A pVII positioning code determines the dynamics of Adenoviral nucleoprotein structure and primes it for early gene activation

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SUMMARY

Inside the capsid adenovirus DNA is associated with the structural protein pVII. However, the viral DNA organisation, pVII positioning and the dynamic changes of viral packaging upon infection, to form a transcriptional active genome, are not known. We combined MNase-Seq and single genome imaging during early infection to provide the structure and time resolved dynamics of viral chromatin changes, correlated with gene transcription. pVII complexes form nucleosome-like arrays, being precisely positioned on DNA, creating a defined and unique adenoviral nucleoprotein-architecture. The structure renders the viral genome transcription competent with lower pVII densities at early gene loci, correlating with viral chromatin decondensation upon infection. Nucleosomes specifically replace pVII at transcription start sites of early genes, preceding transcriptional activation. Our study suggests an underlying regulatory pVII nucleoprotein-architecture, required for the dynamic changes during early infection, including transcription related nucleosome assembly. We suggest that our study provides a basis for the development of recombinant adenoviral vectors exhibiting sustained expression in gene therapy.

INTRODUCTION

Most DNA viruses transport their genomes to the nucleus of uninfected target cells to initiate viral gene expression and genome replication. During cytoplasmic trafficking genomes are highly compacted inside the viral capsid to protect them from nucleic acid sensing molecules. Inside the nucleus genomes are decondensed and transcriptionally activated for viral gene expression and replication. Replicated genomes are then re-condensed for packaging into progeny. While this remarkable and reversible genome plasticity is highly efficient, little is known about the structural organization of the genome and their dynamic changes and regulatory cues of this process during virus entry.

Adenoviruses (Ad) are prototypic DNA viruses that contain a linear double stranded DNA genome of 30-40 kb, being incorporated into a ~ 90 nm non-enveloped icosahedral capsid shell. Ads enter cells by receptor-mediated endocytosis. Entry cues liberate the internal capsid protein VI and permit the virus to escape the endosomal compartment ^{1,2}. Cytoplasmic capsids are transported to the nuclear pore complex (NPC) where they dock, disassemble and release the viral genome for nuclear import ^{3,4}. Within hours after cell entry, nuclear genomes initiate gene expression from the immediate early promoters, as E1A. E1A encodes a transactivator driving early genes responsible for cell and immune modulation (E1B, E3, E4) and replication (E2). In

contrast activation of late genes (L1-L5) encoding most structural proteins, requires prior genome replication^{5,6}.

Packaged genomes are devoid of histones. Instead they associate with basic core proteins VII (pVII), V (pV) and polypeptide μ ^{7,8}. pVII is the most tightly bound and most abundant protein with over 500 copies per virion and packages the viral genome into a structure appearing as irregular “beads-on-a-string” by electron microscopy⁹. pVII, exhibits sequence similarities to protamines and may use its positive charges to neutralise and to compact the viral DNA^{10–12}. No information on genomic binding sites of pVII, its structural organisation on DNA, or its dynamics during infection is available¹³. pVII remains associated with the imported genome while pV dissociates from the viral DNA¹⁴. The fate of the μ -peptide is not known.

Some studies suggest that pVII remains attached throughout the early phase of infection while others argue for a gradual and at least partial removal from the incoming genome^{10,15–18}. pVII was shown to functionally interact with the viral transactivator E1A, pinpointing to a role in transcriptional activation¹⁰. It also was suggested that transcription would subsequently remove pVII from the genome¹⁹. In contrast chromatin immune precipitation (ChIP) analysis proposed that pVII remains associated with viral chromatin for several hours^{10,15–17,20}.

Functional chromatin extraction by using high and low concentrations of micrococcal nuclease (MNase) to digest cellular chromatin, combined with high-throughput sequencing, is a way to analyse protein binding to DNA and to reveal dynamic changes in chromatin structure^{21,22}. In this study we combined time resolved MNase and RNA sequencing and single molecule imaging to correlate chromatin structure changes with transcription at very early time points of adenoviral infection. This approach revealed the specific nucleoprotein architecture of the virus, which dynamically changes during early infection. pVII positioning is driven by sequence and is specifically organized at early genes, correlating with the early de-condensation of these regions upon infection. Defined pVII nucleoprotein complexes are precisely exchanged by nucleosomes on early gene promoters, showing pVII remodeling and nucleosome assembly preceding transcriptional activation and not being its consequence. The time resolved single molecule imaging of transcribing genomes support our findings, identifying the morphological correlate at the single genome level at subcellular resolution. The defined positioning of pVII on the viral genome and its functional involvement in dynamic chromatin rearrangement during infection, suggest a key role in regulating viral life cycle. We propose that knowledge of the adenovirus nucleoprotein architecture could lead to the design of improved Ad vectors for gene-therapy or vaccination, with sustained expression.

RESULTS

Time resolved analysis of the early adenovirus transcriptome by RNA-Sequencing

To investigate the early spatio-temporal reorganization of viral chromatin and transcription activation we infected H1299 lung epithelial cells with a partially E3-deleted but replication competent HAd-C5. The inoculum was replaced with fresh medium after 30 min defining time point zero of the experimental setup (Fig. 1A). On time point 0, 0.5, 1, 2, and 4 hours post infection (hpi) infected cells were subjected to immune fluorescence analysis, RNA-sequencing (RNA-seq) and functional chromatin extraction combined with high-throughput sequencing of protected DNA fragments (MNase-seq) (Fig. 1A). To verify the progression of infection, capsid protein VI (pVI) exposure during endosomal escape was monitored by immunofluorescence²³.

The subsequent import of the nuclear genome, was monitored by immunofluorescence of the genome associated pVII (Fig. 1B)¹⁸. pVI and pVII quantification showed that endosomal escape occurred within 30 min of virus addition (Fig. 1B and 1C) followed by rapid nuclear delivery of viral genomes starting at 30 min post inoculum removal (0.5 hpi), with most genomes being imported 1 hpi (Fig. 1B and 1D). The data confirms a synchronized infection, covering pre- and post-nuclear genome delivery phases resembling previous dynamic entry analysis²⁴.

