HCMV Displays a Unique Transcriptome of Immunomodulatory Genes in Primary Monocyte-Derived Cell Types

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Abstract

Human cytomegalovirus (HCMV) is a betaherpesvirus which rarely presents problems in healthy individuals, yet may result in severe morbidity in immunocompromised patients and in immune-naïve neonates. HCMV has a large 235 kb genome with a coding capacity of at least 165 open reading frames (ORFs). This large genome allows complex gene regulation resulting in different sets of transcripts during lytic and latent infection. While latent virus mainly resides within monocytes and CD34⁺ progenitor cells, reactivation to lytic infection is driven by differentiation towards terminally differentiated myeloid dendritic cells and macrophages. Consequently, it has been suggested that macrophages and dendritic cells contribute to viral spread in vivo. Thus far only limited knowledge is available on the expression of HCMV genes in terminally differentiated myeloid primary cells and whether or not the virus exhibits a different set of lytic genes in primary cells compared with lytic infection in NHDF fibroblasts. To address these questions, we used Illumina next generation sequencing to determine the HCMV transcriptome in macrophages and dendritic cells during lytic infection and compared it to the transcriptome in NHDF fibroblasts. Here, we demonstrate unique expression profiles in macrophages and dendritic cells which significantly differ from the transcriptome in fibroblasts mainly by modulating the expression of viral transcripts involved in immune modulation, cell tropism and viral spread. In a head to head comparison between macrophages and dendritic cells, we observed that factors involved in viral spread and virion composition are differentially regulated suggesting that the plasticity of the virion facilitates the infection of surrounding cells. Taken together, this study provides the full transcript expression analysis of lytic HCMV genes in monocyte-derived type 1 and type 2 macrophages as well as in monocyte-derived dendritic cells. Thereby underlining the potential of HCMV to adapt to or influence different cellular environments to promote its own survival.
Introduction

Human cytomegalovirus (HCMV) is a ubiquitous species-specific betaherpesvirus that infects between 60–100% of the population depending on age, gender, demographics and socio-economic status [1]. The 235kb HCMV genome is characterized by unique long (UL) and unique short (US) regions, each flanked by terminal (TRL and TRS) and internal (IRL and IRS) inverted repeats [2].

Transcription of the HCMV genome is complexly regulated and different sets of transcripts are expressed during productive and latent infection. During latency, the major immediate-early (IE) promotor, which drives IE gene expression, is epigenetically blocked. However, early studies in experimental latency models detected several sense and antisense CMV latency-associated transcripts (CTLs) expressed from the major immediate-early region of the genome, although these CTLs are controversial [3–5]. Since then four transcripts i.e. UL81ast/LUNA [6–10], UL111A [10–12], UL138 [8, 10, 13, 14] and, depending on the virus used and the presence of GATA transcription binding sites, UL144 [15] were characterized thoroughly as latency-associated transcripts. Alternatively, during productive infection, the full array of genes is expressed and gene expression follows a temporal cascade starting with immediate-early gene expression, succeeded by early and late gene expression [16–22]. The transcriptome consists of both polyadenylated protein coding and polyadenylated non-protein coding RNAs. The latter RNA species is comprised of four different long non-coding RNAs (lnc RNAs) which do not overlap with any of the protein coding regions. These four lnc RNAs have been reported to be responsible for 65% of the total polyadenylated RNA pool [23, 24]. In addition, HCMV also expresses non-coding RNA in the form of anti-sense transcripts which often overlap with protein coding genes, and microRNAs which have been attributed a regulatory function [23–25]. Currently, the transcriptome of HCMV has not been entirely unraveled. It has long been assumed that the genome of clinical isolates contained about 170 genes [23, 24]. However, recently a new study revealed an unexpected complexity when Stern-Ginossar et al. identified hundreds of new ORFs through ribosome profiling and transcript analysis in lytically infected fibroblasts [26]. Currently, the functions of these newly discovered transcripts remain largely unknown.

Only recently, transcriptomics technology platforms have been used to study the influence of HCMV infection on host and viral gene expression in different cell lines and cell types. Towler et al. showed that, depending on the cell type, the temporal expression of viral genes varies; but also that, compared to fibroblasts, the transcriptome in epithelial cells and astrocyte cell lines is differentially regulated [27].

CD34+ progenitor cells and CD14+ monocytes are currently the only confirmed sites of latency [28, 29]. Upon differentiation to macrophages or dendritic cells, the virus reactivates and lytic replication is enabled, possibly promoting viral spread after reactivation [17, 30]. The transcriptome in latently infected CD34+ progenitor cells and CD14+ monocytes is slowly being unraveled [8, 10, 12]. However, only one study aimed at delineating the HCMV transcriptome in permissive myeloid cells [31]. It remains unclear how the viral transcriptome adapts to infection in macrophages (MΦs) and dendritic cells (DCs) and if the virus shows a different transcriptional profile in pro-inflammatory DCs or macrophages type 1 (MΦ1) or anti-inflammatory macrophage type 2 (MΦ2). Therefore, we used Illumina next generation sequencing to characterize how HCMV modulates gene expression specifically in terminally differentiated monocyte-derived MΦ1, MΦ2 and DCs in comparison to fibroblasts.

The data reported here is, to our knowledge, the first next generation sequencing transcriptome study investigating HCMV transcription in cell types which are relevant for in vivo viral spread. We show that HCMV follows a specific transcriptional program in primary cells in which genes involved in virion composition, cell tropism or cell-specific replication, viral...
spread and immunomodulation are expressed at different levels compared to fibroblasts. This not only highlights the potential of HCMV to adapt to or influence different cellular environments to promote its own survival. This also reflects the significant differences in gene expression between cell types used in the laboratory environment and in primary cell types.

**Materials and Methods**

**Ethics statement**

Peripheral blood mononuclear cells (PBMCs) and primary monocytes were isolated from whole blood. Blood was obtained via the Antwerp Blood Transfusion Center of the Red Cross (www.redcross.be). Donors gave written consent for their samples to be used for scientific research.

**Cell lines**

ARPE-19 cells (CRL-2302, ATCC) and neonatal NHDF fibroblasts (CC-2509, Lonza) were maintained and propagated in DMEM:F12 with L-glutamine (Biowhittaker) and MEM (Life Technologies) respectively, containing 10% heat-inactivated FCS (HI-FCS; Life Technologies) and 0.04% gentamicine. All infection experiments were done in 24-well plates (Costar).

**Culturing of clinical isolates TB40/E**

High epithelial tropic stocks of the HCMV BAC4-TB40/E [32] were generated in ARPE-19 cells. In brief, ARPE-19 cells were seeded 24 hours before infection to reach 40–60% confluence at the time of infection (3.5 x 10^6 cells/T175). Incubation with TB40/E at MOI 0.02 was done for 3 hours at room temperature on a rocking platform. The virus was harvested until the cells detached from the flasks. Harvests were spun down for 10 minutes at 600xg to remove cellular debris. All harvests were titrated (see below) and only the 6 harvests with the highest titers were used for concentration. Viral stocks were prepared by centrifugation (1 hour at 3000xg) and ultracentrifugation (1 hour at 5800xg). Concentrated virus was resuspended in 5 ml MEM/10% FCS and kept at -80°C for long term storage.

**Virus titration using immediate-early protein immunofluorescent staining**

Virus yield was determined on NHDF and ARPE-19 cells by an immune-fluorescence assay adapted from Chou et al. [33]. NHDF cells and ARPE-19 were used to determine TB40/E titers or epithelial tropism, respectively. In brief, 15000 cells (NHDF or ARPE-19 cells) per well were seeded in a 96-well plate in 75 μl MEM/2% HI-FCS or DMEM:F12/2% HI-FCS. Then, quadruplicates of a ten-fold dilution series of the virus was added to the cells (25μl input). Three days post infection, the viral replication was determined after staining for immediate-early antigen expression in an immune-fluorescence assay (see below). Wells were scored positive if they contained at least one colony of 2–5 IE-positive cells. After scoring, the TCID_{50}/ml was calculated based on the Spearman-Kärber method.

