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Human Cytomegalovirus Infection of Langerhans-Type Dendritic Cells Does Not Require the Presence of the gH/gL/UL128-131A Complex and Is Blocked after Nuclear Deposition of Viral Genomes in Immature Cells

Elvin J. Lauron, Dong Yu, Anthony R. Fehr and Laura Hertel

Human cytomegalovirus (CMV) enters its host via the oral and genital mucosa. Langerhans-type dendritic cells (LC) are the most abundant innate immune cells at these sites, where they constitute a first line of defense against a variety of pathogens. We previously showed that immature LC (iLC) are remarkably resistant to CMV infection, while mature LC (mLC) are more permissive, particularly when exposed to clinical-strain-like strains of CMV, which display a pentameric complex consisting of the viral glycoproteins gH, gL, UL128, UL130, and UL131A on their envelope. This complex was recently shown to be required for the infection of immature monocyte-derived dendritic cells. We thus sought to establish if the presence of this complex is also necessary for virion penetration of LC and if defects in entry might be the source of iLC resistance to CMV. Here we report that the efficiency of LC infection is reduced, but not completely abolished, in the absence of the pentameric complex. While virion penetration and nuclear deposition of viral genomes are not impaired in iLC, the transcription of the viral immediate early genes is a ubiquitous herpesvirus that can cause severe disease in immunocompromised individuals, such as solid organ and bone marrow transplant recipients, AIDS patients, and newborns (7–9). Mucosal DC residing at these anatomical locations are among the first cell types to encounter CMV during entry and can modulate the outcome of infection by contributing to virus dissemination during their migration toward the draining lymph nodes and by stimulating the onset of adaptive immune responses against CMV. Not surprisingly, myeloid DC are major targets of CMV’s immunoevasive strategies aimed at damping and delaying the proper onset of antiviral immune responses until latency is established in hematopoietic progenitor cells (6, 10, 11).

CMV reactivation from latency is also intrinsically linked to the process of CD34+ and CD14+ cell differentiation into DC and macrophages (12, 13), rendering tissue-resident DC an important source of newly produced infectious virus (14–18). By reseeding key tissues such as the salivary glands with reactivated virus, these DC can effectively contribute to the horizontal transmission of CMV.

Langerhans-type DC (LC) account for the totality of innate immune cells residing in the epithelial layer of the oral mucosa, while other types of myeloid DC, displaying surface markers similar to dermal DC, reside in deeper layers, including the lamina propria and submucosa (19–22). DC populations morphologically, immunohistochemically, and ultrastructurally identical to LC can be differentiated in vitro from CD34+ hematopoietic progenitor cells (23–26), while a type of DC considered by some to be analogous to dermal CD14+ DC (25–27) and by others to be inflammatory cells distinct from resident, steady-state dermal DC (28, 29) can be obtained from monocytes (monocyte-derived DC [MDDC]).

These two types of DC differ substantially in their susceptibility to reactivation from latency.
to CMV infection in vitro. While immature LC (iLC) are remarkably resistant regardless of the virus strain (30), immature MDDC (iMDDC) are fully permissive, but exclusively to infection by clinical-strain-like strains, which possess an extended tropism compared to that of laboratory-adapted strains (31–36). Both attenuated strain AD169varATCC and clinical-strain-like strain TB40/E can initiate infection in mature LC (mLC), albeit with different efficiencies (30), while the extent of permissiveness of mature MDDC remains unclear (32, 33, 35, 36).

A pentamer complex that consists of the viral proteins gH, gL, UL128, UL130, and UL131A and is displayed on the envelope of endotheliotropic but not laboratory-adapted strains was recently shown to be required for virion entry into monocytes and endothelial and epithelial cells (37–40). Strains naturally lacking of endotheliotropic but not laboratory-adapted strains was re-...
inoculum was left in contact with cells at 37°C in 5% CO₂ until the time of harvest or for only 4 h. In the latter experiments, after removal of the inoculum, cells were washed twice, treated with citrate buffer (40 mM citric acid, 10 mM KCl, 135 mM NaCl, pH 3) for 1 min (room temperature [RT]) to inactivate residual bound virus (57), washed again twice, and incubated in fresh medium until harvest. At the end of the differentiation (iLC) or maturation (mLC) period, LC were harvested, counted, resuspended in fresh differentiation or maturation medium again, and exposed to CMV at an MOI of 10 or 100 PFU/cell. The virus inoculum was then left in contact with the cells until the time of harvest. In selected experiments, after removal of the inoculum, cells were washed twice in medium prior to treatment with citrate buffer as described above or with 1 mg/ml of proteinase K for 1 h at 4°C to proteolytically remove cell surface virions (58–60). Cells were then washed again twice, pelleted, and stored at −80°C for subsequent analyses or incubated in fresh medium until harvest.