To investigate viral transcription, total RNA was isolated in two independent experiments. A polyA library was prepared to enrich for mRNA transcripts and sequenced on Illumina HiSeq platform yielding on average 26 Mio reads per sample (Table S1). The reads were mapped to an extended version of the human genome amended by the annotation of the HAd-C5 genome. First Ad transcriptional activity could be detected 2 hpi, and strongly increased levels were observed 4 hpi (Fig. 1E and Table S1, whole genome profile shown in Fig. S1). Considering that the first Ad genomes in the nucleus were detected at 0.5 hpi (Fig. 1B and 1C), Ad transcription initiation and mRNA synthesis exhibited a significant lag-time after nuclear entry, suggesting the existence of a remodelling process being required for the assembly of the transcription machinery. Consistent with the spatial organisation of the Ad transcription program^{25,26}, transcription is initiated at both ends of the Ad genome covering the early genes (Fig. S1). We detected E1A as the first viral transcript²⁷, along with significant levels of E3 at 2 hpi (Fig. 1E). At 4 hpi transcription is markedly enhanced with high levels of transcripts originating from all early genes and significant signals arising from the E2A region. Taken together we observed an efficient utilization of the host cell transcription machinery after nuclear entry, revealing a transcriptional competent viral template. Transcription initiates with

a delay of 1 hour after nuclear entry opening the question of which structural changes may be required for assembly of the transcriptional machinery.

Chromatin architecture of the invading Ad genome

To address the Ad pVII nucleoprotein organisation, we partially hydrolysed viral and cellular genomic DNA with the endo-nuclease MNase (Fig. 2A). This enzyme does preferentially hydrolyse DNA in the linker region between nucleosomes and is restricted by stable protein-DNA complexes²⁸. Incubation of cellular chromatin with increasing concentration of MNase resulted in the appearance of a typical MNase ladder, reflected by the monomer and multimers of the nucleosomal DNA fragments (Fig. 2A). MNase concentrations were titrated such that one sample exhibited less than 20% mono-nucleosomal DNA (low digestion) and the other sample was fully hydrolysed to the mono-nucleosomal DNA (high digestion; Fig. S2A). The latter is the common condition used for MNase-seq experiments to date^{29–31}. High and low digestion conditions were used for each time point to isolate the mono-nucleosomal DNA from two replicates. The nucleosomal DNA of 17 individual experiments was used for library preparation and paired-end high-throughput sequencing, yielding in total 3.8 billion paired-end reads (Table S2).

DNA fragments were aligned to the human and the Ad genome, first analysing the read length distribution on both genomes at time point 0 hpi (Fig. 2B). When comparing the virus and host genome digested at the high MNase concentration, we observed a clear difference in the read-length distribution on the two genomes (Fig. 2B). Whereas the majority of DNA fragments exhibited the typical nucleosomal DNA size on the human genome (147 bp), the protected DNA fragments on the Ad genome were significantly shorter with a mean size of 60 to 70 bp. The MNase-seq was capable to reveal a stable adenoviral packaging unit that most probably represents pVII bound to DNA. Previous studies showed that nuclease digestion of Ad nucleoprotein resulted in discrete populations of protected DNA fragments, suggesting a chromatin-like configuration with sizes varying between 180 bp and 200 bp, between 30 and 90 bp, or between 30 and 60 bp depending on the conditions used^{8,10,32}. Assuming that larger DNA fragments represent partial nuclease hydrolysis products, the 60 to 70 bp long Ad-DNA fragment mirrors the previously described pVII-DNA protein complexes. In contrast it is not known if these pVII-DNA complexes are precisely positioned or if they are placed randomly. Detection of the pVII bound DNA allows us for the first time to analyse the occupancy and positioning of Ad pVII. We used the DANPOS2 package³³ to further analyse the viral nucleoprotein structure. We identified defined peaks over the Ad genome, showing non-

random, specific pVII-DNA positions on the viral DNA (Fig. 2D). The MNase profiles and peak calling was highly reproducible between biological replicates and at later time points, suggesting that pVII occupies specific Ad genome positions like the nucleosome on the host genome (Fig. S2C-E). The adenovirus exhibits a defined nucleoprotein architecture that differs along its functional regions. Interestingly, a central region of the viral genome (Late3) exhibits almost no peaks, revealing domains of structural heterogeneity that could reflect functional domains of the virus genome e.g. when packaged inside the virion, or being required for gene activation (Fig. 2D).

pVII exhibits DNA sequence dependent positioning patterns

The observation of a defined nucleoprotein architecture for the adenoviral genome was surprising because it can be *a priori* replaced by unrelated vector DNA of similar length³⁴. We next asked whether intrinsic DNA sequences patterns may establish pVII peaks. Such a sequence dependent positioning code was shown for the nucleosome³⁵. Sequence specific bending of the DNA reduces the energy requirements for wrapping the DNA around the histone octamer surface, revealed by characteristic di-nucleotide repeats in the nucleosomal DNA sequence at 10 bp intervals³⁶. We observed this described di-nucleotide pattern when analysing our nucleosomal DNA fragments annotated to the human genome (Fig. 2C – upper panel).

Interestingly, we also observed a specific di-nucleotide pattern, when computing the di-nucleotide frequency of the mid-point centered pVII fragments, suggesting that a specific DNA code underlies pVII positioning on the Adenovirus genome (Fig. 2C – lower panel). However, to our surprise, a periodic pattern of 5.4 bp was obtained for A/T- and G/C-dinucleotides, much shorter than the nucleosomal DNA repeat pattern, implying a distinct packaging mode. Such a repeat pattern indicates a spring-like winding of the DNA around pVII, similar to protamines (see discussion for details). Even without knowledge of the exact DNA-pVII topology, the sequence pattern and the specific binding sites of pVII on DNA clearly suggest the organisation of the Ad-genome into a specific nucleoprotein-structure that could be essential for its function. On average we detected 238 pVII peaks covering the Ad genome, which was consistent throughout all timepoints (Fig. 2E). Next, we quantified the mean pVII-DNA particle repeat length (pVII-RL) by measuring the center-center distance of adjacent pVII peaks in each sample (Fig. 2F and S2B). The median pVII-RL measures 97.3 bp suggesting that we isolated defined pVII-DNA particles spaced by accessible DNA linkers of 20 to 40 bp in length.

To address the presence of functionally distinct regions, we systematically subdivided the Ad genome into 10 domains of similar size, named after the including gene and classified by the

timing of expression (color code: early to late = light to dark blue) (Fig. 2D). Plotting pVII-RL showed a clear correlation of the repeat length with early expressing genes (Fig. 2G). Increased repeat lengths in the regions of the early genes can serve to increase genome accessibility and to enable the binding of regulatory factors. Strengthening this finding, the pVII occupancy and stability, reflected by the peak height, did anti-correlate with early timing in gene expression (Fig. 2H). In general, less pVII peaks (Fig. 2I) with larger repeat lengths and increased instability were found at early expressed domains indicating an open chromatin conformation (Fig S2D-E).

Our high-resolution analysis of Ad genome packaging provides for the first-time information on pVII-DNA particle positioning and occupancy. The analysis revealed a defined viral nucleoprotein architecture, with an open configuration at early expressed genes potentially enabling immediate access of the host cell transcription machinery.