**Immediate-early protein staining**

The medium of infected cells was removed and the cells were washed once with phosphate buffered saline (PBS). Subsequently, the cells were fixed using 100% ice-cold methanol (10 minutes at -20°C). To remove residual methanol, the cells were washed twice with PBS. Staining with the primary antibody was carried out for 20 minutes at room temperature (RT) with an anti-immediate-early antigen antibody (anti-I.E.A., 11–003, Argene) diluted 1/100 in PBST (PBS containing 0.05% tween-20). Prior to incubation with anti-mouse antibody conjugated to Alexa Fluor 488 (Alexa Fluor 488 goat anti-mouse IgG, A-11001, Life Technologies), the cells
were washed twice with PBS. After 20 minutes incubation at RT, the secondary antibody was removed and the cells were washed twice with PBS. Afterwards, the cells were stored at 4°C in 200μl PBS until readout.

**Isolation of primary CD14+ monocytes and differentiation to HCMV permissive cell types**

Peripheral blood mononuclear cells (PBMCs) were isolated from buffy coats using ficoll-paque. In brief, 60 ml blood was added to 100 ml ice-cold PBS. Twenty-five ml diluted blood was carefully applied on 15 ml ficoll-paque. After centrifugation (800xg, 20 minutes, no brake), the PBMC fraction was collected and divided over four 50 ml tubes. Each tube was filled with ice-cold PBS to a total volume of 50 ml and the cells were spun down for 10 minutes at 300xg. This washing procedure was repeated twice. The total number of PBMCs was determined by manual counting under the light microscope with a counting chamber. Subsequently, monocytes were isolated using magnetic CD14 microbeads (Miltenyi, 130-050-201) as instructed by the manufacturer.

The monocytes were counted manually under a light microscope using a counting chamber, resuspended at 1.5 x 10^6 cells/ml and 1.5 x 10^6 cells were seeded per well in a 24-well plate (tissue treated, Costar). The CD14+ monocytes were allowed to adhere for 3 hours at 37°C/5% CO₂. After attachment, the PBS was removed and 200μl X-Vivo-15 medium (Lonza) without supplements was added and the cells were left overnight at 37°C/5% CO₂. Subsequently, monocytes were differentiated to monocyte-derived dendritic cells (DC), monocyte-derived type 1 macrophages (MΦ1) and monocytes-derived type 2 macrophages (MΦ2) in X-Vivo-15 medium containing 2.5mM L-glutamine and 10% HI-FCS supplemented with 100ng/ml GMCSF and 100ng/ml IL4 (adapted from [34, 35]), 100ng/ml GMCSF or 100ng/ml MCSF (adapted from [34]), respectively, for 7 days (all cytokines from Peprotech). The medium was refreshed on day 4 after the start of differentiation. Activation of terminally differentiated cells was done by adding 500mg/ml LPS (Sigma).

On day 8, the cells were infected for 3 hours at room temperature with TB40/E at MOI 5 in 0.5 ml X-Vivo-15 medium containing 2.5mM L-glutamine and 10% HI-FCS. After infection, the virus inoculum was removed and 1 ml fresh medium without cytokines was added. The cells were incubated for 72 hours at 37°C/5% CO₂ before samples were harvested.

**FACS analysis of surface markers on monocyte-derived cell types**

Monocytes were seeded in a 96-well plate at 3x10^5 cells per well and differentiated to MΦ1, MΦ2 and DCs as described above. After 7 days, the cells were challenged with 500ng/ml LPS (Sigma) and surface markers were determined. Therefore, the cells were detached from the plate using 5mM EDTA (20 minutes at 37°C). The cells were spun down at 300xg, the EDTA was discarded and the cells were incubated with a mix of anti-CD14-APC-Cy7, anti-CD209-PerCP-Cy5.5, anti-CD163-PE and anti-CD80-PE-Cy7 (5μl of each antibody, all antibodies from BD). The antibodies were incubated for 20 minutes at 4°C. After incubation, 200μl PBS was added and the cells were spun down at 300xg. The supernatant was discarded, 200μl PBS was added once more and the cells were collected by centrifugation. Readout was done on a BD FACS CANTO II and subsequent analysis were done using BD DIVA software. Plots were made in Sigmaplot (Systat Software).

**RNA sequencing using the Illumina next generation sequencing platform**

RNA samples were prepared from cells obtained from five 24-wells using a Qiagen RNeasy Plus kit with an additional on-column DNase digestion step according to the manufacturer's
instructions. RNA and DNA concentrations were measured using a ND-1000 spectrophotometer (Nanodrop).

RNA samples were processed for Illumina sequencing using the TruSeq stranded mRNA Library Prep kit (Illumina) according to manufacturer's instructions. In brief, the workflow involved purifying the poly-A containing mRNA molecules using poly-T oligo-attached magnetic beads. Following purification, the mRNA was fragmented into small pieces using divalent cations at elevated temperature. The cleaved RNA fragments were copied into first strand cDNA using reverse transcriptase and random primers. This was followed by second strand cDNA synthesis using DNA Polymerase I and RNase H. These strand-specific cDNA fragments went through an end repair process to a single 'A' base, followed by ligation of the adapters. The products were purified and enriched with PCR to create the final cDNA library.

Finally, samples were pooled and sequenced for 147 cycles on a GAIIx (Illumina) using the TruSeq SBS Kit v5 kit (Illumina). The raw signal was processed into individual sequencing reads per sample with the Illumina software (Casava 1.8.2).

All sequence reads per sample were aligned versus the publicly available reference genome for TB40/E (acc. No. EF999921; 2008). This reference genome is annotated with 168 coding sequences (cds) and was manually updated with 9 additional coding sequences from accession number KF297339 (2013). In addition, also four long non-coding RNAs (RNA1.2, RNA2.7, RNA4.9 and RNA 5.0) were added from accession number KF297339 (2013) [24]. In this study, the TB40/E-BAC4 was used which misses US1-US6 compared to the wild type virus [32], for the analysis the lack of this region was taken into account.

Mapping of the sequence reads and calculating the total read counts per cds were performed using the RNA-seq analysis module of CLC bio's CLC Genomics Workbench using default parameters. All raw data was uploaded to a public repository (http://www.ebi.ac.uk/ena/data/view/PRJEB15199).

For the statistical analysis, the Bioconductor suite was used [36]. Count data were preprocessed (normalized and variance stabilized transformed for visualization purposes) using the DESeq package [37]. For each individual transcript, differential expression between cell types was performed from normalized counts using a generalized linear model hypothesizing a negative binomial distribution. Individual p-values were adjusted for multiple testing with the Benjamini-Hochberg procedure [38]. Finally, a pathway enrichment analysis was performed using the MLP algorithm [39, 40].

Results

HCMV infection of monocyte-derived cell types and percentage HCMV reads

We isolated CD14+ monocytes from six independently processed blood donors (A-F in Fig 1) and differentiated these cells to pro-inflammatory DCs and MΦ1; and anti-inflammatory MΦ2. DCs were characterized as CD14+/CD163-/DCSIGN++ and upon LPS treatment they showed upregulation of CD80; MΦ1 were CD14+/CD163-/DCSIGN+ whereas MΦ2 showed a CD14+/CD163+/DCSIGN+ marker profile with poor upregulation of CD80 upon LPS treatment (S1 Fig).