**IFA.** For immunofluorescence staining analyses (IFA), HFF and ARPE-19 cells were plated on 12-mm-diameter glass coverslips, while iLC and mLC were harvested and deposited on glass slides by centrifugation at 800 rpm for 3 min at RT with a Cytospin 4 (Thermo Shandon). Cells were fixed in 1% formaldehyde for 30 min at RT, permeabilized in 0.5% Triton X-100 for 20 min on ice, and treated with blocking buffer (40% goat serum–4% fetal bovine serum in phosphate-buffered saline [PBS]) for 30 min at RT. Samples were then incubated with mouse monoclonal antibodies against the viral IE1/IE2 proteins (1:50, MAB1810; Chemicon, Temecula, CA) or against pp150 (1:400, originally from W. Britt, Birmingham, AL) for 1 h at RT in a humidified chamber, washed in blocking buffer, and incubated with Alexa Fluor 594-conjugated goat anti-mouse antibodies (1:500; Molecular Probes, Eugene, OR) for 1 h at RT. For dual staining, cells were incubated in blocking buffer containing mouse monoclonal anti-pp150 antibodies and rabbit polyclonal anti-lamin B antibodies (1:50; Santa Cruz Biotechnology, Santa Cruz, CA) for 1 h at RT, washed in blocking buffer, and incubated with Alexa Fluor 594-conjugated goat anti-mouse and Alexa Fluor 488-conjugated goat anti-rabbit antibodies (1:500; Molecular Probes, Eugene, OR) for another hour at RT. Nuclei were labeled with Hoechst 33542 (0.2 mg/ml; Molecular Probes, Eugene, OR) for 3 min at RT. Slides were mounted in 90% glycerol–10% PBS containing 2.5 g/liter of 1,4-diazabicyclo-(2,2,2)-octane (DABCO; Invitrogen, Carlsbad, CA) containing the following DNA fragments from the genome of AD169: nucleotides (nt) 69050 to 77680 (8000 nt), 77680 to 86900 (9220 nt), 86900 to 94700 (7800 nt), 94400 to 102800 (8040 nt), 102800 to 135600 (7100 nt), 135600 to 150200 (6900 nt), 149800 to 158600 (7000 nt), 166400 to 173500 (7100 nt), 173500 to 178000 (46500 nt), and 178000 to 184700 (6700 nt). Two micrograms of a pSC-B plasmid mixture was labeled by nick translation in 150200 (6900 nt), 149800 to 156800 (7000 nt), 166400 to 173500 (7100 nt), or with 1 mg/ml of proteinase K for 30 min at 37°C in a humidified slide hybridization chamber. After the addition of 5 to 10 μl of labeled probe, slides were incubated at 95°C for 2 min to denature DNA. Hybridization was then carried out in a moist hybridization chamber overnight (18 to 20 h) at 37°C. Slides were subsequently washed with 2× SSC twice for 5 min at 60°C and once for 5 min at RT, dried, and stained with Hoechst 33342 (0.2 mg/ml) for 3 min at RT. After three washes in PBS, slides were mounted in ProLong Gold Antifade Reagent (Molecular Probes, Life Technologies, Grand Island, NY) and viewed on a Zeiss LSM710 confocal inverted microscope equipped with Zeiss ZEN image-processing software.

**Real-time quantitative genomic PCR and reverse transcription-PCR.** Genomic DNA was extracted from infected HFF, ARPE-19 cells, iLC, and mLC with the QiAamp DNA minikit (Qiagen, Valencia, CA). mRNA was obtained from infected iLC and mLC with the μMACS mRNA isolation kit (Miltenyi Biotec, Bergisch Gladbach, Germany), treated with DNase (DNA-free kit; Invitrogen, Life Technologies, Grand Island, NY) to remove any contaminating genomic DNA, and reverse transcribed with SuperScript II reverse transcriptase (Invitrogen, Life Technologies, Grand Island, NY). Real-time quantitative PCRs were performed in triplicate with iTaq SYBR green Supermix with ROX (Bio-Rad, Hercules, CA) and an ABI7900 thermocycler (Applied Biosystems, Carlsbad, CA) with primers hybridizing to exon 2 of the viral UL122 and UL123 ORFs (forward primer, 5′-GGCGGAAGAATCTTCAAAAA-3′; reverse primer, 5′-CTGTTGCAATCTCTCGGATCA-3′), to the viral UL50 ORF (forward primer, 5′-AGAACTCTACGATAAGGATAGAAAGGT-3′; reverse primer, 5′-ACTGCGTGACATGGCAGATTGGAACAGGT-3′; reverse primer, 5′-AAACTCTAGGAGCTGCTGTGTT-3′). The following cycling parameters were used: 95°C for 2 min to activate the iTaq polymerase, followed by 40 cycles of template denaturation at 95°C for 15 s, primer annealing at 51°C (UL122/123 and UL50) or 57°C (albumin) for 30 s, and product extension at 72°C for 30 s. Absolute quantifications of viral genome, viral transcript, and cellular genome amounts were obtained by using a standard curve made by serial dilutions of plasmid pON306 (61), a kind gift from E. S. Mocarski (Emory University, Atlanta, GA), of plasmid LNCX UL50-MA (62), and of a plasmid we constructed (pTOPO-albumin) that contains a portion of the albumin gene corresponding to nt 16,000 to 16,500. The number of viral genome copies per cell was then calculated as follows: number of viral genome copies per cell = number of viral DNA copies/ (number of albumin DNA copies/2) (63).