Dynamic changes of Ad genome packaging

Previous studies showed that histones are deposited on the Ad genome³⁷. However, where and whether functional nucleosomes were assembled on the viral DNA is not known. Analysing the MNase-seq pattern of the time points 1, 2 and 4 hpi revealed the appearance of a 147 bp long, protected DNA fragment at 2 and 4 hpi (Fig. 3A and biological replicate in S3A). The nucleosome sized peaks increased in size, within the background of the pVII DNA fragments, revealing a co-occupation of the viral genome by pVII and nucleosomes. To quantify the fraction of nucleosomes on the viral genome, we simulated the peak sizes by computationally adding nucleosomal reads to the pVII fragment data of time point 1 hpi (Fig. S3B). The analysis suggested that at 2 hpi 8% and at 4 hpi 14% of the viral genomes are occupied with nucleosomes. First nucleosomal occupancy is detected at 2 hpi, coinciding with the initiation of viral transcription and indicating the requirement of nucleosome assembly for proper transcription initiation.

In order to reveal the sites of nucleosome assembly, DNA fragments with a size between 137-157 bp were extracted from time points 1, 2 and 4 hpi (Fig. 3A; indicated by dashed lines). The time point 1 hpi was defined as background signal and we calculated the fold-change of nucleosomal DNA arising at 2 hpi and 4 hpi. A cut-off of 5-fold enrichment was used to define the genomic sites of preferential nucleosome assembly (Fig. 3B, raw signal is shown in Fig. S3C). Strikingly, nucleosomal gain on the Ad-genome was not random. At 2 hpi we detected 12 hotspots of nucleosome assembly and additional 17 sites at 4 hpi. All sites showed a gain in enrichment over time, suggesting site specific nucleosome assembly after nuclear entry. The

sites of nucleosome assembly are concentrated within the three regions of the early genes and within the peculiar Late3 region, suggesting a role of nucleosome assembly in transcriptional activation.

Early genes gain a postioned +1 nucleosome

Nucleosome assembly selectively occurs in the genomic regions associated with the early genes, exhibiting a lower pVII protein occupancy and increased pVII repeat-length. An analysis of the incoming nucleosome positions with respect to pVII occupancy, revealed that nucleosomes directly formed over the central positions of the pVII-nucleoprotein particles (Fig. 3C) suggesting a specific mechanism to replace pVII.

A detailed analysis of the nucleosome assembly sites with respect to the early genes revealed that nucleosomes occupy discrete positions. Remarkably nucleosome positions coincide with the 5'-end of the synthesized E1A and E4-orf RNA, resembling the highly positioned +1 nucleosome found in active human genes (Fig. 3D and 3E) ²⁹. Additional nucleosomes are detected in the coding region of the early genes. The +1 nucleosome marks the transcription start sites of genes and flanks the regulatory sequences of the promoter that reside in the upstream linker region of this nucleosome. The Ad-DNA upstream of the +1 nucleosome exhibits either no pVII occupancy or long pVII repeat lengths, allowing binding of regulatory factors for gene activation. The preferential assembly of nucleosomes on the early genes, specifically located at the +1 position, prior to transcription initiation implies the requirement of nucleosome assembly for transcription initiation.

Dynamic changes of viral genome accessibility over time

Assessing the nucleoprotein structure with high MNase concentrations is providing information about stable protein particles occupying the DNA. In contrast, limited MNase concentrations, in analogy to the analysis of regulatory sites with DNase I ³⁸, would reveal genomic regions that de-condense chromatin and identifies regulatory elements ²¹.

As shown below, only a fraction of about 20% of the viruses in the cell are transcriptionally active, explaining why we observed only little variation between samples and the different timepoints treated with high MNase concentrations (Fig. 4A and S4A). In contrast, at low MNase concentration (Fig. 4B) clear changes over time are detected, signifying gross structural changes of the virus during early infection times. The active viruses undergo de-compaction that can be detected by limited MNase hydrolysis. When monitoring the pVII positioning at 0 hpi, the low MNase signal was well correlated with the high MNase signal ($R = 0.91$),

essentially showing the same pVII positions in both conditions (Fig. S4B and S4C). However, when inspecting the viral maps at later time points, we detected specific structural changes, with changes in peak intensities and the appearance of novel peaks in the low MNase signal (Fig. 4C). Structural changes over time, were quantified by a linear regression fit. The average signal intensity obtained in a 300 bp sliding window was quantified for each timepoint. The slope of the regression was used as a score for DNA accessibility changes (accessibility score; Fig. 4C and S5A). Genomic regions with a positive accessibility score are opening up, being more MNase accessible, whereas negative values correlate with DNA remaining condensed. A similar strategy was used in previous studies to measure local chromatin accessibility^{22,39,40}. Correlating with the activity of the virus, we detected Ad-chromatin de-condensation at the early gene regions reflecting gene activation (Fig. 4C and 4D). Interestingly, the score at E1A is not as pronounced as at other early genes (Fig. 4D). As this is the first gene to be transcribed, we suggest that this region already pre-exists in an open chromatin configuration, correlating with the low pVII peak density and therefore not changing over time.

The Late3 domain also exhibited a positive accessibility score, suggesting Ad chromatin de-condensation at this site (Fig. 4D). Additionally, in this domain we observe a discrete peak appearing at low MNase concentrations at 4 hpi (Fig. S5B). This peak is located in a rather MNase insensitive region and was only detected in the low MNase datasets. The occurrence of this local nuclease hypersensitive site suggests the binding of regulatory factors^{39,41,42}. Transcription factor motif analysis revealed potential binding sites for EN1, CREM, C/EBP-alpha and C/EBP-beta in the centre of the peak (Fig. S5B). Interestingly, the transcripts of EN1, CREM and C/EBP-beta in the host cell were upregulated up to twofold upon viral infection (Fig. S5C).

As shown above, we identified the early gene regions and Late3 regions as the sites of preferential nucleosome assembly and of viral chromatin de-condensation (Fig. 4C). To quantify the probability of this co-incidence, we randomly sampled the same number of nucleosome peaks for 10.000 times and calculated the cumulative accessibility score for each run (Fig. 4E). The cumulative scores are normally distributed around zero. The score of the detected nucleosome peaks (red dashed line in Fig. 4E) was very rarely reached by the randomly sampled peaks (p-value: 0.0002), indicating that targeted deposition mechanisms drive nucleosome assembly at the accessible regions of the Ad genome.

