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Integrin β 1 Mediates Vaccinia Virus Entry through Activation of PI3K/Akt Signaling

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Vaccinia virus has a broad range of infectivity in many cell lines and animals. Although it is known that the vaccinia mature virus binds to cell surface glycosaminoglycans and extracellular matrix proteins, whether additional cellular receptors are required for virus entry remains unclear. Our previous studies showed that the vaccinia mature virus enters through lipid rafts, suggesting the involvement of raft-associated cellular proteins. Here we demonstrate that one lipid raft-associated protein, integrin β 1, is important for vaccinia mature virus entry into HeLa cells. Vaccinia virus associates with integrin β 1 in lipid rafts on the cell surface, and the knockdown of integrin β 1 in HeLa cells reduces vaccinia mature virus entry. Additionally, vaccinia mature virus infection is reduced in a mouse cell line, GD25, that is deficient in integrin β 1 expression. Vaccinia mature virus infection triggers the activation of phosphatidylinositol 3-kinase (PI3K)/Akt signaling, and the treatment of cells with inhibitors to block PI3K activation reduces virus entry in an integrin β 1-dependent manner, suggesting that integrin β 1 mediates PI3K/Akt activation induced by vaccinia virus and that this signaling pathway is essential for virus endocytosis. The inhibition of integrin β 1-mediated cell adhesion results in a reduction of vaccinia virus entry and the disruption of focal adhesion and PI3K/Akt activation. In summary, our results show that the binding of vaccinia mature virus to cells mimics the outside-in activation process of integrin functions to facilitate vaccinia virus entry into HeLa cells.

Vaccinia virus is the prototype of the orthopoxvirus genus of the family *Poxviridae*. It has a broad host range, producing multiple infectious forms of virus particles, e.g., mature virus (MV), wrapped virus (WV), and extracellular virus (EV) (14). Vaccinia MV particles are produced in large quantities in infected cells and contain ~80 viral proteins in viral particles (10, 51, 69). The large number of viral proteins in the MV composition contributes to its complex viral entry processes. First, vaccinia MV binds to multiple cell surface components, such as glycosaminoglycans (11, 29, 42), extracellular matrix laminin (9), and sulfatides (49). Furthermore, vaccinia MV contains an entry fusion complex (EFC) of 12 viral envelope proteins that are evolutionarily conserved and essential for membrane fusion (6, 7, 33, 47, 52, 56, 68). Additionally, different strains of vaccinia MV harboring mutations in the viral envelope proteins A25 and A26 enter cells differently (8). Finally, vaccinia MV entry pathways also vary among different cell lines (66), and multiple kinases, such as extracellular signal-regulated kinase (ERK), protein kinase A (PKA), protein kinase C (PKC), and PAK1, have been shown to be activated upon virus entry (18, 43, 45). However, the cellular receptor mediating vaccinia MV entry and signal transduction remains unknown. Our previous work showed that cell-bound MV particles were clustered at the plasma membrane lipid rafts prior to virus entry and that the interruption of lipid raft integrity with *m*- β -cyclodextran significantly reduced vaccinia MV entry into HeLa cells (12). Since lipid rafts on the plasma membrane are known to act as platforms for receptor clustering, endocytosis, and signal transduction, we hypothesized that cellular proteins within plasma membrane lipid rafts mediate vaccinia MV entry into HeLa cells. Therefore, we isolated detergent-resistant domains from HeLa cells upon vaccinia MV infections and extracted proteins for quantitative proteomic analyses (55). Here, we investigated one of the raft-associated proteins, integrin β 1 (ITG β 1), for its role in vaccinia MV entry.

Integrins are a large family of cell surface receptors composed

of 18 different α and 8 different β subunits (35). Integrin β 1 is known to associate with multiple α subunits, including α 1-11 and α V, and is widely distributed in virtually all mammalian cell types (35, 38, 39, 60). Through interactions with the extracellular matrices, integrin β 1 regulates multiple intracellular kinase activation pathways. Our results demonstrate that integrin β 1 mediates MV entry in both mouse embryonic fibroblast (MEF) and HeLa cells and that vaccinia virus-induced phosphatidylinositol 3-kinase (PI3K)/Akt requires integrin β 1.

(R. Izmailyan conducted this research in partial fulfillment of the requirements for a Ph.D. from Academia Sinica, Taipei, Taiwan, Republic of China.)

MATERIALS AND METHODS

Cells and viruses. Two mouse cell lines, GD25 and GD25 β 1A, were kindly provided by Reinhard Fässler (Max Planck Institute of Biochemistry, Germany). The GD25 cell line was derived from integrin β 1 knockout (KO) embryonic stem cells (20). The stably transformed cell line GD25 β 1A resulted from the electroporation of GD25 cells with wild-type human integrin β 1 cDNA (48). HeLa, BSC40, GD25, and GD25 β 1A cells were cultured in Dulbecco's modified Eagle medium (DMEM) supplemented with 10% fetal bovine serum (FBS) and 2% penicillin-streptomycin (Gibco) in a 5% CO₂ incubator at 37°C. The Western Reserve strain of vaccinia MV (WR-VV) was purified through 25 to 40% sucrose gradients as previously described (32, 37). A recombinant WR-VV was also used in this study (8). It was constructed previously to express a dual gene cassette inserted into the thymidine kinase (tk) locus containing the luciferase

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(*luc*) gene driven by a viral early promoter and the *lacZ* gene driven by a viral late promoter (8).

Antibodies and reagents. Anti-integrin $\beta 1$ monoclonal antibodies (MAbs) Ts2/16 and 12G10 were purchased from Santa Cruz Biotechnology and Abcam, respectively, and 9EG7 and Mab13, rat MAbs, were acquired from BD Pharmingen. Anti-transferrin receptor (TfR) antibody (CD71) was obtained from AbD Serotec. Anti-paxillin antibody was purchased from BD Transduction Laboratories. Alexa Fluor 647-phalloidin was purchased from Invitrogen. Anti-phospho-Akt (Ser473) and anti-Akt antibodies were purchased from Cell Signaling Technology. Anti-phospho-focal adhesion kinase (FAK) (pY397) antibody was purchased from Invitrogen. Anti-FAK antibody was purchased from BD Biosciences. Anti-cyclophilin B (CypB) antibody was obtained from Santa Cruz Biotechnology. Anti- β -actin antibody was purchased from Sigma-Aldrich. Anti-A4 and anti-vaccinia MV (anti-VV) rabbit antibodies were previously described (30). Mouse MAb clone 2D5 against the vaccinia virus L1 protein was obtained from Y. Ichihashi (31). Bafilomycin A1 (BFLA), cycloheximide (CHX), and blebbistatin (Bleb) were purchased from Sigma-Aldrich. The PI3K inhibitor (LY294002) and Akt inhibitor (Akt IV) were purchased from Calbiochem. Laminin-1 (LN), fibronectin (FN), and poly-L-lysine (PLL) were purchased from Sigma-Aldrich. The CypB small interfering RNA (siRNA) duplex and the integrin $\beta 1$ siRNA duplex (AAUGUACCAACCGUAGCAUU) were purchased from Dharmacon Inc.

Biological network analysis. Cellular proteins identified in lipid rafts isolated from HeLa cells (55) were subjected to subcellular localization analyses with NCBI Gene Ontology. The “integrin $\beta 1$ (ITG $\beta 1$) signaling network” contains cellular proteins that are known to physically interact with integrin $\beta 1$ and was constructed by using ARIADNE Pathway Studio 7.0 software, which uses automated text-mining engines to extract information (Ariadne Genomics). Plasma membrane proteins identified in lipid rafts were compared with those in the integrin $\beta 1$ signaling network and were displayed in a graphical network by using the open-source software Cytoscape (57).