### RESULTS

**The UL128-131A proteins are not required for CMV infection of iLC and mLC.** We previously showed that CMV strain AD169varATCC can infect mLC, despite lacking the gH/gL/gO trimer (43,46, 47, 64), while TR, BAC4, and BADR particles harbor both the gH/gL/UL128-131A complex. This suggested that the UL128-131A proteins might not be tropism determinants for this type of DC. To test this idea, iLC and mLC populations differentiated from the Blipate Imaris Suite package of Scientific Volume Imaging (Hilversum, The Netherlands) and the Surpass Surfaces and the Surpass Section visualization tools.

**FISH assays.** Probes consisted of a mixture of 10 pSC-B plasmids (StrataClone Ultra Blunt PCR Cloning kit; Agilent Technologies, Santa Clara, CA) containing the following DNA fragments from the genome of CMV strain FIX-BAC (AC146907): nucleotides (nt) 69050 to 77680 (8630 nt in length), 77680 to 86900 (9220 nt), 86900 to 94700 (7800 nt), 94400 to 102800 (8040 nt), 102800 to 135600 (7100 nt), 135600 to 150200 (6900 nt), 149800 to 158600 (7000 nt), 166400 to 173500 (7100 nt), 173500 to 178000 (46500 nt), and 178000 to 184700 (6700 nt). Two micrograms of a pSC-B plasmid mixture was labeled by nick translation in the presence of 0.004 U of Dnase I (New England BioLabs), 10 U of DNA polymerase I (New England BioLabs), and 10 μM ChromaTide Alexa Fluor 594-5-DUTP (Molecular Probes, Eugene, OR) at 15°C for 75 min. Labeled probes were then cleaned with the QiAquick Gel Extraction kit (Qiagen, Valencia, CA) and resuspended in fluorescence in situ hybridization (FISH) buffer (2X SSC [1X SSC is 0.15 M NaCl plus 0.015 M sodium citrate], 50% deionized formamide, 20% dextran sulfate).

iLC and mLC were harvested and deposited on glass slides by centrifugation as described above. Slides were then fixed in cold 95% ethanol–5% glacial acetic acid for 5 min at −20°C, rinsed once in 100% ethanol, and air dried prior to prehybridization in FISH buffer for 30 min at 37°C in a humidified slide hybridization chamber. After the addition of 5 to 10 μl of labeled probe, slides were incubated at 95°C for 2 min to denature DNA. Hybridization was then carried out in a moist hybridization chamber overnight (18 to 20 h) at 37°C. Slides were subsequently washed with 2× SSC twice for 5 min at 60°C and once for 5 min at RT, dried, and stained with Hoechst 33342 (0.2 mg/ml) for 3 min at RT. After three washes in PBS, slides were mounted in ProLong Gold Antifade Reagent (Molecular Probes, Life Technologies, Grand Island, NY) and viewed on a Zeiss LSM710 confocal inverted microscope equipped with Zeiss ZEN image-processing software.
effects of the producer cell type on the tropism and entry modes of the viral progeny (66, 67). HFF were chosen on account of their abilities to support the growth of the AD169, Towne, and UL128mut strains and to produce viral particles with a tropism broader than that of those released by endothelial cells (66).

Second, at least three different stocks of each strain were tested, in order to assess the robustness of each strain’s behavior and to minimize the effect of adventitious mutations potentially arising during CMV amplification in vitro. As the growth of clinical-strain-like strains on HFF is associated with the relatively rapid appearance of mutations in the ORFs encoding the UL128, UL130, and UL131A proteins (68), the nucleotide sequence of the UL128-131A genomic locus of each strain was verified by PCR amplification and sequencing of virion DNA.

Third, infections were carried out in the absence of adsorption-enhancing treatments such as exposure to polyethylene glycol or centrifugation to test the natural capacity of each strain to access cells (59, 66, 69–72). Finally, viruses were left in contact with cells until the time of harvest, in order to avoid the centrifugation step required to remove the virus inoculum from cultures growing in suspension.

Consistent with expectations (39), BAC4 and BADr expressed the IE1/IE2 proteins in the majority of ARPE-19 cells, while AD169, Towne, and UL128mut initiated infection in less than 0.5% of the cells (Table 2). In contrast, and contrary to published data obtained with the TR strain (59), TR-GFP failed to infect cells with high efficiency, despite carrying an intact UL128-131A locus. To verify this finding, we tested the infectivity of a different TR-GFP-derived virus, TRsubUS17, which lacks the ORF encoding US17 (D. Yu, unpublished data). Although the function of US17 is not known, this protein is not expected to be directly involved in CMV tropism. Similar to TR-GFP, TRsubUS17 also inefficiently infected ARPE-19 cells (Table 2), indicating that the observed phenotype was not due to changes occurring during amplification of the TR-GFP strain in our experiments and suggesting that other viral determinants of tropism for epithelial cells are likely to exist in addition to the UL128-131A proteins. Mutations in the ORF(s) encoding this as-yet-unidentified factor(s) and arising within the TR-GFP genome would be maintained in the derived virus TRsubUS17, producing identical phenotypes with respect to these strains’ tropism for epithelial cells.

