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Abstract
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Comments
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Inhibition of "self" engulfment through deactivation of myosin-II at the phagocytic synapse between human cells

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Phagocytosis of foreign cells or particles by macrophages is a rapid process that is inefficient when faced with "self" cells that display CD47—although signaling mechanisms in self-recognition have remained largely unknown. With human macrophages, we show the phagocytic synapse at cell contacts involves a basal level of actin-driven phagocytosis that, in the absence of species-specific CD47 signaling, is made more efficient by phospho-activated myosin. We use "foreign" sheep red blood cells (RBCs) together with CD47-blocked, antibody-opsonized human RBCs in order to visualize synapptic accumulation of phosphotyrosine, paxillin, F-actin, and the major motor isoform, nonmuscle myosin-IIA. When CD47 is functional, the macrophage counter-receptor and phosphatase-activator SIRPα localizes to the synapse, suppressing accumulation of phosphotyrosine and myosin without affecting F-actin. On both RBCs and microbeads, human CD47 potently inhibits phagocytosis as does direct inhibition of myosin. CD47–SIRPα interaction initiates a dephosphorylation cascade directed in part at phosphotyrosine in myosin. A point mutation turns off this motor's contribution to phagocytosis, suggesting that self-recognition inhibits contractile engulfment.

Introduction

A phagocytic cell engulfs another cell or particle that is IgG-opsonized in a coordinated process of adhesion, pseudopod extension, and internalization with phagosome closure. Upon initial binding of IgG, the phagocyte's Fc receptors (FcγRs) activate cytoskeletal assembly with rapid accumulation of phosphopaxillin (Greenberg et al., 1994; Allen and Aderem, 1996) and F-actin (Wang et al., 1984; Greenberg et al., 1991) among other components at a "phagocytic synapse". Nonmuscle myosins also accumulate and suggest a role(s) for contractile motors during particle internalization (Stendahl et al., 1980; Valerius et al., 1981; Diakonova et al., 2002). Signaling activities that influence synapse assembly continue to be clarified (Aderem and Underhill, 1999) and are presumably key to how the macrophage distinguishes foreign cells or particles from autologous cells of "self". Autologous cells are certainly opsonized by Ig (Turrini et al., 1993), and so activation differences are not the complete story. Indeed, based on studies of knockout mice lacking the membrane receptor CD47, this protein on target cells is a phagocytosis-inhibiting "marker of self" (Oldenburg et al., 2000).

CD47 is a ubiquitous member of the Ig superfamily that interacts with the immune inhibitory receptor SIRPα (signal regulatory protein) found on macrophages (Fujisaki et al., 1996; Veillette et al., 1998; Jiang et al., 1999). Although CD47–SIRPα interactions appear to de-activate autologous macrophages in mouse, severe reductions of CD47 (perhaps 90%) are found on human blood cells from some Rh genotypes who show little to no evidence of anemia (Mouro-Chanteloup et al., 2003) and also little to no evidence of enhanced cell interactions with phagocytic monocytes (Arndt and Garratty, 2004). Here, we assess the species-specific inhibition of phagocytosis by CD47–SIRPα interactions at the phagocytic synapse and visualize how the interaction affects cytoskeletal activity.

CD47-SIRPα binding is conserved but species-specific (Subramanian et al., 2007). Based on this divergence and assuming an inhibitory role for human CD47, human macrophages might be expected to efficiently phagocytose opsonized sheep red blood cells (RBCs), which is certainly consistent with years of data showing sheep RBCs are readily phagocytosed by human macrophages (Lowry et al., 1998; Botelho et al., 2000; Cooney et al., 2001). We show below that human-CD47 can inhibit

Supplemental Material can be found at: http://www.jcb.org/cgi/content/full/jcb.200708043/DC1
Fc-receptor mediated phagocytosis by human-derived macrophages and monocytes, but first we map some key components of the phagocytic synapse and identify species-specific differences in localization of SIRPα, phosphotyrosine, and nonmuscle myosin. A microbead system demonstrates the density and affinity dependence of CD47-mediated inhibition and identifies a threshold activity in the inhibitory signaling. These signals propagate downstream and parse cytoskeletal pathways, ultimately implicating a novel phosphotyrosine site on myosin that appears associated with myosin enrichment and contractile function in high efficiency internalization.

## Results

### Affinity, signaling, and cytoskeleton at the phagocytic synapse with and without CD47

Consistent with recent results showing species-specific CD47–SIRPα interactions (Subramanian et al., 2007), soluble human-SIRPα is found here to bind much less to sheep RBCs than to human RBCs (Fig. 1A). Both species of RBC are extensively used below in studies of phagocytosis, but in order to eliminate other potential differences between species we also developed avidin-coated microbeads that display recombinant, biotinylated human CD47. These beads demonstrate a moderate affinity and saturable interaction of CD47 for SIRPα and also establish effective blocking of CD47 (B6H12) with a F(ab')2 made from a monoclonal antibody that is known to inhibit SIRPα binding (Fig. 1B).

To assess the effect of CD47 on protein localization to the phagocytic synapse, we first imaged human-derived THP-1 macrophages incubated with IgG-opsonized human RBCs or sheep RBCs. Briefly, the RBCs were allowed to settle and bind macrophages at 4°C, and then 10 min after warming to 37°C, cells were fixed and immunostained. Imaging by differential interference contrast (DIC) microscopy allowed an unbiased identification of phagocyte–RBC contacts (Fig. 2A). Subsequent immunofluorescence showed that human-RBCs in contact with the human macrophages stimulated accumulation of SIRPα at the synapse. Such localization was lacking both with sheep RBC and after blocking human RBC with a F(ab')2; of the anti-CD47 (Fig. 1B inset; Fig. S2, A–C); the removal of the Fc domain ensured no activating signal. Quantitative intensity analyses of randomly chosen synapses showed a fivefold enrichment of SIRPα with human–RBC contacts when compared with either sheep RBCs or CD47-blocked human RBCs (Fig. 2B, top).