Virus entry assays. Several cell-based biological assays were used to quantify vaccinia MV entry into host cells based on previously established methods (8, 24, 61–63). MV particles bound to cells were quantified by vaccinia MV virion binding assays at 4°C for 60 min with anti-L1 antibody (2D5) (63). Viral core numbers present in the cytoplasm after membrane fusion were quantified by viral core-uncoating assays using an antibody against A4 (62). Luciferase assays driven by a viral early promoter were performed with cell lysates harvested at 2 h postinfection (p.i.), as described previously (8, 61). Acid bypass treatment, which forced cell-bound MV to fuse with the plasma membrane, was performed as previously described (24). In brief, HeLa cells were pretreated with 25 nM bafilomycin A1 or 50 μ M PI3K inhibitor at 37°C for 30 min, cooled at 4°C for 20 min, and subsequently infected with vaccinia MV at a multiplicity of infection (MOI) of 20 PFU per cell for 1 h. After washing, the infected cells were treated with neutral (pH 7.2) or acidic (pH 5) buffer for 5 min, incubated in growth medium, and fixed at 2 h p.i. These infected cells were permeabilized and stained with anti-core A4 antibody for confocal microscopy analyses as described previously (62).

Plaque formation on GD25 $\beta 1$ A and GD25 cells. Freshly confluent cells were infected with WR-VV (approximately 300 PFU per well in a 6-well plate) at 37°C for 1 h; washed and cultured in growth medium containing 1% agarose; fixed at 2 days p.i.; and stained with X-Gal (5-bromo-4-chloro-3-indolyl- β -D-galactopyranoside) as previously described (4). Alternatively, cells were pretreated with dimethyl sulfoxide (DMSO) or LY294002 (25 or 50 μ M) in serum-free DMEM prior to infection, and the inhibitors remained in cultures after infection until cell fixation and X-Gal staining as described above.

Integrin $\beta 1$ siRNA. HeLa cells were either mock transfected (si-control) or transfected with siRNA duplexes (20 nM) targeting either cyclophilin B (si-CypB) or integrin $\beta 1$ (si-ITG $\beta 1$) using the Lipofectamine 2000 reagent (Invitrogen) as described previously (30).

Confocal immunofluorescence microscopy. (i) Copatching experiments. The experiments for the copatching of integrin $\beta 1$ and vaccinia MV were performed as previously described (30, 58). In brief, HeLa cells seeded onto glass coverslips were infected with MV at an MOI of 50 PFU per cell for 1 h at 4°C, washed, and transferred to 12°C, where the cells were incubated with anti-VV antibody (1:500) and anti-integrin $\beta 1$ MAb (12G10) (1:1,000) for 1 h. Tetramethylrhodamine-conjugated goat anti-rabbit IgG (1:1,000) and fluorescein isothiocyanate (FITC)-conjugated goat anti-mouse IgG (1:1,000) were subsequently added for another 1 h of incubation prior to cell fixation for confocal microscopy. Cells were mounted in Vectashield medium (Vector Laboratories, Burlingame, CA), and images were collected with an LSM510 Meta confocal laser scanning microscope (Carl Zeiss) using a 63 \times objective lens.

(ii) Surface staining of integrin $\beta 1$ with virus in GD25 $\beta 1$ A cells. GD25 $\beta 1$ A cells were seeded (1.2×10^5 cells) onto glass coverslips in 12-well plates. The next day, cells were cooled at 4°C for 20 min and subsequently infected with vaccinia MV at an MOI of 60 PFU per cell for 1 h, washed 3 times in phosphate-buffered saline (PBS), fixed, and incubated with primary anti-integrin $\beta 1$ MAb (12G10) (1:1,000) and anti-VV rabbit antibody (1:500) for 1 h. Tetramethylrhodamine-conjugated goat anti-rabbit IgG (1:1,000) and FITC-conjugated goat anti-mouse IgG (1:1,000) were subsequently added for 30 min, and cells were analyzed by confocal microscopy.

Outside-in integrin signaling activation assay. Outside-in integrin signaling activation assays were performed as follows. (i) Glass coverslips in 24-well plates were coated with FN (10 μ g/ml), LN (20 μ g/ml), or PLL (100 μ g/ml) in PBS at 4°C. After 24 h, the dishes were blocked with 1% bovine serum albumin (BSA) in PBS at 37°C for 1 h. Serum-starved HeLa cells were seeded at a density of 8×10^4 cells into each well in serum-free DMEM; incubated at 37°C for 20, 30, 45, and 90 min; and lysed in cold lysis buffer (10 mM Tris [pH 7.4], 150 mM NaCl, 0.5% NP-40, 1% sodium deoxycholate, 0.1% SDS, 1% Triton X-100, 1 mM EGTA, and 1 mM EDTA with 1 \times protease inhibitor cocktail [tablets purchased from Roche Applied Science]). Samples were then prepared for immunoblot analyses with anti-phospho-Akt (1:1,000), anti-phospho-FAK (1:1,000), anti-Akt (1:1,000), and anti-FAK (1:1,000) antibodies. Alternatively, cells were fixed after 90 min of plating, permeabilized, and stained with anti-paxillin antibody (1:1,000). (ii) HeLa cells seeded for 90 min, as described above, were subsequently infected with vaccinia MV at an MOI of 5 PFU per cell at 37°C for 1.5 h and harvested for luciferase assays. (iii) HeLa cells were seeded, as described in above, for 15 min at 37°C. DMSO or blebbistatin (25 and 50 μ M) was added to cells, and the cells were incubated for another 45 min. The cells either were fixed, permeabilized, and stained with anti-paxillin antibody (1:1,000) or were infected with vaccinia MV at an MOI of 5 PFU per cell for 1.5 h and harvested for luciferase assays.

Flow cytometry. Cells were detached from the dishes with 20 mM EDTA, washed, and stained with anti-integrin $\beta 1$ MAb (9EG7) (1:1,000) in PBS containing 150 mM NaCl and 1% BSA for 1 h at 4°C. Samples were then washed 3 times, followed by incubation with FITC-conjugated goat anti-rat secondary antibody for 1 h at 4°C. Following a final wash, the cells were analyzed by fluorescence-activated cell sorting (FACS).

Vaccinia MV infections activated Akt phosphorylation. HeLa cells (7.5×10^4 cells/well) or GD25 $\beta 1$ A cells (1×10^5 cells/well) were cultured in 12-well plates for 24 h. Cells were serum starved at 37°C for 2 h (for HeLa cells) or 18 h (for GD25 $\beta 1$ A cells) and then stimulated with medium containing 20% FBS or purified vaccinia MV (at MOIs of 40 PFU per cell for HeLa cells and 60 PFU per cell for GD25 $\beta 1$ A cells) at 37°C for various times. Cell lysates were prepared with cold lysis buffer (10 mM Tris [pH 7.4], 150 mM NaCl, 0.5% NP-40, 1% sodium deoxycholate, 0.1% SDS, 1% Triton X-100, 1 mM EGTA, and 1 mM EDTA with 1 \times protease inhibitor cocktail [tablets purchased from Roche Applied Science]) for immunoblot analysis.

Statistical analysis. Statistical analyses were performed by using Student's *t* test with Prism software (GraphPad).