Subsequently, these cell-types were infected with highly epithelial tropic TB40/E HCMV strain at MOI 5. In this study, the infection was carried out for 3 hours without subsequent inactivation of the remaining surface-bound virus e.g. with citrate buffer to avoid any damage or any changes in host transcriptome of the cells thereby attempting to keep the cell-virus interplay as authentic as possible. In contrast to fibroblasts where 100% of the cells could be infected, 3.11% (2.91–4.12%) of the DCs, 1.20% (1.01–1.36%) of the MΦ1 and 6.27% (3.5–8.16%) of the MΦ2 could be infected (S2 Fig).
Illumina RNA sequencing resulted in an average of almost 23.3 million \((9.4-36.4 \times 10^6)\) sequences per sample covering both HCMV and host mRNA. In order to perform a comparative analysis of HCMV gene expression between fibroblasts and monocyte-derived cells, we argue that a more representative control is provided by infecting NHDF cells at a lower MOI thereby mimicking the actual percentage of HCMV reads in DCs and MΦs (Fig 1). Infection of DCs, MΦ1 and MΦ2 resulted in a median percentage of mapped reads of 0.63%, 0.04% and 0.67%, respectively. There was some variability in infectivity between the cells derived from different donors. We infected NHDF cells at three different MOIs and show that infection of NHDF fibroblasts at MOI 0.16 reflects better the percentage of reads mapped to the HCMV transcriptome (2.97%) in primary cells compared to MOI 0.5 (10.62%) and MOI 5 (32.12%) (Fig 1).

First, we evaluated the expression levels of each HCMV gene in NHDF cells at MOI 0.16. Expression levels are represented by the reads per kilobase of transcript per million (RPKM) value, which represents the number of mapped sequences per gene normalized for the gene length and the total amount of sequences from the experiment (accounting for experimental variability). We plotted the binned logs of these RPKM values on the HCMV genome annotation (acc.no. EF999921) used for this study (Fig 2). The highest expressing genes in this study during lytic infection at low MOI in NHDF fibroblasts were RNA4.9, RNA2.7, RNA1.2, UL22A, US18, RL12, RL13-RL14, UL4, UL5, UL40, UL41A, UL42, UL132 and UL148. On the other end of the spectrum UL12, UL38, UL80.5, UL121, UL123, UL127, truncated UL141, and vMIA are only expressed at low levels (10–100).

In contrast, no reads were mapped for UL60 and UL90 in this study.

Differentially regulated genes and gene clusters in primary cell types

Next, we compared HCMV gene expression in monocyte-derived DCs, MΦ1 and MΦ2 with HCMV gene expression in NHDF cells (Fig 3). In general, the regulation of the US region was striking in primary cell types. Especially US26, US33 (not in MΦ1), US33A, US34 and US34A were found to be expressed at higher levels in primary cells \((p<0.01)\). Also in the US region, US7-US9, US10 (only significant in MΦ1), US11 (only significant in DCs), US23 (only significant in MΦ2) and US23 (only significant in MΦ2) were expressed to higher levels in primary cells. Striking was the significantly lower expression of US28 (in DCs) and US29 (in MΦ2) compared to NHDF cells.
In particular in DCs, the regulation of HCMV genes was clustered. RL11-RL12-RL13, UL41A-UL42 and (UL1)UL4-UL11 were present in lower amounts in primary cells while, besides the aforementioned genes in the US region, UL120-UL121 and UL148D-UL149 were found to be expressed higher. In MΦ1, except for the US region, this clustering was not as clear. Individual genes such as UL25, UL100, UL141, RNA5.0, UL80, UL83 and UL86 were detected at higher expression levels in contrast with the lower expression of UL17, UL20, UL78, UL105 and UL123. UL80, UL83 and UL86 may be part of a cluster. In MΦ2, RL10-RL11 and a cluster from UL82 to UL86 (p>0.05 for UL84) as well as individual genes such as RNA5.0, UL1, UL8, UL42, UL52, UL100, UL115 and UL146 were found to be lower expressed than in NHDF fibroblasts. Also genes in MΦ2 were more difficult to cluster, only UL2-UL2 and UL148A-UL149 were significantly higher expressed neighboring genes in addition to individual genes as UL12, UL17, UL20, UL29, UL78 and UL137.

Fig 2. Expression levels of the HCMV genes transcriptome in fibroblasts (MOI 0.16). Log(10) of the RPKM values per gene were binned in six categories and mapped on the reference genome of TB40/E.

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In a head to head comparison between the different primary cell types, the only significant difference were the lower amounts of UL4-UL5, US7 and US9 in DCs compared to MΦ1. This suggests that the virus regulates its gene expression similarly in both macrophage subtypes, despite their different inflammatory character and infectability.

Functional analysis of differentially regulated genes

Each HCMV gene has a specific function during the lytic or latent part of the viral life cycle. By annotating each gene with its function [40], we were able to explore if any functional groups were significantly enriched (S3 Fig). This means that the p-values of each functional group were compared between two cell types. The enrichment of a functional groups can be driven either by the differential expression of many genes with p-values between 0.05 and 0.01; or by a small number of highly significant genes (p<0.001).

This pathway analysis showed that in DCs and MΦ1, compared to NHDF fibroblasts, the most affected functional group consisted of genes involved in immunomodulation. In DCs, the second and third functional groups which were enriched contained genes which have not been annotated with a function or genes involved in cell tropism and cell-dependent replication, respectively. In MΦ1, genes with a function in viral spread and entry contributed to the
enrichment of the similarly named functional classes. Finally, in MΦ2 only two functional groups were significantly affected, being genes involved in cell tropism and cell-specific replication; and genes involved in viral spread (Fig 4).

The group of immune-modulatory genes is the largest (38 tested genes) and the presence of many significant p-values (p<0.05) demonstrates the importance of the regulation of genes in this group in all examined monocyte-derived cells. In DCs, RL11-RL13, UL141, UL11, UL111A and UL20 were the most important differentially regulated genes involved in immunomodulation (p<0.01). To a lesser extent (0.01<p<0.05), TRS1, UL123, UL21A, UL78, US10-US11, US2 and US28 were contributing to the enrichment of this functional group. In MΦ1, functional analysis also revealed a shift in immunomodulatory genes, but this was driven by slightly different genes (based on significance) i.e. UL123, UL20, UL78, UL83, US10, US7,
US8 and US9 (p<0.01) and to a lesser extend RL11–RL12, UL141, UL11, UL111A, UL146, UL74, UL82 and US11 (0.01<p<0.05).

In both DCs and MΦ2, the second most significant functional annotation is comprised of genes with a function in cell tropism and cell-specific replication. This was due to the modulation of different genes in DCs and MΦ2. In DCs, the p-values of RL13, UL1 and UL111A were the main contributors whereas in MΦ2 the identification of this pathway was driven by UL24, UL29, UL78 and UL83 (p<0.01). In MΦ1 and MΦ2, viral spread was also found to be an enriched functional group mainly influenced by the highly significant p-values of UL74 and US9. Finally, in MΦ1, UL110 and UL74 were the main contributors to the significant p-value of the functional group involved in entry processes.

The same functional distributions were generated when primary cell types were compared to each other. Genes which were differentially regulated between DCs and macrophages mostly code for virion proteins (i.e. RL12, RL13, TRL14, US23, UL4–UL5, UL24–UL25, UL29, UL82–UL83, UL100 and UL115) which underlines the adaptability of the virion to the specific cell type where they are produced (Fig 5). When comparing DCs to MΦ1, US9 shows the highest significance in the group of factors involved in viral spread (Fig 5). Further, many differentially when comparing DCs to MΦ2 expressed genes were described in non-permissive cells and may be involved in latency (Fig 5). However, the relevance of most of these genes in lytic infection has not yet been established. Finally, when comparing DCs and MΦ2, genes contributing to cell tropism and cell-dependent replication such as UL132 and UL148 were responsible for the significance of this functional group. Between pro- and anti-inflammatory macrophage types, UL93 drives the significant enrichment of cellular trafficking factors (Fig 5).

