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Multiple stiffening effects of nanoscale knobs on human red blood cells infected with *Plasmodium falciparum* malaria parasite

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During its asexual development within the red blood cell (RBC), *Plasmodium falciparum* (Pf), the most virulent human malaria parasite, exports proteins that modify the host RBC membrane. The attendant increase in cell stiffness and cytoadherence leads to sequestration of infected RBCs in microvasculature, which enables the parasite to evade the spleen, and leads to organ dysfunction in severe cases of malaria. Despite progress in understanding malaria pathogenesis, the molecular mechanisms responsible for the dramatic loss of deformability of Pf-infected RBCs have remained elusive. By recourse to a coarse-grained (CG) model that captures the molecular structures of Pf-infected RBC membrane, here we show that nanoscale surface protrusions, known as “knobs," introduce multiple stiffening mechanisms through composite strengthening, strain hardening, and knob density-dependent vertical coupling. On one hand, the knobs act as structural strengtheners for the spectrin network; on the other, the presence of knobs results in strain inhomogeneity in the spectrin network with elevated shear strain in the knob-free regions, which, given its strain-hardening property, effectively stiffens the network. From the trophozoite to the schizont stage that ensues within 24–48 h of parasite invasion into the RBC, the rise in the knob density results in the increased number of vertical constraints between the spectrin network and the lipid bilayer, which further stiffens the membrane. The shear moduli of Pf-infected RBCs predicted by the CG model at different stages of parasite maturation are in agreement with experimental results. In addition to providing a fundamental understanding of the stiffening mechanisms of Pf-infected RBCs, our simulation results suggest potential targets for antimalarial therapeutics.

*The most virulent human malaria parasite, *Plasmodium falciparum* (Pf), causes ~700,000 deaths each year (1, 2). Following entry into red blood cells (RBCs), the parasite matures through the ring (0–24 h), trophozoite (24–36 h), and schizont stages (40–48 h). This intraerythrocyte maturation is accompanied by striking changes in the surface topography and membrane architecture of the infected RBC (3–5). A notable modification is the formation of nanoscale protrusions, commonly known as knobs, at the RBC surface during the second half (24–48 h) of the asexual cycle. These protrusions mainly comprise the knob-associated histidine-rich protein (KAHRP) and the membrane-embedded cytoadherence protein, Pf-erythrocyte membrane protein 1 (PfEMP1). KAHRP binds to the fourth repeat unit of the spectrin α-chain, to ankyrin, to spectrin–actin–protein 4.1 complexes, and to the cytoplasmic domain of PfEMP1 (6–9). These attachments enhance the vertical coupling between the lipid bilayer and the spectrin network. Another striking modification in the Pf-infected RBC membrane is the reorganization of the cytoskeletal network caused by parasite-induced actin remodeling (10). As a result of these molecular-level modifications, the Pf-infected RBC exhibits markedly increased stiffness [the shear modulus increases on average from ~4–10 μN/m in normal/uninfected RBCs, to ~40 μN/m at the trophozoite stage, and to as high as 90 μN/m at the schizont stage (11–13)] and cytoadherence to the vascular endothelium, which enable sequestration from circulation in vasculature, and evasion from the surveillance mechanisms of the spleen. Although in vitro experimental studies have revealed roles of particular parasite-encoded proteins in remodeling the host RBC (14–22), the mechanism by which Pf-infected RBCs gain dramatically increased stiffness has remained unclear. Indeed, uncertainty remains as to whether the loss of deformability arises from the structural reorganization of the host membrane components or from the deposition of parasite proteins. That is, it is not clear whether the stiffening is due to remodeling of the spectrin network, or to the formation of the knobs, or both. As experimental studies alone have heretofore not been able to determine the molecular details, numerical modeling, combined with a variety of experimental observations and measurements, offers an alternative approach to reveal the underlying mechanisms.