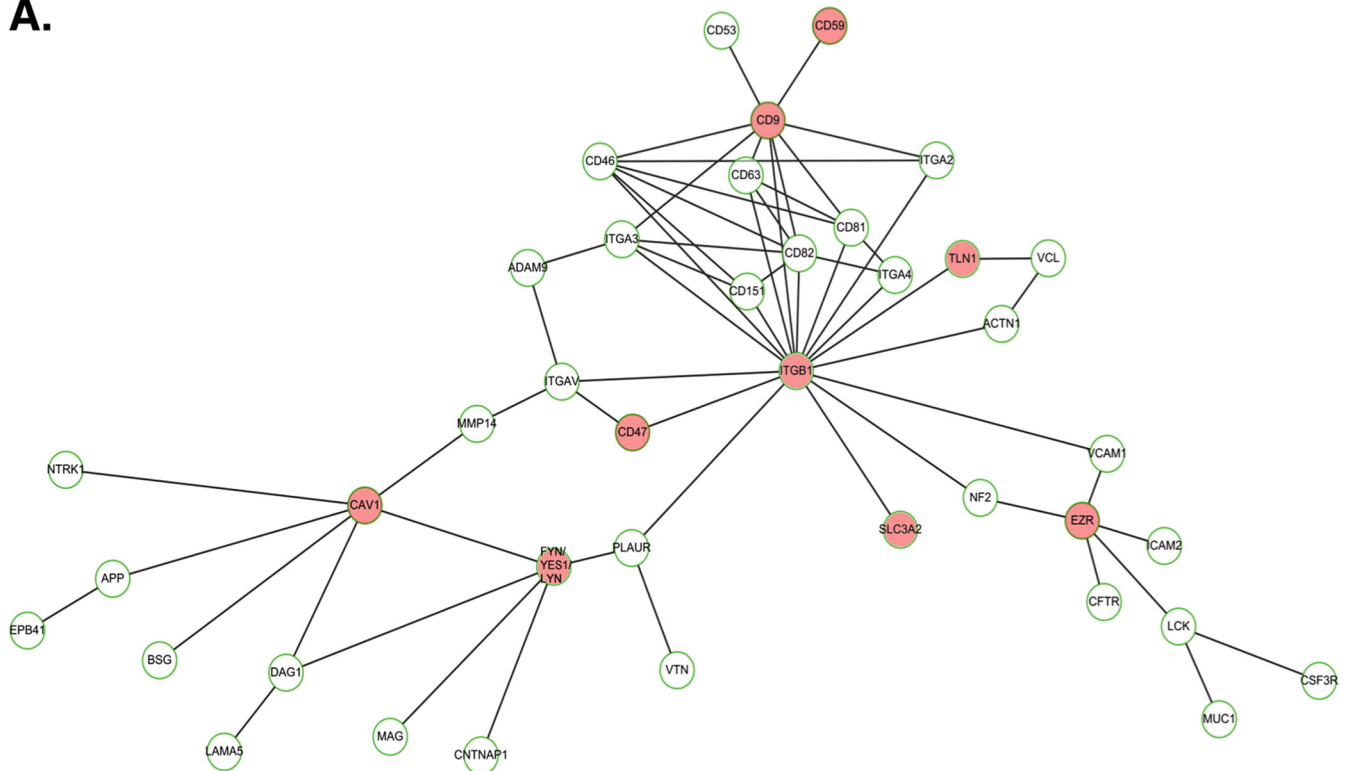
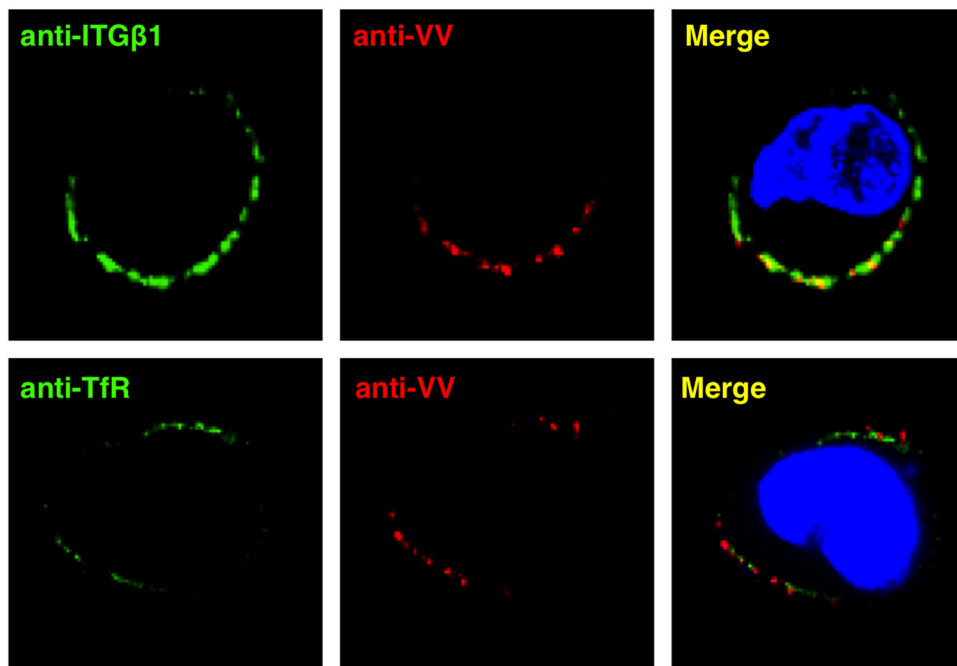
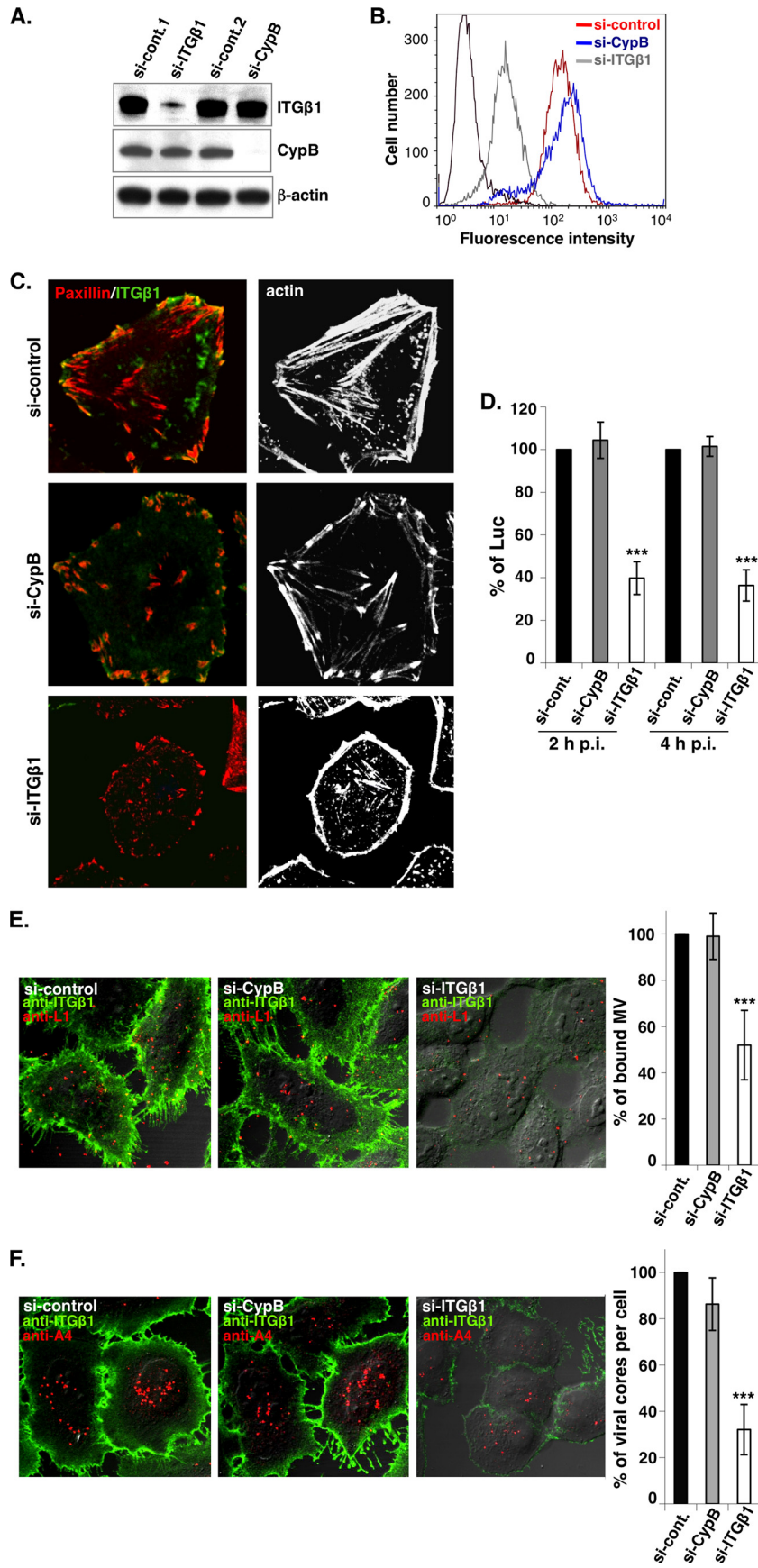
A.**B.**