***In situ* detection of E1A mRNA identifies transcribing Ad genomes.**

Time resolved analysis using MNase-seq and RNA-seq showed strong correlation between localized viral chromatin remodelling and subsequent gene activation. We next used RNAscope technology for the direct detection of individual viral transcripts at the single cell level⁴³. A specific set of probes for the E1A mRNA was designed (ACDBio) and validated on cells infected with either HAd-C5 virus or an E1-deleted vector (Ad5-GFP). At 4 hpi specific E1A transcripts could be readily detected in virus- but not in vector-infected cells (Fig. 5A), nor in non-infected control cells (not shown). We next repeated the time course of infection (Fig. 1A) and detected at each time point viral capsids (not shown) and viral genomes using specific antibodies as well as E1A transcripts using RNAscope (Fig. 5B). Most genomes were imported until 1 hpi with no significant additional import after this time point. Normalization with cell associated capsids revealed that only ~40% of genomes were imported into the nucleus (Fig. 5C). While the nuclear import of genomes was finished 1 hpi, E1A transcripts only appeared at 2 hpi and further increased at 4 hpi (Fig. 5D). E1A transcript accumulation could be related to the continued expression from a stable pool of transcriptionally active genomes or by an increasing number of genomes initiating transcription over time. We quantified the colocalization of nuclear genomes with nuclear E1A transcripts to discriminate between both possibilities. Colocalizations resulting from association by probability would augment over time when active genome numbers remain constant and E1A transcript numbers increase over time. However, our results clearly showed that E1A positive nuclear genomes rapidly plateau at ~20% despite strong accumulation of nuclear E1A transcripts (Fig. 5E). It is shown that only a relatively small subpopulation of imported genomes contributes to early viral gene expression until 4 hpi.

***In vivo* detection of E1A mRNA identifies a subpopulation of transcribing Ad genomes.**

The RNAscope analysis confirmed the RNA-seq data and shows that the fraction of active viral genomes first reconstitute nucleosomes and only then activate early gene expression. To confirm this observation and further characterize transcribing and non-transcribing genomes we developed an imaging system for living cells. To mark E1A transcripts in living cells we inserted 24x MS2 binding sites into the 3'UTR of the E1A gene of a replication competent virus. MS2 binding sites are short stem-loop structures derived from the MS2 bacteriophage and can be detected by binding of multiple copies of fluorophore tagged MS2 binding protein (i.e. MS2BP-NLS-GFP) forming a fluorescent spot *in situ*⁴⁴. Production and amplification of the virus resulted in normal yields with unaltered infectivity or protein composition compared to the parental virus (data not shown). To detect Ad genomes in living cells we used our

previously established life cell imaging system based on fluorescent TAF-I β , which associates with incoming genomes also generating a fluorescent spot *in situ*¹⁸. A stable mCherry-TAF-I β expressing cell line was transduced with an expression vector for MS2BP-NLS-GFP to generate double fluorescent stable cell lines. Cells were left either uninfected, or infected with untagged control virus, or infected with E1A-MS2 tagged replicative virus and at ~4 hpi analysed by life cell imaging at high temporal resolution (1 frame/sec) (Movies S1-S4). To display the results in a single image we superimposed individual movie frames (60 sec t-projections, Fig. 6A). In non-infected control cells both, mCherry-TAF-I β and MS2BP-NLS-GFP localized to the nucleoplasm, with some unspecific enrichment of the MS2 signal in nucleoli (Fig. 6A top row, taken from Movie S1). When cells were infected with non-MS2 containing control virus, genomes formed mCherry-TAF-I β fluorescent spots in the nucleoplasm with limited confined motion as previously described¹⁸ while the MS2 signal remained unchanged compared to non-infected cells (Fig. 6A 2nd row taken from Movie S2). In contrast cells infected with virus expressing MS2-repeat-tagged E1A mRNA formed mCherry-TAF-I β spots indicating viral genomes but also rapidly moving GFP spots visible inside the nucleoplasm and in the cytoplasm (Movie S3/S4). GFP spots accumulated in most cells around 3-5 hpi in agreement with the RNAscope experiment. Upon merging the mCherry-TAF-I β channel (viral genomes) with the MS2BP-NLS-GFP channel (viral E1AmRNA) using t-projections, some genomes were clearly double positive suggesting transcribing genomes (Fig. 6A 3rd and 4th row, taken from Movie S3 and S4). Similar observations were made when we used a different MS2BP construct (MS2BP-NES-GFP) predominantly localizing to the cytosol or when we lowered the input MOI resulting in fewer (<10) genome copies per imaging field (Fig. S6). To quantify the number of E1AmRNA positive genomes we performed kymograph analysis using dual-channel t-projections of individual cells. This approach was possible because the genome position was stable over time allowing the identification of MS2 positive (transcribing) genomes (Fig. 6B). Kymograph analysis of several cells showed that ~20% of detected genomes inside a given nucleus are positive for MS2BP-NLS-GFP (Fig. 6C), which was very similar to the RNAscope approach (Fig. 6D). Taken together the analysis of transcriptionally active genomes in living cells supports our findings that early in infection transcription is restricted to a subpopulation of imported genomes that actively remodel the viral chromatin for transcription onset.

DISCUSSION

The adenovirus exhibits a defined nucleoprotein architecture

Early studies suggested that pVII binds adenoviral DNA at a 1:1 ratio like histones and protects DNA from endonuclease cleavage^{45,46}. However, no clearly defined pVII-DNA particle has been identified, as the protected DNA had a variable size range of 30 to 250 bp depending on experimental conditions and the degree of endonuclease digestion^{8,13,32}. In addition, it was not known whether pVII binds to specific sites on the viral genome, creating a defined viral nucleoprotein architecture, like the cellular nucleosomes. Here, our MNase-seq approach revealed the 147 bp long DNA fragments of the host nucleosomes and clearly identified a shorter, protected DNA species with a mean size of about 60 to 70 bp originating from the viral sequences. With pVII being the major DNA binding protein of the adenovirus, we suggest that we observe DNA fragments protected by pVII. The pVII digestion pattern is not as clearly defined as the nucleosomal DNA fragments, suggesting a heterogeneity in pVII-DNA complexes. The rather broad peak of protected DNA fragments (50 to 150 bp) is mirroring the variable results of the early studies, suggesting a rather heterogenous association of pVII with DNA. Alternatively, such a digestion pattern could also be the result of discrete pVII core complex (60 bp) that tightly associates with neighbouring pVII complexes, partially protecting the interconnecting linker DNA from endonuclease cleavage. Structures that are visible in early electron microscopic studies¹³.

pVII organizes the viral DNA into a specific nucleoprotein structure

Annotation of the pVII protected DNA fragments to the viral genome revealed for the first time a specific positioning of pVII on the genome. We show that the Ad genome is covered by about 238 stable pVII particles located at defined positions that are spaced by a DNA linker of about 20 to 40 bp. Precise pVII positioning suggest a highly organized and defined nucleoprotein architecture shared by all virus particles. DNA structure determines histone binding preferences, as shown by the DNA sequence dependent nucleosome positioning code³⁵. DNA bending allows wrapping of the DNA around a protein core, requiring site specific A/T and G/C di-nucleotide enrichments. A/T and G/C repeats differ in the wideness of the DNA major/minor groove thereby bending the DNA³⁵. If the sequence dependent DNA shape follows the protein surface path it increases binding affinity and thereby determines binding site preferences³⁶. Whereas the nucleosomal di-nucleotide repeats are spaced by 10.3 bp, pVII exhibits a much shorter repeat length of 5.4 bp. Nucleosomes and pVII-DNA complexes are organized differently and we suggest that the DNA associates in form of a spiral around the pVII core (Fig. 2C). This toroidal structure was proposed for protamines, to which pVII exhibits sequence similarities⁴⁶.