**Discussion**

The organization of regulatory motifs in different promotor regions on the HCMV genome shows the potential for HCMV to intricately regulate its transcription under specific circumstances [41]. This was recently confirmed with rat CMV (RCMV) which followed an alternative transcriptional profile in commonly used cell lines in the laboratory and cell types which are relevant in vivo [42].

Macrophages (MΦ) and dendritic cells (DCs) can be infected with HCMV in vivo [43–47] and in vitro [44, 48]. Both cell types play a role in latency and reactivation since their progenitors i.e. CD14⁺ monocytes and CD34⁺ cells support latency whilst terminal differentiation to DCs or MΦ results in reactivation [8, 18, 30, 49–52]. Because of the limited knowledge about transcription in these cell lines and their potentially vital role in vivo, we set up a next generation sequencing study to investigate the profile of the HCMV transcriptome in monocyte-derived pro-inflammatory MΦ1, anti-inflammatory MΦ2 and DC compared to lytic infection in NHDF fibroblasts.

We infected pro-inflammatory CD14⁺/CD163⁺/DCsign⁺/CD80⁺/+LPS monocyte-derived DCs, pro-inflammatory CD14⁺/CD163⁺/DCsign⁺/CD80⁺/+LPS MΦ1 and anti-inflammatory CD14⁺/CD163⁺/DCsign⁺/CD80⁺/+LPS MΦ2 [34, 53–62]. As reported before, MΦ2 were easier to infect than MΦ1 [44] resulting in respectively 6.27% (3.5–8.16%) and 1.2% (1.01–1.36%) infection. The infectivity of DCs was intermediate between MΦ1 and MΦ2 (3.11%, 2.91–4.12). These percentages were reflected in the number of mapped reads on the HCMV genome. A study by Bayer et al. [44] reports infection rates up to 75% in MΦ2; however in none of the donors used in this study we obtained infection rates above 10%. While in literature the polarization in macrophages is perceived distinct, the differences between these cell types is much more complex and subtle [58, 60, 63, 64]. It has been shown that a plethora of monocyte-derived MΦs exists under the influence of cytokines [65], but also under influence of FCS on its own.
distinct MΦ subtypes are produced [66, 67]. Therefore, it is not unlikely to assume that small technical differences such as the method of monocyte isolation, plate type, the source of the cytokines, differences in the culture medium or the handling of the buffy coat pre-isolation have an impact on the type of monocyte-derived cell and thus the rate of infection [68, 69].

For the RNA sequencing analysis, we updated the sequence of the TB40/E-BAC4 (EF999921; 2008) with later gene annotations from KF297339 (2013). Of note, the complexity of the transcriptome may still increase as the clinical Merlin strain was recently annotated with hundreds of additional ORFs [26]. However, because genetic differences between strains may influence the transcriptome [23], we limited our analysis to the available TB40/E sequence information.

Fig 5. Enrichment of functional groups when comparing DCs with both types of macrophages and when comparing MΦ1 with MΦ2 macrophages. The HCMV was annotated and a mathematical model was used to determine which functional groups were significantly enriched in the differentially regulated genes. Based on this algorithm, genes involved in cell tropism, viral spread and immunomodulation are the most enriched groups in DCs, MΦ2 and MΦ1.
We used infection in NHDF fibroblasts as a reference cell type for lytic infection to compare HCMV gene expression in M\(\Phi\)1, M\(\Phi\)2 and DCs. It has been reported that M\(\Phi\)s support productive lytic infection characterized by the shedding of virus [44]. In agreement with these published results, all viral transcripts except UL60 and UL90, including late genes were detected in this study. We, and others, observe that the majority of the HCMV reads in NHDF, DCs, M\(\Phi\)1 and M\(\Phi\)2 is dedicated to RNA4.9, RNA2.7 (36% of all HCMV specific reads), RNA1.2, UL4, UL5 and UL22A transcripts [24]. These similarities between NHDF and the primary cell types suggest a lytic program and not a latency transcriptional profile as in CD34\(^+\) progenitor cells which only express a restricted set of transcripts and do not follow the immediate-early, early, late gene expression cascade [8, 70]. It has to be noted that the presence of late transcripts does not always rule out the presence of some degree of abortive infection as late virion-associated genes have been identified as well [71]. However, only one of these virion-associated genes (RL13) was of any significance in this study. Taken together, we conclude that in this study primary cells displayed a lytic transcriptional program.

In primary cell types, differentially regulated genes were often found in clusters suggesting that these clustered genes may have similar functions. In DCs, the downregulation of RL11-RL13-RL14-UL1 and UL4-UL11; and the upregulation of UL120-UL121, UL148D-UL149 and US33-US34A are striking. In M\(\Phi\)2, RL11-RL12 and UL82-UL86 (UL84 not significant) were upregulated and besides UL148A-UL149 and UL2-UL3, very strong clustering was seen in the US region where US7-US9 and US33-US34A were upregulated. A very similar observation was made for M\(\Phi\)1 but the downregulated cluster of UL80-UL86 was interrupted because the lower expression of UL81-UL82 and UL85 did not reach significance. Although the functions of many of the HCMV genes remain unknown, pathway analysis reveals that the functional group of immunomodulatory genes is significantly enriched in DCs, M\(\Phi\)1 and to a lesser extend in M\(\Phi\)2 compared to NHDF fibroblasts. These observations were made before in several focused studies which showed a profound effect of HCMV on the host immune-transcriptome and an active modulation of viral transcripts via viral proteins such as pp65 [31, 72–75] but recently also a transcriptomics study with RCMV showed an enrichment of immunomodulatory genes in relevant in vivo cell types [42]. Especially the regulation of the US region contributed to the enrichment of this functional group. Of note is that TB40/E-BAC4 used in this study misses US1-US6 compared to the wild type virus [32]. Future research with other strains such as Merlin may elucidate if US1-US6 are also part of this unique transcriptional program.

The US region was associated with immunomodulation before [72, 76–88]. More specifically, differentially expressed genes such as US8, US10 and US11 have a role in the modulation of MHCI molecules [76–84, 89], US9 was reported to be involved in the evasion of NK activation [90], US28 interferes with cyto- and/or chemokine production or responsiveness [72, 87, 88] and based on sequence similarity the miRNA regulated US7 [91] and the largely unknown US9 are predicted to have similar roles [92]. Other studies such as a study using superSAGE describing the transcriptome in DCs indicated the importance of immunomodulatory genes in HCMV infection of DCs [31]. Further, also in in vivo cell types infected with RCMV it was suggested that the differential expression of immunomodulatory genes is a unique way for the virus to evade the immune system during infection [42]. Also, several groups evaluated the host’s transcriptome in monocytes which, under the influence of HCMV, are driven towards a M\(\Phi\)1 phenotype. These groups found that in the host’s transcriptome, most genes that were differentially regulated were involved in anti-viral responses, inflammatory response, viral spread and apoptosis.

The US region is not fully annotated with functions and some of the most significantly regulated genes in this study such as US26 and US33-34A have unknown functions [24]. US34 was...
identified before as an abundantly expressed transcript in DCs [31] and US34A was identified as a SUMOylation target [93]. The relevance of these observations is not clear but based on this study a role in immunomodulation would not be surprising.

Next to genes involved in immunomodulation, we found that also factors involved in cell tropism and cell-specific replication were enriched, especially in DCs and MΦ2 compared to NHDF fibroblasts. The genes responsible for this enrichment are different for both cell types, i.e., RL13-RL14, UL1 and UL111A in DCs versus UL24, UL29, UL78 and UL83 in MΦ2. The differential regulation of cell-tropic factors but also of viral spread factors (driven by UL74, US27 and US9) suggests that the virion may adapt to optimally infect surrounding cells of the same type as previous research already suggested [27, 94].

