Primary human leukocyte subsets differentially express vaccinia virus receptors enriched in lipid rafts

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Abstract

Poxviruses including vaccinia virus (VV) and canarypox virus (ALVAC) do not indiscriminately infect all cell types of primary human leukocytes (PHLs) they encounter, but instead demonstrate an extremely strong bias towards infection of monocytes and monocyte-lineage cells. We studied the specific molecular events that determine the VV tropism for major PHL subsets including monocytes, B cells, neutrophils, NK cells, and T cells. We found that VV exhibited an extremely strong bias of cell surface protein-dependent binding to monocytes, B cells, and activated T cells to a similar degree, and to neutrophils to a much lesser extent. Resting T cells and resting NK cells exhibited only trace amounts of VV binding. Activated T cells, however, became permissive to VV binding, infection, and replication, while activated NK cells still resisted VV binding. VV binding strongly co-localized with lipid rafts on the surfaces of all VV binding-susceptible PHL subsets, even when lipid rafts were relocated to cell uropods upon cell polarization. Immunosera raised against detergent-resistant membranes (DRMs) from monocytes or activated T cells, but not resting T cells, crossly and effectively blocked VV binding to and infection of PHL subsets. CD29 and CD98, two lipid raft-associated membrane proteins that had been found to be important for VV entry into HeLa cells, had no effect on VV binding to and infection of primary activated T cells. Our data indicates that PHL subsets express VV protein receptors enriched in lipid rafts, and that receptors are crossly presented on all susceptible PHLs.
Introduction

In general, all viruses must bind to their receptors on the surface of target cells to initiate infection. Virus–receptor interactions determine the cell type, organ specificity, and host range, and therefore constitute an interspecies barrier. Poxviruses are a family of large, complex, enveloped DNA viruses that show species specificities (1, 2), i.e., variola virus is a strict human-specific pathogen that causes smallpox in humans only (1), and myxoma virus is a rabbit-specific poxvirus that causes a lethal disease (myxomatosis) in rabbits only (1, 2). However, the molecular basis underlying the strict species barrier for poxviruses remains mysterious. In particular, no specific cellular receptor for any poxvirus has yet been identified. Poxviruses infect a wide variety of cell lines in culture, leading to the presumptions that either specific receptors for these viruses may not be required, or that conserved and ubiquitous receptors may be widely distributed on the surface of diverse cell types (1). These conjectures may have impeded attempts to identify cellular receptor(s) that mediate poxvirus binding and infection. However, recent reports from our group and others have shown that vaccinia virus (VV), the prototypical member of the poxvirus family, and canarypox virus (ALVAC) do not indiscriminately infect all cell types of primary human hematopoietic cells they encounter, but instead demonstrate an extremely strong preference for infection of monocytes among peripheral blood mononuclear cells (PBMCs) and monocyte-lineage cells in the bone marrow (3, 4). Significantly, expression of VV receptor(s) can be induced de novo on primary human T cells upon T cell activation (3). As a consequence, activated T cells
become susceptible to VV binding, infection and replication, in contrast to resting T cells that are non-permissive to VV binding (3, 4). These receptors are likely proteins because inhibitors of transcription (actinomycin D), protein synthesis (cycloheximide) and intracellular protein transport (brefeldin A) significantly reduce VV binding to activated primary human T cells, and also treatment of primary human monocytes or activated T cells with trypsin or pronase completely diminishes VV binding and infection (3).

Poxviruses not only bind to and infect monocytes but also use these cells to initiate a systematic infection. A recent report using high doses of variola virus, the most virulent member of poxvirus family, to infect Cynomolgus macaques in an attempt to develop an animal model of smallpox has demonstrated that variola virus is disseminated by means of monocytic cell-associated viremia (5), suggesting that monocytes play a significant role in the initiation of systematic infection. Monocytes may use putative viral receptors to grab infectious variola virus particles and then disseminate them to uninfected cells and tissues, resulting in a generalized infection. However, the specific molecular events that determine poxvirus bias towards monocyte binding and infection remain unclear. In the present study, we investigated the susceptibility of major subsets of primary human leukocytes (PHLs) to VV binding and infection. Our data demonstrate that PHL subsets express and share protein VV receptors that are enriched in lipid rafts on the cytoplasmic membrane, and that VV receptors are induced de novo on certain but not all PHL subsets.

Materials and methods
Antibodies and flow cytometric analysis

The following anti-human monoclonal antibodies (mAbs) or polyclonal Abs (pAbs) conjugated with fluorochrome were purchased from BD PharMingen (San Diego, CA): anti-CD3^APC, anti-CD4^PerCP, anti-CD8^PE, anti-CD14^APC, anti-CD16^PE, anti-CD19^PE, anti-CD56^PE, and matched-isotype control Abs conjugated with FITC, PE, PerCP, or APC. Anti-human neutrophil lipocalin (HNL) and anti-human CD66b^PE Abs were purchased from Novus Biologicals (Littleton, CO) and Biolegend (San Diego, CA), respectively. Rabbit pAbs against full-length human Integrin beta-1 (CD29) and rabbit pAbs against human amino acid transporter SLC3A2 (CD98) were purchased from Abnova (Taipei, Taiwan) and Thermo Fisher Scientific (Pittsburgh, PA), respectively. Isolated PHL subsets including monocytes, B cells, T cells, neutrophils, and NK cells were subjected to VV binding and surface staining with different combinations of Abs, followed by flow cytometric analysis (FACS) using a BD FACSCalibur (BD Biosciences, San Diego, CA). The data were analyzed using FlowJo software (TreeStar, San Carlos, CA). Appropriate isotype controls were used at the same protein concentration as the test Abs and control staining was performed during every FACS.

