Localized glucose and water influx facilitates Cryptosporidium parvum cellular invasion by means of modulation of host-cell membrane protrusion

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Dynamic membrane protrusions such as lamellipodia and filopodia are driven by actin polymerization and often hijacked by intracellular microtubules to enter host cells. The overall rate of membrane protrusion depends on the actin polymerization rate and the increase of localized cell volume. Although the signaling pathways involving actin polymerization are well characterized, the molecular mechanisms regulating local cell volume associated with membrane protrusion are unclear. Cryptosporidium parvum, an intracellular parasite, depends on host-cell membrane protrusion to accomplish cell entry and form the parasitophorous vacuole. Here, we report that C. parvum infection of cholangiocytes recruits host-cell SGLT1, a Na+/glucose cotransporter, and aquaporin 1 (AQP1), a water channel, to the attachment site. SGLT1-dependent glucose uptake occurs at the attachment site. Concordantly, the region of attachment displays localized water influx that is inhibited by either suppression of AQP1 by means of AQP1-small interfering RNA (siRNA) or inhibition of SGLT1 by a specific pharmacologic inhibitor, phlorizin. Inhibition of SGLT1 does not affect actin accumulation but decreases the membrane protrusion at the attachment site. Moreover, functional inhibition of host-cell AQP1 and SGLT1 hampers C. parvum invasion of cholangiocytes. Thus, glucose-driven, AQP-mediated localized water influx is involved in the membrane protrusion during C. parvum cellular invasion, phenomena that may also be relevant to the mechanisms of cell membrane protrusion in general.

Localized glucose and water influx facilitates Cryptosporidium parvum cellular invasion by means of modulation of host-cell membrane protrusion in general.

actin | aquaporin | cryptosporidiosis | Na+/glucose cotransporter 1 | bile ducts

C. parvum, an intracellular parasite within the protist phylum Apicomplexa, infects gastrointestinal epithelia and other mucosal surfaces in humans, causing self-limited diarrhea in immunocompetent subjects and potentially life-threatening syndromes in immunocompromised individuals, primarily those with AIDS (13). Once ingested, C. parvum oocysts excyst in the gastrointestinal tract, releasing infective sporozoites. Mediated by poorly characterized but specific ligands on the sporozoite surface and unidentified receptors on the host cell, the sporozoite attaches to the apical membrane of the epithelial cell, inducing host-cell membrane protrusion that encapsulates the parasite in a parasitophorous vacuole (14–16). The parasite is then retained on the apical surface of the cell, and a unique “electron-dense band” of unknown composition is established that separates the organism from the host-cell cytoplasm and may facilitate the uptake of nutrients from the host. Thus, the parasitophorous vacuole membrane and dense band keep the internalized parasite intracellular but extracytoplasmic (14–16). Despite intensive efforts over the past decades, there is currently no effective therapy for cryptosporidiosis in humans (17).

Host-cell membrane protrusion plays an active role in C. parvum entry of epithelial cells. Inhibition of host-cell actin polymerization by pharmacological inhibitors, such as cytochalasin B and cytochalasin D (18, 19), or by cellular expression of specific inhibitory fragments of actin-associated proteins, such as Scar-WA (20), blocks C. parvum cellular invasion. Several intracellular signaling pathways are involved in C. parvum-induced actin polymerization, including the c-Src/cortactin and phosphatidylinositol 3-kinase (PI3-K)/Cde42 pathways (21–23). Functional inhibition of those actin-associated signal pathways results in an inhibition of host-cell membrane protrusion and, subsequently, of parasite cellular invasion (21–23).

In work described here, we found that C. parvum recruits host-cell AQP1 and SGLT1 to the attachment site during C. parvum infection of cultured cholangiocytes. Recruitment of SGLT1 and AQP1 results in a localized increase of glucose uptake and a subsequent AQP1-mediated water influx at the attachment site. The localized water influx seems to enhance C. parvum-induced host-cell membrane protrusion and thus facilitates parasite cellular invasion. Thus, localized glucose-driven, AQP-mediated water influx is involved in membrane protrusion during C. parvum invasion.
a phenomenon that is also relevant to the general mechanisms of cell membrane protrusion.

Materials and Methods

C. parvum and Cholangiocyte Cell Lines. C. parvum oocysts of the Iowa strain were purchased from a commercial source (Bunch Grass Farm, Deary, ID). H69 cells (a gift of D. Jefferson, Tufts University, Medford, MA) are SV40-transformed normal human cholangiocytes (24). 603B cells are immortalized normal mouse cholangiocytes (a gift from Y. Ueno, Tohoku University School of Medicine, Sendai, Japan) (25).