In addition, there are also discrete differences when comparing the different primary cell types head to head. Between DCs and MΦ1, there were four genes (UL4, UL5, US7 and US9) significantly lower expressed in DCs. While US7 and US9 only have proposed roles, UL4 and UL5 are virion proteins and their differential regulation may once more suggest the adaptability of the virion when infecting different cell types. Despite their different inflammatory character and infectability no genes were found to be differentially expressed (p < 0.05) when comparing MΦ1 and MΦ2 or when comparing DCs and MΦ2. In a previous study investigating HCMV infection in both MΦ subtypes, it was reported that IE1-2 (UL122-123), pp65 (UL83) and pp150 (UL32) were more abundantly present in MΦ2 [44], however in this study this difference did not reach significance. Perhaps, if a sequencing study was carried out on different time points, these differences could reach significance.

The HCMV genome is foreseen of several regulatory motifs and can replicate in a variety of cell types suggesting that in each of these cell types, a specific transcriptional program is followed [42, 95]. This HCMV study is the first transcriptomics study which compares gene expression in commonly used NHDF fibroblasts and DCs, MΦ1 and MΦ2, all cell types which are considered relevant for in vivo viral spread. By using pathway analysis, we show that compared to fibroblasts, a unique transcriptional profile is observed in primary cells in which genes involved in immunomodulation genes, cell tropism and replication, viral spread and virion proteins are enriched. While the differential regulation of these specific pathways is still consistent with lytic infection, it resembles more closely a lytic low level persistent infection with careful adaptations to the cellular environment rather than with a fulminant lytic infection as observed in fibroblasts. These observations not only show the adaptability of the virus, but also explain any discrepancies between what may happen in vivo and what is observed in the laboratory.

As in the RCMV study, many questions remain unanswered. We report a number of genes that are strongly upregulated in the US region, but defined functions for these genes have not been described and functional research is needed to delineate their actual role in the viral life cycle. Especially, the high expression of the US33-US34 cluster warrants further research as it is the first time that alterations in gene expression levels are observed for these genes. It would be interesting to attribute these genes with critical functions in primary cells. In addition, it is unclear whether cellular factors in immune cells such as DC and MΦs push the virus into the production of specific transcripts or if it is the virus that manipulates the cell by producing transcripts that aid with immune-evasion. Most likely it is a combination of both. Several groups evaluated the host's transcriptome in monocytes which, under the influence of HCMV, are driven towards a MΦ1 phenotype. These groups found that in the host's transcriptome, most genes that were differentially regulated were involved in anti-viral responses, inflammatory response, viral spread and apoptosis which is consistent with the differential regulation of viral genes involved in these processes reported in this study [96–98]. Yet, it is an intriguing observation that MΦ1 and MΦ2, which display distinct polarization, show a similar viral
transcription profiles. Also, it may be warranted to dedicate future research to investigate regular and short lived HCMV transcripts using specialized methods preferably in a kinetic time course set up [99]. Finally, since expression levels of transcripts are not always reflected in actual protein expression profiles, it would be interesting to correlate the findings presented here to proteomics data once these data are available.

Supporting Information

S1 Fig. Surface marker analysis of monocyte-derived cell types. Monocytes were seeded and differentiated to MΦ1, MΦ2 and DCs or left untreated. After 7 days, the cells were characterized using FACS. The boxplots are based on 11 different and independently processed donors. The boxes represent all counts within the 25th and 75th percentile; error bars show the 10th and 90th percentile. All values outside the 10th and 90th percentiles are considered outliers (•), the mean is indicated as a horizontal line. A one way ANOVA (Holm-Sidak method) was used to test for significant differences between cell types. (Panel A) CD14 surface expression was significantly different (p > 0.05) between all cell types (Mo vs MΦ1, Mo vs MΦ2, Mo vs DC; MΦ1 vs DC; MΦ2 vs DC), except between MΦ1 and MΦ2 (p > 0.05). (Panel B) DC marker DCSign was expressed significantly higher on DCs and all other cell types (p < 0.05; Mo vs DC, MΦ1 vs DC, MΦ2 vs DC). DCSign was also expressed more on both types of MΦ compared to monocytes (p < 0.05). No statistical difference was observed between both types of macrophages (p > 0.05). (Panel C) CD163 was only expressed on MΦ2 and poorly on all other cell types (p < 0.05; MΦ2 vs Mo, MΦ2 vs MΦ1, MΦ2 vs DC). (Panel D) To assess CD80 response, all cell types were challenged with 500ng/ml LPS for 24h. MΦ2 macrophages showed a significantly lower response in C80 upregulation compared with the other cell types (p < 0.05; MΦ2 vs Mo, MΦ2 vs MΦ1, MΦ2 vs DC). CD80 expression between all other cell types was comparable (p > 0.05).

S2 Fig. A representative image (donor A) of the number of IE expressing cells in TB40/E and mock (UV-inactivated TB40/E) infected primary cell types. In each representative image, we indicated the average number and the range of IE expressing cells for three independently processed donors.

S3 Fig. Distribution of functions of DCs, MΦ1 and MΦ2 compared to NHDF fibroblasts and of primary cells compared to each other. The most enriched functions are represented by the highest bars. The cut off to determine the most significantly enriched functions was based on the point where the box plot curves flatten out (marked by a red box).

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References
1. Cannon MJ, Schmid DS, Hyde TB. Review of cytomegalovirus seroprevalence and demographic characteristics associated with infection. Rev Med Virol. 2010; 20(4):202–13. Epub 2010/06/22. doi: 10.1002/rmv.655 PMID: 20564615.
2. Murphy EA, Shenk T. Human cytomegalovirus Genome. In: Shenk T, Stinski MF, editors. Human cyto- megalovirus. Heidelberg: Springer Berlin Heidelberg; 2008. p. 1–20.
3. Kondo K, Kaneshima H, Mocarski ES. Human cytomegalovirus latent infection of granulocyte-macro- phage progenitors. Proceedings of the National Academy of Sciences of the United States of America. 1994; 91(25):11879–83. PMID: 7991550; PubMed Central PMCID: PMC453339.
4. Kondo K, Mocarski ES. Cytomegalovirus latency and latency-specific transcription in hematopoietic progenitors. Scandinavian journal of infectious diseases Supplementum. 1995; 99:63–7. PMID: 8668945.
5. Slobedman B, Mocarski ES. Quantitative analysis of latent human cytomegalovirus. J Virol. 1999; 73(6):4806–12. Epub 1999/05/11. PMID: 10233941.
6. Bego M, Maciejewski J, Khaiboullina S, Pari G, St Jeor S. Characterization of an antisense transcript spanning the UL81-82 locus of human cytomegalovirus. J Virol. 2005; 79(17):11022–34. Epub 2005/08/17. 79/17/11022 [pii] doi: 10.1128/JVI.79.17.11022-11034.2005 PMID: 16103153.
7. Bego MG, Keyes LR, Maciejewski J, St Jeor SC. Human cytomegalovirus latency-associated protein LUNA is expressed during HCMV infections in vivo. Arch Virol. 2011; 156(10):1847–51. Epub 2011/06/01. doi: 10.1007/s00705-011-1027-7 PMID: 21625978.
8. Goodrum FD, Jordan CT, High K, Shenk T. Human cytomegalovirus gene expression during infection of primary hematopoietic progenitor cells: a model for latency. Proc Natl Acad Sci U S A. 2002; 99(25):16255–60. Epub 2002/11/29. doi: 10.1073/pnas.252630899 252630899 [pii] PMID: 12456880.
9. Keyes LR, Hargett D, Soland M, Bego MG, Rossetto CC, Almeida-Porada G, et al. HCMV Protein LUNA is Required for Viral Reactivation from Latently Infected Primary CD14(+) Cells. PLoS One. 2012; 7(12):e52827. Epub 2013/01/10. doi: 10.1371/journal.pone.0052827 PONE-D-11-22739 [pii] PMID: 23300789.
10. Rossetto CC, Tarrant-Elorza M, Pari GS. Cis and Trans Acting Factors Involved in Human Cytomegalovirus Experimental and Natural Latent Infection of CD14 (+) Monocytes and CD34 (+) Cells. PLoS Pathog. 2013; 9(5):e1003366. Epub 2013/05/30. doi: 10.1371/journal.ppat.1003366 PPATHOGENS-D-12-02509 [pii] PMID: 23717203.
11. Avdic S, Cao JZ, Cheung AK, Abendroth A, Slobedman B. Viral interleukin-10 expressed by human cytomegalovirus during the latent phase of infection modulates latently infected myeloid cell differentiation. J Virol. 2011; 85(14):7465–71. doi: 10.1128/JVI.00088-11 PMID: 21593144; PubMed Central PMCID: PMC3126599.
12. Cheung AK, Abendroth A, Cunningham AL, Slobedman B. Viral gene expression during the establishment of human cytomegalovirus latency in myeloid progenitor cells. Blood. 2006; 108(12):3691–9. Epub 2006/08/26. blood-2005-12-026682 [pii] doi: 10.1182/blood-2005-12-026682 PMID: 16931631.
13. Goodrum F, Reeves M, Sinclair J, High K, Shenk T. Human cytomegalovirus sequences expressed in latently infected individuals promote a latent infection in vitro. Blood. 2007; 110(3):937–45. doi: 10.1182/blood-2007-01-070078 PMID: 17440050; PubMed Central PMCID: PMC1924770.
14. Petruccelli A, Rak M, Grainger L, Goodrum F. Characterization of a novel Golgi apparatus-localized latency determinant encoded by human cytomegalovirus. J Virol. 2009; 83(11):5615–29. doi: 10.1128/JVI.01989-08 PMID: 19297488; PubMed Central PMCID: PMC2681962.
15. Poole E, Walther A, Raven K, Benedict CA, Mason GM, Sinclair J. The Myeloid Transcription Factor GATA-2 Regulates the Viral UL144 Gene during Human Cytomegalovirus Latency in an Isolate-
Reeves MB, Lehner PJ, Sissons JG, Sinclair JH. An in vitro model for the regulation of human cytomegalovirus latency and reactivation in dendritic cells. J Gen Virol. 2005; 86(Pt 11):2949–54. Epub 2005/10/18. 86/11/2949 doi: 10.1099/vir.0.81161-0 PMID: 16227215.