FIG 1 Integrin $\beta 1$ was identified in lipid rafts and is associated with vaccinia MV on HeLa cells. (A) The hierarchical ITGB1 network, constructed by ARIADNE Pathway Studio 7.0 software (see Materials and Methods), shows integrin $\beta 1$ and its direct and indirect interacting membrane proteins. The nodes are labeled with gene symbols, and the red nodes represent the cellular proteins identified in the lipid raft fractions upon vaccinia MV infections (55). ITGB1, integrin $\beta 1$; TLN1, talin-1; EZR, ezrin; CAV1, caveolin-1; SLC3A2, CD98. (B) Vaccinia MV colocalizes with ITGB1 at the surface of HeLa cells. HeLa cells were infected with vaccinia MV at an MOI of 50 PFU per cell for 1 h at 4°C, washed, and transferred to 12°C, where the cells were incubated with rabbit anti-VV antibody (1:500) (red) and anti-integrin $\beta 1$ MAb (12G10) (1:1,000) (green) for 1 h for copatching as previously described (30). Alternatively, anti-VV antibody was incubated with anti-transferrin receptor (TfR) (green) MAb as a control. Tetramethylrhodamine-conjugated goat anti-rabbit IgG (1:1,000) and FITC-conjugated goat anti-mouse IgG (1:1,000) were subsequently added, and cells were fixed for confocal microscopy. DNA was visualized by 4'-6-diamidino-2-phenylindole (DAPI) staining (blue).



RESULTS

Integrin β 1 associates with vaccinia MV on HeLa cells. In order to identify cellular proteins within lipid raft microdomains, we previously performed stable isotope labeling for quantitative proteomic analyses and identified 570 cellular proteins. Those proteins with altered levels after vaccinia virus infection constitute about 3% of the total candidates and were described elsewhere previously (55). To complement the above-mentioned study, here we analyzed the remaining 97% of the “constitutive” raft-associated proteins whose quantifications fell within the range of a ratio of 1.0 ± 0.5 , i.e., a ratio that was considered not significantly changed by virus infection. These proteins were analyzed through biological network analyses in order to identify specific signaling complexes resident in the rafts. As described in Materials and Methods, biological network analyses (Fig. 1A) revealed the presence of integrin β 1 and its associated proteins, such as CD9, CD47, CD59, CD98, talin, ezrin, and Fyn/yes/lyn, suggesting a possibility that integrin β 1-mediated biological signaling may participate in vaccinia MV entry. We thus investigated whether integrin β 1 associates with vaccinia MV on the surface of infected HeLa cells. HeLa cells were infected with the vaccinia MV WR strain at an MOI of 50 PFU per cell for 60 min at 4°C and then washed and transferred to a temperature of 12°C. Anti-integrin β 1 (ITG β 1) and anti-vaccinia MV (VV) antibodies were subsequently added to cells for copatching (Fig. 1B), as previously described (30, 58). An anti-transferrin receptor (TfR) antibody was also included as a negative control. The results show that vaccinia MV on HeLa cells copatched with cell surface integrin β 1 but not with the transferrin receptor on HeLa cells (Fig. 1B), suggesting a role of integrin β 1 in vaccinia MV entry.

Vaccinia virus entry is reduced upon integrin β 1 knockdown in HeLa cells. To test whether integrin β 1 mediates vaccinia MV entry into HeLa cells, control siRNA (si-control) or siRNA targeting integrin β 1 (si-ITG β 1) or cyclophilin B (si-CypB) was transfected into HeLa cells, followed by harvesting for immunoblot analyses (Fig. 2A) and FACS analyses (Fig. 2B). The si-ITG β 1 and si-CypB constructs specifically knocked down (KD) the total amounts of integrin β 1 and cyclophilin B proteins, respectively (Fig. 2A). In addition, integrin β 1 expression on the cell surface was effectively reduced in si-ITG β 1 KD HeLa cells but not in si-control or si-CypB KD cells (Fig. 2B). si-ITG β 1 also affected integrin β 1-mediated cell adhesion, resulting in alterations in cell morphology and a disorganization of focal adhesions, as deter-

mined by actin and paxillin staining in si-ITG β 1 KD HeLa cells (Fig. 2C), confirming the specificity of si-ITG β 1. We then infected these HeLa KD cells with vaccinia MV at an MOI of 5 PFU per cell and harvested the cells at 2 or 4 h p.i. for viral early luciferase activity assays. The results showed that the rates of vaccinia virus infection of si-ITG β 1 KD HeLa cells were reduced to 40% (at 2 h p.i.) and 38% (at 4 h p.i.) of the infection rates seen for the si-control and si-CypB KD HeLa cells (Fig. 2D), suggesting that the reduction was not due to delayed kinetics. Finally, we performed vaccinia MV binding assays and virus core-uncoating assays as previously described (62, 63). The results showed that the rate of MV attachment was reduced to 52% in si-ITG β 1 KD HeLa cells (Fig. 2E) and that the rate of MV penetration was reduced to 32% (Fig. 2F), indicating that integrin β 1 is important for vaccinia virus entry at both the attachment and penetration steps.

Vaccinia virus entry is reduced in mouse cells lacking integrin β 1 expression. In addition to HeLa cells, we obtained a mouse cell line, GD25 β 1A, derived from an integrin β 1 KO cell line, GD25 (20), which expresses only human integrin β 1 (48). Indeed, integrin β 1 was detected on the surface of GD25 β 1A cells but not on the surface of GD25 cells by FACS analyses (Fig. 3A). GD25 β 1A and GD25 cells were infected with vaccinia MV at an MOI of 60 PFU per cell at 4°C for 60 min, washed, and fixed for immunofluorescence analyses. Abundant MV particles bound to GD25 β 1A cells and colocalized with surface integrin β 1 concentrated at cellular protrusions, whereas fewer MV particles (~34%) bound to GD25 cells (Fig. 3B). In addition, both GD25 β 1A and GD25 cells were infected with vaccinia WR-VV at different MOIs from 10 to 40 PFU per cell and were harvested to measure the early luciferase activity at 2 h p.i. or the late β -galactosidase (β -Gal) activity at 8 h p.i. (Fig. 3C). Data from both enzymatic activity assays and the core-uncoating assay (data not shown) revealed that vaccinia MV entry into GD25 cells was less efficient than entry into GD25 β 1A cells. We also performed plaque assays on GD25 β 1A and GD25 cells and stained the plaques with X-Gal. Consistently, vaccinia MV produced fewer plaques (~30%) in GD25 cells than in GD25 β 1A cells (Fig. 3D). Interestingly, plaques formed in GD25 cells also appeared smaller than those formed in GD25 β 1A cells, implying a role of integrin β 1 in vaccinia virus spreading among cells, although we did not pursue it further. Taken together, the results show that cell surface integrin β 1 mediates vaccinia MV infections.

