Human papilloma virus E7 oncoprotein abrogates the p53-p21-DREAM pathway

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High risk human papilloma viruses (HPV) are oncogenic DNA viruses that can cause cancer of the cervix uteri, oropharynx, penis, vagina, vulva and anus1-4. The primary transforming capacities of HPV stem from the E6 and E7 proteins. These two oncoproteins cooperate in silencing the anti-proliferative control of the cell. The best-known target of E7 is the cell cycle regulator pRB5. Direct binding of E7 to the retinoblastoma tumor suppressor protein pRB impairs its function6-8. Furthermore, association of E7 with pRB leads to an increase in p53 level9. The tumor suppressor p53 can either trigger checkpoints causing cell cycle arrest or lead to the induction of apoptosis. In the context of HPV infection, the E6 oncoprotein initiates degradation of p5310, 11.

Moreover, the E7 oncoprotein has additional functions aside from targeting pRB12, 13. These include means to impair p53 function even in the absence of E6. Thus, HPV E7 is sufficient to block cell cycle checkpoint control by p5314-16. The cyclin-dependent kinase (CDK) inhibitor p21 (CDKN1A) is a central mediator of p53 checkpoint control19, 20, and its function can be impaired by E718, 21, 22. This finding is particularly interesting given that p21 is required for downregulation of genes in response to p5323, 24.

Recently, another mechanism to impair p53 function that is independent of E6 was discovered. HPV E7 was found to disrupt the pRB-related transcriptional repressor complex DREAM (DP, RB-like, E2F4 and MuvB)25-27. The DREAM protein complex consists of E2F4, DP1 and p130/p107 in addition to RBBP4 and the LIN proteins LIN9, LIN37, LINS2 and LINS4 that form the MuvB core38-39. DREAM binds promoters through cell cycle-dependent elements (CDEs), cell cycle genes homology regions (CHRs), CHR-like elements (CLEs) and E2F sites3-13. In response to p53, DREAM is recruited to promoters of cell cycle genes, leading to their repression36, 37. While p53 itself is solely an activator of transcription, the p53-p21-DREAM pathway mediates indirect gene downregulation by p5338, 39. For example, Polo-like kinase 4 is an important target of this pathway. The mitotic kinase PLK4 is repressed through the p53-p21-DREAM-CDE/CHR pathway40, and its p53-dependent repression can be abrogated by HPV E741. Importantly, CDE/CHR elements are required for p53-dependent repression of PLK4, and the expression of HPV E7 impairs DREAM binding to the CDE/CHR elements in the PLK4 promoter41. As genome-wide expression profiling datasets of E7-expressing IMR90 lung fibroblasts42 and
NIKS keratinocytes43 became recently available, we asked whether targets of the p53-p21-DREAM pathway are generally deregulated by HPV E7 on a genome-wide level.

Here, we integrate these new data with earlier genome-wide datasets that were derived from comparing HPV-16/18-infected cervical tumor samples with normal tissue44,45, from CaSkii cells expressing HPV E2C, a potent transcriptional repressor of E6 and E746, or from HeLa cells in which E6 and E7 were downregulated by RNAi47. Our analysis identifies genes that were observed as E7-regulated in most datasets, and we compared the results with lists of DREAM and pRB-E2F target genes43. We found that many DREAM targets are upregulated by E7 and that DREAM targets are the main class of genes deregulated by E7. Most importantly, p53-dependent down-regulation of DREAM target genes is abrogated in HPV E7-expressing cells. In summary, our analysis provides a genome-wide high-confidence list of genes deregulated by HPV E7, most of which are DREAM targets. This study reveals the importance of E7-mediated DREAM disruption that interferes with p53-dependent gene down-regulation. Thus, in HPV-infected cells, p53 function can be impaired by E7 independently of E6.

Materials and Methods
Computational analysis. A step-wise meta-analysis approach was employed to integrate multiple datasets43. This approach enables the integration of pre-analyzed datasets and does not require re-analysis of the raw data. Publicly available HPV E7 gene expression profiling datasets were curated42–47 and mapped to a collection of protein-coding genes47. Expression values of the analyzed genes were compiled and classified into downregulated (−1), upregulated (+1) and non-regulated genes (0).