Reeves MB, MacAr PA, Lehner PJ, Sissons JG, Sinclair JH. Latency, chromatin remodeling, and reactivation of human cytomegalovirus in the dendritic cells of healthy carriers. Proc Natl Acad Sci USA. 2005; 102(11):4140–5. Epub 2005/03/02. 0408994102 doi: 10.1073/pnas.0408994102 PMID: 15738399.

Stenberg RM, Witte PR, Stinski MF. Multiple spliced and unspliced transcripts from human cytomegalovirus immediate-early region 2 and evidence for a common initiation site within immediate-early region 1. J Virol. 1985; 56(3):665–75. Epub 1985/12/01. PMID: 2994923.

Stinski MF, Thomsen DR, Stenberg RM, Goldstein LC. Organization and expression of the immediate early genes of human cytomegalovirus. J Virol. 1983; 46(1):1–14. Epub 1983/04/01. PMID: 6298447.

Wathen MW, Stinski MF. Temporal patterns of human cytomegalovirus transcription: mapping the viral RNAs synthesized at immediate early, early, and late times after infection. J Virol. 1982; 41(2):462–77. Epub 1982/02/01. PMID: 6281461.

Dolan A, Cunningham C, Hector RD, Hassan-Walker AF, Lee L, Addison C, et al. Genetic content of wild-type human cytomegalovirus. J Gen Virol. 2004; 85(Pt 5):1301–12. Epub 2004/04/24. PMID: 15105547. doi: 10.1099/vir.0.79888-0

Gatherer D, Seirafiyan S, Cunningham C, Holton M, Dargan DJ, Baluchova K, et al. High-resolution human cytomegalovirus transcriptome. Proc Natl Acad Sci U S A. 2011; 108(49):19755–60. Epub 2011/11/24. 1115861108 doi: 10.1073/pnas.1115861108 PMID: 22109557.

Hook L, Hancock M, Landais I, Grabski R, Britt W, Nelson JA. Cytomegalovirus microRNAs. Current opinion in virology. 2014; 7C:40–6. doi: 10.1016/j.coviro.2014.03.015 PMID: 24769092.

Stern-Ginossar N, Weisburd B, Michaels A, Le VT, Hein MY, Huang SX, et al. Decoding human cytomegalovirus. Science. 2012; 338(6110):1088–93. doi: 10.1126/science.1227919 PMID: 23180859; PubMed Central PMCID: PMC3817102.

Towler JC, Ebrahim B, Lane B, Davison AJ, Dargan DJ. Human cytomegalovirus transcriptome activity differs during replication in human fibroblast, epithelial and astrocyte cell lines. J Gen Virol. 2012; 93(16):2013–2019. Epub 2012/12/01. vir.0.010538-0 PMID: 22258857.

Sissons JG, Bain M, Wills MR. Latency and reactivation of human cytomegalovirus. J Infect. 2002; 44(2):73–7. Epub 2002/06/22. doi: 10.1053/jinf.2001.0948 S016344530190948X [pii]. PMID: 12076064.

Springer KL, Weinberg A. Cytomegalovirus infection in the era of HAART: fewer reactivations and more immunity. J Antimicrob Chemother. 2004; 54(3):582–6. Epub 2004/07/30. doi: 10.1093/jac/dkh396 dhk396 [pii]. PMID: 15282241.

Soderberg-Naucler C, Streiblow DN, Fish KN, Allan-Yorke J, Smith PP, Nelson JA. Reactivation of latent human cytomegalovirus in CD14(+) monocytes is differentiation dependent. J Virol. 2001; 75(16):7543–54. Epub 2001/07/20. doi: 10.1128/JVI.75.16.7543-7554.2001 PMID: 11462026.

Raftery MJ, Moncke-Buchner E, Matsumura H, Giese T, Winkelmann A, Reuter M, et al. Unravelling the interaction of human cytomegalovirus with dendritic cells by using SuperSAGE. J Gen Virol. 2009; 90(Pt 9):2221–33. doi: 10.1099/vir.0.010538-0 PMID: 19439557.

Sinzger C, Hahn G, Digez M, Katona R, Sampaio KL, Messerle M, et al. Cloning and sequencing of a highly productive, endothe liotropic virus strain derived from human cytomegalovirus TB40/E. J Gen Virol. 2008; 89(Pt 2):359–68. Epub 2008/01/17. 89/2/359 [pii] doi: 10.1099/vir.0.038286-0 PMID: 18198366.

Chou SW, Scott KM. Rapid quantitation of cytomegalovirus and assay of neutralizing antibody by using monoclonal antibody to the major immediate-early viral protein. J Clin Microbiol. 1988; 26(3):504–7. Epub 1988/03/01. PMID: 2833529.

Verreck FA, de Boer T, Langenberg DM, van der Zanden L, Ottenhoff TH. Phenotypic and functional profiling of human proinflammatory type-1 and anti-inflammatory type-2 macrophages in response to microbial antigens and IFN-gamma- and CD40L-mediated costimulation. J Leukoc Biol. 2006; 79(2):285–93. Epub 2005/12/07. jlb.0105015 [pii] doi: 10.1189/jlb.0105015 PMID: 16330536.
35. HARGETT D, SHENK TE. Experimental human cytomegalovirus latency in CD14+ monocytes. Proc Natl Acad Sci U S A. 2010; 107(46):20039–44. Epub 2010/11/03. 1014509107 [pii] doi: 10.1073/pnas.1014509107 PMID: 21041645.