Consistent with our previous findings (30), mLC were overall more permissive to infection than iLC were (Fig. 1A and B). Similar to ARPE-19 cells, BAC4 and BADr were more efficient than UL128mut and AD169 at initiating infection in mLC, while TR-GFP yielded the lowest percentage of IE1/IE2+ cells (Fig. 1B). As the extent of infection was highly dependent on the donor of the progenitor cells used to differentiate LC, the infectivities of strains possessing or not possessing the gH/gL/UL128-131A complex were compared in side-by-side infections of mLC derived from the same donors (Fig. 1C and D). The percentage of infected mLC observed after exposure to BAC4 was virtually identical to that detected after exposure to UL128mut in cultures derived from 3 of 11 donors and 2- to 5-fold higher in the remaining cultures (Fig. 1C), with an overall median BAC4/UL128mut ratio of 2.3 ± 1.3. Similarly, the ratio of the percentages of BADr- and AD169-infected cells was ~1 for mLC derived from two of nine donors and ranged between 1.4 and 2.6 for the rest of the cultures (Fig. 1D), with a median value of 1.8 ± 0.5.

Both TR-GFP and TRsubUS17 did not effectively enter mLC, yielding percentages of IE1/IE2+ cells 3.5-fold (TR-GFP) and

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**Table 1 Name and envelope complex composition of the CMV strains used in this study**

<table>
<thead>
<tr>
<th>Full name</th>
<th>Name used in text</th>
<th>ORF(s) deleted for BAC insertion</th>
<th>Origin</th>
<th>gH gL gO</th>
<th>Reference(s) for virion complex</th>
</tr>
</thead>
<tbody>
<tr>
<td>AD169 var ATCC</td>
<td>AD169</td>
<td>None</td>
<td>Plaque-purified derivative of ATCC stocks</td>
<td>Yes</td>
<td>No</td>
</tr>
<tr>
<td>Towne GFP-IE2</td>
<td>Towne</td>
<td>None</td>
<td>GFP-IE2-expressing derivative of Townevar RIT1</td>
<td>Yes</td>
<td>No</td>
</tr>
<tr>
<td>TR-GFP</td>
<td>TR-GFP</td>
<td>None</td>
<td>GFP-expressing derivative of TR-BAC containing a full-length genome</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>TRsubUS17</td>
<td>TRsubUS17</td>
<td>None</td>
<td>TR-GFP derivative carrying the GalK/kanamycin cassette in lieu of the US17 ORF</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>BAC4</td>
<td>BAC4</td>
<td>US2-US6</td>
<td>Highly endotheliotropic BAC clone of TB40/E</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>TB40-BAC4</td>
<td>UL128mut</td>
<td>US2-US6</td>
<td>BAC clone of TB40/E with the HSLTR amino acid sequence of UL128 mutated to ASLTA</td>
<td>Yes</td>
<td>No</td>
</tr>
<tr>
<td>TB40-BAC4-UL128ctta82-86</td>
<td>UL21.5</td>
<td>BADr</td>
<td>BAC clone of AD169 with the UL21.5 ORF replaced with the GFP and kanamycin resistance genes and the original UL131A ORF replaced with the UL131A ORF from strain TR</td>
<td>Yes</td>
<td>Yes</td>
</tr>
</tbody>
</table>

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**Table 2 Percentages of IE1/IE2+ ARPE-19 cells after infection at an MOI of 10**

<table>
<thead>
<tr>
<th>CMV strain</th>
<th>Median % of IE1/IE2+ ARPE-19 cells (SD)</th>
<th>No. of stocks tested</th>
<th>No. of replicate tests</th>
</tr>
</thead>
<tbody>
<tr>
<td>BAC4</td>
<td>48 (5.4)</td>
<td>3</td>
<td>3</td>
</tr>
<tr>
<td>UL128mut</td>
<td>0.02 (0.1)</td>
<td>2</td>
<td>5</td>
</tr>
<tr>
<td>BADr</td>
<td>100 (0)</td>
<td>2</td>
<td>6</td>
</tr>
<tr>
<td>AD169</td>
<td>0.24 (0.4)</td>
<td>3</td>
<td>18</td>
</tr>
<tr>
<td>Towne</td>
<td>0.12 (0.4)</td>
<td>3</td>
<td>5</td>
</tr>
<tr>
<td>TR</td>
<td>1.81 (1.3)</td>
<td>4</td>
<td>8</td>
</tr>
<tr>
<td>TRsubUS17</td>
<td>2.53 (2)</td>
<td>3</td>
<td>5</td>
</tr>
</tbody>
</table>
2-fold (TRsubUS17) lower than those observed after infection with BAC4 (six donors). Similar to the situation in ARPE-19 cells, TRsubUS17 was slightly more efficient than TR-GFP at initiating infection of mLC, with a median TRsubUS17/TR-GFP ratio of 1.7 ± 0.3 (nine donors).

Aside from being more resistant to infection, iLC responses to each strain were largely similar to those of mLC, with BADr being better than AD169, TRsubUS17 better than TR-GFP, and BAC4 better than both TR-GFP and TRsubUS17 at initiating infection (Fig. 1A). In contrast to mLC, however, exposure to BAC4 or UL128mut yielded virtually identical proportions of infected cells, with a median BAC4/UL128 ratio of 0.9 ± 1.6 in parallel infections of cultures derived from six donors.