Our results show that pVII folds Ad genomes into an organized higher order structure, driven by its specific binding sites encoded by the Ad-DNA sequence. Taken together with the active remodelling of individual pVII molecules at early genes, this findings have to be taken into account for the optimal (sequence-)design of therapeutic DNAs packed into adenoviral vectors to achieve high and sustained transgene expression. In analogy to eukaryotic chromatin organisation, specific assembly factors and ATP dependent remodelling enzymes may be required to adjust the precise positioning of pVII on DNA, in addition to the DNA assembly code. Like nucleosomes, pVII does not bind as isolated protein to DNA but is forming homotypic complexes. The number of pVII proteins per Ad genome was estimated to be about 500 copies⁹ and early biochemical studies suggested the occurrence of pVII dimers^{47,48}. Taking into account the median number of 238 pVII positions we identified on the Ad genome (plus 19 on average for the 1871 bp deleted E3 region) our data would be compatible with two pVII molecules forming dimers at 238 discrete sites. This number would also come close to the ~200 “beads-on-string” estimated for purified viral cores⁴⁹.

Viral chromatin dynamics drives early gene expression

Here we show that nuclear delivery of viral genomes takes about 30 min but requires an additional hour to initiate early gene expression. In agreement with earlier observations full activation of early gene expression is reached after 4 h (reviewed in⁵⁰). Correlating gene expression with our differential MNase digestion conditions, we show that early genes exhibit reduced pVII density and increased pVII repeat lengths. Direct detection of genomes and of the E1A mRNA showed that genomes imported to the nucleus require about two hours to activate gene expression (i.e. E1A transcription). In this lag phase pVII is selectively replaced by nucleosomes at early genes, preceding gene activation. We identified that 80% of the imported Ad genomes are transcriptionally inactive, but the remaining 20% do actively rearrange the chromatin structure to allow transcription factor and RNA Polymerase binding. Once entering the nucleus, we showed that the active viral genomes do selectively de-condense the two early gene regions and curiously a specific domain in the Late3 gene. The overlap of open viral chromatin with early transcription, suggests an organisation that allows binding of regulatory factors for gene activation. For the Late3 region we observe increased DNA accessibility overlapping with binding motifs for transcriptional regulators that may serve as enhancing factors.

pVII is exchanged for nucleosomes at early genes

Genome decompaction correlates with low pVII occupancy and coincided with a site-specific exchange of pVII protein for histone octamers. pVII molecules were exchanged for positioned nucleosomes at the +1 site of actively transcribed genes (i.e. the E1A and E4 gene, Fig. 4D and 4E). The calculated number of nucleosomes and their positions on DNA, as determined by the DNA fragment length of 147 bp, corresponds to the number of ~20% transcribing genomes, exhibiting single positioned nucleosomes at the +1 position of early genes. As nucleosome exchange is almost complete at the time point 2 hpi when transcription activation starts, we suggest that nucleosome positioning is a prerequisite for gene activation. At this point we can only speculate why only a subpopulation of genomes is transcriptionally active. Reasons could be that the few genomes undergoing nucleosome assembly and active transcription produce the replication enzymes whereas the bulk of genomes enters replication without activation as an elegant way to avoid repeated chromatinization. Alternatively, cellular defence mechanisms could suppress transcription.

Interestingly, pVII removal to place nucleosomes at early genes preceded transcription, suggesting that pVII removal is driving transcription and not vice versa as previously reported¹⁹. Future work has to reveal which machinery drives the pVII nucleosome exchange and determines specificity. Generally, active promoters are free of nucleosomes (nucleosome depleted region, NDR) and the transcription start site is typically located upstream of a well-positioned '+1' nucleosome influencing recruitment of the transcriptional machinery and initiation of transcription^{51,52}. In case of the Ad genome our data show that the E1A promoter region is already pVII free and exists in an accessible nucleoprotein conformation when entering the cell, potentially allowing fast assembly of the +1 nucleosome and enabling immediate gene activation. This organization might thus be defined in the producer cell during genome packaging, which occurs in a sequential packaging sequence driven manner^{53,54}. In contrast, other early regions were actively de-compacted with nuclear entry of the virus, correlating with slightly delayed transcriptional onset. Whether this requires E1A expression is not known.

Taken together our analysis provides unprecedented insight into the adenoviral nucleoprotein structure and its dynamic reorganization during the onset of viral gene expression. We reveal genome sequence, pVII-DNA occupancy and nucleosome positioning as driving forces for transcriptional activation. Reproducing such organization in adenoviral vectors could result in efficient and sustained transgene expression.

EXPERIMENTAL PROCEDURES

RNA library preparation and sequencing

Total cellular RNA was isolated from infected cells at 0, 0.5, 1, 2, 4 hpi using TRI Reagent (Molecular Research Center, Inc.). Two time series sets of total RNA were prepared from two independent infections. Library preparation, comprising rRNA depletion, polyA enrichment, fragmentation to ~270 nucleotide length, reverse transcription to cDNA using random hexamer primers and adapter ligation, was accomplished using TruSeq RNA Sample Preparation Kit v2 (Illumina) according to the manufacturers protocol by EMBL GeneCore facility in Heidelberg. Libraries were sequenced on Illumina HiSeq2000 platform resulting in 18-75 Mio 50 bp single-end reads per sample.