Genes identified as significantly differentially regulated in HPV-16 E7 expressing NIKS cells were retrieved from Table 2 in Zhou et al.43. The pre-analyzed dataset of Rozenblatt-Rosen et al. from HPV-18 E7-expressing IMR90 cells was retrieved from the deposited Supplementary Table 19 in Rozenblatt-Rosen et al.45 and a gene was considered significantly differentially regulated if it passed the thresholds of adj. p-value ≤ 0.05 and absolute log2 (fold-change expression) ≥0.5. Genes identified as significantly differentially regulated in HPV-16/18 infected early stage cervical cancers compared to normal cervical epithelium were retrieved from the deposited Tables 2 and 3 in Santin et al.45. The pre-analyzed dataset from HeLa cells in which endogenous HPV-18 E6 and E7 expression was silenced by RNAi displays significantly differentially expressed genes and was retrieved from the Supplementary Table S1 in Pang et al.46. Of note, datasets by Rosty et al. and Pang et al. exclusively reported upregulated genes.

It is generally agreed that gene expression data from different experimental platforms are not directly comparable, and thus we used the stepwise meta-analysis approach instead that ranks genes by the number of datasets that find them significantly differentially regulated. Given that raw data were not re-analyzed, the approach does not include data points that were below the thresholds set in the individual studies. Differences in unprocessed data acquisition between several studies may reduce reproducibility, yet it minimizes the bias that would be introduced by using one particular analysis approach for all datasets. Following the stepwise meta-analysis approach43,44, genes were ranked by the number of datasets finding the gene to be significantly upregulated minus the number of datasets that find the gene to be downregulated (Supplementary Table S1).

Cell culture and drug treatment. HCT116 cells were grown in Dulbecco’s modified Eagle’s medium (DMEM; Lonza, Basel, Switzerland) supplemented with 10% fetal calf serum (FCS) (Biochrom, Berlin, Germany) and penicillin/streptomycin and maintained at 37 °C and 10% CO₂. Stably transfected HCT116 cells were generated by transfection with pCMV-HPV16-E7 wt (kindly provided by Karl Münger48), and selection with G418/Geneticin (PAA Laboratories, Pasching, Austria) at a concentration of 0.5 mg/ml41. Wild-type mouse keratinocytes were isolated from C57Bl6 mouse embryos as described previously49. Cells were grown on plates coated with collagen (Invitrogen, Darmstadt, Germany) and maintained at 10% CO₂ and 32 °C in DMEM/Ham’s F12 (3.5:1.1) (PAN Biotech, Aidenbach, Germany). Cells were treated with doxorubicin (0.2 μg/ml; Medac, Wedel, Germany) or Nutlin-3a (10 μM; Cayman Chemicals, Ann Arbor, MI, USA) for 24 h. For cell sorting of transiently transfected wild-type mouse keratinocytes, pEGFP plasmid (Clontech, Mountain View, CA, USA) was co-transfected with pCMV-HPV16-E7 wt plasmid at a 1:3 ratio using GeneJuice (Merck, Darmstadt, Germany). Fluorescence-activated cell sorting was carried out on a FACS Aria SORP instrument (Becton Dickinson Biosciences, Franklin Lakes, NJ, USA).

RNA extraction, reverse transcription and semi-quantitative real-time PCR. Total cellular RNA was isolated using TRIzol reagent (Invitrogen, Carlsbad, CA, USA) following the manufacturer’s protocol. One-step reverse transcription and quantitative real-time PCR were performed with an ABI 7300 Real-Time PCR System (Applied Biosystems, Forster City, CA, USA) using QuantiTect SYBRGreen PCR Kit (Qiagen, Hilden, Germany) as described previously41. Primer sequences have been published previously34,40,41,50,51.

Sodium dodecyl sulphate-polyacrylamide gel electrophoresis and immunoblot. Sodium dodecyl sulphate-polyacrylamide gel electrophoresis and western blot were performed following standard protocols40. The following antibodies were used: E2F1 (sc-193, Santa Cruz Biotechnology, Santa Cruz, CA, 1:500 dilution), KIF23 (sc-136473, Santa Cruz Biotechnology, 1:200), CDC25C (sc-327, Santa Cruz Biotechnology, 1:1000), B-MYB (LX015.1, kindly provided by Roger Watson35, hybridoma media 1:5) and β-actin (A5441, Sigma-Aldrich, Munich, Germany, 1:5000).
Results