In Vitro Models and Infection Assay. An attachment model and an attachment/invasion model were used as described (18). Before infecting cells, oocysts were excysted to release infective sporozoites (18). Infection was done in a culture medium consisting of DMEM-F12, 100 units/ml penicillin, 100 µg/ml streptomycin, and freshly excysted sporozoites (1 × 10⁶ sporozoites per slide well or culture plate). Inactivated organisms (treated at 65°C for 30 min) were used for sham infection controls. Infection assays (attachment rate or attachment/invasion rate) were carried out after incubation of 1 h with the parasite employing an indirect immunofluorescent approach (18). Briefly, the number of parasites and cells was counted randomly in each group, and results were expressed as attachment or attachment/invasion rate [(the number of parasites per total number of cells) × 100]. A total of >2,000 cells were counted for each slide, and all assays were performed in triplicate. For the inhibitory experiments, one specific inhibitor of SGLT1, phlorizin (26), was added in the medium at the same time as C. parvum. A concentration of 0.2 mM phlorizin, which showed no cytotoxic effects on 603B cells or on C. parvum sporozoites, was selected for the study.

AQP1-Small Interfering RNA (siRNA) and Plasmid Constructs. Generation of AQP1-siRNA and subsequent suppression of AQP1 expression have been described (27). A nonspecific siRNA containing the same nucleotides but in irregular sequence (i.e., scrambled siRNA) was used as the control. The AQP1-siRNA was labeled with Cy3 by using the Silencer siRNA labeling kit (Ambion, Austin, TX). Cells were transfected with AQP1-siRNA by using the siPORT lipid transfection agent (Ambion). pEGFP-N1-AQP1 was generated by inserting the complete rat AQP1 into the expression vector pEGFP-N1 (Invitrogen). The pEGFP-N1 empty vector and pEGFP-N1-GFP-Dynamin2 (a gift of M. A. McNiven, Mayo Clinic College of Medicine) were used as the controls. Cells were transfected with GFP constructs with the Lipofectamine Plus reagent kit (Invitrogen). 603B cells, which don't express endogenous AQP1, and H69 cells, were transfected with the GFP-AQP1 construct into H69 cells and then exposed to infection medium containing 10 µM calcein-acetoxyethyl ester (calcine-AM, Molecular Probes) at 37°C for 30 min. After extensive washing with culture medium, cells were then exposed to freshly excysted C. parvum sporozoites for 1 h at 37°C, followed by real-time confocal microscopy as described above.

Visualization of Glucose Uptake. A fluorescent glucose analogue, 2-[N-(7-nitrobenz-2-oxa-1,3-diazol-4-yl) amino]-2-deoxy-glucose (2-NBDG) (Molecular Probes), was used to measure localized glucose uptake in single cells (29). Briefly, 603B cells were grown on cover-glass-bottomed 3-cm dishes to subconfluence. Cells were then exposed to freshly excysted C. parvum sporozoites in the infection medium containing 1% glucose with or without 0.2 mM phlorizin for 50 min at 37°C. 2-NBDG was then added to the culture medium (at a final concentration of 200 µM). 2-NBD, a control probe provided by the manufacturer (Molecular Probes), was also used. After incubation at 37°C for 10 min, cells were assessed by real-time confocal microscopy. Fluorescence intensity at 520–560 nm wavelength (excitation wavelength 488 nm) was measured and calculated with an analysis system of the LSM 510 provided by Carl Zeiss. A total of 500 parasite attachment sites were randomly selected and measured for each condition. Results represent three independent experiments.

Visualization of Sites with Water Influx. A calcine fluorescence quenching method was used to assess the water influx as described (12, 30). A water influx changes the cytosolic protein concentration in the region and thus alters calcine quenching, resulting in an increase of calcine fluorescence (30). Briefly, 603B cells were grown on cover-glass-bottomed 3-cm dishes to subconfluence. Cells were then exposed to the infection medium containing 10 µM calcine-acetoxyethyl ester (calcine-AM, Molecular Probes) at 37°C for 30 min. After extensive washing with culture medium, cells were then exposed to freshly excysted C. parvum sporozoites for 1 h at 37°C, followed by real-time confocal microscopy as described above.