Thus, absence of the gH/gL/UL128-131A complex from the virion envelope effectively abolished viral entry into ARPE-19 cells and had essentially no impact on iLC infection but reduced the efficiency of mLC infection by approximately 2-fold. These data therefore indicate that, in contrast to epithelial cells, the presence of the UL128, UL130, and UL131A proteins is irrelevant for viral entry into iLC and advantageous but not essential for entry into mLC. The low infectivity of TR-GFP for iLC, mLC, and ARPE-19 cells, moreover, implies that other viral proteins besides UL128, UL130, and UL131A may have a role in mediating CMV tropism for epithelial cells and LC. Finally, the small but consistent difference between the efficiencies of infection of TR-GFP and TRsubUS17 may hint at a potential role for US17 in restraining infection onset in LC and ARPE-19 cells, identifying this protein as a potential temperance factor (73).

**CMV virion entry is not blocked in iLC.** The fact that no viral strain with the ability to initiate productive infection in iLC has been found hints at the presence of host defense mechanisms that cannot be overcome by any viral protein encoded by the strains tested so far.

To identify the step in the viral cycle that is blocked in iLC, the progress of viral particles was tracked by staining infected iLC and mLC for pp150, a tegument phosphoprotein that remains strongly associated with capsids during entry and has been used as a capsid marker by us and others (60, 74, 75). Cytospin preparations of iLC and mLC exposed to BAC4 at an MOI of 10 PFU/cell and harvested at 0.25, 1, 5, 20, and 30 hpi were fixed and permeabilized prior to the addition of anti-pp150 antibodies. The efficacy of permeabilization was verified by staining cells with antibodies against the nuclear envelope protein lamin B. While no signal corresponding to pp150 was observed in mock-infected, permeabilized iLC (Fig. 2A to C) or mLC (not shown), viral particles were clearly visible as bright dots (Fig. 2D and G) in infected samples. At each of the times tested, the overall abundances of viral particles appeared to be similar in infected iLC and mLC (Fig. 2, compare D and G; data not shown). To precisely quantify these data, the number of particles per cell observed in at least 200 cells from at least five different fields per time point was determined and the median number of virions per iLC versus the mLC...
ratio was calculated (Fig. 2). At 15 min postinfection, iLC and mLC populations appeared to harbor similar numbers of particles/cell, with mean values of 4 ± 1 for iLC and 4.6 ± 1 for mLC (Fig. 2, 0.25 hpi), indicating that the efficiencies of virion attachment and initial entry into the two cell types were highly comparable, albeit lower than expected for an MOI of 10 PFU/cell. At both 1 and 5 hpi, mLC contained approximately 40% more particles than iLC did, but by 20 hpi, these differences had waned and both cell types again contained similar number of particles. No disparity was observed at 30 hpi either, although the total amount of virions was greatly reduced from the beginning of infection, potentially because of capsid disassembly after viral genome delivery into the nucleus. Thus, while mLC appeared to contain slightly larger amounts of virions than iLC did at early times, similar number of particles were found in each cell type at later time points, suggesting that virion entry into iLC is not dramatically hindered.

As these assays did not allow us to distinguish between virions attached to the cell surface and those that had already penetrated the cell, we next treated infected cells with citrate buffer, which was expected to inactivate and detach nonpenetrating virions from the cell surface (57, 76–78), or with proteinase K, which has been reported to proteolytically remove residual extracellular particles (58–60). Real-time quantitative PCR was then used instead of IFA to determine the number of viral genomes per cell under each condition. iLC and mLC populations derived from the progenitor cells of four different donors were exposed to BAC4 at an MOI of 10 PFU/cell for 15 min, 1 h, or 4 h. At each time point, cells were collected, washed, and stored (untreated) or washed and treated with citrate buffer for 1 min at room temperature or with proteinase K for 1 h at 4°C. After additional washes, cells were pelleted and subjected to real-time quantitative PCR with primers hybridizing to the viral UL122/123 ORF (79) or to the cellular albumin gene (63) and the number of viral genomes/cell was calculated.

At each of the time points tested, iLC contained slightly but consistently larger amounts of viral DNA than mLC did (Fig. 3A, white bars), with iLC/mLC ratios ranging from 2 ± 1.5 at 15 min postinfection to 3.8 ± 1 at 4 h postinfection (Fig. 3B, white bars). These values are higher than those obtained by IFA (Fig. 2), likely because of the superior precision of the quantification method used (real-time PCR instead of manual counting), the testing of cell populations derived from four different CD34+ cell donors instead of one, and the tracking of viral genomes instead of viral particles.
capsids. Treatment of infected cells with citrate buffer did not change the viral genome content of iLC and had only modest effects (25, 10, and 17% decrease from untreated samples at 0.25, 1, and 4 h, respectively) on mLC (Fig. 3A, gray bars), suggesting that, contrary to what was reported for other cell types (57, 76–78), exposure to a low pH did not completely remove nonpenetrating virions from infected LC. Identical results were obtained after iLC treatment with proteinase K, while enzymatic digestion of mLC resulted in the removal of 55 to 64% of the bound virions (Fig. 3A, black bars), consistent with what was previously reported for monocytes, fibroblasts, and epithelial cells (58, 59). Viral particle attachment and internalization may thus occur more efficiently in iLC than in mLC. Already at 15 min postinfection, virtually all of the virions bound to iLC are protected from proteolysis, presumably on account of their localization within the cell’s interior. In contrast, only about 36 to 44% of the attached particles are internalized by mLC at each of the time points tested, suggesting that entry into mLC may be slower or less effective than entry into iLC. Because of these differences, the intracellular content of viral particles was consistently higher (6.5-fold ± 2-fold to 9-fold ± 0.7-fold) in iLC than in mLC at each time point postinfection (Fig. 3B, black bars).