We present here a coarse-grained (CG) molecular dynamics (MD) RBC membrane model to correlate structural modifications at the molecular ultrastructure level with the shear responses of the Pf-infected RBC membrane, focusing on the second half of the parasite’s intra-RBC asexual cycle (24–48 h), i.e., the

Significance

Our coarse-grained molecular dynamics (CGMD) simulations show that the deposition of nanoscale knobs, rather than spectrin network remodeling, is the primary cause of the dramatically increased stiffness of the *Plasmodium falciparum* (Pf)-infected red blood cell (RBC) membranes. Our analyses further reveal that the knobs stiffen the RBC membrane in a unique manner by simultaneously harnessing composite strengthening, strain hardening, and knob density-dependent vertical coupling effects. In addition to providing a fundamental understanding of the stiffening mechanism of Pf-infected RBCs, our simulation results suggest potential targets for antimalarial therapeutics.


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trophozoite and schizont stages. The CG model is computationally efficient, and able to capture the molecular structures of the RBC membrane in both normal and infected states. CGMD simulations reveal that spectrin network remodeling accounts for a relatively small change in shear modulus. Instead, the knobs stiffen the membrane by multiple mechanisms, including composite strengthening, strain hardening, and knob density-dependent vertical coupling. Our findings provide molecular-level understanding of the stiffening mechanisms operating in Pf-infected RBCs and shed light on the pathogenesis of falciparum malaria.

A Composite Model for Healthy and Pf-Infected RBC Membranes

The human RBC is one of the simplest cell types, comprising a lipid bilayer and an underlying spectrin network, coupled by integral membrane proteins. During its 120-d life span, the disk-shaped RBC (resting long diameter, \( \sim 8 \) \( \mu \)m) exhibits pronounced deformability and durability. It circulates approximately a million times in the human body, squeezing through narrow capillaries and the 1- to 2-\( \mu \)m interendothelial slits that separate the splenic cords and venous sinuses (23). Mechanically, the lipid bilayer of an uninfected RBC contributes to the out-of-plane bending resistance and helps maintain the membrane surface area, but cannot sustain in-plane static shear. The spectrin network is responsible for the shear modulus but possesses negligible bending rigidity. As such, many existing membrane models (24–33) reduce computational complexity by simulating only the lipid bilayer or the spectrin network, depending on the membrane deformation mode of interest. Such a simplification is well justified for the uninfected RBC membrane, but not for the Pf-infected RBC membrane, given its increased vertical coupling between the lipid bilayer and the spectrin network. Therefore, a composite RBC membrane model comprising both lipid bilayer and spectrin network with high computational efficiency is required.

Here, we formulate a composite CG RBC membrane model by integrating a one-agent–thick lipid bilayer model and a chain-of-bead spectrin network model via vertical connections that represent integral proteins (Fig. 1). Similar composite models have been developed, with variations in the lipid and spectrin representations (34–37). In the lipid bilayer model of a normal (uninfected) RBC membrane, a small patch of lipids in the lateral direction is coarse-grained into a single agent (Fig. 1A1). The agent carries not only translational but also orientational degrees of freedom. The interagent interaction is described by a simple pairwise potential that is dependent on both the interagent distance and orientation, which drives the self-assembly of the agents into a 2D fluid phase (38). This lipid bilayer model significantly improves the computational efficiency compared with the chain-of-bead lipid models, while faithfully capturing both the in-plane viscosity and out-of-plane bending rigidity of the lipid bilayer. In the spectrin network model (Fig. 1A2), spectrin tetramers and actin junctional complexes form a hexagonal spectrin network in the normal RBC membrane (39). Each spectrin tetramer is modeled as a chain of 39 beads (40) (corresponding to the 39 triple-helical segments in a spectrin tetramer (41)), connected by unbreakable springs. Each vertex in the hexagonal network is an actin junctional complex at which an actin oligomer is linked to the beads at the end of six spectrin tetramers. The association between actin oligomers and spectrin tetramers is breakable and represented by a Lennard–Jones-type potential (40). To couple the lipid bilayer and the spectrin network, actin junctional complexes (representing protein 4.1–glycophorin C linkages) and the middle beads of spectrin tetramers (representing spectrin–ankyrin–band 3 linkages) are bonded to their closest lipid agents (Fig. 1D1). Further details of the models are provided in SI Text.