Tyrosine phosphorylation is known to be strongly enhanced when macrophages phagocytose IgG-opsonized targets. Compared with SIRPα, phosphotyrosine showed the opposite trends after immunostaining: synapses with human RBCs showed only a small increase of phosphotyrosine above cytoplasmic levels and at a level that was 3.2-fold lower than sheep RBCs or CD47-blocked human RBCs (Fig. 2B, bottom). IgG interactions with Fc receptors (FcγRs) are well known to initiate Src family phosphorylation of the immunoreceptor tyrosine-based activating motif (ITAM) that then propagate a phosphorylation cascade (Huang et al., 1992; Ghazizadeh et al., 1994; Greenberg et al., 1994). In contacts with human RBCs, however, the CD47-induced accumulation of SIRPα at the synapse is expected to phospho-activate SIRPα’s immunoreceptor tyrosine-based inhibiting motif (ITIM) (Kharitonenkov et al., 1997) with subsequent recruitment of inhibitory tyrosine phosphatases, particularly SHP-1 (Tsuda et al., 1998; Veillette et al., 1998; Vernon-Wilson et al., 2000; Kant et al., 2002). Phosphatase activation is consistent with the relatively large and dominant decrease found here for phosphotyrosine.

Pseudopod extension and phagocytic cup formation around the target predictably requires extensive remodeling of the actin cytoskeleton. Surprisingly, SIRPα–CD47 interactions did not exert any statistically significant effect (P = 0.3) on
Recruitment of NMM IIA to the synapse formed with sheep RBCs was seen to take minutes and to persist typically for at least 15 min (Fig. 3 A). Macrophages were stably transfected with GFP-NMM IIA, and real-time fluorescence imaging was used to follow the contact and engulfment process. Both sheep RBCs and human RBCs adhered and engaged the macrophages, initiating the formation of a phagocytic cup, but NMM IIA did not enrich adjacent to the human RBCs (Fig. 3, B and C). RBC contacts per macrophage additionally showed no major F-actin localization to the synapse (Fig. 2, C and D, top). Previous reports have also suggested a role for nonmuscle myosin in phagocytosis (Mansfield et al., 2000), and immunofluorescence for the dominant myosin isoform, nonmuscle myosin IIA (NMM IIA), shows clear recruitment to both sheep RBCs and CD47-blocked human RBCs. However, localization of myosin to the synapse with human RBCs is minimal, with a 3.8-fold decrease relative to sheep RBCs and CD47-blocked human RBCs (Fig. 2, C and D, bottom).
Figure 3. **Time-lapse image of phagocytic synapse formation.** Human-derived THP-1 macrophages stably transfected with GFP-NMM IIA were incubated at 37°C with IgG-opsonized sheep RBCs (A) or human-RBCs (B) for 10 min after an initial 4°C incubation. Time-lapse images in DIC and fluorescence microscopy were taken upon identification of a target cell adhered to the macrophage. Arrows indicate the site of target cell contact with magnified images of the phagocytic synapse. Bars, 10 μm. (C) Phagocytic synapses fixed after 10 min. Cells were immunostained for total NMM IIA and protein localization to the phagocytic synapse was quantified for randomly chosen GFP+ cells by normalization to cytoplasmic intensity of 1.0 (n = 5, ± SD).
Myosin recruitment to the phagocytic synapse is inhibited by blebbistatin
Nonmuscle myosin has long been known to be expressed in macrophages (Stendahl et al., 1980), and although recent studies have implicated myosin in FcyR-mediated phagocytosis (Swanson et al., 1999; Titus, 1999; Mansfield et al., 2000; Diakonova et al., 2002) through inhibition studies with 2,3-butanedione monoxime (BDM), this is a relatively nonspecific drug compared with blebbistatin (Ostap, 2002; Limouze et al., 2004). This recently synthesized, membrane-permeable drug inhibits the ATPase activity of myosin types II and VI (Straight et al., 2003; Kovacs et al., 2004). Consistent with a significant role for myosin in phagocytosis, we found that incubation of macrophages with blebbistatin, inhibited enrichment of NMM IIA at synaptic contacts with sheep RBCs targets (Fig. 4, A and B). CD47 on human RBCs inhibited phospho-paxillin localization to the synapse with sheep RBCs in the presence or absence of the myosin inhibitor as well as for CD47-blocked human RBCs.

Paxillin is also known from past studies to accumulate at the phagocytic synapse (Greenberg et al., 1994) and to be phosphorylated (Hall, 1998). Phospho-paxillin Y118 localized to the synapse with sheep RBCs and this persisted with myosin inhibition by blebbistatin (Fig. 4 A). In contrast, human RBCs inhibited phospho-paxillin localization through the effects of CD47. Note the double-labeling studies here consistently show statistically similar levels of F-actin localization, whereas phospho-paxillin varies. Phosphatase activity initiated by CD47–SIRPα interaction likely has multiple downstream targets that directly or indirectly include phospho-paxillin and—as elaborated below—a novel phosphorylation site in myosin.