Together, these data indicate that virion entry into iLC occurs with similar or better efficiencies than virion entry into mLC, suggesting that defects in binding and penetration are not the main source of iLC resistance to infection.

Viral genome losses are similar in iLC and mLC. To determine if any difference might exist in the way viral particles are processes by each LC type after penetration, the fate of viral genomes over time was tracked in HFF and ARPE-19 cells, used as reference, and in iLC and mLC infected with BAC4, UL128mut, BADr, and AD169 at an MOI of 10 PFU/cell. Cells were exposed to each strain for 4 h before being washed with medium, treated with citrate buffer to inactivate residual nonpenetrating virus, and washed extensively again. Real-time quantitative PCR was then used to determine the number of viral genomes present in each cell population at 4, 8, 24, and 48 hpi and to calculate the proportion of genomes remaining at each time point relative to that at 4 hpi (Fig. 4), when the entry of noninternalized virions was halted by low-pH treatment. The calculated average number of viral genomes per cell found in each population at 4 hpi averaged 2.6 ± 1.8 in iLC, 1.3 ± 1.1 in mLC, 2.7 ± 2 in ARPE-19 cells, and 3.1 ± 3 in HFF. Again, at this early time postinfection, iLC appeared to hold more viral genomes than mLC did, while the viral DNA contents of HFF and ARPE-19 cells were almost identical.

As expected, all of the strains could efficiently infect HFF. The abundance of UL122/UL123 and UL50 transcripts is significantly lower in iLC and mLC (Fig. 5B to E, arrows), suggesting that capsids could successfully dock to the nuclear envelope in both cell types. To determine if viral genomes were subsequently deposited into the nucleus, confocal microscopy imaging further revealed the existence of what appeared to be points of contact between some viral particles and the outer surface of the nuclei of both iLC and mLC (Fig. 5D to G), with no apparent difference between the two cell types with respect to genome localization or the number of genomes per nucleus. In contrast, iLC and mLC samples could not be distinguished in blinded analyses of FISH slides, suggesting that nuclear deposition of viral genomes was occurring with similar efficiencies in both LC types.

The abundance of UL122/UL123 and UL50 transcripts is significantly lower in iLC. Finally, we sought to determine if transcription of the UL122/123 and UL50 ORFs was similarly efficient in iLC and mLC. The UL122/123 ORF is expressed with immedi-
ate early kinetics and encodes the IE1 and IE2 proteins essential for infection onset (80–82), while the UL50 ORF is expressed with early kinetics and encodes the nuclear egress complex component UL50 (83–85). Real-time, quantitative reverse transcription-PCR was used to determine the numbers of UL122/123 and UL50 transcript copies present in each cell type at 24, 48, and 72 h postexposure to AD169, BADr, BAC4, or UL128mut and to calculate the ratio of the number of cDNA copies of each gene present in the mLC to the number of copies in the iLC of each donor (Fig. 6). Although the UL122/123 ORF was expressed in both cell types, mLC consistently contained more transcript copies than iLC at each time point. Already at 24 hpi, the number of UL122/123 mRNA copies in mLC was 6 times (AD169 and BAC4), 8 times (UL128mut), and 12 times (BADr) greater than that in iLC.

**FIG 4** Viral genome contents of HFF, ARPE-19 cells, iLC, and mLC at later times postinfection. Cells were exposed to BAC4, UL128mut, BADr, or AD169 virions at an MOI of 10 PFU/cell for 4 h, washed in medium to remove the inoculum, and treated with citrate buffer to prevent any further virion penetration. A quarter of the cells were then harvested immediately (4 hpi), while the rest were plated and harvested at 8, 24, and 48 hpi. Real-time quantitative PCR was used to determine the number of viral genome copies per cell at each time point and to calculate the percentage of genomes remaining at 8, 24, and 48 hpi relative to that at 4 hpi (% 4 hpi). (A, C, E, and G) Genomes remaining in ARPE-19 cells and HFF from one representative experiment, with the 48-hpi values shown on a log10 scale. (B, D, F, and H) Mean percentages of genomes remaining in iLC and mLC derived from four different CD34+/H11001 cell donors with standard deviations. (I and J) Mean percentages of genomes remaining in each population after infection with BAC4, UL128mut, BADr, and AD169 with standard deviations. Values above the columns at 48 hpi are the percentages of IE1/IE2+ cells obtained in IFA analyses of each cell type at 48 hpi.
Most genome copies were present in the equivalent of 1 donor iLC and mLC from different donors at each time point, while a horizontal bar marks the median value of each group. Ratios were obtained for UL50 in LC populations derived from the progenitor cells of three or four (UL122/123, A) and two (UL50, B) different CD34+ progenitor cell donors. Each floating vertical bar extends from the minimum to the maximum ratio obtained for the iLC and mLC from different donors at each time point, while a horizontal bar marks the median value of each group.