Two main modifications need to be taken into account in the model to simulate the Pf-infected RBC membrane: remodeling of the spectrin network due to the loss of actin oligomers, and the formation of the knobs and associated increase in vertical linkages to the lipid bilayer. Cryoelectron tomography experiments suggest substantive reorganization of the RBC cytoskeleton in the infected RBCs, with convergence on a reduced number of junction points, resulting in reorganization of the spectrin network. To examine the likely consequences of actin mining, we examined the effect of having the same total number of spectrin tetramers converging on a reduced number of junction points, resulting in an enhanced spectrin network (Fig. 1B2). We also considered the effect of removing up to 20% of the junctional complexes, resulting in holes in the spectrin network (Fig. 1C2). Although other possibilities exist, these represent the extreme scenarios that give rise to the upper and lower bounds of the shear modulus, respectively. In the enhanced spectrin network, the network still maintains a perfect hexagonal arrangement. However, the distance between neighboring actin complexes (hence the end-to-end distance of the spectrin tetramers) is increased by 10% to keep the covered area unchanged from the normal network. Meanwhile, to keep the total number of spectrin heterodimers unchanged, 20% of spectrin tetramers are replaced by spectrin octamers. In the deficient spectrin network, some spectrin tetramers are disconnected in accordance with the loss of actin oligomers. According to a recent atomic force microscopy study of the RBC cytoskeleton (43), the enhanced spectrin network resembles the cytoskeleton of the Pf-infected RBCs at the trophozoite stage (Fig. 1B2), whereas the deficient spectrin network resembles that of the schizont stage (Fig. 1C2).

Another critical modification in the RBC membrane that needs to be incorporated into the model is the presence of the knobs. As the knob is enriched with proteins (KAHRP and PIEMP1), it can be regarded as a stiff nanosized domain with one end linked to the spectrin network and the other immersed in the lipid bilayer. The cross-sectional area and stiffness of the nanosized domain and its...
connectivity to the spectrin network are key parameters governing its mechanical properties. In our simulations, instead of modeling the knobs as independent structures, we establish a biomechanically equivalent and computationally convenient model for the knobs. Specifically, each knob is modeled as a circular, relatively stiff lipid bilayer region (Fig. 1 B1 and C1, yellow regions) that connects, whenever reachable, to the fourth beads of the nearest spectrin tetramers, to actin junctional complexes, and to ankyrins (Fig. 1D2), using harmonic springs. The potential wells between a pair of lipid agents within the knob region is set to be deeper than that in the normal lipid bilayer so as to effectively stiffen it to capture the effective stiffness of the knob (see SI Text for details). Because the lipid bilayer region representing the knob has the same area and connectivity to the spectrin network and the replaced lipid domain contributes negligibly to the shear modulus, it is mechanically faithful to the real structure of the knob. In the course of parasite maturation, the density, size, and distribution of the knobs on the cell membrane dynamically change. Previous microstructural studies evidenced that in general the knob density increases, whereas the knob radius decreases from the trophozoite (Fig. 1C1) to the schizont (Fig. 1C1) stage (44, 45).

**CG Simulations**

To compute the shear modulus of the RBC membranes, a piece of RBC membrane (~600 nm × 600 nm) as shown in Fig. 1E was subjected to simple shear. Unless otherwise mentioned, the shear rate applied to the membrane was defined as \(\gamma = 2.97 \times 10^5 \text{s}^{-1}\), where \(\gamma\) is the engineering shear strain. As shall be seen later, such a shear rate is chosen so that the lipid bilayer in uninfected RBCs contributes negligibly to the shear stress (\(\tau\)). Fig. 2A depicts the simulated configuration of the normal cell membrane at \(\gamma = 1\). We found that only a small portion of the actin junctional complexes is broken. Fig. 2D shows the shear stress-strain responses of the uninfected RBC membrane. As expected, shearing generates negligible viscous stress in the lipid bilayer owing to its fluid nature, and the shear resistance is mainly contributed by the spectrin network. Notably, the spectrin network exhibits a strong strain-hardening effect, with a shear modulus of ~10 \(\mu\)N/m at a small shear strain (<0.3) but of ~18.4 \(\mu\)N/m at a high shear strain (>0.8). The extracted shear modulus of the healthy RBC thus falls within the range of existing experimental data (~4-10 \(\mu\)N/m at small shear strain (<0.4); 14-20 \(\mu\)N/m at large shear strain (<1.0) (46, 47)).