36. GENTLEMAN RC, CAREY VJ, BATES DM, BOLSTAD B, DETTLING M, DUDOIT S, et al. Bioconductor: open software development for computational biology and bioinformatics. Genome biology. 2004; 5(10):R80. doi: 10.1186/gb-2004-5-10-r80 PMID: 15461798; PubMed Central PMCID: PMC545600.

37. ANDERS S, HUBER W. Differential expression analysis for sequence count data. Genome biology. 2010; 11(10):R106. doi: 10.1186/gb-2010-11-10-r106 PMID: 20979621; PubMed Central PMCID: PMC3218662.

38. BENJAMINI Y, HO. Controlling the False Discovery Rate: A Practical and Powerful Approach to Multiple Testing. Journal of the Royal Statistical Society. 1995; 57:pp. 289–300.

39. ARRODE G, DAVRINCHE C. Dendritic cells and HCMV cross-presentation. Curr Top Microbiol Immunol. 2003; 276:277–94. Epub 2003/06/12. PMID: 12797453.

40. BAYER C, VARANI S, WANG L, WALTHER P, ZHOU S, STRASCHEWSKI S, et al. Human cytomegalovirus infection of M1 and M2 macrophages triggers inflammation and autologous T-cell proliferation. J Virol. 2013; 87(1):67–79. Epub 2012/10/12. JVI.01585-12 [pii] doi: 10.1128/JVI.01585-12 PMID: 23055571.

41. CHAMBERS J, ANGULO A, AMARUTANGA D, GUO H, JIANG Y, WAN JS, et al. DNA microarrays of the complex human cytomegalovirus genome: profiling kinetic classes with drug sensitivity of viral gene expression. J Virol. 1999; 73(7):5757–66. Epub 1999/06/11. PMID: 10364327.

42. BOLSTAD B, DETTLING M, DUDOIT S, et al. Bioconductor: open software development for computational biology and bioinformatics. Genome biology. 2004; 5(10):R80. doi: 10.1186/gb-2004-5-10-r80 PMID: 15461798; PubMed Central PMCID: PMC545600.

43. SCHNEIDER-NAUCLR C, FISH KN, NELSON JA. Efficient human cytomegalovirus reactivation is maturation dependent in the Langerhans dendritic cell lineage and can be studied using a CD14+ experimental latency model. J Virol. 2012; 86(16):8507–15. Epub 2012/06/01. JVI.00598-12 [pii] doi: 10.1128/JVI.00598-12 PMID: 22647696.

44. SODERBERG-NAUCLR C, FISH KN, NELSON JA. Reactivation of latent human cytomegalovirus by alloimmune stimulation of blood cells from healthy donors. Cell. 1997; 91(1):119–26. PMID: 9335340.

45. BULLWINKEL J, LUDERMANN A, DEBARRY J, SINGH PB. Epigenotype switching at the CD14 and CD209 genes during differentiation of human monocytes to dendritic cells. Epigenetics. 6(1):45–51. Epub 2010/09/08. 13314 [pii] doi: 10.4161/epi.6.1.13314 PMID: 20818162.
54. Schmitte A, Scheibenbogen C, Keilholz U. Lipopolysaccharide effectively up-regulates B7-1 (CD80) expression and costimulatory function of human monocytes. Scand J Immunol. 1995; 42(6):701–4. Epub 1995/12/01. PMID: 8552895.

55. Reis e Sousa C, Hiery S, Scharton-Kersten T, Jankovic D, Charest H, Germain RN, et al. In vivo microbial stimulation induces rapid CD40 ligand-independent production of interleukin 12 by dendritic cells and their redistribution to T cell areas. J Exp Med. 1997; 186(11):1819–29. Epub 1998/01/07. PMID: 9362881.

56. de Saint-Vis B, Fugier-Vivier I, Massacrier C, Gaillard C, Vanbervliet B, Alt-Yahia S, et al. The cytokine profile expressed by human dendritic cells is dependent on cell subtype and mode of activation. J Immunol. 1998; 160(4):1666–76. Epub 1998/02/20. PMID: 9469432.

57. Steinbach F, Krause B, Blass S, Burmester GR, Hiepe F. Development of accessory phenotype and function during the differentiation of monocyte-derived dendritic cells. Research in immunology. 1998; 149(7–8):627–32. PMID: 9851513.

58. Mantovani A, Sica A, Sozzani S, Allavena P, Vecchi A, Locati M. The chemokine system in diverse forms of macrophage activation and polarization. Trends Immunol. 2004; 25(12):677–86. Epub 2004/11/09. S1471-4906(04)00295-9 [pii] doi: 10.1016/j.it.2004.09.015 PMID: 15530839.

59. D’Andrea A, Aste-Amezaga M, Valiante NM, Ma X, Kubin M, Trinchieri G. Interleukin 10 (IL-10) inhibits human lymphocyte interferon gamma-production by suppressing natural killer cell stimulatory factor/IL-12 synthesis in accessory cells. J Exp Med. 1993; 178(3):1041–8. Epub 1993/09/01. PMID: 8102386.

60. Elgini S, Brioschi M, Fiorelli S, Tremoli E, Banfi C, Colli S. Human monocytic-derived macrophages are heterogeneous: Proteomic profile of different phenotypes. Journal of proteomics. 2013; 124:112–23. doi: 10.1016/j.jprot.2013.03.026 PMID: 25857278.

61. Vogt G, Nathan C. In vitro differentiation of human macrophages with enhanced antimycobacterial activity. J Clin Invest. 2011; 121(10):3889–901. doi: 10.1172/JCI57235 PMID: 21911938; PubMed Central PMCID: PMC3195467.

62. Goodrum F, Jordan CT, Terhune SS, High K, Shenk T. Differential outcomes of human cytomegalovirus infection in primitive hematopoietic cell subpopulations. Blood. 2004; 104(3):687–95. Epub 2004/04/20. doi: 10.1182/blood-2003-12-4344 2003-12-4344 [pii] PMID: 15090458.

63. Bresnahan WA, Shenk T. A subset of viral transcripts packaged within human cytomegalovirus particles. Science. 2000; 288(5475):2373–6. PMID: 10879294.

64. Gurczynski SJ, Das S, Pellet PE. Deletion of the human cytomegalovirus US17 gene increases the ratio of genomes per infectious unit, and alters regulation of immune and ER stress response genes at early and late times after infection. Journal of virology. 2013. doi: 10.1128/JVI.02704-13 PMID: 24335296.
73. Mezger M, Bonin M, Kessler T, Gebhardt F, Einsele H, Loeffler J. Toll-like receptor 3 has no critical role during early immune response of human monocyte-derived dendritic cells after infection with the human cytomegalovirus strain TB40E. Viral immunology. 2009; 22(6):343–51. doi: 10.1089/vim.2009.0011 PMID: 19951172.

74. Renneson J, Dutta B, Gorily S, Danis B, Lecomte S, Laes JF, et al. IL-12 and type I IFN response of neonatal myeloid DC to human CMV infection. European journal of immunology. 2009; 39(10):2789–99. doi: 10.1002/eji.200939414 PMID: 19637227.

75. Zhu H, Cong JP, Shenk T. Use of differential display analysis to assess the effect of human cytomegalovirus infection on the accumulation of cellular RNAs; induction of interferon-responsive RNAs. Proceedings of the National Academy of Sciences of the United States of America. 1997; 94(25):13985–90. PMID: 9391139; PubMed Central PMCID: PMC284119.

76. Hesse J, Ameres S, Besold K, Krauter S, Moosmann A, Plachter B. Suppression of CD8+ T-cell recognition in the immediate-early phase of human cytomegalovirus infection. J Gen Virol. 2013; 94(Pt 2):376–86. doi: 10.1099/vir.0.045662-0 PMID: 23100361.