Phagocytosis increases upon blocking CD47 on human RBCs
Past results from mouse cells certainly suggest that CD47 inhibits phagocytosis, but evidence has been lacking with human phagocytes interacting with any type of target cell or particle. CD47-blocking of human RBCs with the F(ab’)2 was shown above to perturb the phagocytic synapse, which made the synapse look like that of sheep RBCs interacting with macrophages (Fig. 2). Human CD47’s interaction with SIRPα was certainly blocked (Fig. 1 B inset; Fig. S2, A–C, available at http://www.jcb.org/cgi/content/full/jcb.200708043/DC1), and so we expected that CD47-blocked human RBCs would be engulfed by human phagocytes at a greater frequency than untreated cells.

Phagocytosis by the human-derived THP-1 macrophages of IgG-opsonized RBCs was studied by imaging in DIC microscopy: after 45 min at 37°C the non-engulfed RBCs were hypotonically lysed (Fig. 5 A) and the number of ingested RBCs per macrophage was counted. Phagocytosis by fresh human peripheral blood monocytes used fluorescence to visualize RBCs, all of which had membranes prelabeled with PKH26 (red), and then the non-engulfed RBCs were labeled with a fluorescein-labeled (green) antibody against the Fc of the IgG-opsonin (Fig. 5 B). For either type of human phagocyte, human RBCs were internalized at a small fraction of the frequency of sheep RBCs (Fig. 5, C and D), but a small number of human RBCs was always engulfed for either type of phagocyte. All dependencies on IgG-opsonization fit well to a saturation binding process, consistent with specific activation of the FcR phagocytosis pathway. Most important, blocking of CD47 on human RBCs produced major increases in phagocytosis (blue arrows) with two- to threefold more engulfment that approached or slightly exceeded the phagocytosis of sheep RBCs.
Nonmuscle myosin IIA contributes to FcγR-mediated phagocytosis, but is not essential

In light of the myosin differences seen by fluorescence at the various synapses (Figs. 2–4), we also examined the effect of blebbistatin-inhibited myosin on phagocytosis. Engulfment of sheep RBCs by THP-1 macrophages was found to be inhibited by blebbistatin to an extent similar to that of CD47 on human RBCs (Fig. 5 C, inset bar graph). Moreover, a dose–response gave a $K_{\text{Blebb}} = 5$ μM for blebbistatin inhibition of phagocytosis (Fig. S3, available at http://www.jcb.org/cgi/content/full/jcb.200708043/DC1), which is in excellent agreement with the $K_{\text{ATPase}}$ for inhibition of NMM IIA’s ATPase by blebbistatin (Limouze et al., 2004).

To directly confirm the implied correlation between synaptic myosin and the extent of phagocytosis, fluorescence imaging was repeated on the enrichment of myosin and F-actin at the synapse for both human and sheep RBCs treated with or without blebbistatin. As demonstrated in Figs. 4 and 5, when synaptic NMM IIA is at or near background cytoplasmic levels, phagocytosis is low but nonzero. The comparatively rapid enrichment of F-actin at the phagocytic synapse described here and by others (Strzelecka et al., 1997) proved here to be dependent primarily on the initial synaptic contact rather than the ultimate level of phagocytosis. The result highlights a basal level of pseudopod extension and engulfment that is independent of myosin.

CD47 on microbeads is sufficient to inhibit phagocytosis

RBCs are common targets in phagocytosis, but RBC membranes are complex and blocking results above could have other interpretations. We therefore tested whether CD47 alone on synthetic microbeads could also inhibit phagocytosis. The extracellular immunoglobulin-like domains of human CD47 (hCD47) was recombinantly expressed with a spacer domain...
In the absence of functional CD47, phagocytosis is proportional to myosin

Blebbistatin’s inhibition of phagocytosis functionally implicated NMM II because it inhibits all three isoforms (A–C), and so to more directly assess the function of the IIA isoform, the effects of both knockdown and overexpression were functionally studied. Transfection of GFP-NMM IIA added ~50% more total IIA isoform, based on an immunoblot comparison to wild type (Fig. S4, available at http://www.jcb.org/cgi/content/full/jcb.200708043/DC1), and the overexpression also resulted in ~50% more phagocytosis of sheep RBCs (Fig. 7). Similarly, knockdown of NMM IIA with Lentivirus gave ~50% less myosin, and this resulted in nearly 50% less phagocytosis. Transfection of GFP-NMM IIA into the knockout cells (driven by a powerful CMV promoter) returned expression to near wild-type levels, and also recovered full phagocytic function. Additionally, the GFP-NMM IIA was inhibited ~50% by 5 μM blebbistatin (Fig. 7, inset), which again indicates that this chimeric construct has normal, myosin-like activity. A linear fit (R² = 0.99) through all of these phagocytosis results for sheep RBCs not only emphasizes the proportional role for myosin in efficient engulfment, but also yields a nonzero intercept at “zero” myosin activity that is nearly the same as the low rate for “actin only” engulfment of human RBCs (~0.2–0.3 RBC per macrophage). These results thus suggest that human CD47 parses pathways and primarily signals inhibition of NMM IIA’s contribution to efficient phagocytosis.

**CD47 signals through SIRPα and ultimately to myosin**

SIRPα localization to the phagocytic synapse with targets presenting CD47 (Fig. 2, A and B) is consistent with ligand-receptor interactions as well as SIRPα ITIM activation and subsequent SHP-1 phosphatase induction (Brown and Frazier, 2001; Latour et al., 2001). Immunoprecipitation of SIRPα followed by Western blot analysis of phosphotyrosine showed a clear and saturable signaling difference with hCD47 (Fig. 8 A). The term denoted as $K_p$ is defined as the normalization of CD47 densities to the phagocytosis inhibition constant for human CD47 (Fig. 6 C) and normalization of phosphotyrosine levels to SIRPα intensities gave an effective signaling constant $K_p$ that approximated the $K_i$.