Preparation of human primary leukocyte subsets

Whole blood samples were collected from healthy donors. Written consent was obtained from each participant and all investigational protocols were approved by Institutional Review Boards for Human Research at the Indiana University School of Medicine (Indianapolis, IN). PBMCs were separated from blood samples by gradient centrifugation on Ficoll-Hypaque (Amersham Pharmacia Biotech AB, Uppsala, Sweden).
Monocytes, B cells, and NK cells were then purified by negative isolations using Ab-conjugated magnetic beads in the Monocyte, B cell, and NK cell Negative Isolation Kits (Dynal, Oslo, Norway). The resulting cell preparations contained more than 95% desired cell types assessed by CD14, CD3, CD4, CD8, CD19, or CD56 staining and FACS.

Resting T cells were isolated from the PBMCs using the Pan T Cell Isolation Kit II (Miltenyi Biotec, Auburn, CA), which yielded >95% purity of CD3+ T cells. CD3+ T cells were activated by incubating with anti-CD3/anti-CD28 Abs-coated magnetic beads (Life Technologies, Carlsbad, CA), and allowed to culture in RPMI 1640 medium supplemented with 10% fetal bovine serum (FBS) and 2 mM L-glutamine (complete RPMI 1640 medium). Neutrophils were isolated from whole blood from healthy donors by density gradient separation in Lympholyte-Poly solution (Cedarlane Labs, Hornby, ON) to isolate polymorphonuclear cells, followed by treatment with a hypotonic solution to lyse red blood cells. Neutrophil purity was >98% as determined by FACS analysis of HNL+CD66b+ cells.

**VV preparation, binding and infection**

EGFP-VV and vA5L-YFP, two recombinant viruses of the Western Reserve (WR) VV strain, were obtained from Dr. Yewdell JW. and Dr. Moss B. (NIH, Bethesda, MD). EGFP-VV contains a chimeric gene encoding the influenza virus nucleoprotein, the ovalbumin SIINFEKL peptide, and enhanced green fluorescence protein (EGFP) regulated by the P7.5 early-late promoter(6). EGFP expression was used in this study to monitor VV infection. The vA5L-YFP virions were constructed with VV core protein A5L fused with yellow fluorescence protein (YFP), and can be directly visualized by
fluorescence microscopy or FACS(7). Viral stocks including EGFP-VV and vA5L-YFP were grown and titrated in primary chicken embryo fibroblasts (Charles River Laboratories, Wilmington, MA) or the monkey kidney cell line CV-1 (ATCC, Manassas, Virginia) in Dulbecco's minimal essential medium (DMEM) supplemented with 10% FBS and 2 mM L-glutamine. VV intracellular mature virions (IMV) were purified by 24 – 40% sucrose gradient as previously described(8), and the viral titers were determined by a plaque assay in CV-1 cells(8).

To determine the susceptibility of individual PHL subsets to VV binding, vA5L-YFP virions were incubated with isolated monocytes, B cells, resting T cells, activated T cells, neutrophils, resting or activated NK cells at an multiplicity of infection (MOI) of 10 at 4°C for 30 min in complete RPMI 1640 medium, a condition that permits virus binding but not entry(3, 8, 9). After extensive washing with ice-cold phosphate-buffered saline (PBS), cells were fixed with 1% paraformaldehyde (PFA) and then subjected to confocal microscopy analysis and FACS to determine VV binding at single cell and whole cell population levels, respectively. TA3 cells (a mouse B cell hybridoma cell line) that were previously shown not to bind with VV were used as a negative control of VV binding(8, 10).

VV-EGFP was used to monitor the susceptibility of individual PHL subsets to VV infection as previously described(4). Infection intensity was calculated by the percentage of EGFP-positive cells or the median fluorescence intensity (MFI).
Cells were incubated with cholera toxin subunit B (CTB) conjugated with Alexa Fluor 647 (Life Technologies, Carlsbad, CA) at 4°C for 20 min to stain ganglioside M1 (GM1) lipid rafts. For CTB-patching, cells were treated with 1:100 diluted goat anti-CTB pAbs (Millipore, Darmstadt, Germany) in 2% FBS/PBS for 30 min on ice, and then incubated at 37°C for 20 min as previously described(11). After washing, cells were then incubated with 10 - 30 pfu/cell of vA5L-YFP for 30 min at the binding condition as described above. Cells were fixed with 2% PFA, permeabilized with 0.1% saponin, and then incubated with phalloidin conjugated with Alexa Fluor 546 (Life Technologies, Carlsbad, CA) to stain F-actin. Cells were then adhered to poly-L-lysine-coated coverslips, mounted onto glass slides using ProLong Gold Antifade Reagent (Life Technologies, Carlsbad, CA) containing 4',6-diamidino-2-phenylindole (DAPI) dye for fluorescent staining of DNA content and nuclei. Cells were analyzed using an Olympus FV1000-MPE confocal/multiphoton microscope fitted with a 60x objective. For each field, a Z-series of images was collected and the number of virions per cell was counted on a 3D image of cells constructed using the open source software FluoRender 2.7 (University of Utah) and ImageJ 1.44p (NIH, Bethesda, MD).

**Polarization of primary human leukocyte subsets**

Individual PHL subsets were treated with various agents to induce membrane polarization and lipid raft relocation. Briefly, isolated monocytes were incubated with 100 ng/mL granulocyte/macrophage colony stimulating factor (GM-CSF) (BioVision, Milpitas, CA) for 24 h(12), B cells were incubated with 100 ng/mL SDF-1 (Biolegend, San Diego, CA) over rhICAM-2 (fc)-coated coverslips for 30 min(13), activated T cells were adhered
to anti-CD44 coated coverslips for 30 min(14)(15), and neutrophils were treated with 10
nM of the bacterial peptide fNLPNTL (Bachem, Torrence, CA) for 5 min(16). After
treatment, all cell types were fixed with 2% PFA, and subjected to VV binding and the
fluorescence microscopy assay.