The HPV E7 oncoprotein deregulates cell cycle genes targeted by the DREAM complex. Two recently published datasets identified genes deregulated upon expression of the HPV E7 oncoprotein on a genome-wide basis. Combined, those two datasets identified 753 genes deregulated by E7, including 453 upregulated and 300 downregulated genes. A fraction of these genes was identified in both datasets, 66 upregulated and 2 downregulated genes. The small number of genes downregulated upon E7 expression indicates that expression of E7 primarily causes gene upregulation. When comparing the overlap of the 66 upregulated genes with recently published lists of cell cycle genes and targets of the DREAM, MMB-FOXM1 and pRB-E2F complexes, it becomes evident that most E7-upregulated genes are cell cycle genes and DREAM targets.

Next, we integrated additional datasets that identified genes deregulated by HPV E6 and E7. Employing tools and data from a recent meta-analysis, this approach yields reliable target identification. In total, these six datasets identified 1,783 genes as deregulated by HPV E7. No gene was identified in all datasets as downregulated, further supporting the notion that E7 expression primarily results in target gene induction. Fourteen genes were identified as upregulated by E7 in at least five of the six datasets. Remarkably, all of these genes are cell cycle genes and DREAM targets. Furthermore, when looking at the 49 genes identified in at least four of the six datasets, 34 are cell cycle genes and DREAM targets. Only one gene from this group, CDKN2A (p16), is not a DREAM target.

Table 1. HPV E7 deregulates DREAM target genes. 68 genes overlap in two datasets of genes deregulated upon HPV E7 expression. Two genes described in both datasets as downregulated are underlined. Annotation of genes as DREAM targets or cell cycle genes, including the cell cycle phase of peak expression, were extracted from Fischer et al. UNKN, timing of peak expression of the cell cycle gene is unknown; X, cell cycle-dependent expression was not reported in the datasets.
| Gene Symbol | Identified as upregulated by E7 in No. of datasets | Cell cycle gene | DREAM target | MMB-FOXM1 target | pRB-E2F target |
|-------------|-----------------------------------------------|----------------|--------------|--------------------|----------------|
| MCM2        | 6                                             | G1/S           | ✓            | ×                  | ✓              |
| ZWINT       | 6                                             | G1/S           | ✓            | ×                  | ×              |
| APOBRC3B    | 5                                             | G2/M           | ✓            | ×                  | ×              |
| CDC6        | 5                                             | G1/S           | ✓            | ×                  | ✓              |
| KIF2C       | 5                                             | G2/M           | ✓            | ✓                  | ×              |
| LMNB1       | 5                                             | G2/M           | ✓            | ✓                  | ✓              |
| MCM4        | 5                                             | G1/S           | ✓            | ×                  | ✓              |
| MYBL2       | 5                                             | G1/S           | ✓            | ×                  | ✓              |
| NUSAP1      | 5                                             | G2/M           | ✓            | ✓                  | ✓              |
| PRC1        | 5                                             | G2/M           | ✓            | ✓                  | ×              |
| RRM2        | 5                                             | G1/S           | ✓            | ✓                  | ✓              |
| SMC4        | 5                                             | G2/M           | ✓            | ✓                  | ✓              |
| STIL        | 5                                             | G2/M           | ✓            | ✓                  | ✓              |
| TOP2A       | 5                                             | G2/M           | ✓            | ✓                  | ×              |
| APOBEC3B    | 5                                             | G1/S           | ✓            | ✓                  | ✓              |
| ASFM        | 5                                             | G2/M           | ✓            | ✓                  | ×              |