CD47’s activation of SIRPα (about twofold here) and downstream phosphatase activity is certainly known but downstream phospho-targets have not yet been identified. Studies of SIRPα-knockout macrophages engulfing IgG-opsonized mouse RBCs (Okazawa et al., 2005) suggested no major perturbation of either phospho-FcyR, downstream phospho-Syk or phospho-Cbl, although it had been reported that SHP-1 can dephosphorylate Cbl and thereby moderate Rac activity in FcγR-mediated phagocytosis (Kant et al., 2002). Through immunoblotting, changes in the phosphorylation state of these activator/effector proteins ranged from 1.1- (FcγR) to 1.5-fold (Cbl) (Fig. 8 B). Considerably larger down-regulation of phosphotyrosine by CD47 at the synapse (3.2-fold) was clear in our imaging studies (Fig. 2 B) as was CD47-induced delocalization of myosin and phospho-paxillin (3.8- and 3.0-fold, respectively; Fig. 2 D, Fig. 4 B).
in the present context of CD47 function. Anti-phosphotyrosine immunoblots were made from whole cell lysates of THP-1 cells phagocytosing opsonized human or sheep RBCs (Fig. 8C). Three band regions (a, b, c) showed the greatest differences in phosphotyrosine levels compared with THP-1 cells alone. Bands a–c each

The only report of phosphotyrosine regulation of NMM IIA suggests it to be a direct target of SHP-1 in B-cells (Baba et al., 2003), with the speculated site of interaction appearing in the N-terminal head domain. Phosphotyrosine regulation of NMM IIA therefore seemed an intriguing possibility to pursue

Figure 6. Human-CD47 is sufficient to inhibit phagocytosis. (A) Streptavidin beads coated with both anti-streptavidin IgG as the opsonin and biotinylated human CD47. Phagocytosis of beads (red) by THP-1 cells was assessed in DIC and fluorescence microscopy with non-ingested beads (green) visible with rabbit anti-streptavidin plus a second, goat anti–rabbit antibody. Bars, 10 μm. Beads ± hCD47 (at normal RBC density) were engulfed by THP-1 cells or human monocytes (B), demonstrating inhibition of phagocytosis by hCD47 unless the beads are blocked with anti-CD47. Inset bar graph compares to inhibited phagocytosis of uncoated beads after 10 min pretreatment of THP-1 cells with blebbistatin (50 μM). (C) Inhibition of phagocytosis depends on density of human CD47 on beads. All assays were conducted and analyzed as in Fig. 5. Phagocytosis inhibition occurs with an effective $K_i \approx 20$ molecules/μm², which considerably exceeds the relevant dissociation constant for a 10-nm gap of $K_{d,10 \text{ nm}} \approx 1$ molecule/(10 μm)² (Fig. 1B). This ratio of $\approx 1,000$ as well as the known cell surface densities imply that almost all SIRPα and CD47 that diffuse into the gap will bind and thus enrich in the synapse [see Fig. 2, A and B]. (D) Bar graph compares hCD47 inhibition of phagocytosis in peripheral blood monocytes.

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showed 3.3- to 1.6-fold less phosphotyrosine with human RBCs versus sheep RBCs (Fig. 8 C, plot), which was similar in range to the down-regulation of phosphotyrosine imaged at the phagocytic synapse (3.2-fold in Fig. 2 B). In fact, band “a” had a MW in the range of nonmuscle myosins in addition to a similarly high level of down-regulation.

To identify the major phosphotyrosine bands a–c, these bands were excised from Coomassie-stained polyacrylamide gels, digested in-gel by trypsin, and the peptide fragments were characterized by liquid chromatography-mass spectrometry/mass spectrometry (LC-MS/MS) (Fig. 8 C). Within band “a”, 45 tryptic peptides matched sequences of NMM IIA (23% coverage), including one fragment from myosin’s self-assembling tail region with a novel phosphorylation at Y1805 (Fig. 8 C, bottom). Additional cytoskeletal proteins that were detected in the macrophages and might be targeted in the CD47-SIRPα phosphotyrosine pathway include Talin-1 (Turner et al., 1989; Greenberg et al., 1990), mDIA1 (Meng et al., 2004), and α-actinin (Crowley and Horwitz, 1995; Izaguirre et al., 1999). Capabilities of MS to identify phosphotyrosine are currently limited, as here, by low sequence coverage and sub-stoichiometric phosphorylation (McLachlin and Chait, 2001). However, the one solid lead from MS to identify phosphotyrosine is currently limited, as here, by low sequence coverage and sub-stoichiometric phosphorylation (McLachlin and Chait, 2001). However, the one solid lead from MS to identify phosphotyrosine in NMM IIA that is ultimately regulated by CD47 interactions (Fig. 8 D).

**Phosphotyrosine activation of NMM IIA in phagocytosis**

In light of the proportionality between phagocytosis and myosin expression (Fig. 7), several mutants of the GFP-NMM IIA construct were made and studied in their effects on phagocytosis of sheep RBCs and localization to the synapse. One mutant truncated the C terminus by 170 residues and had been found previously to compromise myosin filament assembly and function (Wei and Adelstein, 2000). Two Y→F point mutations were made at the putative phospho-sites in the head (Y277) and tail (Y1805). Although the wild-type GFP-NMM IIA construct showed gain-of-function phagocytosis that was statistically distinct from untransfected wild type (P < 0.03), all three mutants appeared the same as wild type (Fig. 9 A). Similar results with even greater statistical significance (P < 0.01) were found upon transfection of these constructs into the knockdown macrophages (Fig. 9 A).