**The Effect of Spectrin Network Remodeling.** We next investigate the role of spectrin network remodeling in the shear responses of Pf-infected RBC membranes. The simulation model comprises the normal lipid bilayer (Fig. 1A1) coupled with a spectrin network that incorporates the proposed Pf-induced modifications (Fig. 1B2 or C2). For the deficient spectrin network (Figs. 1C2 and 2B), the nonbonded tetramers do not sustain shear, and therefore, the shear modulus of the RBC membrane decreases to ~8.1 and ~15.9 \(\mu\)N/m at low and high shear strains, respectively (Fig. 2E). This reduction in the membrane shear modulus by the deficient spectrin network (i.e., with holes) might facilitate rupture of the RBC membrane during egress (48, 49). In contrast to the deficient spectrin network, the enhanced spectrin network maintains a perfect hexagonal topology upon shear (Fig. 2C). The shear modulus of the RBC membrane with the enhanced spectrin network is stiffened to ~14.9 and ~28.2 \(\mu\)N/m at low and high shear strains, respectively, as shown in Fig. 2E. The stiffness of each individual spectrin tetramer derived from the worm-like chain model (27) is positively correlated to its end-to-end distance. In addition, a spectrin octamer can be considered equivalent to two tetramers in parallel and consequently has a higher stiffness than a single tetramer. Thus, both the elongated end-to-end distance of spectrin tetramers and the emergence of the spectrin octamers contribute to the increase in shear modulus. However, the shear modulus (19.6 \(\mu\)N/m, averaged over the range of the shear strain from 0 to 1) caused by enhanced spectrin network is much less than that measured for Pf-infected RBC membranes (~40 \(\mu\)N/m at trophozoite stage and ~60 \(\mu\)N/m at schizont stage on average (11, 13)). Thus, mechanisms other than the spectrin network remodeling must be present for stiffening the Pf-infected RBC membrane.

**The Effect of Knobs.** The contribution of knobs to the stiffening of Pf-infected RBC membranes has been experimentally investigated by genetically eliminating KAHRP expression so as to prevent knob formation (50). It was shown that knobs account for ~50% of the overall increase in the shear modulus in Pf-infected RBCs (50). However, it is possible that knocking out KAHRP also disrupts the transport of other parasite-encoded proteins. Additional proteins that might also contribute to the increased shear modulus include PIEMP1, RESA, MESA, Pf332, and STEVOR (17, 18, 20, 22, 51, 52). Our CG model is capable of separating the effect of the knobs from other factors. To this end, we establish a simulation model that combines the normal spectrin network with an overlying lipid bilayer that is modified with knobs. The knob density has been shown to vary between different human hosts (45), but the knob density increases as the intraerythrocytic parasite matures from the trophozoite to the schizont stage. Based on an ultrastructural study (44), we modeled the knob density to increase by 2.5 times as the intraerythrocytic parasite matures from the trophozoite to the schizont stage, from
20 knobs per µm² (Fig. 1B1) to 50 knobs per µm² (Fig. 1C1). The average knob radius was modeled to decrease by 1.5 times, from 75 to 50 nm, and the ratio of the total knobby area to RBC membrane area slightly increased from 35% to 39% (44). From atomic force microscopy imaging (43), the center of each knob is roughly located on the top of an actin junctional complex but otherwise randomly positioned. Each knob forms vertical links by connecting to the actin junctional complex, and whenever reachable, to the fourth beads of the nearest spectrin tetramers, and to ankaryins. Following this connecting topology, each knob connects to the spectrin network by 8–11 vertical links for the knobs of size r = 75 nm and 6–8 vertical links for r = 50 nm; this gives rise to 73 total vertical links for the low-density large knobs and 147 for the high-density small ones. Note the twofold increase in the total number of vertical links due to the increase of the knob density, despite the small change of the total knob area.