77. Noriega VM, Hesse J, Gardner TJ, Besold K, Plachter B, Tortorella D. Human cytomegalovirus US3 modulates destruction of MHC class I molecules. Molecular immunology. 2012; 51(2):245–53. doi: 10.1016/j.molimm.2012.03.024 PMID: 22497807; PubMed Central PMCID: PMC3367378.

78. Oresic K, Noriega V, Andrews L, Tortorella D. A structural determinant of human cytomegalovirus US2 dictates the down-regulation of class I major histocompatibility molecules. J Biol Chem. 2006; 281(28):19395–406. doi: 10.1074/jbc.M601026200 PMID: 16687410.

79. Oresic K, Tortorella D. Endoplasmic reticulum chaperones participate in human cytomegalovirus US2-mediated degradation of class I major histocompatibility complex molecules. J Gen Virol. 2008; 89(Pt 5):1122–30. doi: 10.1099/vir.0.83516-0 PMID: 18420789; PubMed Central PMCID: PMC2634742.

80. Wiertz EJ, Jones TR, Sun L, Bogyo M, Geuze HJ, Ploegh HL. The human cytomegalovirus US11 gene product dislocates MHC class I heavy chains from the endoplasmic reticulum to the cytosol. Cell. 1996; 84(5):769–79. PMID: 8625414.

81. Park B, Spooner E, Houser BL, Strominger JL, Ploegh HL. The HCMV membrane glycoprotein US10 selectively targets HLA-G for degradation. J Exp Med. 2010; 207(9):2033–41. doi: 10.1084/jem.20091793 PMID: 20713594; PubMed Central PMCID: PMC2931171.

82. Tirabassi RS, Ploegh HL. The human cytomegalovirus US8 glycoprotein binds to major histocompatibility complex class I products. Journal of virology. 2002; 76(15):6832–5. PMID: 12050396; PubMed Central PMCID: PMC1362586.

83. Furman MH, Dey N, Tortorella D, Ploegh HL. The human cytomegalovirus US10 gene product delays trafficking of major histocompatibility complex class I molecules. Journal of virology. 2002; 76(22):11753–6. PMID: 12308773; PubMed Central PMCID: PMC136774; PubMed Central PMCID: JVI.76.22.11753-11756.2002

84. Tirosh B, Ikawaishi NN, Lilley BN, Lee AH, Glimcher LH, Ploegh HL. Human cytomegalovirus protein US11 provokes an unfolded protein response that may facilitate the degradation of class I major histocompatibility complex products. Journal of virology. 2005; 79(5):2768–78. doi: 10.1128/JVI.79.5.2768-2779.2005 PMID: 15708995; PubMed Central PMCID: PMC548438.

85. Dugan GE, Hewitt EW. Structural and Functional Dissection of the Human Cytomegalovirus Immune Evasion Protein US6, J Virol. 2008; 82(7):3271–82. doi: 10.1128/JVI.01705-07 PMID: 18199642; PubMed Central PMCID: PMC2268496.

86. Kim Y, Park B, Cho S, Shin J, Cho K, Jun Y, et al. Human cytomegalovirus UL18 utilizes US6 for evading the NK and T-cell responses. PLoS Pathog. 2008; 4(8):e1000123. doi: 10.1371/journal.ppat.1000123 PMID: 18688275; PubMed Central PMCID: PMC2483941.

87.Arnolds KL, Lares AP, Spencer JV. The US27 gene product of human cytomegalovirus enhances signaling of host chemokine receptor CXCR4. Virology. 2013; 439(2):122–31. doi: 10.1016/j.virol.2013.02.006 PMID: 23490053; PubMed Central PMCID: PMC3639318.

88. Stropes MP, Schneider OD, Zagorski WA, Miller JL, Miller WE. The carboxy-terminal tail of human cytomegalovirus (HCMV) US28 regulates both chemokine-independent and chemokine-dependent signaling in HCMV-infected cells. J Virol. 2009; 83(19):10016–27. doi: 10.1128/JVI.00354-09 PMID: 19605482; PubMed Central PMCID: PMC2748033.

89. van den Boomen DJ, Lehner PJ. Identifying the ERAD ubiquitin E3 ligases for viral and cellular targeting of MHC class I. Molecular immunology. 2015; 68(2 Pt A):106–11. doi: 10.1016/j.molimm.2015.07.005 PMID: 26210183; PubMed Central PMCID: PMC4678111.

90. Seidel E, Le VT, Bar-On Y, Tsukerman P, Enk J, Yamin R, et al. Dynamic Co-evolution of Host and Pathogen: HCMV Downregulates the Prevalent Allele MICA *008 to Escape Elimination by NK Cells. Cell reports. 2015. doi: 10.1016/j.celrep.2015.01.029 PMID: 25683719; PubMed Central PMCID: PMC4641326.
91. Tirabassi R, Hook L, Landais I, Grey F, Meyers H, Hewitt H, et al. Human cytomegalovirus US7 is regulated synergistically by two virally encoded microRNAs and by two distinct mechanisms. J Virol. 2011; 85(22):11938–44. doi: 10.1128/JVI.05443-11 PMID: 21900172; PubMed Central PMCID: PMC3209316.

92. Apweiler R, Bairoch A, Wu CH, Barker WC, Boeckmann B, Ferro S, et al. UniProt: the Universal Protein knowledgebase. Nucleic Acids Res. 2004; 32(Database issue):D115–9. doi: 10.1093/nar/gkh131 PMID: 14681372; PubMed Central PMCID: PMC308865.

93. Kim ET, Kim YE, Kim YJ, Lee MK, Hayward GS, Ahn JH. Analysis of human cytomegalovirus-encoded SUMO targets and temporal regulation of SUMOylation of the immediate-early proteins IE1 and IE2 during infection. PLoS One. 2014; 9(7):e103308. doi: 10.1371/journal.pone.0103308 PMID: 25050850; PubMed Central PMCID: PMC4106884.

94. Gibson W. Structure and Formation of the Cytomegalovirus Virion. In: Shenk T, Stinski MF, editors. Human Cytomegalovirus. Berlin Heidelberg: Springer-Verlag; 2008. p. 187–204.

95. Kreklywich CN, Smith PP, Jones CB, Cornel A, Orloff SL, Streblow DN. Fluorescence-based laser capture microscopy technology facilitates identification of critical in vivo cytomegalovirus transcriptional programs. Methods Mol Biol. 2014; 1119:217–37. doi: 10.1007/978-1-62703-788-4_13 PMID: 24639226; PubMed Central PMCID: PMC4347879.

96. Chan G, Nogalski MT, Yurochko AD. Human cytomegalovirus stimulates monocyte-to-macrophage differentiation via the temporal regulation of caspase 3. J Virol. 2012; 86(19):10714–23. Epub 2012/07/28. JVI.07129-11 pii doi: 10.1128/JVI.07129-11 PMID: 22837201.

97. Smith MS, Bentz GL, Smith PM, Bivins ER, Yurochko AD. HCMV activates PI(3)K in monocytes and promotes monocyte motility and transendothelial migration in a PI(3)K-dependent manner. J Leukoc Biol. 2004; 76(1):65–76. Epub 2004/04/27. doi: 10.1189/jlb.1203621 jlb.1203621 pii. PMID: 15107461.

98. Stevenson EV, Collins-McMillen D, Kim JH, Cieply SJ, Bentz GL, Yurochko AD. HCMV reprogramming of infected monocyte survival and differentiation: a Goldilocks phenomenon. Viruses. 2014; 6(2):782–807. doi: 10.3390/v6020782 PMID: 24531335; PubMed Central PMCID: PMC3939482.

99. Dolken L, Ruzsics Z, Radle B, Friedel CC, Zimmer J, Mages J, et al. High-resolution gene expression profiling for simultaneous kinetic parameter analysis of RNA synthesis and decay. Rna. 2008; 14(9):1959–72. doi: 10.1261/rna.1136108 PMID: 18658122; PubMed Central PMCID: PMC2552961.