Consistent with the lack of a functional contribution to phagocytosis, the GFP mutants exhibited no significant localization to the synapse (Fig. 9 B). Conveniently, the transfections of these GFP constructs into wild-type macrophages also allowed immunostaining for total NMM IIA. Quantitative ratio imaging (see Fig. 2) showed that the endogenous myosin accumulated at phagocytic synapses, whereas the mutant constructs did not. These results contrast with the clear localization of the wild-type GFP-NMM IIA to the phagocytic synapse (Fig. 3 A) and point to the role for phospho-regulation of myosin-II in phagocytosis.

**Discussion**

Human CD47 has been shown here to communicate marker of self signals to human primary and immortalized monocytes/macrophages, and the mechanism for species-specific inhibition of phagocytosis shows that CD47 parses pathways by selectively down-regulating the cytoskeleton’s contractile contributions to engulfment. Our results were obtained with IgG-opsonized RBCs from various species that are common in phagocytosis studies (e.g., sheep RBCs) and also with synthetic beads bearing recombinant CD47. The signaling and remodeling processes are highlighted in Fig. 10. IgG activation of the Fcγ receptor on the phagocytes (Cambier, 1995) is known to induce actin cytoskeleton assembly (Araki et al., 1996;
CD47 in mouse has been described as a marker of self that can inhibit phagocytosis by mouse macrophages (Oldenborg et al., 2000; Gardai et al., 2005). Blocking of CD47 on human RBCs with an anti-CD47 F(ab')2 increased uptake by both human macrophages and monocytes. Given the low densities of Crowley et al., 1997; Caron and Hall, 1998; Lowry et al., 1998; May and Machesky, 2001), and this F-actin assembly was proven here to be independent of CD47's signaling pathway(s) (Fig. 2). In the absence of CD47 on the target, paxillin assembles at the phagocytic synapse, consistent with past results for IgG-opsonized sheep RBCs (Greenberg et al., 1994; Coppolino et al., 2001), and so does NMM IIA. By contrast, human RBCs with functional CD47 show strong localization of the counter-receptor SIRPα to the synapse (fivefold), and also show strongly reduced levels of both phosphorysine and NMM IIA (3.2- and 3.8-fold, respectively; Fig. 2). We find that human CD47 and the myosin inhibitor blebbistatin block synaptic localization of NMM IIA and phagocytosis to the same extent (Fig. 4 B, Fig. 5 C). The proportionality between NMM IIA levels and efficient phagocytosis is also clear from knockdown and overexpression studies (Fig. 7). Although CD47 signals locally, we conclude that it ultimately has a similar, but downstream effect in inhibiting the major myosin isoform's contractile contributions to engulfment.

Figure 8. Species-specific signaling through SIRPα and ultimately to cytoskeletal proteins. (A) hCD47 was bound at varying densities to opsonized beads and phagocytosed by THP-1 macrophages. From macrophage lysates, SIRPα was immunoprecipitated and immunoblotted [inset] for quantitation of phosphorysine and total SIRPα for normalization. Fits of the data gave an effective signaling constant $K_S$ for each species that depends on the CD47 density; all densities are scaled by hCD47’s inhibitory constant ($K_i$) as determined in Fig. 6 at the same opsonization. (B) Phosphotyrosine decreases in Cbl, Syk, and FcγR within THP-1 during phagocytosis of IgG-opsonized human RBCs normalized to sheep RBCs. Whole-cell lysates were immunoblotted and densitometry was used to quantify suitable MW bands (also see Fig. S5, available at http://www.jcb.org/cgi/content/full/jcb.200708043/DC1). (C) Major phosphotyrosine differences in THP-1 during phagocytosis of IgG-opsonized human RBCs versus sheep RBCs. Whole-cell lysates were immunoblotted and densitometry was used to identify bands (a–c) that showed the largest relative differences in intensity for sheep or human RBC targets compared with THP-1 cells. For bands a–c, the plot shows the intensity obtained with phagocytosis of human RBCs relative to sheep RBCs (triplicate experiments ± SEM). The list from MS analyses of bands a–c indicates top candidate phospho-proteins or else a directly detected sequence (for myosin). (D) Immunoprecipitation of NMM IIA from lysates followed by immunoblot for pTyr, confirming the major decrease in phosphorysine NMM IIA when THP-1 phagocytose human RBCs compared with sheep RBCs.
CD47 on cells, this ubiquitous membrane protein seems less likely to mediate cell adhesion than function as an extremely potent signaling receptor, functional at densities of a just a few dozen molecules per μm² (Fig. 6 C). Major co-deficiencies of CD47 on human cells (Mouro-Chanteloup et al., 2003) can therefore be tolerated without compromising marker of self function, and because CD47-null humans have yet to be reported, our results suggest an important role for CD47 in the innate immunity of humans.

Regardless of species, normal macrophages should be expected to engulf “self” cells at a considerably lower frequency than typical targets—apoptotic and foreign cells or particles. The decision of a macrophage to phagocytose is in part made by the extent of target opsonization, which activates assembly of the F-actin cytoskeleton at the phagocytic synapse; however, Ig is high in bodily fluids and almost certainly binds or adsorbs at some level to all cells (Turrini et al., 1993), and so activating signals seem unavoidable. CD47 signaling probably occurs in parallel because IgG binding to FcγR (Duchemin et al., 1994) promotes intimate adhesion and thereby promotes hCD47 interactions with SIRPα within the synapse (Fig. 2 A). It will be interesting to visualize the effects of SIRPα polymorphisms in human (as well as mouse) that have been recently described (Takenaka et al., 2007).