**Immunosera raised against cell membrane extracts or whole cells**

All animal experimentation was conducted following the NIH guidelines for
housing and care of laboratory animals and performed in accordance with Indiana
University Institutional regulation after review and approval by the institutional Animal
Care and Use Committee at Indiana University (approved IACUC protocol number: MD-
0000003309-R). Female BALB/c mice, 6-8 weeks of age, from the Jackson
Laboratories (Bar Harbor, ME) were subjected to immunization through intraperitoneal
injections (i.p.). Mice were divided into nine groups with 3 mice per group, and then
subjected to immunization with: (1) detergent-resistant membranes (DRMs), (2) crude
membrane extracts (CMEs), or (3) whole cells. These immunogens were prepared from
either 40x10^6 monocytes, resting T cells, or activated T cells from the same blood
donors. DRMs and CMEs were prepared from each type of these cells as previously
described(17). Briefly, 40x10^6 cells were lysed with 1% Triton X-100/PBS plus 1x
Protease Inhibitor Cocktail (Fisher Scientific, Pittsburgh, PA) at 4°C for 1 h. Lysates
were clarified with centrifugation at 1,000 x g for 10 min, and the resulting supernatants
were mixed with sucrose to make 45% sucrose solutions which were added to the
bottom of an ultracentrifuge tube. Equal volumes of 35% and 5% sucrose/PBS were
sequentially added to the tube to make discontinuous gradients. The tube was
centrifuged at 166,000 x g at 4°C for 18 h in an Optima LE-80K ultracentrifuge (Beckman Coulter, Brea, CA). The light-scattering band near ~20% sucrose (DRMs) was harvested, diluted in PBS containing 1x Protease Inhibitor Cocktail, and then centrifuged at 166,000 x g at 4°C for 2 h. The DRM pellet was homogenized in PBS using a Dounce homogenizer. For CME preparation, 40x10^6 cells of each subset in PBS with 1x Protease Inhibitor Cocktail were disrupted with a Dounce homogenizer. After clarification, the resulting supernatant was centrifuged at 120,000 x g, and the pellet was resuspended in PBS. Primary immunizations were followed by two immunization boosts on day 14 and day 28. Two weeks after the last immunization boost, animals were anesthetized with isoflurane and sacrificed by bleeding. Sera were collected from mice in each group.

An ELISA assay was developed to titrate Abs against human CD55, a glycosylphosphatidylinositol (GPI) protein anchored in cell lipid rafts(18), to evaluate the immunization efficacy. Briefly, microplates were coated with recombinant CD55 protein at 0.1 ug/mL (R&D Systems, Minneapolis, MN). After washing and blocking with 5% FBS/PBS, plates were incubated with serially diluted immunosera, followed by addition of anti-mouse IgG mAb conjugated with horseradish peroxidase (HRP). Pooled pre-immunization mouse sera were used as Ab-negative controls. Absorption was read at a wavelength of 450 nm by a plate spectrophotometer (BioTek, Winooski, VT).

Knockdown of CD29 and CD98 in HeLa cells and activated T cells

Dharmacon Smartpool Accell siRNA constructs against human CD29 and CD98 were purchased from Thermo Fisher Scientific (Pittsburgh, PA). These siRNA
constructs were transfected into cells using the Amava Nucleofector system (Lonza, Basel, Switzerland) according to the manufacturer’s instructions. Briefly, 150 - 300 nM of each siRNA mixture was used per 5 x 10^6 activated T cells or per 1 x 10^6 HeLa cells. Transfected cells were allowed to culture for 48 h, and then subjected to Western blot or FACS using rabbit pAbs against human CD29 or CD98 to analyze knockdown of human CD29 or CD98. These cells were also subjected to VV binding and infection to determine the effects of CD29 and CD98 on VV binding and entry.

Statistical analysis

Data were analyzed using Tukey’s Post-hoc ANOVA test and the Student t-test. p < 0.05 was considered statistically significant.

Results

VV differentially binds to primary human leukocyte subsets

The majority of studies investigating the entry of VV into host cells have focused on single-enveloped VV IMV particles since they are the most abundant (>98%) and maintain their membrane integrity after freezer storage (19, 20). The IMV particles of vA5L-YFP or EGFP-VV were therefore used in this study. Isolated monocytes, B cells, neutrophils, resting T cells, and NK cells were incubated with vA5L-YFP particles at the binding condition (4°C for 30 min) to study VV binding profiles for these PHL subsets. At an MOI of 10, vA5L-YFP bound to 76 ± 10%, 71 ± 9%, 28 ± 2%, 3 ± 2% and 2 ± 2% of monocytes, B cells, neutrophils, resting T cells, and NK cells, respectively (Fig. 1A, 1B).
These values were the results of mean ± standard deviation (SD) from six healthy blood donors. VV binding to monocytes, B cells, and neutrophils was not affected by soluble heparan sulfate (HS) at 10 μg/mL (Fig. 1A, 1B), an optimal concentration that completely blocks VV non-specific binding to HS glycosaminoglycan (GAG) side chains of cell surface proteoglycans of BSC40 cell line(21). In contrast, HS at 10 μg/mL completely eliminated the trace amount of VV binding to resting T cells and NK cells (Fig. 1A, 1B), suggesting that the binding is GAG-dependent. VV binding to monocytes, B cells and neutrophils was markedly reduced by trypsin treatment (Fig. 1A, 1B). Next, we investigated whether activated T cells become sensitive to VV binding to confirm that VV receptor(s) can be induced de novo upon T cell activation(3). We found that VV binding was correlated to the degree of T cell activation (Fig. 1C, 1D). Similar to monocytes, B cells and neutrophils, VV binding to activated T cells was markedly reduced by treatment with trypsin, but not HS (Fig. 1E). In contrast, activated NK cells remain non-permissive to VV binding (Fig. 1F). These results indicate that VV binding to monocytes, B cells, neutrophils, and activated T cells is mediated by protein VV receptors independent of HS GAGs, and that these receptors are induced upon activation of T cells, but not NK cells.