FIG 2 Vaccinia virus entry is reduced in si-ITG β 1 KD HeLa cells. (A) Immunoblots of lysates prepared from mock-KD (si-Cont.1 and 2), si-ITG β 1 KD, and si-CypB KD HeLa cells using anti-ITG β 1 (Mab13), anti-CypB, or anti- β -actin antibodies. (B) Cell surface expression of ITG β 1 determined by flow cytometry. The KD HeLa cells described above (A) were stained with anti-ITG β 1 Mab (9EG7), followed by FITC-conjugated goat anti-rat secondary antibody, and analyzed by FACS. si-control is shown in red, si-CypB is shown in blue, and si-ITG β 1 is shown in gray. The background staining (shown in black) represents the si-control cells that were stained with the secondary antibody only. (C) Immunofluorescence analysis of ITG β 1, paxillin, and actin proteins in KD HeLa cells. The KD HeLa cells described above (A) were fixed and stained with anti-ITG β 1 Mab (Mab13), followed by FITC-conjugated goat anti-rat secondary antibody (green). These cells were subsequently permeabilized and stained with Alexa Fluor 647-phalloidin (white) and anti-paxillin antibodies, followed by tetramethylrhodamine-conjugated goat anti-mouse IgG (red). (D) Viral early luciferase assays. The KD HeLa cells described above (A) were infected with VV-WR at an MOI of 5 PFU per cell and harvested at 2 and 4 h p.i. for luciferase assays as described in Materials and Methods. The luciferase activity in the si-control cells was defined as 100%. The bars represent the standard deviations from three independent experiments. (E) Immunofluorescence analyses of vaccinia virus binding assay. The KD HeLa cells described above (A) were infected with purified vaccinia MV particles at an MOI of 20 PFU per cell for 60 min at 4°C, fixed, and stained with anti-ITG β 1 Mab13 (green) and anti-L1 Mab 2D5 (red). The cell-bound virions were quantified as described previously (63). (F) Immunofluorescence analyses of vaccinia virus core-uncoating assays. The KD HeLa cells described above (A) were infected as described above, cultured in the presence of cycloheximide (10 μ g/ml) for an additional 2 h at 37°C, fixed, and stained with anti-ITG β 1 Mab Mab13 as described above. These cells were subsequently permeabilized and stained with anti-A4 antibody, and internalized viral cores were quantified as described previously (62). Particle numbers counted in the si-control HeLa cells were used as 100%. The bars represent the standard deviations from five independent experiments. Statistical analyses in panels D to F were performed by using Student's *t* test in Prism software (GraphPad). The *P* value is shown (***, *P* < 0.0001).

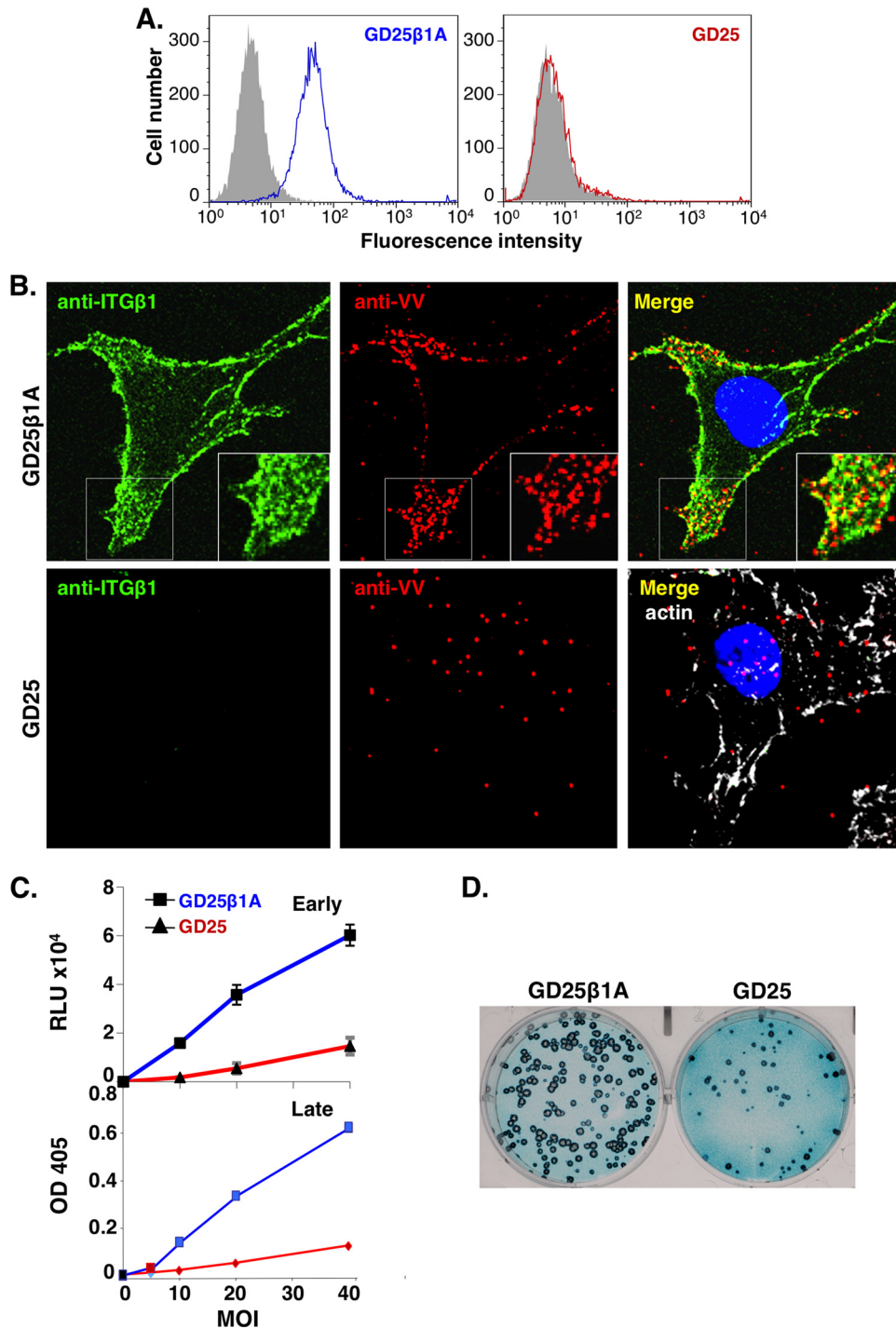


FIG 3 Vaccinia virus entry is reduced in mouse cells lacking integrin β 1 expression. (A) Flow cytometry analyses of cell surface expression of ITG β 1. GD25 β 1A (blue) and GD25 (red) cells were stained with anti-ITG β 1 MAb (9EG7), followed by FITC-conjugated goat anti-rat secondary antibody, and analyzed by FACS. The background staining (in gray) represents cell staining with the secondary antibody only. (B) Confocal immunofluorescence of vaccinia MV particles colocalized with integrin β 1 in GD25 β 1A and GD25 cells. GD25 β 1A and GD25 cells were infected with purified vaccinia MV particles at an MOI of 60 PFU per cell at 4°C for 1 h, washed, and stained with anti-ITG β 1 (12G10) (green) and anti-VV (red) antibodies, followed by FITC-conjugated goat anti-mouse IgG and tetramethylrhodamine-conjugated goat anti-rabbit IgG. GD25 cells were negative for ITG β 1 staining and were subsequently permeabilized and stained with Alexa Fluor 647-phalloidin (white) to mark the cell body. (C) GD25 β 1A and GD25 cells were infected with WR-VV at MOIs of 10, 20, and 40 PFU per cell and harvested at 2 h p.i. for luciferase assays (Early) and 8 h p.i. for β -Gal activity assays (Late). RLU, relative light units; OD 405, optical density at 405 nm. (D) Vaccinia MV formed more plaques in GD25 β 1A cells than in GD25 cells. Both cell types were infected with \sim 300 PFU of WR-VV, fixed, and stained for X-Gal at 2 days p.i. to visualize the blue plaques.