Synaptic activation of SHP-1 phosphatase by CD47-SIRPα has been clear, but downstream targets and mechanisms in phagocytosis inhibition have remained unknown. Deactivation could have been restricted to membrane receptors, particularly FcγR, but our phosphorylation data indicates otherwise (Fig. 8). Conceivably, F-actin could have been found to be the primary target of CD47’s signaling to macrophages because polymerization alone is sufficient to drive pseudopod extension and particle internalization (May and Machesky, 2001). We find, however, that F-actin establishes a basal level of phagocytosis regardless of CD47. From our sheep versus human comparison of phosphotyrosine and MS analysis of dephosphorylated proteins, NMM IIA emerged as the major down-regulated species (3.3-fold) (Fig. 8 C). This regulation of the myosin heavy chain differs from the usual regulation through the light chain (Conti and Adelstein, 2008). Talin, α-actinin, and the actin nucleator mDia1 also appeared regulated (to a lesser extent), and all have been implicated previously in phagocytosis (respectively: [Turner et al., 1989; Greenberg et al., 1990]; [Crowley and Horwitz, 1995; Izaguirre et al., 1999]; [Meng et al., 2004; Brandt et al., 2007])). Although phospho-paxillin was down-regulated in our imaging studies, it was not prominent in our immunoblots and was unaffected by blebbistatin, which otherwise mimicked CD47’s inhibitory effects (Figs. 5 C and 6 B, insets). Nonetheless, focal adhesion-type cytoskeletal assemblies of paxillin, talin (Niewohner et al., 1997), or even α-actinin (Sampath et al., 1998) could also contribute to CD47 function, as these phospho-proteins are also implicated in phagocytosis.
FcγR-induced tyrosine phosphorylation leads to the downstream activation of Rac/Cdc42 (Olazabal et al., 2002), which then signals the recruitment and assembly of F-actin and a focal adhesion type of structure. Syk is not required because macrophages from Syk-deficient mice will extend actin-rich pseudopods around target particles even though FcγR-mediated phagocytosis is ultimately defective (Crowley et al., 1997).

In comparison, SIRPα-null macrophages engulf IgG-opsonized mouse RBCs more readily than wild-type macrophages (Okazawa et al., 2005) and show no major differences in either phospho-FcγR, or downstream phospho-Syk or phospho-Cbl. Likewise, in our phagocytosis studies of human versus sheep RBCs, no significant difference in phospho-FcγR was detected, and, relative to phospho-SIRPα and cytoskeletal changes (Fig. 8 C), only small decreases in phospho-Syk or phospho-Cbl were detected (Fig. 8 B: 1.3- and 1.5-fold, respectively). Such differences upstream could in principle reflect a classical amplification of signaling, but our finding of slight decreases in phospho-Cbl due to CD47 is perhaps most consistent with the activity of SHP-1 (Kant et al., 2002). The small effect would suggest perturbations to actin assembly; indeed, with CD47–SIRPα interactions (Fig. 2 D and Fig. 4 B) all suggest a slight tendency toward decreased actin polymerization (based on staining with rhodamine-phalloidin), but the effects on myosin are statistically certain in comparison.

Past studies of antibody-opsonized beads as targets have shown that FcγR-mediated phagocytosis is independent of Rho or ROCK activity while dependent on myosin-II for particle internalization but not for actin cup formation (Olazabal et al., 2002). With CD47 on the target, activation of the SIRPα→SHP-1 phosphatase pathway (Vernon-Wilson et al., 2000) could in principle block engulfment by regulating various kinase or phosphatase activities (such as Cbl) or perhaps even by regulating signaling to NMM IIA. Our immunoblot analysis followed by LC-MS/MS indeed identified direct phosphorylation of Y1805 in this myosin’s self-assembling tail as an ultimate target for CD47-SIRPα interaction. Mutation of this site disables this major myosin from contributing to phagocytosis (Fig. 9). Moreover, initial studies of NMM IIA in primary human mesenchymal stem cells (that are also marrow derived) identify the same site as being phosphorylated (unpublished data), and since we have shown NMM’s contribute in key ways to differentiation of these adult stem cells (Engler et al., 2006), a general mechanism of regulation is perhaps emerging. Detailed study of purified point mutants would certainly help clarify broader effects on myosin assembly/activity, but regulatory roles for specific residues in the myosin tail region already appear reasonable because: (1) nearby sequences (aa 1666–1728 and 1914–1961) are involved in isoform-specific assembly of functional myosin filaments (Sato et al., 2007), and (2) disease-causing mutations occur nearby with the charge changes E1841K and R1933X (Heath et al., 2001). The head and tail of at least some myosin-Is can interact to form a compact, diffusible monomer and regulatory interactions then open up this conformation for polymerization into filaments (Kudryashov et al., 2002); pY might regulate such a conformational switch. The only other report of a phosphotyrosine in NMM IIA suggested pY277 is a direct target for SHP-1 (Baba et al., 2003).

Both phosphotyrosine sites identified here in human myosin are found in mouse, for which CD47’s “self” signaling was first described (Oldenborg et al., 2000). Both phospho-sites in NMM IIA are found in other mammalian species as well as in other proteins (i.e., eight myosin heavy chain proteins share the head motif and one non-myosin, coiled-coil protein has the tail motif). In addition, the MS data here indicated that NMM IIA dominates the B and C myosin isoforms by at least 10-fold (Table S1, available at http://www.jcb.org/cgi/content/full/jcb.200708043/DC1), and mouse knockouts of the B and C isoforms do not exhibit immune defects whereas the NMM IIA-null mouse is embryonic lethal as an undifferentiated mass of cells (Conti et al., 2004). The results here with human cells have implicated NMM IIA for good reason it seems and thus suggest that recognition of “self” entails blocking engulfment by turning off this ubiquitous and essential motor.