To visualize VV binding at the single cell level, we incubated monocytes, resting T cells, activated T cells and TA3 cells with vA5L-YFP particles at the binding condition, and then used confocal microscopy to examine VV binding. At an MOI of 10, vA5L-YFP bound to monocytes at ~39 virions per cell (mean from 100 cells counted) (Fig. 1G, 1H), whereas VV barely bound to resting T cells (0.05 virions per cell, mean from 100 cells counted) which has no difference with the results of vA5L-YFP binding to TA3 cells that
were previously shown not to bind with VV(8, 10). After activation with anti-CD3/anti-
CD28 mAbs-coated magnetic beads for 3 days, activated T cells became sensitive to
VV binding with a similar binding degree to that of monocytes (Fig. 1G, 1H).

To determine whether VV binding was correlated to its infection tropism, we
infected PHL subsets with EGFP-VV at an MOI of 10 for various durations. We
confirmed that VV preferentially infected monocytes where 65 ± 8% (n=6) cells became
EGFP-positive 6 h post-infection, whereas 15 ± 5% (n=6) of activated T cells became
EGFP-positive 24 h post-infection (Fig. 1I, 1J). In contrast, only 4 ± 2% (n=6) B cells
were infected, and neutrophils and resting T cells resisted VV infection as only trace
amounts of these cells were EGFP-positive 24 h post-infection (Fig. 1I, 1J). VV infection
of monocytes and activated T cells was significantly reduced by trypsin treatment, but
not HS (Fig. 1I, 1J). Further analysis of monocyte subpopulations revealed that
"classical" CD14<sup>high</sup>CD16<sup>-</sup> monocytes received the vast majority of bound virus in
binding experiments and were sensitive to VV infection when compared to "patrolling"
CD14<sup>low</sup>CD16<sup>+</sup> monocytes (Fig. 1K). Two previous studies have demonstrated that VV-
infected primary human monocytes do not produce either viral late-gene products or
viral DNA copies, indicating that VV undergoes abortive infection in primary human
monocytes(3, 22). Our data together with these results indicate that: (1) monocytes are
the most sensitive PHL subset to VV binding and infection, but the infection is abortive;
(2) B cells and neutrophils are sensitive to VV binding, albeit at different degrees, but
non-permissive to VV infection; (3) NK cells (both resting and activated states) and
resting T cells resist VV binding and infection; and (4) activated T cells are the only cell
type among PHLs to permit VV to complete the whole cycle of binding, infection and replication.

**VV strongly binds to lipid rafts on the surface of all susceptible PHL subsets**

Lipid rafts (also called DRMs) play a critical role in VV entry into cell lines (23). VV particles colocalize with lipid rafts on HeLa cells (11), and VV uncoating, but not attachment, is inhibited by treatment of HeLa cells with methyl-β-cyclodextrin (mβCD), a drug that disrupts lipid rafts by depleting cellular cholesterol (23). To test whether VV-binding molecules are enriched in lipid rafts of primary human cells, we studied colocalization of VV binding with lipid rafts on the cell surface of all susceptible PHL subsets. Colocalization of VV with lipid rafts on PHL subsets including monocytes, B cells, neutrophils, and activated T cells was observed in both patched and unpatched states, while CXCR4 on monocytes, neutrophils, and activated T cells, and CD19 on B cells did not colocalize with VV binding (Fig. 2A-2D).

**VV binds to lipid rafts enriched in uropods of polarized leukocytes**

To further analyze the association of VV binding with lipid rafts, we polarized monocytes, B cells, neutrophils and activated T cells, and then conducted lipid raft staining and VV binding. As previously reported, GM-CSF, SDF-1, bacterial peptide fNLPNTL, and anti-CD44-coated coverslips effectively induced polarization of monocytes (24), B cells (25), neutrophils (26), and activated T cells (15), respectively, as 80 ± 8% (n=6) of monocytes, 65 ± 6% (n=6) of B cells, 75% ± 11% (n=6) of neutrophils, and 35% ± 4% (n=6) of activated T cells displayed elongated phenotypes and uropod
formation. In all polarized cell types, vA5L-YFP strongly colocalized with CTB-stained lipid rafts enriched in polarized cell uropods (Fig. 3A). In contrast, VV did not colocalize with F-actin molecules in lamellipodia in the leading edge of polarized cells (Fig. 3A). If monocytes were continually cultured for 7 days in GM-CSF-containing complete RPMI 1640 medium to differentiate into macrophages in vitro, these newly differentiated macrophages also maintained a polarized phenotype with bound VV strongly colocalized with lipid rafts in the uropods (Fig. 3B). These results indicate that the VV receptors are strongly associated with lipid rafts in PHL subsets in both ex vivo and polarized states.

During PHL migration and/or polarization in vivo and in vitro, GM1-stained lipid rafts and raft components move to the uropod ends of cells(24-27). Many cell surface proteins also move in and out of lipid rafts during these processes to fulfill certain physiological roles. Thus, our polarization study not only provides another way to demonstrate colocalization of VV with lipid rafts, but also presents a unique characteristic about the location of VV receptors during cell migration and polarization.