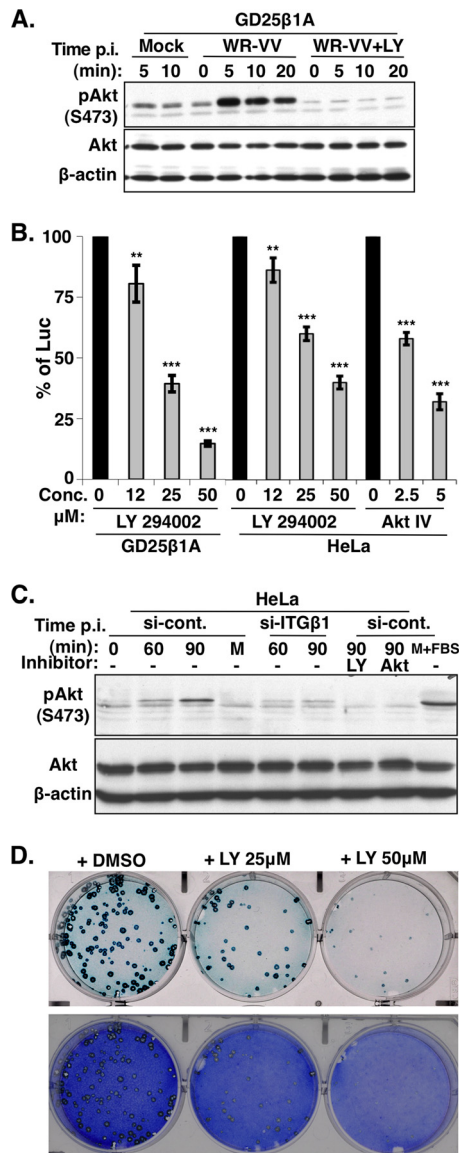


FIG 4 PI3K/Akt activation induced by WR-VV is required for virus entry. (A) WR-VV induced phosphorylation of Akt in GD25 β 1A cells. GD25 β 1A cells were serum starved, pretreated with DMSO or the PI3K inhibitor (LY294002), and either mock infected or infected with medium containing purified MV particles (WR-VV) at an MOI of 60 PFU per cell at 37°C. Cells were harvested at 0, 5, 10, and 20 min after the addition of virus for immunoblot analyses with anti-phospho-Akt (pAkt) (S473), anti-Akt, and anti- β -actin antibodies. (B) Viral early luciferase activity in GD25 β 1A and HeLa cells was blocked by PI3K/Akt inhibitors. Cells were pretreated with DMSO; the PI3K inhibitor (LY294002) at a concentration of 12, 25, or 50 μ M; or the Akt inhibitor (Akt IV) at a concentration of 2.5 or 5 μ M and subsequently infected with WR-VV at MOIs of 10 PFU (for GD25 β 1A cells) and 5 PFU (for HeLa cells) per cell and harvested at 2 h p.i. for luciferase assays as described above. The luciferase activity present in the DMSO-treated samples was used as 100%. The bars represent the standard deviations from three independent experiments. Statistical analyses were performed by using Student's *t* test in Prism software (GraphPad). *P* values are shown (**, *P* < 0.001; ***, *P* < 0.0001). (C) WR-VV-induced phosphorylation of Akt requires integrin β 1 in HeLa cells. The si-control and si-ITG β 1 KD HeLa cells, as described in the legend of Fig. 2A, were serum starved, pretreated with or without inhibitors of PI3K (LY294002) or Akt (Akt IV), and subsequently infected with purified MV particles (WR-VV) at an MOI of 40 PFU per cell at 37°C. Cells were harvested at 0, 60, and 90 min after the addition of virus for immunoblot analyses with anti-phospho-Akt (pAkt) (S473), anti-Akt, and anti- β -actin antibodies. The si-control HeLa

Vaccinia MV binds to integrin β 1 to induce PI3K/Akt activation, which leads to virus endocytosis in HeLa cells. Integrin β 1 was shown previously to mediate multiple inside-out and outside-in signaling pathways that play crucial roles in cell-to-cell and cell-to-matrix communications (28, 38, 46). The clustering of integrin at the cell membrane leads to kinase activation, which is followed by alterations to the cytoskeleton and receptor endocytosis. Since it was shown previously that PI3K and Akt are critical regulators downstream of integrin β 1 on GD25 β 1A cells (15, 48, 65), we wanted to test whether vaccinia MV binding to integrin β 1 activates PI3K/Akt signaling in GD25 β 1A cells and whether such kinase activation is critical for vaccinia MV entry.

GD25 β 1A cells were serum starved and subsequently infected with vaccinia MV at an MOI of 60 PFU per cell at 37°C for 5, 10, and 20 min and then harvested for immunoblot analyses with anti-phospho-Akt (Ser473) antibody. As shown in Fig. 4A, vaccinia MV stimulated the robust phosphorylation of Akt in GD25 β 1A cells as early as 5 min after the addition of virus, compared to that in medium alone. The pretreatment of GD25 β 1A cells with the PI3K inhibitor LY294002 completely abolished MV-induced Akt phosphorylation, showing that vaccinia MV infection triggered the activation of Akt through PI3K. To determine whether PI3K/Akt activation was required for vaccinia virus entry, we pretreated both GD25 β 1A and HeLa cells with inhibitors that blocked PI3K (LY294002) and Akt (Akt IV) activities, subsequently infected the cells with vaccinia MV, and measured viral early promoter luciferase activity at 2 h p.i. As shown in Fig. 4B, PI3K as well as Akt inhibitors readily reduced vaccinia MV infections in both GD25 β 1A and HeLa cells in a dose-dependent manner, showing that PI3K/Akt signaling is crucial for vaccinia MV entry into both cell lines. It is worth noting that GD25 cells were not suited to kinase signaling studies because it was previously shown that this cell line contains altered kinase regulation that compensates for the loss of integrin β 1 and that multiple tyrosine kinases are activated without integrin β 1 (3). Thus, in order to demonstrate that MV-induced PI3K activation requires integrin β 1, we infected si-control and si-ITG β 1 KD HeLa cells with vaccinia MV and harvested the cells at 0, 60, and 90 min for phospho-Akt immunoblot analyses. When these cells were infected with vaccinia MV, the phosphorylation of Akt was also induced in si-control HeLa cells although with slower kinetics than in GD25 β 1A cells (Fig. 4C). Most importantly, the phosphorylation of Akt was significantly reduced in si-ITG β 1 KD HeLa cells. Finally, GD25 β 1A cells were pretreated with 25 and 50 μ M LY294002 and infected with vaccinia MV for plaque formation assays, which showed a dosage-dependent reduction of plaque numbers with LY294002 and not the DMSO control (Fig. 4D), consistent with the data from the luciferase assays (Fig. 4B). Taken together, these results demonstrate that vaccinia MV-induced PI3K/Akt activation is mediated through integrin β 1 and that

cells stimulated with medium containing 20% fetal bovine serum (M+FBS) for 30 min were used as a control. (D) Vaccinia virus plaque formation was reduced with the PI3K inhibitor LY294002 in GD25 β 1A cells. GD25 β 1A cells were pretreated with DMSO or the inhibitor LY294002 at concentrations of 25 and 50 μ M for 1 h at 37°C and subsequently infected with WR-VV (~300 PFU/well), cultured in medium containing inhibitors, fixed at 2 days p.i., and stained with X-Gal to visualize the blue plaques. After photography, cells were subsequently stained with crystal violet to reveal the monolayer of cells.