Materials and methods

**Chemicals**

Dulbecco’s phosphate-buffered saline (DPBS) without Ca²⁺ or Mg²⁺ (Invitrogen) was supplemented with either 1% BSA or 1% BSA + 0.05% Tween 20 (Sigma-Aldrich). PKH26 (Sigma-Aldrich) hydrophobic dye was used for...
red cell labeling. TBS (Tris-buffered saline) and TTBS (TBS with Tween 20) were used in Western blotting.

Antibodies
The fluorescein-labeled antibody B6H12-FITC was used against human CD47 (BD Biosciences). Opossonizing antibodies against human and sheep RBCs included intact anti-human RBCs, and anti-rabbit–sheep RBCs (Sigma-Aldrich); these were used as IgG opsonin in phagocytosis assays. Antibodies against NMM IIA, actin was obtained from Sigma-Aldrich. Antibodies against streptavidin coated polystyrene beads (Spherotech) included rabbit anti-streptavidin (Sigma-Aldrich) and rabbit anti-streptavidin conjugated with FITC (Rockland Immunocoulomicals) used also as IgG opsonin in phagocytosis assays. Secondary antibodies used for detecting opsonin-labeled beads included goat anti-rabbit–FITC or goat anti-rabbit F(ab')2, R-PE (Sigma-Aldrich). Secondary antibodies used for detecting SRPs binding included anti-GST Alexa 488 (Invitrogen). Antibodies against Syk (SYK-KO) were purchased from AbCam, and Phospho-Syk (Tyr525/526), Phospho c-Cbl (Tyr527), and c-Cbl (2747) were purchased from Cell Signaling Technology.

Plasmid construction
The plasmid vector containing a cytomegalovirus promoter in pEGFP-C3 containing the DNA fragment encoding aa 1–1961 from NMHC II A were obtained from Addgene. The original construct was prepared as described (Wai and Adelstein, 2000). To generate pCMVGST-ΔC170 encoding aa 1–1701, the original NMHC II A construct from Addgene was digested with AlliI–Spel and the purified larger fragment was rendered blunt and self-ligated. pCMVGST-NMHC II A point mutants Y277F and Y1805F were modified by using primers 5'-CCGTCCAGTCATCCTTTCTGCTCTTG–3' and 5'-TCAGTCCGAACGGGCTTCATTCTGC-3', respectively; done using the QuikChange Site-Directed Mutagenesis kit (Stratagene). For convenience pCMVGST-NMHC II A, pCMVGFP-ΔC170, and pCMVGFP-NMHC II A point mutants Y277F and Y1805F will be referred to as GFP-NMHC II A, GFP-ΔC170, GFP-Y277F, and GFP-Y1805F, respectively. All constructs were confirmed by sequencing.

Lentiviral knockdown of NMM IIA in THP-1 cells
SRNA lentiviral supernatants to NMM IIA purchased from Sigma-Aldrich were targeting MYH9 (TRC#. TRCN00000029467) encoding NMHC IIA. Further details of these clones are available from the Sigma-Aldrich website. Target THP-1 cells were infected with lentiviral supernatants at a multiplicity of infection (MOI) of 10, and then cells with integrated viral sequence were selected using puromycin (Sigma-Aldrich). The degree of NMM IIA silencing was regularly monitored by both immunofluorescence and Western blotting. Control cell cultures were generated with control lentiviruses in parallel.

Phagocytosis assay
Peripheral blood monocytes from human donors were obtained from Cell Signaling Technology. Differentiation of THP-1 cells was achieved in 100 ng/ml phorbol myristate acetate (PMA) for 3 days. Differentiated THP-1 cells were cultured in DMEM, MEM (Thermo Fisher Scientific), or RPMI 1640 media (Invitrogen) at 37 °C. These cells were used in phagocytosis assays. CD47 was attached to streptavidin-coated beads at room temperature for 5 min. Beads were washed and blocked 3x in PBS plus 0.4% BSA. Biotinylated human CD47 was attached to streptavidin-coated beads at room temperature for 30 min, washed 3x, and resuspended in PBS plus 0.4% BSA.

Preparation and quantification of CD47 density on polystyrene beads
Streptavidin-coated polystyrene beads of 2.1 μm diameter (Spherotech) were washed and blocked 3x in PBS plus 0.4% BSA. Biotinylated human CD47 was attached to streptavidin-coated beads at room temperature for 30 min, washed 3x, and resuspended in PBS plus 0.4% BSA.

Production of recombinant human CD47
Plasmids encoding the extracellular domain of human CD47 were PCR amplified, digested with XbaI and SalI (New England Biolabs, Inc.), and ligated to similarly digested vector, pEF-BO5-XB (Vernon-Wilson et al., 2000), which results in an in-frame fusion of CD4d3 + 4-biotin at the C terminus of the extracellular domain of CD47. The above vector containing the extracellular domain of CD47 was transfected into CHO (- K1) cells using Lipofectamine 2000 (Invitrogen). Secreced CD47-CD4d3 + 4 was concentrated using a 10 K MWCO Amicon (Millipore) and biotinylated at the C terminus using a biotin–protein ligase (AviDity, LLC) and dialyzed against PBS (Invitrogen). The protein was affinity purified using a monoclonal avidin (Promega) and dialyzed against PBS (Invitrogen).