Immunosera raised against DRMs strongly block VV binding and infection

Since VV receptors are strongly associated with lipid rafts in PHLs and are likely proteins, we hypothesized that DRMs from susceptible PHL subsets were able to induce Abs that would block VV binding. To this end, we immunized BALB/c mice with DRMs fractionated from monocytes, activated T cells, or resting T cells. Immunosera against DRMs from resting T cells would not block VV binding as resting T cells do not express VV-binding receptors. Immunosera against whole cells or CMEs from
monocytes, activated T cells or resting T cells were also raised to be used as VV-blocking comparisons as a previous study has reported that immunosera against whole monocytes or activated T cells effectively blocked VV binding to activated T cells(3). We found that all immunogens from all cell types effectively elicited Abs against CD55 protein (Fig. 4A). Because CD55 is associated with lipid rafts, DRMs from all cell types induced the highest titers of anti-CD55 Abs when compared with immunogens of whole cells or CMEs (Fig. 4A). Immunosera raised against DRMs, whole cells, or CMEs from monocytes or activated T cells crossly and effectively blocked VV binding to monocytes (Fig. 4B), B cells (Fig. 4C), and activated T cells (Fig. 4D). In contrast, immunosera raised against DRMs, whole cells, or CMEs from resting T cells did not affect VV binding to any of these cell types, which is similar to the results observed from pre-immunization sera (Fig. 4B-4D). Anti-DRM immunosera exhibited the strongest blockage activity, followed by immunosera against CMEs and then whole cells (Fig. 4B-4D). Concordantly, infection of monocytes was drastically reduced when cells were pretreated with anti-DRM immunosera, and to a lesser extent by anti-CME immunosera, but not by immunosera against DRMs or CMEs from resting T cells (Fig. 4E). Notably, anti-DRM immunosera did not affect endocytosis of latex beads (Fig. 4F), whereas cytochalasin D, a known endocytosis inhibitor, effectively blocked endocytosis of latex beads (Fig. 4F). Thus, immunosera raised against DRMs from monocytes or activated T cells crossly and effectively blocked VV binding to and infection of VV-susceptible PHL subsets. The blocking activity is significantly higher than that mediated by immunosera raised against CMEs or whole cells. These results suggest that VV receptors are
enriched in DRMs, and these receptors are shared by the virus-susceptible subsets of PHLs.

**Immunosera depleted with VV-susceptible PHL subsets lose their blocking activity against VV binding**

To further confirm that monocytes, activated T cells, and B cells share VV receptors, we incubated anti-DRM immunosera with monocytes or activated T cells to deplete Abs in these immunosera. We found that anti-DRM immunosera depleted with either monocytes or activated T cells, but not resting T cells, profoundly reduced their activity in blocking VV binding to all cell types examined including monocytes (Fig. 5A), B cells (Fig. 5B), and activated T cells (Fig. 5C). PHL subsets from 6 healthy donors were used in the Ab depletion. In some cases, Ab-depleted anti-DRM immunosera totally lost their ability in blocking VV binding (data not shown). These results further indicate that VV receptors are proteins enriched in lipid rafts, and that efforts to identify poxvirus receptors and to study interactions of individual poxvirus proteins with viral receptors should be focused on DRMs instead of soluble membrane proteins extracted from target cells by non-ionic detergent lysis methods.

These results add to the knowledge of characteristics of VV receptors on PHLs, where: (1) receptors are mainly expressed on monocytes, B cells, and neutrophils among PHL subsets; (2) receptors are induced de novo following T cell activation; (3) receptors are upregulated on CD14^{high}CD16^{-} monocytes versus CD14^{low}CD16^{+} monocytes; (4) receptors are lipid raft-associated. Using these criteria, a list of putative VV receptors was made according to mass spectrometry or RNA-seq data available
from previous studies (Table 1). These proteins were selected based on the presence in monocyte DRMs\(^{(28)}\), the upregulation on CD14\(^{\text{high}}\)CD16\(^{-}\) monocytes versus CD14\(^{\text{low}}\)CD16\(^{+}\) monocytes\(^{(29)}\), and the upregulation on activated T cells versus naïve T cells\(^{(30)}\).

Lipid raft-associated proteins CD29 and CD98 are not directly involved in VV binding

The lipid raft-associated proteins CD29 and CD98 in HeLa cells and mouse embryonic fibroblasts (MEFs) play a critical role in VV entry into these cells, as knockdown or knockout of these proteins significantly reduces VV entry\(^{(31, 32)}\). In addition, VV entry into GD25 cells (a mouse cell line that is deficient in CD29 expression) was less efficient than entry into GD25\(^{\beta1A}\) cells (GD25 cells expressing human CD29)\(^{(31)}\).

We therefore wanted to determine whether these two lipid raft-associated proteins also play a role in VV binding to and infection of PHLs. We found that knockdown of either CD29 or CD98 in HeLa cells (Fig. 6A) or activated T cells (Fig. 6B) did not affect VV binding to these cells (Fig. 6C, 6D). However, knockdown of CD29 or CD98 in HeLa cells reduced VV infection, as MFI of EGFP was significantly reduced in HeLa cells transfected with siRNA constructs against human CD29 or CD98 (Fig. 6C), which is consistent with the previous reports as described above\(^{(31, 32)}\). In contrast, knockdown of CD29 or CD98 in activated T cells had no effect on VV infection (Fig. 6D). In fact, CD29 expression on the surface of HeLa cells pre- or post-knockdown of CD29 had no correlation with VV binding, as both the CD29-negative and CD29-positive population...
did not show any difference in VV binding (Fig. 6E). These results are in agreement with previous reports that CD29 and CD98 are important for VV infection in HeLa cells through mediating VV entry, but not attachment(31, 32). However, these two proteins have no effect on VV binding, entry and infection of primary human T cells although they are highly expressed on these cells.