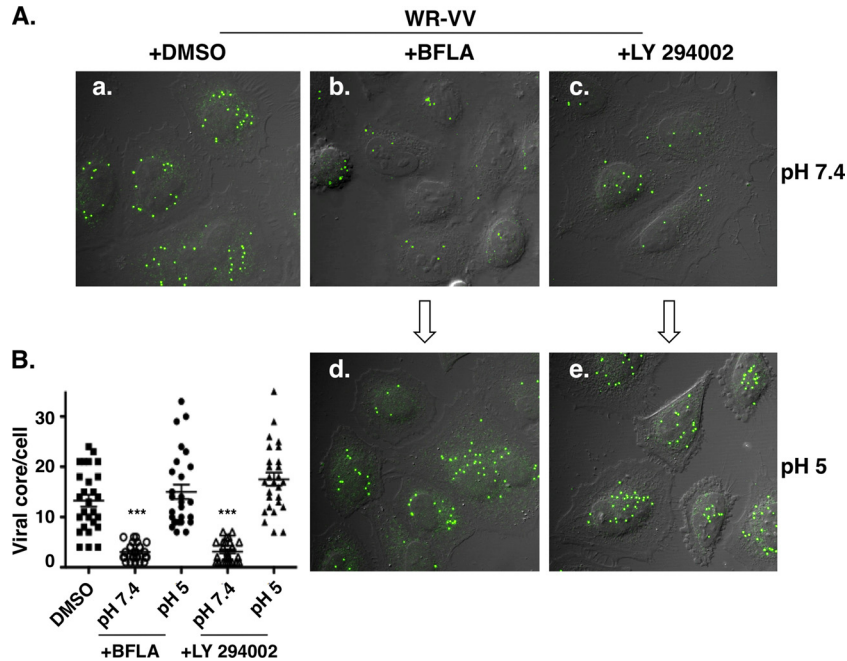


FIG 5 PI3K/Akt activation is required for vaccinia MV endocytosis. (A) Immunofluorescence analyses of virus-uncoating assays with HeLa cells. HeLa cells were pretreated with DMSO (a), 25 nM BFLA (b and d), or 50 μM PI3K inhibitor (LY294002) (c and e) and subsequently infected with purified MV particles at an MOI of 20 PFU per cell for 60 min at 4°C. Cells were then treated with neutral (pH 7.4) (a to c) or acidic (pH 5) (d and e) buffer for 5 min, washed, cultured at 37°C in medium containing 10 μg/ml cycloheximide for an additional 2 h, and analyzed by virus core-uncoating assays using anti-A4 antibody, as described in the text. (B) Quantification of viral core numbers per cell from each group (30 cells/group) described above (A). Statistical analyses were performed by using Student's *t* test in Prism software (GraphPad). The *P* value is shown (***, *P* < 0.0001).

such an activation is required for vaccinia virus entry and plaque formation.

To determine whether the PI3K inhibitor specifically blocks viral endocytic entry, we performed acid bypass experiments in which the cell-bound viruses were briefly treated with acidic buffer to force viral entry through plasma membrane fusion, as described previously (24). HeLa cells were pretreated with DMSO, bafilomycin A (a drug known to inhibit endosomal acidification), or the PI3K inhibitor LY294002 and infected with vaccinia MV at 4°C for 1 h. The unbound virions were washed, and the infected cells were briefly exposed to either neutral (pH 7.4) or acidic (pH 5) buffer for 5 min at 37°C. The cultures were then maintained for another 2 h in growth medium prior to fixation for viral core-uncoating assays as previously described (8). As shown in Fig. 5A, control HeLa cells treated with DMSO (Fig. 5Aa) were successfully infected, and abundant viral cores were detected in the cytoplasm. BFLA pretreatment significantly reduced viral core numbers in cells, confirming virus entry through a low-pH-dependent endocytic process (Fig. 5Ab). The pretreatment of HeLa cells with LY294002 also reduced numbers of viral cores in cells, suggesting that PI3K/Akt is important for vaccinia virus uncoating (Fig. 5Ac). Most importantly, the exposure of these cells to a low-pH buffer rendered MV entry through plasma membrane fusion resistant to inhibition by BFLA (Fig. 5Ad) and LY294002 (Fig. 5Ae). Quantifications of viral uncoated cores in each sample are shown in Fig. 5B. Taken together, these results demonstrate that PI3K/Akt activation induced by vaccinia MV is important for virus endocytosis in HeLa cells.

Outside-in activation of integrin β1 facilitates vaccinia MV entry. The fact that the integrin network is essential for cell viability,

migration, and growth suggests that vaccinia MV exploits the integrin/PI3K/Akt signaling pathway to modulate cellular environments preferable for viral entry and growth. We thus performed outside-in activation experiments to turn on the integrin-dependent signaling pathway in HeLa cells. HeLa cells were plated onto dishes precoated with the extracellular matrix proteins fibronectin (FN) and laminin (LN) for only a short time so that few extracellular matrix proteins were secreted from cells to induce integrin-mediated cell adhesion. As a negative control, HeLa cells were plated onto dishes precoated with poly-L-lysine (PLL), which mediates cell attachment through electrostatic interactions and is independent of integrins (53, 54). Cells were allowed to adhere and spread for 20, 30, 45, and 90 min before they were harvested for immunoblot analysis. As shown in Fig. 6A, the integrin-mediated phosphorylation of FAK and Akt was detected as early as 20 min after cell plating onto FN and LN but not onto PLL, showing a specific activation of integrin/PI3K/Akt through cell-matrix interactions. As expected, immunofluorescence staining with anti-paxillin antibody revealed the formation of focal adhesions in cells plated onto FN and LN but not onto PLL (Fig. 6B). When these cells were infected with vaccinia MV, the early luciferase activity level was higher in HeLa cells plated onto FN and LN than in those plated onto PLL (Fig. 6C). Altogether, these results demonstrate that the outside-in activation of integrin β1-mediated PI3K signaling is important for vaccinia MV entry into HeLa cells.

Integrin adhesome formation induced by outside-in activation was shown previously to be disrupted upon blebbistatin treatment (53). When the above-described outside-in activation procedures were performed with HeLa cells in the presence of 25 and 50 μM blebbistatin, the formation of focal adhesions was completely dis-