Isolation of mono-nucleosomes, library preparation and sequencing

H1299 cells infected with Ad were permeabilized with the detergent IGEPAL-CA630 and digested at 0, 0.5, 1, 2 and 4 hpi with 100U/600U of MNase for 4/5 min at 37°, denoted as ‘low’/‘high’ MNase. Mnase treatment was performed as described in ²¹. After fragmented DNA has been separated on 1.3% agarose gels, mono-nucleosomal fractions were isolated. Paired-end sequencing on Illumina HiSeq2000 platform (EMBL GeneCore facility, Heidelberg), yielding about 220 Mio read pairs per sample, was accomplished following library preparation using NEBNext DNA library prep Master Mix Kit (New England Biolabs)

Viruses and virus production

Bacterial artificial chromosome (BAC) mutagenesis was used to generate recombinant adenoviruses following the protocol described in ⁵⁵ and ²³. Briefly, bp 1-3513 of the HAdV-C5 genome including the left inverted terminal repeat (ITR) and the whole E1 region was replaced by an FRT (Flippase Recognition Target) site and maintained as BAC (termed B12) to perform lambda red recombination essentially as described. Plasmid pO6-Ad5-E1-FRT (encoding the left ITR and the complete E1 region followed by an FRT recombination site) was used to clone 24 copies of the MS2 binding site repeats into the 3’UTR of the E1A gene. Subsequently flp-recombination was used ⁵⁵ to recombine plasmids pO6-Ad5-E1-FRT-MS2 and BAC B12 to reconstitute BxAd5-E1-MS2 a BAC encoding the whole AdV genome including the E1A gene tagged with 24 copies of the MS2 binding site. To generate the virus the BxAd5-E1-MS2 BAC was digested by *PacI* restriction enzyme to release the viral genome and transfected into HEK293 cells to reconstitute the recombinant viruses. Two rounds of plaque purification were performed followed by large scale production in HEK293 cells and purification through double CsCl₂ gradients banding. Replicative E3-deleted HAd-C5 virus or E1/E3-deleted HAd-C5 GFP

expressing control virus were essentially produced in the same way. Purified viruses were extensively dialyzed against PBS/10% glycerol and stored in aliquots at -80°C and quantified as physical particles per μ L following the method described in ⁵⁶.

Cell culture and Infection

U2OS (ATCC #HTB-96) and H1299 (ATCC #CRL-5803) and HEK293 (kindly provided by G. Nemerow, TSRI, La Jolla USA) cells were maintained at 37°C and 5% CO₂ in Dulbecco's modified Eagle's medium (DMEM)-Glutamax supplemented with 10% fetal calf serum (FCS) and 1% of Penicillin/Streptavidin. U2OS cells stably expressing mCherry-TAF-I β ⁵⁷ were transduced with lentiviruses expressing MS2BP-NLS-GFP and double positive cells were sorted by fluorescence activated cell sorting (FACS). The resulting cell population was maintained without further selection. For control experiments using cytosolic MS2BP (Fig. S6) stable mCherry-TAF-I β cells were transfected with an expression vector for MS2BP-NES-GFP. For RNAscope and immunofluorescence (IF) analysis H1299 cells were grown on 15mm glass coverslips. Subsequently cells were infected with 3000 physical particles/cell of replicative E3-deleted HAd-C5 virus or E1/E3-deleted HAd-C5 GFP expressing control virus, washed once with PBS and fixed at different time points with paraformaldehyde 4%/PBS for 10min at room temperature. Fixed cells were washed 3 times with PBS and slowly dehydrated in 50%/70%/100% ethanol for 5 min each and stored at -20°C in 100% ethanol. For the RNAscope protocol cells were rehydrated for 2 min in 70%/50% ethanol and equilibrated for 10 min in 1x PBS.

RNAscope

RNAscope experiment were performed using reagents and protocols from the RNAscope® Multiplex Fluorescent Assay essentially as indicated by the supplier (ACD Bio). Briefly, cells were incubated with Protease III (freshly diluted 1:30 with PBS) for 15 min at room temperature (RT) and washed 3 times with PBS. Hybridisation with the E1A RNAscope probe was performed for 2h at 40° in a humidified chamber. Hybridized cells were washed twice for 2 min and hybridized with signal amplifier probes (final amplifier Amp 4 AltB-FL) following the manufacturer's instructions. Hybridized cells were washed twice in PBS followed by incubation in IF buffer (PBS/10% FCS/0.1% Saponin) for 15 min at room temperature to saturate unspecific binding sites followed by IF analysis.

Immunofluorescence

Cells (infected or controls) grown on cover slips were fixed in 4% EM-grade PFA at indicated time points and stored in cold PBS until use. For IF, cover slips were initially incubated in IF

buffer (PBS/10% FCS/0.1% Saponin) for 15min and then replaced by antibody solution. Primary antibodies (mouse anti-pVI, mouse anti-pVII and rabbit anti Ad5- capsids) were diluted in IF buffer and incubated with cells for 1h at RT. Antibody was removed by washing in PBS followed by incubation with secondary antibodies (mouse Alexa 647 and rabbit Alexa 594) diluted in IF buffer for 1h at RT. Cells were washed with PBS, ddH₂O and absolute ethanol, air-dried and mounted in DAKO mixed with DAPI (1:1000).

Live-cell imaging and image analysis

For live-cell imaging, U2OS cells stably expressing mCherry-TAF-I β and MS2BP-NLS-GFP were seeded in ibidi μ -slide VI^{0.4} (Ibidi). The next day, the medium in the ibidi μ -slide was replaced by imaging medium (FluoBrite DMEM supplemented with Prolong antifade reagent, both Life technologies) containing 3000 physical particles/cell of replicative BxAd5-E1-MS2 for 30min followed by washing with imaging medium to remove the inoculum. Infection and subsequent cultivation were performed at 37°C. Cells were imaged using a Leica spinning-disk microscopy system (x100 objective) equipped with an environmental chamber at a frame rate of 1 frame per second using MetaMorph software. To generate kymographs, movies were processed using the ImageJ KymographBuilder plugin (https://imagej.net/KymographBuilder:_Yet_Another_Kymograph_Fiji_plugin) on confocal sections. Line width was adapted to cover all viral genomes throughout the length of the movie and double positive genomes were scored as transcribing genomes. Mounted cells for IF and RNAscope analysis were imaged using a SP8 confocal microscope (Leica) at the Bordeaux Imaging Center (BIC) platform. Stacks (7x) were taken every 0,3 μ m using a pinhole of 1 at a pixel size of 85 nm. Quantifications were done using an adapted semi-automated macro in ImageJ (provided upon request). Briefly, channels were split and stacks were Z-projected and the cell periphery was outlined manually. A threshold was applied to every channel and single objects or colocalized objects exceeding a defined area of pixels were counted. Statistical analysis was done using one-way ANOVA multicomparison test.

QUANTIFICATION AND STATISTICAL ANALYSIS

Virus annotation

The reference genome sequence and gene annotation of Human Adenovirus C serotype 5 (accession: AY339865.1) was downloaded from NCBI. The annotation was adapted to the used Adenovirus strain and nucleotides 28,593-30,464 were excised from the full version (available at github project page).