**Discussion**

In the present study, we systemically studied the patterns of VV binding to and infection of major PHL subsets including monocytes, B cells, neutrophils, NK cells, and T cells. We found that VV exhibited an extremely strong bias towards binding to and infection of monocytes in PHLs (Fig. 1A-1B, 1G-1J) and this result is in agreement with previous reports(3, 4, 33). VV also bound to primary B cells to a similar degree as that of monocytes and to neutrophils to a much lesser extent, but barely infected these cells (Fig. 1A-1B, 1I-1J). These results suggest that monocytes not only express VV receptors on the cell surface but also have all pathways necessary for viral uptake, entry, transmembrane trafficking and ultimately penetration to the cytoplasm where the whole process of poxvirus replication takes place. However, VV infection of primary human monocytes is abortive because either viral late-gene products or viral DNA copies are not significantly generated in infected monocytes(3, 22). Primary B cells and neutrophils also express VV receptors on their surface, albeit at different degrees, but they likely lack cellular pathways for VV entry or other downstream events. It is also possible that VV may require more than one molecular species as receptors, and monocytes have all these molecular species, whereas B cells and neutrophils only have
one of these molecular species. Many viruses such as poliovirus use a single molecular
species as its receptor (34), whereas other viruses such as human immunodeficiency
virus 1 (HIV-1) use more than one molecular species including CD4 and CCR5 (34).
Different outcomes of VV binding to monocytes versus B cells or neutrophils imply that
these cell types can be used as cell models to dissect the molecular mechanisms of
poxvirus binding, penetration, entry, and infection, eventually leading to a better
understanding of poxvirus tropism and species specificities. These results may also
provide a better understanding of why live VV-based vaccines were so effective against
smallpox that we continue to use these same vaccines today. This knowledge is
requisite for the rational development of safer and more effective poxvirus-based
vaccines against other infectious pathogens and tumors. Currently, over a dozen viral
vaccines based on live poxvirus vectors are licensed in veterinary medicine (35). In
humans, a combination of a poxvirus-based HIV-1 vaccine with HIV-1 envelope (Env)
demonstrates a promising protective effect in the HIV-1 vaccine efficacy trial, known as
RV144 clinical trial (36). The success of these poxvirus-based vaccines greatly renews
research interests in poxvirus biology and virology. Because VV replication is
dependent on epidermal growth factor receptor (EGFR)/Ras pathway signaling which is
commonly active in epithelial cancers (37), VV has been developed as a promising
oncolytic agent to kill tumor cells, and has been engineered as a vehicle for the
intravenous delivery and expression of anti-tumor siRNA and peptides (37-39).
Therefore, characterization of VV binding and infection tropism will also advance the
concept of using live viruses to treat cancer.
The VV envelope consists of approximately 25 surface membrane proteins and several have been proposed as receptor-binding proteins (RBPs)(40). However, none of them has been validated as a RBP because they failed to pull down a specific ligand from soluble membrane proteins extracted from VV permissive cells by conventional detergent lysis methods, implying that VV receptors may not exist in soluble membrane extracts, but in detergent-resistant lipid rafts. We found that VV colocalized with lipid rafts on the surface of all major PHL subsets (monocytes, B cells, and neutrophils) that are susceptible to VV binding (Fig. 2A-2D). Activated T cells become sensitive to VV binding and infection because VV receptors are induced de novo upon T cell activation(3). The VV receptors newly induced on activated T cells were also found to colocalize with lipid rafts (Fig. 2C). Strikingly, these receptor molecules move together with lipid rafts as VV binding is concentrated in lipid rafts even they are relocated to uropods of polarized cells (Fig. 3A), and VV binding is dispersed on the cell surface when lipids rafts are dispersed by MβCD treatment (data not shown). Since VV receptors are strongly associated with lipid rafts, efforts to identify poxvirus receptors and to study interactions of individual poxvirus proteins with viral receptors should be focused on DRMs instead of soluble membrane proteins extracted from target cells by non-ionic detergent lysis methods. If detergents are to be used in an attempt to isolate possible VV receptor proteins, ionic detergents should be preferred, and lysates should not be cleared to avoid losing detergent-insoluble materials.

It is pertinent to note that lipid rafts of cell lines in culture versus PHLs may play different roles in VV binding and infection. In HeLa cells, membrane lipid rafts are important for VV penetration, but not for VV binding as MβCD treatment significantly
inhibits VV uncoating without affecting virion attachment(23). In addition, HeLa cell
surface CD29 and CD98, two lipid raft-associated proteins, are important for VV
entry(32)(41), although not for VV binding (Fig. 6C, 6E). However, these two proteins
have no effect on VV binding to and infection of primary human T cells, as knockdown
of their expression on the surface of activated T cells does not affect viral binding and
infection (Fig. 6), and pAbs against full-length human CD29 did not block VV binding to
PHLs (data not shown). Furthermore, primary human NK cells express high levels of
CD29 together with many other adhesion molecules(42), but these cells are resistant to
VV binding and infection(3, 43)(also Fig. 1A, 1B). These data indicate that VV receptors
are strongly associated with lipid rafts, but not CD29 and CD98, on the cell surface of
PHLs.