Fig. 3A and B display the shear morphologies of membranes with the knobs at trophozoite and schizont stages, respectively. Notably, as shown in Fig. 3C, the stiff lipid bilayer regions representing the knobs undergo finite shear stress, in contrast to the vanishing shear stress in the fluid lipid bilayer. Shear stresses in both the spectrin network and the knobby regions first rise slowly at low shear strains, and then sharply at large shear strains, corresponding to a pronounced stiffening effect. Our result indicates that the lipid domain representing the knobs no longer behaves as a Newtonian fluid, but more like a solid, which makes a considerable contribution to the shear modulus increase, as shown by the black curves in the figures. At γ = 1, the shear stress in the lipid bilayer makes up about 30% of the total shear stress, which is in clear contrast to the negligible viscous stress in the uninfected RBC membrane. The shear stress in the spectrin network also rises in the presence of knobs because of the enhanced vertical connections. At the trophozoite stage, the presence of knobs elevates the shear modulus of the RBC membrane to ~16.0 and ~44.9 µN/m at low and high shear strains, respectively.

Fig. 3E displays the shear stress–strain responses of the RBC membrane with knobs at the schizont stage, showing that the knob density significantly affects the shear modulus of the RBC membrane. The shear modulus is accordingly increased to ~22.2 and ~77.9 µN/m at low and high shear strains, respectively.

**Combined Effects.** We next compute the shear moduli of *P.f-*infected RBC membranes at different asexual stages, combining the physiologically relevant spectrin network remodeling with different knob characteristics. Specifically, we combine the knob-stiffened membrane bilayer with large knob size (Fig. 1B1) and the enhanced spectrin network (Fig. 1B2) to model the *P.f-*infected RBC membrane at the trophozoite stage. We also combine the lipid bilayer with small knob size (Fig. 1C1) and the deficient spectrin network (Fig. 1C2) to model the membrane at the schizont stage. Fig. 4 shows the shear stress–strain responses of the *P.f-*infected RBC membranes at the two later asexual stages, along with that of the uninfected membrane for comparison. At small shear strains, the shear modulus of the *P.f-*infected RBC membrane increases by twofold compared with the normal membrane. At large shear strains, the shear moduli of the *P.f-*infected RBC membrane at trophozoite and schizont stages are increased to ~48.1 and ~77.2 µN/m, respectively. Compared with the cases with physiologically relevant knob radii and densities but with normal spectrin network (Fig. 3D and E), the shear moduli of the *P.f-*infected RBC membranes change negligibly, further demonstrating the insignificant role of the spectrin network remodeling on stiffening. The shear moduli predicted by this model at both small and large shear strains for these two stages fall in the range of experimental measurements [20–60 µN/m at trophozoite stage; 25–90 µN/m at schizont stage (11, 13)].
Analysis

We view the knobby RBC membrane as a composite with two regions: one with the spectrin network only (knob-free region) and the other with the spectrin network with attached knobs. Because the fluid lipid bilayer does not contribute to shear resistance, it is not considered in the analysis. According to the rule of mixtures, the shear strain applied to the composite γc is distributed in the two domains,

$$\gamma_c = \alpha \gamma_K + (1 - \alpha) \gamma_F, \quad [1]$$

where γF and γK are the shear strains in the knob-free spectrin network region and the knobby region, respectively, and α is the areal fraction of the knobby region. Because these two regions are subjected to the same shear stress τ,

$$\tau = \gamma_K G_K = \gamma_F G_F(\gamma_F), \quad [2]$$

where GF and GK are the shear moduli of the knob-free and the knobby regions, respectively. To emphasize the material nonlinearity and strain-hardening effects of the spectrin network, the shear modulus of the spectrin network is expressed here as a function of the shear strain γF. Substituting Eq. 2 to Eq. 1 and noting that the protein-enriched knobby region is much more rigid than the spectrin network (G_K ≫ G_F), one arrives at the conclusion that the shear strain in the knob-free region is amplified by a factor of approximately 1/(1 − α) from the applied shear strain:

$$\gamma_F = \frac{\gamma_c}{[\alpha G_F(\gamma_F)/G_K + (1 - \alpha)]} \approx \gamma_c/(1 - \alpha). \quad [3]$$

This suggests that the knob-free region experiences an elevated shear strain (γF) compared with the apparent, applied shear strain (γC).