Electron Microscopy. For scanning electron microscopy (SEM) and transmission electron microscopy (TEM), cells were prepared as described in the product literature (Technical Bulletins 405 and 406, Electron Microscopy Sciences, Hatfield, PA). The length of membrane protrusions was determined by measuring the distance from tip to base on digitized TEM images. A total of ~200 parasite attachment sites were measured for each condition, and results represent three independent experiments. For immunogold labeling, cells were fixed and processed according to the protocol as described (21, 22). The relative distribution of AQP1 was determined by counting gold particles over cell profiles. Dense-band and parasitophorous vacuole membrane-associated labeling was quantitated by counting the gold particles in the area within a 0.2-µm distance along the dense-band or vacuole membrane; totals were described as gold particles per µm². Gold particles at randomly selected cytoplasm and apical membrane areas (within a 0.2-µm distance along membrane surface) were also counted in both sham and C. parvum-infected cells. A total of ~100 parasite attachment sites and >2,000 gold particles were counted randomly for each condition; data represent three independent experiments.

Results

C. parvum Recruits Cholangiocyte AQP1 to the Attachment Sites. We chose two cell lines for the study: 603B cells, which express endogenous AQP1, and H69 cells, which don’t express endogenous AQP1 as confirmed by both Western blotting and immunofluorescent staining (data not shown). When 603B cells were exposed to C. parvum, very strong staining of AQP1 was observed at the attachment site by immunofluorescence (Fig. L4). No accumulation of AQP1 was detected in sham-infected control cells, and no positive staining of AQP1 was found in the parasite itself (data not shown). To confirm the host-cell origin of AQP1, we transfected a GFP-AQP1 construct into H69 cells and then exposed the cells to C. parvum. A similar expression pattern of AQP1-GFP was found in transfected H69 cells compared with the antibody staining in
603B cells, and a strong accumulation of GFP-AQP1 was found at the attachment site (Fig. 1B). In contrast, no accumulation of fluorescence was found at the parasite attachment sites in cells transfected with the GFP-dynamin 2 (Fig. 1C), a GFP fusion protein that is not associated with C. parvum cellular invasion (21), and the GFP empty vector (Fig. 1D). To further localize the accumulation of AQP1 in infected cells, 603B cells exposed to C. parvum sporozoites were assessed by immunohistochemistry using an antibody to AQP1. Colloidal gold particles specific for AQP1 accumulated along the dense-band region. Some labeling was also observed at the parasitophorous vacuole membrane (Fig. 1E–G), whereas fewer gold particles were found in the cytoplasm not directly adjacent to the parasite in infected cells (solid arrow in Fig. 1F). Sham-infected cells showed no membrane accumulation of gold particles but a scattered distribution in the cytoplasm (arrows in Fig. 1H). Quantitative analysis showed a significant increase of gold particles for AQP1 along the dense-band and at the parasitophorous vacuole membrane compared with other areas not directly adjacent to the parasite or in sham infection controls (Fig. 1J).

C. parvum Recruits Cholangiocyte SGLT1 to the Attachment Sites and Induces a Localized Glucose Uptake. 603B cells were exposed to C. parvum followed by immunofluorescent labeling for SGLT1, a Na+/glucose cotransporter on the apical membrane domain. Strong staining of SGLT1 was observed at the C. parvum attachment site (arrowheads in Fig. 2A and B). To test whether SGLT1 is activated, we used a real-time fluorescent-based approach using 2-NBDG, a fluorescent glucose analogue, to measure the uptake of glucose in infected cells. An increase of fluorescence was found at the C. parvum attachment site in 603B cells (Fig. 2C), and no increase of fluorescence was found over the parasite attachment site in the presence of phlorizin (Fig. 2D). No increase of fluorescence to the control probe, 2-NBD, was found at the attachment site (arrowhead). (C–E) Quantitative analysis of 2-NBD and 2-NBDG uptake at the attachment site. *, P < 0.05; ANOVA, C. parvum vs. control (fluorescence at the nearby region); **, P < 0.05 vs. C. parvum. (Scale bars: 5 μm.)
ing approach. Strong fluorescence was observed at the attachment site in infected 603B cells (Fig. 3A), reflecting localized water influx (30). No labeling of calcein-AM was found in the parasite itself with the approach used (data not shown). To test whether the localized water influx at the attachment site is AQP1-mediated, we first transfected 603B cells with Cy3-tagged AQP1-siRNA to suppress AQP1 expression and then measured water influx at the attachment site in infected cells after exposure to C. parvum. As shown in Fig. 3B, no obvious increase of fluorescence was found at the attachment site in siRNA-transfected 603B cells.

To test whether SGLT1-associated glucose uptake is associated with C. parvum-induced AQP1-mediated localized water influx, we tested the effect of phlorizin on C. parvum-induced water influx. As shown in Fig. 3C, no obvious increase of calcein-AM fluorescence was found at the attachment site in the presence of phlorizin. Quantitative analysis showed: (i) a significant increase of water influx-associated fluorescent intensity at the attachment site in 603B cells (Fig. 3D); and (ii) a significant decrease of water influx-associated fluorescent intensity at the attachment site in 603B cells in the presence of phlorizin or in cells transfected with AQP1-siRNA (Fig. 3D).