Both LC types, albeit at different levels (median mLC/iLC ratios of 11 in AD169, 12 in BADr, 10 in BAC4, and 7 in UL128mut infections, Fig. 6B, white bars). While the number of UL50 mRNA copies in mLC remained at least 5-fold higher than that in iLC at each time point after infection with AD169 or BADr, differences were somewhat lower at 72 h after infection with BAC4 and UL128mut (Fig. 6B, dark gray bars). Considering that at both 24 and 48 h after infection with each strain, iLC contain more viral genomes than mLC (Fig. 4, A, B, C, D, F, and H) but substantially less UL122/123 and UL50 mRNA copies (Fig. 6), transcription of viral genes in iLC appears to be highly restricted. Together, these data identify viral gene transcription as the first severely impaired step in the regular progression of the viral cycle in iLC.
DISCUSSION

Efficient CMV penetration of specific cell types requires fusion of the virion envelope with cellular membranes in a process controlled by both viral and cellular factors. Capsid release into the cytoplasm can indeed occur at the periphery of the cell or from intracellular vesicles, depending on the complement of glycoprotein complexes displayed on the virion surface and on the cell type.

The gH/gL/UL128-131A complex was shown to promote entry by endocytosis and to be essential for the infection of epithelial cells, endothelial cells, monocytes, and iMDDC (38, 40, 45, 59, 67, 86). Exposure to strains carrying only the gH/gL/gO complex was not followed by viral gene expression in these cells, implying that endocytosis is likely the predominant, if not the only, entry pathway conducive to productive infection onset.

Consistent with this, we found that viral DNA maintenance and replication, as well as IE1/IE2 protein expression, in ARPE-19 epithelial cells occurred only after infection with strains possessing the pentameric complex and presumably entering by endocytosis (Table 2 and Fig. 4).

In contrast to the infection of epithelial cells, that of HFF does not require expression of the UL128, UL130, and UL131A proteins but depends on the presence of the gH/gL/gO complex to support fusion of the viral envelope with the plasma membrane (46, 72, 87, 88). In our assays, penetration of HFF by virions from strains possessing or lacking the gH/gL/UL128-131A complex progressed equally well and led to the onset of productive infections (Fig. 4A, C, E, and G), suggesting that both entry modes may be allowed to occur in this cell type. Alternatively, virions of strains harboring both complexes on their envelope, such as BAC4 and BADr, may selectively use the trimeric complex to access HFF.

While the intracellular release of naked capsids at the cell periphery is considered to be the main entry route of CMV into these cells (88), incoming virions were also detected within phagolysosome-like vacuoles in electron microscopic images of infected cells (89), suggesting that endocytosis of viral particles can also take place in HFF.

Here we show that the pentameric complex is not required for infection of iLC and mLC, although in its presence, the efficiency of mLC infection was increased by about 2-fold (Fig. 1). Similar to the situation in HFF, strains containing both complexes or just gH/gL/gO were equally able to bind to and likely enter both types of LC, but their genomic DNA was subsequently lost (Fig. 4B, D, F, and H), in a fashion comparable to that of UL128mut and AD169 in ARPE-19 cells (Fig. 4C and G). In contrast to the latter, however, the rate and extent of viral genome loss were similar for all strains (Fig. 4I) and were not accompanied by a complete absence of IE1/IE2 gene expression in mLC (Fig. 4J and 1B). It is thus possible that, akin to HFF, both endocytosis and fusion with the plasma membrane are viable routes of entry into iLC and mLC and may be used interchangeably by virions possessing both complexes. When only the gH/gL/gO complex is present, particles may still use both pathways to access cells, as suggested by the lack of difference between the intracellular contents of viral DNA of different strains. Because the endocytic pathway is impracticable in the absence of the pentameric complex, a portion of these virions may fail to reach the nucleus, leading to a reduction in the percentage of IE1/IE2 mLC.

Our finding that mLC are consistently more permissive to infection than iLC are (Fig. 1) (30) is in stark contrast to the situation with MDDC, whereby immature cells, and not their mature counterparts, are the most susceptible to CMV infection (33–36). Despite being related and sharing many of the features and functions typical of antigen-presenting cells, LC and MDDC respond to CMV in opposite ways, reflecting the heterogeneity and functional specialization of different DC populations. If these differences observed in vitro are also mirrored in vivo, then dermal DC and epidermal LC are likely to play dramatically different roles in the development of CMV pathogenesis and in the generation of antiviral immune responses, a concept that may be relevant for the development of live-attenuated vaccines intended for delivery via mucosal surfaces.

None of the CMV strains tested so far has the ability to initiate productive infections in iLC. This strongly supports the existence of some iLC-specific defense mechanism that cannot be overcome by any viral countermeasure. Intriguingly, while entry is the target of multiple cellular antiviral strategies restricting infection in different cell types, CMV penetration of iLC is not blocked. Quite to the contrary, data from our particle neutralization/detachment experiments indicate that virion internalization is more efficient in iLC than in mLC (Fig. 3A), leading to higher, instead of lower, intracellular contents of virions at early times postinfection (Fig. 3B).