This amplified shear strain in the knob-free region is confirmed by our CG simulations, as shown in Fig. 5, where γF = 1.44 for γC = 1 and α = 0.35. Rule of mixtures gives rise to the shear modulus of the composite as follows:

$$\frac{1}{G_C(\gamma_C)} = \frac{\alpha}{G_K} + \frac{1 - \alpha}{G_F(\gamma_F/(1 - \alpha))}. \quad [4]$$

The expression of the shear modulus of the Pf-infected RBC membranes thus manifests as a dual stiffening mechanism of the knob-membrane complexes. First, the knobs act as structural strengtheners of the spectrin network, as represented by the first term on the right side of Eq. 4. Second, the presence of rigid knobs results in inhomogeneous shear strain distribution. In particular, the shear strain in the knob-free regions is amplified compared with the externally applied shear strain (γF > γC). Owing to the strain-hardening property of the spectrin network, G_F(γ_F) > G_F(γ_C), the knob-free regions are also effectively stiffened, as indicated in the second term of Eq. 4. For an applied shear strain γ = 1, the expected shear modulus of the knob-free region is ~50 μN/m, compared with ~20 μN/m without the strain-hardening effect (see Fig. S1 for shear modulus at high shear strains). The dual role of the knobs explains the significantly increased shear modulus of Pf-infected RBC membranes.

Our simulations further revealed that the strong knob density (or size) effect originates from the increase in the total number of vertical links from the knobs to the spectrin network through a comparative study, as detailed in SI Text. In the case of the same total knob area and the same total number of vertical links but different knob size [thus different knob density (Fig. S2)], the shear modulus changes negligibly. Whereas for the same knob size but with the increase in the total knob area and hence in the total number of vertical links, the shear stiffness markedly increases (Fig. S3). These comparative studies demonstrate that the total number of vertical links, rather than the apparent knob size or the total knob area, is a primary stiffening factor. From the trophozoite to the schizont stage, the total knob area remains roughly the same, but the knob density increases. As a result, the total number of vertical links increases and a much stiffer membrane develops. It is interesting to consider how the parasite could stiffen the membrane by remodeling the knob size and density so as to increase the vertical coupling to the spectrin network to stiffen the membrane.

Discussion and Concluding Remarks

We have developed an efficient composite RBC membrane model of the lipid bilayer and the spectrin cytoskeleton network. This model enables a systematic study of the effects of changes in the molecular structures on the shear elasticity of Pf-infected RBC membranes. Our modeling results show that the deposition of knobs, rather than spectrin network remodeling, is a primary cause of the dramatically increased stiffness of the Pf-infected RBC membranes. Our analysis further revealed that the knobs stiffen the RBC membrane in a unique manner by simultaneously harnessing composite strengthening, strain hardening, and knob density-dependent vertical coupling effects. First, the knobs stiffen the spectrin network as structural strengtheners, commonly seen in composites reinforced by stiff inclusions. Second, the presence of knobs significantly elevates the shear strain in the knob-free regions of the spectrin network; given the strain-hardening property of the spectrin network, the knob-free regions are also effectively stiffened. Third, during maturation from the trophozoite to the schizont stage, the knob density increases, resulting in increased number of the knob-associated vertical links, which further increases the membrane stiffness. The predicted shear moduli of Pf-infected RBC membranes by the CG model fall in the range of the experimental measurements [20–60 μN/m at trophozoite stage; 25–90 μN/m at schizont stage (11, 13)].

The present study suggests possible targets for the development of effective antimalarial agents. The cytoadherence protein, PfEMP1 (53, 54), has been previously suggested (55) as...
promising target molecules for developing medication against malaria. Disrupting the PfEMP1 proteins would undoubtedly reduce the cytoadherence, but it might not be effective for changing the high stiffness of the membrane. In addition to the stiffening role, the knobs also act as platforms for the presentation of PfEMP1. The high stiffness of the knob region together with the knob-associated vertical connections facilitate the distribution of the tensile forces imposed on PfEMP1 to the entire knob region and to the cytoskeleton. In light of these multifunctional roles, disrupting the vertical proteins would potentially reduce both the stiffness and the strength of adhesion to the endothelium. Molecules that block the binding of KAHRP (or other parasite-encoded proteins) to RBC spectrin network could be promising candidates as antimalarial agents.

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6. Pei X, et al. (2005) Structural and functional studies of interaction between Plasmo-
20. Boal DH (1994) Computer simulation of a model network for the erythrocyte cyto-