| ASPM        | 5                                             | G2/M           | ✓            | ✓                  | ✓              |
| ATAD2       | 4                                             | G1/S           | ✓            | ✓                  | ✓              |
| BIRC5       | 4                                             | G2/M           | ✓            | ✓                  | ×              |
| BRCA1       | 4                                             | G1/S           | ✓            | ×                  | ✓              |
| CCNA2       | 4                                             | G2/M           | ✓            | ✓                  | ×              |
| CCNB1       | 4                                             | G2/M           | ✓            | ✓                  | ✓              |
| CCNB2       | 4                                             | G2/M           | ✓            | ✓                  | ✓              |
| CDC20       | 4                                             | G2/M           | ✓            | ✓                  | ×              |
| CDC25C      | 4                                             | G2/M           | ✓            | ✓                  | ×              |
| CDC45       | 4                                             | G1/S           | ✓            | ×                  | ✓              |
| CDRN2A      | 4                                             | ×              | ×            | ×                  | ×              |
| CENPF       | 4                                             | ×              | ×            | ×                  | ×              |
| DTL         | 4                                             | ×              | ×            | ×                  | ×              |
| E2F1        | 4                                             | G1/S           | ✓            | ×                  | ✓              |
| FANCI       | 4                                             | G1/S           | ✓            | ×                  | ✓              |
| FKBP5       | 4                                             | UNKN           | ✓            | ×                  | ×              |
| FOXM1       | 4                                             | G2/M           | ✓            | ×                  | ×              |
| Gins2       | 4                                             | G1/S           | ✓            | ×                  | ✓              |
| KIF20A      | 4                                             | G2/M           | ✓            | ✓                  | ×              |
| KIF23       | 4                                             | G2/M           | ✓            | ✓                  | ×              |
| MELK        | 4                                             | G2/M           | ✓            | ×                  | ×              |
| NCAPG2      | 4                                             | G1/S           | ✓            | ×                  | ✓              |
| NEK2        | 4                                             | G2/M           | ✓            | ✓                  | ×              |
| POLQ        | 4                                             | G2/M           | ✓            | ×                  | ×              |
| PRIM1       | 4                                             | G1/S           | ✓            | ×                  | ✓              |
| PTG1        | 4                                             | G2/M           | ✓            | ×                  | ✓              |
| RAD51AP1    | 4                                             | G1/S           | ✓            | ×                  | ✓              |
| RFC3        | 4                                             | G1/S           | ✓            | ×                  | ✓              |
| SPAG5       | 4                                             | G2/M           | ✓            | ✓                  | ×              |
| TMPO        | 4                                             | G2/M           | ✓            | ×                  | ✓              |
| TRIP13      | 4                                             | G2/M           | ✓            | ×                  | ×              |
| TTK         | 4                                             | G2/M           | ✓            | ✓                  | ×              |
| WDHD1       | 4                                             | G1/S           | ✓            | ×                  | ✓              |
| WDR87       | 4                                             | G1/S           | ✓            | ×                  | ✓              |