This finding may not be that surprising in view of the well-known superior antigen-capturing abilities of iLC, compared to mLC (25, 90). While all three antigen internalization pathways, namely, macropinocytosis, endocytosis, and phagocytosis, are fully operational in iLC, two (macropinocytosis and phagocytosis) are downregulated during maturation (91, 92), leaving endocytosis as the predominant means of extracellular material acquisition by mLC. While our data suggest that CMV entry into mLC may occur independently of vesicle-mediated transport pathways (Fig. 1), the loss of two prominent entry routes following maturation may negatively impact the efficiency of CMV penetration of these cells, at least in the case of pentamer–containing strains such as BAC4.

Subsequent to entry, no major differences were detected between iLC and mLC in the kinetics of viral genome loss (Fig. 4I) or in the number of viral genomes reaching the nucleus, arguing against enhanced virion degradation or inefficient nuclear deposition of viral genomes as the main cause of iLC resistance to infection. In contrast, the abundance of UL122/123 and UL50 ORFs transcripts was considerably lower in iLC than in mLC at each of the times tested (Fig. 6), pointing at viral gene transcription as the first severely restricted step in the regular progression of the viral cycle.

The most intriguing question now is how viral gene transcription is blocked in iLC. It is, of course, possible for iLC to express higher levels of specific transcriptional repressors, whose binding to the major immediate early promoter (MIEP) is followed by the severe inhibition of UL122/123 expression. Conversely, mLC may contain higher levels of transcriptional activators, whose positive activity on the MIEP may promote the onset of lytic infection. The resistance to CMV infection of undifferentiated human monocyteic THP-1 cells, for instance, was found to be partially due to the presence of the MBF1 and MRF transcriptional repressors in large amounts (93, 94). Upon THP-1 differentiation into macrophages, the abundance of both factors was substantially reduced, with a concomitant increase in MIEP activity. This allowed the expres-
sion of the IE1/IE2 proteins and initiation of the viral replication cycle to occur (93, 94). Epigenetically acting proteins such as histone acetyltransferases, histone deacetylases, and histone methyltransferases may also be potential players. In latently infected CD34+ progenitor cells, the transcriptionally inactive UL122/123 promoter was indeed found to be associated with deacetylated H4 histones and with the transcriptional repressor heterochromatin protein 1 (HP1), while in mLC derived from latently infected progenitors and harboring reactivating virus, the now active UL122/123 promoter was associated with acetylated H4 histones and had lost HP1 binding (15–17). Interestingly, and similar to our findings with directly infected iLC, no expression of the viral UL122/123 ORFs was detected in iLC differentiated from latently infected CD34+ cells, indicating that maturation is crucial for both infection onset and viral reactivation to occur. Thus, although CMV reactivation in the CD34+-LC model was proposed to be “differentiation dependent” (95), we believe that it may be more tightly linked to the process of maturation (i.e., “maturation dependent”). Interestingly, while differentiation and maturation of latently infected CD34+ progenitors into mLC trigger viral reactivation (15, 16), maturation of directly infected iLC has not yet been reported to yield infectious virus. This suggests that the mechanisms mediating silencing of the MIEP in iLC derived from latently infected progenitors may be different from those acting after direct infection of these cells.

Finally, mLC susceptibility to direct infection, as well as the establishment of viral latency in CD34+ cells, was also proposed to be regulated by the activity of nuclear domain 10 (ND10) bodies (96), whose negative effects on MIEP transcription (97–99) can be partially reversed by the nuclear delivery of the viral tegument protein pp71 (100–105). Although no evidence was provided by Saffert et al. as to the actual presence of viral genomes within the ND10 bodies of infected CD34+ cells or mLC (96), pp71 appeared to reach the nucleus in at least a few infected mLC. If incoming viral genomes are indeed sequestered within the ND10 bodies of iLC and if pp71 is subsequently prevented from reaching the nucleus, then silencing of the MIEP may remain in effect, preventing the expression of the UL122/123 ORFs in these cells. On the other hand, depletion of the ND10 component and the strong transcriptional repressor hDaxx from nonpermissive human teratocarcinoma T2 cells did not render them susceptible to infection (106), and neither was their loss of resistance (triggered by exposure to vasoactive intestinal peptide) accompanied by hDaxx degradation (107), suggesting that the role played by hDaxx and pp71 in controlling CMV infection onset may be cell dependent and that they may not be active in iLC.

Altogether, we postulate that iLC resistance to infection is highly unlikely to depend on the activity of single cellular proteins. Instead, multiple host defense effectors may cooperate in generating an unfavorable environment for viral gene transcription. Conversely, mLC susceptibility to CMV infection and reactivation may depend on the assembly of an intranuclear milieu conducive to MIEP activation and be characterized by the presence of selected transcription factors, the absence of others, and the occurrence of specific epigenetic modifications of the viral and cellular genomes (95, 108). The relative contributions of some of these elements to the regulation of LC susceptibility to infection are under investigation.

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