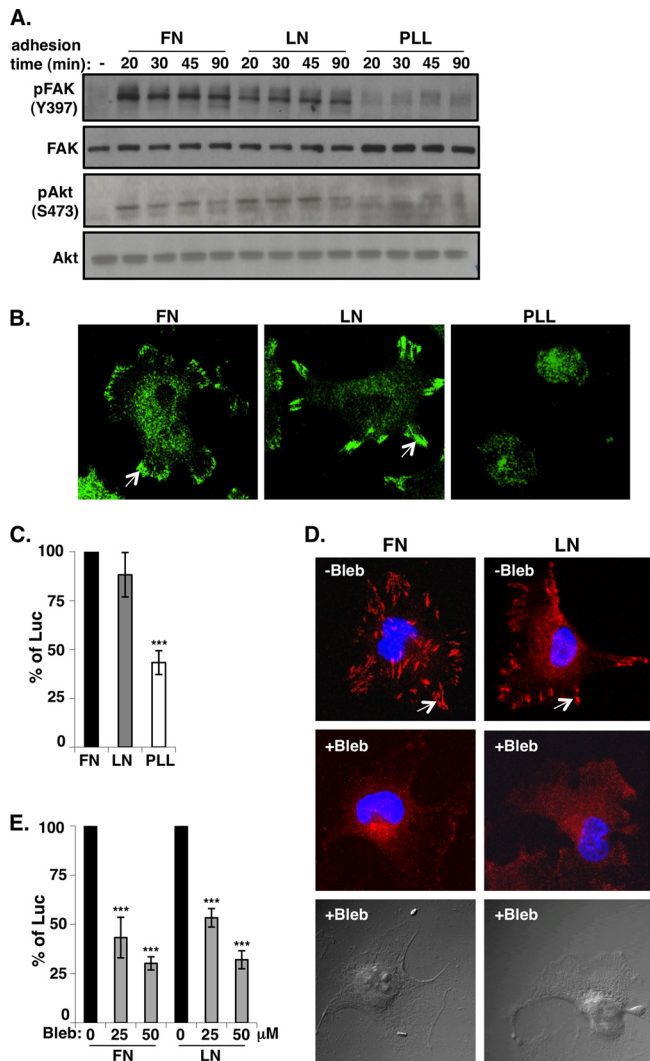


FIG 6 Outside-in integrin activation enhances vaccinia MV entry. (A) Cell adhesion to the extracellular matrix activates FAK and Akt phosphorylation. Serum-starved HeLa cells in suspension (–) were seeded onto fibronectin (FN), laminin-1 (LN), and poly-L-lysine (PLL) at 37°C and harvested at 20, 30, 45, and 90 min for immunoblot analysis with anti-phospho-FAK (pFAK) (Y397), anti-phospho-Akt (pAkt) (S473), anti-FAK, and anti-Akt antibodies. (B) Immunofluorescence staining of paxillin in HeLa cells seeded onto a different extracellular matrix. HeLa cells were seeded onto FN, LN, and PLL, as described above, for 90 min; fixed; permeabilized; and stained with anti-paxillin antibody, followed by FITC-conjugated goat anti-mouse IgG. (C) Viral early luciferase assays. HeLa cells were seeded as described above (B), infected with WR-VV at an MOI of 5 PFU per cell, and harvested at 1.5 h p.i. for luciferase assays as described in Materials and Methods. The luciferase activity in HeLa cells seeded onto FN was defined as 100%. The bars represent the standard deviations from three independent experiments. (D) Immunofluorescence staining of paxillin in HeLa cells treated with blebbistatin. HeLa cells were seeded onto FN and LN as described above (A) and incubated with 50 μ M blebbistatin (+Bleb) or DMSO (–Bleb) (see Materials and Methods) for 60 min. Cells were fixed and permeabilized, and the focal adhesions were stained with anti-paxillin antibody (red). Cell nuclei were stained with DAPI (blue). In panels B and D, white arrows identify areas of focal adhesion. (E) HeLa cells seeded as described above were treated with DMSO or blebbistatin (25 and 50 μ M) and subsequently infected with WR-VV at an MOI of 5 PFU per cell and harvested at 1.5 h p.i. for luciferase assays. The luciferase activity in DMSO-treated HeLa cells was defined as 100%. The bars represent the standard deviations from three independent experiments. Statistical analyses in panels B, C, and E were performed by using Student's *t* test in Prism software (GraphPad). The *P* value is shown (***, *P* < 0.0001).

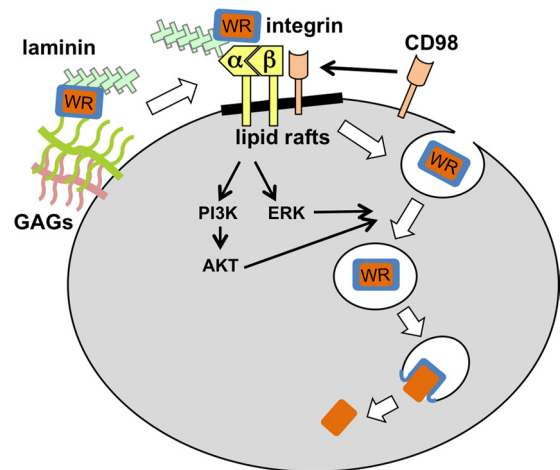


FIG 7 Current model of integrin $\beta 1$ in vaccinia MV entry. The vaccinia MV WR strain binds to cell surface glycosaminoglycans (GAGs) and laminin, which induce further interactions between MV and cellular surface receptor integrin in lipid rafts. The subsequent recruitment of the cellular membrane protein CD98 and the activation of downstream kinases such as PI3K/Akt and ERK lead to the endocytosis of vaccinia MV.

persed (Fig. 6D). These drug-treated HeLa cells were subsequently infected with vaccinia MV, and a dosage-dependent reduction of viral entry was observed, compared with the mock-treated cells (Fig. 6E). These results suggested that the recruitment of cellular proteins to form integrin adhesomes, which was blocked by blebbistatin treatment, is important for vaccinia MV entry.

DISCUSSION

In this study, we demonstrated the role of the cellular protein integrin $\beta 1$ in vaccinia MV entry by several experimental criteria. First, we showed that integrin $\beta 1$ and vaccinia MV colocalized on plasma membrane lipid rafts and that the knockdown of integrin $\beta 1$ in HeLa cells and the knockout of integrin $\beta 1$ in mouse cells reduced vaccinia MV attachment. Second, we showed that the requirement of integrin $\beta 1$ is not limited to virus attachment, since integrin $\beta 1$ also mediated the activation of PI3K/Akt for subsequent MV endocytosis. We further showed that adhesome formation activated by the interaction of the extracellular matrix and integrin is also critical for vaccinia MV entry. Taken together, these data demonstrated that cell surface integrin $\beta 1$ in lipid rafts serves as a signaling platform which activates downstream kinases and cellular adhesome formation, both of which are important for vaccinia MV entry.

Vaccinia virus infection is known to activate multiple kinases, including ERK (18), PKC, and PAK1 (43, 45). However, the virus-induced upstream signal that leads to kinase activation remains unknown. Our study shows that the binding of vaccinia MV to integrin $\beta 1$ activated the PI3K/Akt kinase pathway. We also found that the KD of integrin $\beta 1$ led to a reduction of ERK activity in HeLa cells and that the addition of the ERK inhibitor PD98059 interfered with vaccinia MV entry (R. Izmailyan, unpublished results). Although we did not investigate PAK1 activation in this study, interestingly, PAK1 was previously shown to be activated by integrin $\beta 1$ (50). Taking these findings together, we conclude that vaccinia MV interacts with integrin $\beta 1$ to activate a downstream kinase network to induce viral endocytosis (Fig. 7). At this point,

we have not been able to identify any viral envelope protein specifically associated with integrin $\beta 1$ in coimmunoprecipitation experiments. It could be that vaccinia MV interacts with multiple cellular receptor proteins during virus entry and that each receptor interaction is transient and dynamically regulated (see below).

Although our study focused primarily on integrin $\beta 1$, the significance of the involvement of integrin in vaccinia virus entry goes beyond this molecule. Blebbistatin, an inhibitor of myosin II light-chain ATPase (41), was recently shown to interrupt integrin adhesome formation (53), and in our experiments, blebbistatin-treated HeLa cells lost the outside-in activation of integrin and reduced vaccinia virus endocytosis. This finding implies that other components of the integrin adhesome may be involved in vaccinia virus endocytosis. Indeed, a recent study conducted in our laboratory by Schroeder et al. (55) demonstrated that CD98, a type II membrane protein found in lipid rafts (55) and a component of the integrin adhesome (53), is required for vaccinia MV endocytosis (55). The KD of CD98 does not affect vaccinia MV binding to cells but reduces virus endocytosis and core uncoating (55), suggesting that it participates in vaccinia virus entry at a step subsequent to integrin $\beta 1$. It is worth noting that CD98 was reported previously to promote integrin-dependent signaling, leading to the activation of FAK, PI3K, Akt, and Rac (22, 23, 70). Currently, our model proposes that vaccinia MV binds to integrin $\beta 1$, recruits CD98 in rafts, activates kinases, and induces cytoskeleton rearrangements to trigger MV endocytosis in cells (Fig. 7). Finally, we noticed that vaccinia MV formed smaller plaques on GD25 cells than those on GD25 $\beta 1A$ cells. The KD of integrin $\beta 1$ in HeLa cells also reduced MV and EV production in a low-MOI infection, implying that vaccinia virus spreading among cells may also require integrin $\beta 1$ (data not shown). Further studies are needed to explore this possibility.

Integrins are involved in the entry of many viruses (2, 5, 19, 21, 26, 34, 36, 44, 59, 64) and bacteria (16, 27, 40). The pathogen-host interaction may promote the clustering of integrins and focal adhesion formation but subvert the downstream signaling network to modify membrane traffic and cytoskeletal dynamics to meet their own needs (1, 17). Several unique properties of integrins may explain why they are so frequently targeted for bacterial and viral invasions. Integrins are ubiquitously expressed on virtually all cells. They regulate both outside-in and inside-out signaling to maintain important cell functions related to cell adhesion, migration, and survival. Furthermore, integrins act as global regulators of endocytosis, affecting the intracellular trafficking of growth factor receptors and endosome localization in cells (67). A genome-wide screen of endocytosis regulators identified adhesion-mediated molecules that support cross talk between cell adhesion regulation and endocytic activity (13). The fate of endocytosed vaccinia MV is not well understood. Recently, it was shown that integrin internalization is mediated through macropinocytosis. Internalized integrins were found to transit through early endosomes to recycling endosomes and then back to the cell surface to form new adhesions (25). Currently, we are investigating whether integrin $\beta 1$ regulates the intracellular trafficking of vaccinia virions through the same route prior to virus-cell membrane fusion. More studies will be needed in the future to dissect the structure-function relationship of integrins in vaccinia virus entry.

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