Preparation and quantification of CD47 density on polystyrene beads
Streptavidin-coated polystyrene beads of 2.1 μm diameter (Spherotech) were washed and blocked 3x in PBS plus 0.4% BSA. Biotinylated human CD47 was attached to streptavidin-coated beads at room temperature for 30 min, washed 3x, and resuspended in PBS plus 0.4% BSA.

Binding isotherm for soluble hSRPα for CD47 coated beads
Human CD47 was attached to the identical density as described above. The binding isotherm of soluble hSRPα was performed for different species over a range of concentrations using flow cytometry. Forward scatter, side scatter, and fluorescence (FL1, FL2, FL3, FL4 channels in logarithmic mode) were measured for a least 10⁴ events using a FACScan or FACSCalibur (BD Immunocytometry Systems). Data points from flow cytometry were plotted and fitted: the Ki values for human CD47-coated beads was 0.12 μ M and 6.9 μ M, respectively (Fig. 1 B).

Phagocytosis assay
For phagocytosis assays, macrophages were plated in 4-cm² laboratory-Tek II chambered coverslips (Nalge Nunc International) on 10⁵ cells/cm². Streptavidin polystyrene beads or RBCs were added to macrophages at a ratio of 20:1 and allowed to incubate at 37 °C for 45 min. Non-phagocytosed beads or RBCs were washed with PBS. Cells were fixed with 5% formaldehyde (Thermo Fischer Scientific) for 5 min, followed by immediate replacement with PBS. For differentiation of non-internalized beads, beads were labeled with a primary antibody, rabbit anti-streptavidin (Sigma-Aldrich) at 1:1,000 in PBS for 20 min at 25 °C. A second antibody, anti-rabbit R-PE (Sigma-Aldrich) was added at 1:1,000 in PBS to the cells and incubated for an additional 20 min at 25 °C. Cells were then washed with PBS/0.4% BSA and then quantified by light and fluorescent microscopy. At least 200 cells were scored per well and experiments were repeated at least three times. Assays using RBCs as targets, lysis of unengested RBCs was performed by adding deionized H₂O for 60 s, followed by immediate replacement with PBS/0.4% BSA and fixing with 5% formaldehyde for 5 min.

For stimulated phagocytosis assays, beads with or without CD47 were incubated with rabbit anti-streptavidin serum and for sheep and human RBCs with rabbit anti-sheep RBCs and rabbit anti-human RBCs, respectively, as the opsonin. Beads or RBCs were opsonized at the respective concentration for 30 min at room temperature. In some experiments, human CD47 was blocked with a Fab ', monoclonal antibody against CD47. The Fab' fragments of mAb B6H12 were removed by Ficin digestion and separated by affinity purification by protein A chromatography after 10 min of the addition of opsonin. Opsonized beads and RBC were washed 2x and resuspended in 50 μl of PBS/0.4% BSA. Phagocytosed beads were washed with PBS and unengested RBC was lysed and unengested beads were labeled as described above.

For cytoskeletal involvement at the phagocytic synapse, sheep and human RBCs were opsonized with rabbit anti-sheep RBCs or rabbit anti-human RBCs, respectively. Opsonized RBCs were added to IP-1.
cells, PMA treated, and immediately placed at 4°C for 30 min to synchronize phagocytosis. The temperature of the cells was then immediately increased to 37°C for 10 min and then fixed with 5% formaldehyde for immunofluorescence. For studies involving blebbistatin (EMD Biosciences), macrophages were treated for 10 min at 4°C before temperature increase to 37°C or for 45 min and 37°C. Macrophages treated with DMSO were used to verify no solvent effects. Adhesion studies were done as above with incubations at 4°C for 30 min, and then the cultures were fixed and gently washed in PBS to eliminate non-adherent cells. RBCs per phagocyte were then counted.

Quantification of fluorescent intensity
Immunostaining was performed by permeabilization with 0.1% Triton X-100 in PBS for 20 min before blocking for 1 h with 5% BSA in PBS. Staining with primary antibodies was performed for 1 h at room temperature in PBS. After washing, samples were incubated with appropriate PE-conjugated secondary antibodies [1:1,000]. To label cytoskeletal proteins at the phagocytic synapse, primary antibodies prepared with Zenon Alexa 488 or Zenon Alexa 647 Fab Labeling Kits (Invitrogen) were labeled for 45 min at room temperature. Cells were immediately fixed using 5% formaldehyde for 20 min and washed with PBS. Samples were analyzed by differential interference contrast (DIC) and fluorescence microscopy.

Images were acquired with an inverted microscope [IX71; Olympus] with a 60x (oil, 1.4 NA) objective using a Cascade CCD camera [Photometrics]. Image acquisition was performed with Image Pro software (Media Cybernetics, Inc.). Time-lapse imaging was performed using a heated stage [Ibid Gmbh] at 37°C in normal growth medium. Intensity analysis of the phagocytic synapse was performed using Image with a 10 x 2 μm box. The synapse was aligned at the center of the 10μm length box and the peak intensity designated as the zero in the distance scale on the plots. Fluorescence intensity was normalized by setting the cytoplasmic signal to 1 as the base signal, and five randomly selected cells were averaged. Two-tailed t-tests were performed to assess the significance.

Immunoprecipitation and Western blotting
Human phagocytes, THP-1 cells (2 x 10^6) were cultured and differentiated in 6-well plates for 48 h after PMA differentiation. Human CD47 was at-

tered as the base signal, and five randomly selected cells were averaged. Two-
tailed t-tests were performed to assess the significance.

Cytoskeletal proteins during Fcgamma receptor-mediated phago-
cytosis controlled by different Rho GTPases.

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