RNA-seq analysis

STAR⁵⁸ was used to align the RNA-seq reads simultaneously to the human and the adenoviral genome. Therefore, a STAR index was generated based on the hg19 reference, the modified adenoviral genome (see section *Virus annotation* for details) and the GENCODE human transcript annotation version 25. Read mapping was conducted using following parameters:

```
--outSAMmultNmax 1
--outMultimapperOrder Random
--outFilterType BySJout
--outFilterIntronMotifs RemoveNoncanonical
--outSAMstrandField intronMotif
--outFilterMultimapNmax 20
--alignSJoverhangMin 8
--alignSJDBoverhangMin 1
--alignIntronMin 20
--alignIntronMax 1000000
```

To quantify the RNA abundance for each gene the number of uniquely mapped reads per gene was counted using the featureCounts function of the subread package⁵⁹. The read counts were normalized to transcripts per million (TPMs).

MNase-seq analysis

A bowtie2 index comprised of the human reference genome hg19 and the sequence of the used Adenovirus strain (see section *Virus annotation* for details) was built (code available at⁶⁰) and paired-end reads were mapped to the index using bowtie2-aligner⁶¹ with following parameters:

```
--very-sensitive-local
--no-discordant
```

Reads were filtered to mapping quality ≥ 30 and only reads mapped in proper pair were kept using samtools⁶². Reads aligned to the Adenovirus genome were extracted for further analysis. The fragment size distribution was plotted using the CollectInsertSizeMetrics function from the Picard toolkit (GitHub Repository. <http://broadinstitute.github.io/picard/>; Broad Institute).

pVII positioning analysis

Fragments mapped to the Adenovirus reference with a maximum insert size of 139 bp were selected and used for pVII positioning analysis. pVII positions were called using the DANPOS2 toolkit³³. First, a pVII occupancy profile for each sample was generated and normalized to 20,000 reads using the dpos function with following parameters: -u 1e-5; -a 1; -z 5; -p 0.001, --extend 35 and --mifrsz 0. Next, the profiles were normalized by a genome wide quantile normalization to the first replicate at 0 hpi of each MNase condition (high/low) using the wiq script. Finally, pVII positions were obtained from the normalized pVII occupancy profiles using the dpos function with following parameters: -jd 50; -q 10; -z 5; -a 1.

Rotational positioning analysis

49 to 51 bp sized fragments mapped to the Adenovirus reference of all high MNase samples were selected and pooled together. Frequencies of A/T or G/C di-nucleotides around the fragment center were calculated using the annotatePeaks.pl script from the HOMER suit⁶³. Noise was filtered using the Fast Fourier Transform build in the filterFFT function of the nucleR Bioconductor package⁶⁴. The cutoff was set with the option pcKeepComp=0.2. Finally, to reflect the symmetry of the pVII binding the reverse di-nucleotide frequency was added, and the total frequency was divided by 2. The same procedure was conducted to derive the di-nucleotide periodicity of nucleosomal fragments, except that 147 bp fragments of the first replica at 0 hpi were selected.

Sites of nucleosome assembly

The R Bioconductor package nucleR⁶⁴ was used to call sites of nucleosome assembly onto the Ad genome. First, fragments specified to the Ad genome were loaded into the R environment. Fragments between 137 to 157 bp from time points 1, 2 and 4 hpi of the high MNase condition were extracted for further analysis. The coverage was normalized to reads per million Ad mapped reads and noise was removed using the built-in Fast Fourier Transform (filterFFT with option pcKeepComp=0.007). Potential nucleosome positions were called based on the normalized profile using the peakDetection function with the option width=147. Peaks showing at least a 5-fold enrichment compared to the background signal at 1 hpi were considered as nucleosome positions.

Ad genome accessibility

DNA fragments released under low MNase conditions were used to identify structural changes of Ad chromatin over time. Therefore, a chromatin accessibility score was calculated

along the Ad genome using a 300 bp bin sliding along the Ad genome by 20 bp. The accessibility score was obtained for each bin in the genome by fitting a linear regression on the normalized read frequencies computed for each time point (average of both replicates). The slope of the linear fit was used as a measure of dynamic changes of viral genome accessibility.

Comparison of nucleosome assembly sites and Ad genome accessibility

The cumulative accessibility score at the sites of nucleosome assembly was calculated. To compare how likely this score is obtained by chance, the nucleosome peaks were randomly shuffled along the Ad genome and the cumulative accessibility score was computed. This process was iterated 10.000 times.

Transcription factor motif identification

The Ad DNA sequence from position 19,890 to 19,940 was extracted and uploaded to the PROMO web server ⁶⁵. Motif search was performed against known human and mouse motifs stored in TRANSFAC data base using the SearchSites tool. The maximum dissimilarity rate was set to 2 %. The obtained motifs were further filtered to have at least a recognition sequence of 5 nucleotides and that the corresponding transcription is expressed in the infected H1299 cells.

ACCESSION NUMBERS

The Gene Expression Omnibus accession number for the MNase-seq, and RNA-seq data reported in this paper is GSE136550. The code used to analyze the sequencing data is available at GitHub: <https://github.com/uschwartz/AdVir>

SUPPLEMENTAL INFORMATION

Supplemental Information includes 6 figures, 4 movies, 2 tables, and can be found with this article online at <http://xxxxx>

AUTHOR CONTRIBUTIONS

Conceptualization H.W. and G.L.; Methodology F.L., T.K., E.S., C.H., E.B., H.W. and G.L.; Formal Analysis F.L., U.S. and H.W.; Investigation T.K., C.H., F.L., E.S., H.W. and G.L., Data Curation U.S.; Visualization F.L., U.S. and H.W.; Writing – Original Draft U.S. G.L. and H.W.; Writing – Review & Editing E.B., T.K., U.S., G.L. and H.W., Resources E.B., H.W. and G.L.; Supervision H.W. and G.L.; Funding Acquisition G.L. and H.W.

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FIGURES AND LEGENDS

Figure 1. Spatio-temporal analysis of adenovirus infection. **A)** Experimental design. Virus was added (+Ad) and removed (Wash) after 30 min and cell were processed as indicated (hpi = hours post infection) using immunofluorescence (IF), RNA sequencing (RNA-seq) and nucleosome sequencing (MNase-seq). **B)** Images show representative confocal images at indicated time points stained with anti-pVI (left column, cyan signal) and anti-pVII (middle column, red signal) antibodies and merged with DAPI detection of the nucleus (right column, grey signal). Scale bar is 10 μ m. **C)** Quantification of the pVI signal over time as indicated on the x-axis. The distribution of pVI foci per cell are shown as Box and Whiskers plot ($n > 44$ cells per time point). **D)** As in (C) showing the quantification of pVII foci per nucleus from the same experiment ($n > 44$ nuclei per time point). **E)** Time resolved analysis of Ad transcription using RNA-seq. TPMs (Transcripts Per Kilobase Million) for early expressed genes were calculated at each timepoint. The barplots show the average of two replicates and each replicate is indicated by a black dot.