It has been reported that immunosera raised against whole monocytes or
activated T cells effectively block VV binding to these cells(3). If VV receptors were
enriched in lipid rafts, immunosera raised against DRMs would be more effective than
immunosera raised against whole cells in blocking VV binding. In fact, anti-DRM
immunosera significantly blocked VV binding to the highest degree, whereas
immunosera raised against whole cells or CMEs also blocked VV binding but at much
lesser degrees (Fig. 4B-4D). Blockage of VV binding by these immunosera appears to
be specific as immunosera raised against either DRMs, CMEs, or whole cells of resting
T cells did not affect VV binding (Fig. 4B-4D). In addition, these immunosera lost their
blocking activity if they were depleted against monocytes or activated T cells, but not
resting T cells (Fig. 5A-5C). Importantly, immunosera raised against monocyte DRMs,
CMEs, or whole cells were able to crossly block VV binding to B cells and activated T
cells (Fig. 4B-4D). Similarly, immunosera raised against activated T cell DRMs, CMEs, or whole cells were able to crossly block VV binding to B cells and monocytes (Fig. 4B-4D). These data strongly suggest that monocytes, B cells, and activated T cells share one or more unique protein receptors for VV. Notably, the consequences of VV binding to these cell types are different. Activated T cells are permissive to VV binding, infection and replication. In contrast, primary B cells and neutrophils are only sensitive to VV binding, but not permissive to VV infection. VV binds to and enters monocytes to initiate virus infection, but the infection is abortive as no viral late gene products have been detected(3) and viral DNA copies have not been found to increased in infected monocytes(22), which indicates that the state of certain cellular pathways in monocytes is not permissive for VV replication. However, monocytes bound with VV may help VV dissemination from initially infected sites to distant organs and tissues as variola virus has been found to be disseminated by monocyctic cell-associated viremia(5). It is possible that monocytes use putative viral receptors to grab infectious variola virus particles and then disseminate them to uninfected cells and tissues via filopodial extensions, a major mechanism that HIV-1 uses to disseminate virus infection from dendritic cells to other cell types(44). Cell-associated VV spread by filopodial extensions greatly reduces the time needed to infect neighboring cells in culture, and this process only requires VV early gene transcriptions that monocyte-lineage cells are known to express during VV infection(45). It is also possible that VV binds to the uropods of phagocytes such as monocytes, macrophages, neutrophils in PHLs to not only spread to tissues, but also to resist phagocytic activity. A recent report has demonstrated that *Neisseria meningitidis* binds to the uropods of migrating neutrophils to resist neutrophil
phagocytic activity, and to spread the bacteria during cell migration through epithelial cell layers(46). This report also showed that uropod-bound bacteria were resistant to phagocytosis, which only occurs at the pseudopod end (leading edge)(46). Our results also showed that VV would preferentially bind to the uropod ends of phagocytes, which may also provide protection from phagocytosis and assist in viral dissemination leading to generalized VV infections.

Statement

The authors have no conflict of interest to declare.

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Figure Legends
**Fig. 1. VV binding and infection of PHL subsets.** (A) Major PHL subsets including monocytes, B cells, neutrophils, resting T cells and NK cells were treated or untreated with trypsin or HS, and then subjected to vA5L-YFP binding at an MOI of 10. VV binding was measured by YFP intensity using FACS. (B) Pooled data represented mean ± SD of VV binding (% of YFP-positive cells) to PHL subsets from 6 blood donors. (C) A representative sample showed the kinetics of VV binding to activated T cells. (D) Pooled data represented mean ± SD (n=6) of VV binding to activated T cells during 0 – 72 h activation periods. (E) VV binding to activated T cells on day 3 of activation was affected by treatment with trypsin, but not HS. (F) Both resting NK cells (CD69-negative) and IL-2-activated NK cells (CD69-positive) were not permissive to VV binding. (G) Confocal microscope analysis of VV binding to monocytes versus resting T cells versus activated T cells at the single cell level. (H) Pooled data represented mean ± SD (n=6) of VV binding to monocytes versus resting T cells versus activated T cells. (I) VV infection profiles of PHL subsets and the effects of trypsin on VV infection of these cell types. (J) Pooled data represented mean ± SD of VV infection (% of EGFP-positive cells) to PHL subsets from 6 blood donors. (K) Isolated monocytes were subjected to VV binding, and then surface staining with CD14 and CD16. Cells were then subjected to FACS to determine VV binding efficacy of different monocyte subsets. Data represented VV binding to monocytes of 6 blood donors. Data were compared using Tukey’s ANOVA assay. NS, no significant; *p<0.05; **p<0.01; HS, heparan sulfate, histogram in solid gray, virus binding control; histogram in dotted line, vA5L-YFP binding.
Fig. 2. VV binding to lipid rafts on the surface of PHL subsets. Isolated monocytes (A), B cells (B), activated T cells on day 3 of activation (C), and neutrophils (D) were patched or unpatched, and then subjected to staining with CTB conjugated with Alexafluor 647 (red), anti-human CXCR4 Ab conjugated with Alexafluor 647 (red) for monocytes, activated T cells and neutrophils, or anti-human CD19 Ab conjugated with Alexafluor 647 (red) for B cells. All cell types were then incubated with an MOI of 10 vA5L-YFP (green) at the binding condition, fixed with 2% PFA, and adhered to poly-l-lysine coated coverslips. Coverslips were mounted with mounting medium containing DAPI (blue), and analyzed with confocal microscopy. Scale bars represent 10 μM. The data represents the results of VV binding to lipid rafts on PHL subsets from 6 blood donors.

