative data are expressed as the mean ± SD of triplicate samples and are representative of at least three separate experiments. The difference between two groups was statistically analyzed by t test or an analysis of variance (one-way ANOVA). A p-value of <0.05 was considered significant. Survival data were analyzed using Log-Rank and Wilcoxon tests using JMP5 software (SAS Institute, Inc.).

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