**Functional Manipulation of AQP1 and SGLT1 Affects C. parvum Invasion of Cholangiocytes.** To further determine whether accumulation of AQP1 and SGLT1 and subsequent water influx are involved in parasite cellular invasion, we manipulated the function of AQP1 and SGLT1 in host cells and tested the effects on C. parvum invasion. 603B cells were first transfected with AQP1-siRNA and, 24 h later, exposed to C. parvum. Phlorizin was also used to test the role of SGLT1 in C. parvum entry of host cells. When cholangiocytes were prefixed and then exposed to C. parvum sporozoites, a model in which the organism can both attach to and enter into host cells, significantly fewer parasites were detected in cells transfected with AQP1-siRNA or in the presence of phlorizin (Fig. 4A). Attachment assay in prefixed cells. (B) Attachment/invasion assay in nonprefixed cells. (C and D) Representative confocal images of attachment/invasion of C. parvum by nonprefixed cells treated with scrambled siRNA (C) or AQP1-siRNA (D). Representative confocal images of attachment/invasion of C. parvum by nonprefixed cells in the absence (E) or presence (F) of phlorizin.

**Inhibition of SGLT1 by Phlorizin Does Not Affect Localized Actin Accumulation but Hampers C. parvum-Induced Cholangiocyte Membrane Protrusion.** To test how SGLT1, as well as subsequent AQP1-mediated water influx, affects C. parvum cellular invasion, we observed host-cell actin accumulation and membrane protrusion in 603B cells during C. parvum invasion in the presence or absence of phlorizin by confocal microscopy and electron microscopy. Phalloidin, which binds to host-cell F-actin but not C. parvum actin (19), revealed accumulation of actin at the attachment sites both in the presence and absence of phlorizin (Fig. 6A and B). In the absence of phlorizin, cells showed obvious membrane protru-
sion enveloping the organism at the parasite attachment site (arrowheads in Fig. 6C). In contrast, much less host-cell membrane protrusion was found in the presence of phlorizin (arrowhead in Fig. 6D). Quantitative analysis showed a significant decrease of length of membrane protrusions at the attachment site in 603B cells in the presence of phlorizin compared with the controls (Fig. 6E).

Discussion

The key findings in this report are as follows: (i) C. parvum infection of cholangiocytes recruits host-cell SGLT1 and AQP1 to the attachment site, resulting in a localized increase of active glucose uptake by SGLT1 and subsequent passive water influx by means of AQP1; and (ii) localized glucose and water influx seem to be involved in host-cell membrane protrusion and subsequent parasite cellular invasion. Thus, C. parvum recruits host-cell SGLT1 and AQP1 to the attachment site, generating a localized water influx, thereby facilitating parasitic cellular invasion by means of modulation of host-cell membrane protrusion.

Cellular invasion by C. parvum is characterized by the formation of a tunnel connection between the anterior vacuole in the apical region of the sporozoite and the host-cell cytoplasm (31) and by host-cell membrane protrusion to encapsulate the invading parasite. At the base of each vacuole, a unique “electron-dense band” is established (13–16). A variety of molecules from both the parasite and the host cell accumulate at the host–parasite interface, including CpABC, a recently identified C. parvum ATP-binding cassette protein (32), and various host-cell actin-associated proteins, such as cortactin (21) and Cdc42 (22). It has been postulated that recruitment of transporters/exchangers/channels to the host–parasite boundary can supply the parasite with various nutritional requirements (14). Here, we have shown that C. parvum infection of cholangiocytes recruits host-cell SGLT1 and AQP1 to the attachment site. Accumulation of AQP1 was found both at the parasitophorous vacuole membrane and along the dense band region. Interestingly, functional inhibition of host-cell SGLT1 by a specific inhibitor, phlorizin, or suppression of AQP1 expression in host cells by a specific AQP1-siRNA, significantly decreased parasite cellular invasion. In addition, overexpression of a GFP-AQP1 construct in H69 cells increased C. parvum cellular invasion. Thus, it seems that, during the early infection stage, C. parvum recruits host-cell SGLT1 and AQP1 to the attachment site, a process that is directly associated with parasitic cellular invasion.