Fig. 3. VV binding to lipid rafts enriched in uropods of polarized PHL subsets. (A) VV binding to lipid rafts enriched in uropods of polarized monocytes, B cells, activated T cells on day 3 of activation, and neutrophils. These cell types were treated with GM-CSF, SDF-1, anti-CD44 coated plates, and bacterial peptide fNLPNTL, respectively, to induce cell polarization. Polarized cells were subsequently fixed with 2% PFA and stained with CTB conjugated with Alexafluor 647 (red), phalloidin conjugated with Alexa Fluor 546 to stain actin filaments (pink), and DAPI (blue). Cells were then subjected to VV binding with vA5L-YFP (green), and confocal microscopy analysis. (B) VV binding to lipid rafts enriched in uropods of polarized macrophages that were differentiated from monocytes after 7 days of culture in vitro in the presence of GM-CSF. Scale bars...
represent 10 μM. The data represents the results of VV binding to these polarized cell types derived from 6 blood donors.

**Fig. 4. Blockage of VV binding and infection of PHL subsets by immunosera.** (A) Titers of anti-human CD55 Abs in immunosera raised against DRMs, CMEs, or whole cells from monocytes, resting T cells or activated T cells were determined using an ELISA assay. Diluted immunosera at 1:10 with PBS were used to block VV binding to monocytes (B), B cells (C), and activated T cells (D). (E) Diluted immunosera at 1:10 with PBS were used to block VV infection of monocytes. (F) Effects of diluted immunosera at 1:10 with PBS and cytochalasin D on monocyte endocytosis of FITC-latex beads. The MFI of YFP (FITC for latex beads) or EGFP represented VV binding or infection intensity to PHL subsets of 6 blood donors. Data were compared using Tukey’s ANOVA assay. Mono W, whole monocytes; Mono CME, monocyte crude membrane extracts (CMEs); Mono DRM, monocyte detergent-resistant membranes (DRMs); ActT W, whole activated T cells on day 3 of activation; ActT CEM, activated T cell CMEs, ActT DRM, activated T cell DRMs, ResT W, whole resting T cells, ResT CME, resting T cell CMEs, ResT DRM, resting T cell DRMs, Pre, preimmunization sera. Statistical analysis was used to compare each experimental condition with medium (control), NS: no significant, *p<0.05, and **p<0.01.

**Fig. 5. Immunosera depleted with VV-susceptible PHLs subsets reduced their blocking activity against VV binding.** Immunosera raised against DRMs from monocytes were diluted 1:10 in PBS, and then treated with activated T cells (40x10^6) or...
immunosera raised against DRMs from activated T cells were treated with monocytes (40x10^6) to deplete Abs. Ab-depleted immunosera were then used to block VV binding to monocytes (A), B cells (B), and activated T cells (C). The MFI of YFP represented immunoserum-mediated blockage of VV binding to PHL subsets of 6 blood donors. Data were compared using Tukey’s ANOVA assay. Anti-M DRMs, immunosera raised against DRMs of monocytes were incubated with activated T cells to deplete Abs; Anti-Act T DRMs, immunosera raised against DRMs of activated T cells were incubated with monocytes to deplete Abs; and *p<0.05.

Fig. 6. The lipid raft-associated proteins CD29 and CD98 were not directly involved in VV binding to activated T cells. HeLa cells and primary human activated T cells were transfected with siRNA constructs against human CD29 and CD98, and knockdown was measured by Western blot on HeLa cells (A) and surface expression on activated T cells (B). VV binding with vA5L-YFP and infection with VV-EGFP were performed on HeLa (C) and activated T cells (D) to analyze the effects on CD29 and CD98 on VV binding and infection. (E) A representative FACS plot of HeLa cell CD29 surface staining versus vA5L-YFP binding shows the relationship between CD29 expression and VV binding. NS, no significant, *p<0.05.

Table 1. Partial list of the lipid raft-associated proteins cross-presented on different PHL subsets.

Note: The list is compiled from previous reports available using these criteria: (1) lipid raft-association on monocytes according to mass spectrometry data for monocyte DRM
proteins (28), (2) upregulation on CD14^{high} \text{CD16}^{-} monocytes versus CD14^{low} \text{CD16}^{+} monocytes according also to a mass spectrometry study (29), (3) upregulation on activated T cells versus naïve T cells according to an RNA-seq analysis (30). Cell type-specific expression was confirmed using the BioGPS microarray library from the Scripps Research Institute (La Jolla, CA). *These factors were further tested for their roles as VV receptors on PHLs in this study. X, represents proteins expressing on the surface of cells.

References

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Fig. 3

A) CTB, vA5L-YFP, F-actin, Merge

- Monocyte
  - CTB: 5 μM
  - vA5L-YFP: 5 μM
  - F-actin: 5 μM
  - Merge: 6 μM

- B cell
  - CTB: 5 μM
  - vA5L-YFP: 5 μM
  - F-actin: 5 μM
  - Merge: 5 μM

- Activated T
  - CTB: 5 μM
  - vA5L-YFP: 5 μM
  - F-actin: 5 μM
  - Merge: 5 μM

- Neutrophil
  - CTB: 5 μM
  - vA5L-YFP: 5 μM
  - F-actin: 5 μM
  - Merge: 5 μM

B) CTB, vA5L-YFP, F-actin, Merge

- Macrophage
  - CTB: 5 μM
  - vA5L-YFP: 5 μM
  - F-actin: 5 μM
  - Merge: 5 μM
Fig. 5

A) VV binding to monocytes

B) VV binding to B cells

C) VV binding to activated T cells

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<th></th>
<th>VV</th>
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<th>Anti-Act T DRM</th>
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<tr>
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<td>Depleted with monocytes or activated T cells</td>
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Table 1. Partial list of the lipid raft-associated proteins cross-presented on different PHL subsets

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