Figure 2. Characterization of Ad DNA packaging by pVII. **A)** Schematic view of MNase digestion of infected cells. Agarose gel shows nucleosome ladders of a MNase titration. Mono-, di- and tri-nucleosomes are indicated at the right. The mono-nucleosomal DNA fraction was isolated after MNase digestions and subjected to high-throughput sequencing. **B)** Fragment size distribution of paired-end reads either mapped to the human genome (red) or the Ad genome (purple). **C)** Average A/T- and G/C-di-nucleotide frequency of nucleosomal (upper panel) and pVII fragments (lower panel). Size-selected fragments were centred at the midpoint and the di-nucleotide frequency was calculated. Every 10.3 (nucleosome) / 5.4 (pVII) bp a dashed line highlights the periodicity of the signal. **D)** Profile illustrates Ad genome coverage by MNase-seq reads. pVII positions, which have been called with the DANPOS2 package³³, are represented by grey boxes. Partitioning of the genome into functional domains and the timing of their transcription is indicated by the color code (lightblue = early, darkblue=late). Genomic location and underlying gene annotation of HAd-C5 (NCBI accession: AY339865.1) are indicated at the bottom. **E)** Boxplot showing the number of detected pVII peaks for each sample. **F)** Boxplot showing the genome-wide pVII repeat length (pVII-RL) for each sample. **G)** and **H)** Boxplots showing the distribution of pVII center-center distances / peak heights in each domain. Individual replicates are shown side by side. **I)** Barplot showing the number of pVII peaks per kb detected in the domains. Individual replicates are shown as black dots.

Figure 3. Nucleosome assembly onto the Ad genome. **A**) Fragment size distribution of paired-end reads either mapping to the human genome (red) or the Ad genome (shades of purple). Nucleosome sized fragments are highlighted between two dashed lines at 137 bp and 157 bp. **B**) Enrichment of nucleosome sized fragments across the Ad genome. Enrichment was calculated as fold-change of 137-157 bp sized fragments at 2 hpi / 4 hpi to a background level at 1 hpi. Dashed lines indicate the threshold used for histone peak calling. Nucleosome positions are represented by grey boxes. H3 ChIP-qPCR verified regions³⁷ are shown as green boxes. **C**) Average pVII occupancy at 0 hpi around predicted nucleosome positions. Dashed lines indicate nucleosome boundaries. **D**) and **E**) Genome browser tracks showing pVII occupancy at 0 hpi (upper panel), the abundance of nucleosome sized fragments (137-157 bp, middle track) from 1 to 4 hpi, predicted sites of nucleosome assembly (grey boxes) and expression levels of early genes (bottom panel).

Figure 4. Early decondensation of Ad chromatin. **A)** and **B)** Sample-wise Pearson correlations of **A)** high MNase or **B)** low MNase profiles. **C)** Profile illustrates Ad genome coverage by low MNase-seq fragments. The average of two replicates is shown, except at timepoint 0 hpi where only one replicate was available. The accessibility score was calculated as the slope of a linear regression fit of the average signal intensity across each timepoint. The score was assessed in 300-bp sliding windows. Nucleosome positions at 4 hpi derived from Fig. 4B are shown. **D)** Barplot showing the average accessibility score in each domain. **E)** Density distribution of the cumulative accessibility score of sampled peaks. The cumulative accessibility score of the estimated nucleosome positions at the Ad genome is highlighted by a dashed red line.

Figure 5. E1A transcription at the single cell level. **A)** Representative confocal images of cells infected with HAd-C5 (Ad5, top row) or an E1-deleted HAd-C5 vector (Ad5-GFP, bottom row). Detection of E1A transcript with specific RNAscope probes (left column, green signal), genomes using anti-pVII antibodies (middle column, magenta signal) and merged with DAPI detecting the nucleus (right column, grey signal). Scale bar is 10 μ m. **B)** Representative confocal images as in (A) of cells infected with HAd-C5 at different time points as indicated. Scale bar is 10 μ m. **C)** Quantification of the ratio of nuclear pVII foci normalized by cell associated capsid signals over time as indicated on the x-axis. The distribution of normalized pVII foci at each time point is shown as scatter plot (Mean $\pm$ SD, n = 30 cells per time point). **D)** Graph as in (C) showing the distribution of E1A transcript foci per cell as scatter plot (Mean $\pm$ SD, n = 30 cells per time point). **E)** Graph as in (C) showing the distribution of pVII foci per nuclei (top graph) and E1A transcript foci per nuclei (middle graph) as scatter plot (Mean $\pm$ SD, n = 30 nuclei per time point). The bottom graph was generated by normalizing the number of E1A positive genomes by the total number of genomes per nucleus shown as scatter plot (Mean $\pm$ SD, n = 30 nuclei per time point). Statistical analysis for C-E via one-way ANOVA multi-comparison test.

Figure 6. E1A transcription in living cells. **A)** Representative images showing t-projections of 60 frames (frame rate 1F/sec) imaged at ~4 hpi. Cells express mCherry-TAF-I β showing viral genomes (first column, red signal in merge) and MS2BP-NLS-GFP showing MS2-repeat tagged E1AmRNAs (second column, green signal in merge) and an overlay (third column, merge) of both channels. The first row shows non-infected control cells, the second row cells infected with untagged control virus and the third and fourth row examples of cells infected with virus expressing E1AmRNA tagged with MS2-repeats. Black arrows point at non-

transcribing and white arrows at transcribing genomes. Details are shown as magnifications of the boxed area. Scale bar is 10µm. **B)** Kymograph analysis of the bottom two cells shown in A and as described in the material and methods section. Cell overview to the left, kymograph separated into the individual channels and a superimposition of both channels to the right. Genome position over time appears as red vertical line, E1AmRNA signal as green vertical line. The boxed area in the overview corresponds to the boxed area in the kymograph. Green arrows point at the E1AmRNA signal, orange arrows at transcribing (double positive) and red arrow at non-transcribing (single positive) genomes. **C)** Quantitative kymograph analysis. Scatter plot representation of the percentage of MS2-BP positive genomes per cell (n = 11 cells) vs. control cells infected with viruses expressing untagged E1AmRNA (n = 5 cells) determined by kymograph analysis. **D)** Scatter plot representation comparing the percentage of E1AmRNA positive genomes per cell determined with the MS2-BP method (left, n = 11 cells) vs. the RNAscope method (right, n = 22 cells). Unpaired t-test was used for statistical analysis in C and D.