Cellular invasion by C. parvum depends on host-cell actin polymerization and subsequent membrane protrusion at the host–parasite interface. Inhibition of host-cell actin polymerization inhibits C. parvum-induced membrane protrusion and reduces parasite cellular invasion (18–20). We found that inhibition of SGLT1 did not affect actin-accumulation at the host–parasite interface. Instead, we detected a localized increase of glucose uptake at the attachment site during C. parvum invasion. A specific inhibitor of SGLT1, phlorizin, completely blocked glucose uptake by 603B cells. Cells loaded with calcein-AM displayed an increase

![Fig. 5. Expression of AQP1 in H69 cells increases C. parvum invasion. H69 cells were transfected with a AQP1-GFP construct for 24 h and then exposed to C. parvum for 1 h. (A) Attachment assay in prefixed cells. (B) Attachment/invasion assay in nonprefixed cells. (C) A representative confocal image of attachment/invasion of C. parvum (arrowheads) by nonprefixed cell culture transfected with the AQP1-GFP construct. Cells transfected with AQP1-GFP were identified with the GFP expression. *, P < 0.05, t test. (Scale bar: 5 μm.)](image)

![Fig. 6. Inhibition of SGLT1 by phlorizin does not affect localized actin accumulation but hampers C. parvum-induced cholangiocyte membrane protrusion in 603B cells. (A and B) Strong actin accumulation was found at the attachment site both in the absence (A) and presence (B) of phlorizin. (C and D) Host-cell membrane protrusion at the parasite–host interface as assessed by scanning electron microscopy (SEM). (E) Quantitative analysis of length of membrane protrusion based on transmission electron microscopy (TEM) images. *, P < 0.05, t test. (Scale bars: 1 μm.)](image)
of fluorescence at C. parvum attachment sites compared with nearby regions, suggesting localized water influx (30). Inhibition of either SGLT1 or AQPs blocked the localized water influx, indicating that the water-influx is AQPi-mediated and likely requires the osmotic gradient generated by the uptake of glucose by means of SGLT1. Moreover, inhibition of SGLT1 by phlorizin significantly hampered C. parvum-induced cholangiocyte membrane protrusion as assessed by electric microscopy. Therefore, we postulated that C. parvum recruits host-cell SGLT1 and AQPi to the attachment site, generating a glucose-driven localized inward osmotic gradient followed by regional influx of water, processes that facilitate the localized host-cell membrane protrusion required for parasitic cellular invasion.

Pollard and Borisy (1) have recently depicted the dendritic-nucleation model to explain the formation of membrane protrusions based on actin polymerization. It proposes that actin filaments growing by actin assembly at the barbed ends produce enough force to push the plasma membrane outward. What this model does not address is how cells quickly accommodate the resultant increase of localized cell volume accompanying membrane protrusion. Cells regulate their volume mainly by means of water and ion transport, and AQPs are water channels that allow rapid passive water movement across membranes in response to gradients established by the transport of osmotically active molecules such as glucose (7, 8). Our data support the notion that C. parvum recruits functionally active host-cell SGLT1 to the attachment site to induce a localized uptake of glucose, generating an inward osmotic gradient to drive regional inward water movement by means of AQPi. AQPi-mediated localized water influx accommodates the increase of cell volume accompanying localized membrane protrusion, facilitating C. parvum cellular invasion.

Loitto et al. (12) recently reported that AQPi9 is recruited to the motile regions of neutrophil leukocytes. It has been postulated that, during membrane protrusion, localized water influx generates a gap between the filament ends and the membrane, thereby allowing for rapid polymerization of actin. Thus, a regulated water influx will allow for gel-to-sol transitions, osmotic forces, and subsequent dendritic nucleation, indicating that AQPs may be involved in the regulation of water movement in a localized region such as membrane protrusion. Coupling our current and previous studies with the work of others, the results suggest that C. parvum activates host-cell intracellular signaling pathways to induce actin polymerization at the attachment site, generating a force for membrane protrusion. C. parvum also recruits host-cell SGLT1 and AQPi to the attachment site, generating localized glucose and water influx to accommodate the increase of cell volume accompanying membrane protrusion. Actin polymerization and rapid increase of local cell volume result in efficient localized membrane protrusion, thereby facilitating parasite cellular invasion (Fig. 7).

In conclusion, our data indicate that a pathogen, C. parvum, recruits host-cell SGLT1 and AQPi to the attachment site, generating localized glucose and water influx that are required for efficient membrane protrusion and parasite invasion. It will be of interest to extend these studies to determine the mechanisms by which C. parvum recruits SGLT1 and AQPi at the attachment site. Our studies not only provide insight into the mechanism of C. parvum entry of host epithelial cells, but also are relevant to the general molecular mechanisms of cell membrane protrusion.

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