Table 2. HPV E7-deregulated genes with an identification-overlap of at least four out of six datasets. 49 genes were identified in at least 4 of the 6 datasets as being deregulated by HPV E7 (compiled from Table S1). Annotation of cell cycle genes, including the phase of peak expression and DREAM, MMB-FOXM1 or pRB-E2F targets were extracted from Fischer et al. UNKN, timing of peak expression of the cell cycle gene is unknown; X, cell cycle-dependent expression was not reported in the datasets.
| Gene Symbol | Identified as upregulated by E7 in No. of datasets | Cell cycle expression | DREAM target | MMBOFOX1 target | pRB-E2F target | Gene Symbol | Identified as upregulated by E7 in No. of datasets | Cell cycle expression | DREAM target | MMBOFOX1 target | pRB-E2F target |
|-------------|-----------------------------------------------|----------------------|--------------|-----------------|---------------|-------------|-----------------------------------------------|----------------------|--------------|-----------------|---------------|
| ANLN        | 3                                             | G2/M                 | ✓            | ✓              | ✓             | KNTC1       | 3                                             | G1/S                 | ✓            | ×               | ✓             |
| ANP32E      | 3                                             | G2/M                 | ✓            | ✓              | ✓             | MAD2L1      | 3                                             | G2/M                 | ✓            | ✓              | ✓             |
| ATAD5       | 3                                             | UNKN                 | ✓            | ✓              | ✓             | MASTL       | 3                                             | G1/S                 | ✓            | ×               | ×             |
| AURKA       | 3                                             | G2/M                 | ✓            | ✓              | ✓             | MCM10       | 3                                             | G1/S                 | ✓            | ×               | ✓             |
| BRCA2       | 3                                             | G1/S                 | ✓            | ✓              | ✓             | MCM3        | 3                                             | G1/S                 | ✓            | ✓              | ✓             |
| BRIP1       | 3                                             | G1/S                 | ✓            | ✓              | ✓             | MCM5        | 3                                             | G1/S                 | ✓            | ×               | ×             |
| BUB1        | 3                                             | G2/M                 | ✓            | ✓              | ✓             | MCM6        | 3                                             | G1/S                 | ✓            | ×               | ✓             |
| BUB1B       | 3                                             | G2/M                 | ✓            | ✓              | ✓             | MCM7        | UNKN                                              | G1/S                 | ✓            | ×               | ✓             |
| CCNE2       | 3                                             | G1/S                 | ✓            | ×              | ✓             | MKI67       | 3                                             | G2/M                 | ✓            | ✓              | ×             |
| CCNF        | 3                                             | G2/M                 | ✓            | ✓              | ✓             | MMS2L       | 3                                             | G1/S                 | ✓            | ×               | ✓             |
| CDC7        | 3                                             | G1/S                 | ✓            | ×              | ✓             | MSH2        | 3                                             | G1/S                 | ✓            | ×               | ✓             |
| CDC40       | 3                                             | G2/M                 | ✓            | ✓              | ✓             | MSH6        | 3                                             | G1/S                 | ✓            | ×               | ✓             |
| CENPA       | 3                                             | G2/M                 | ✓            | ✓              | ✓             | MTPB        | 3                                             | G1/S                 | ✓            | ×               | ✓             |
| CENPE       | 3                                             | G2/M                 | ✓            | ✓              | ✓             | MTHFD1      | 3                                             | G1/S                 | ✓            | ×               | ✓             |
| CENPF       | 3                                             | G1/S                 | ✓            | ✓              | ✓             | MBTBL1      | UNKN                                              | ✓                     | ✓            | ×               | ×             |
| CEP50       | 3                                             | UNKN                 | ✓            | ✓              | ✓             | MBTBL1      | 3                                             | UNKN                 | ✓            | ×               | ×             |
| CEP54       | 3                                             | G1/S                 | ✓            | ×              | ✓             | NAP1        | 3                                             | G1/S                 | ✓            | ×               | ✓             |
| CEP60       | 3                                             | G1/S                 | ✓            | ×              | ✓             | NAP1        | 3                                             | G1/S                 | ✓            | ×               | ✓             |
| CHAF1A      | 3                                             | G1/S                 | ✓            | ×              | ✓             | NDC1        | 3                                             | G2/M                 | ✓            | ×               | ×             |
| CHAF1B      | 3                                             | G1/S                 | ✓            | ×              | ✓             | NEMF1       | 3                                             | G1/S                 | ✓            | ×               | ✓             |
| CKS1B       | 3                                             | G2/M                 | ✓            | ✓              | ✓             | OIP5        | 3                                             | G2/M                 | ✓            | ✓              | ✓             |
| DDMA        | 3                                             | G1/S                 | ✓            | ✓              | ✓             | ZC3H1       | 3                                             | G1/S                 | ✓            | ×               | ✓             |
| DHFR        | 3                                             | G1/S                 | ✓            | ✓              | ✓             | PCNA        | 3                                             | G1/S                 | ✓            | ×               | ×             |
| DLGAP5      | 3                                             | G2/M                 | ✓            | ✓              | ✓             | POL2        | 3                                             | G2/M                 | ✓            | ×               | ×             |
| DNA2        | 3                                             | G1/S                 | ✓            | ✓              | ✓             | PML         | 3                                             | G1/S                 | ✓            | ×               | ×             |
| DONSON      | 3                                             | G1/S                 | ✓            | ×              | ✓             | PLK1        | 3                                             | G2/M                 | ✓            | ×               | ×             |
| DSN1        | 3                                             | G1/S                 | ✓            | ×              | ✓             | POLA1       | 3                                             | G1/S                 | ✓            | ×               | ✓             |
| EMPL2       | 3                                             | G1/S                 | ✓            | ×              | ✓             | POLA2       | 3                                             | G1/S                 | ✓            | ×               | ✓             |
| EXO1        | 3                                             | G1/S                 | ✓            | ×              | ✓             | POLD1       | G1/S                                              | ✓                     | ✓            | ×               | ×             |
| EZH2        | 3                                             | G1/S                 | ✓            | ✓              | ✓             | POLD3       | 3                                             | G1/S                 | ✓            | ×               | ×             |
| FAM111B     | 3                                             | G1/S                 | ✓            | ×              | ✓             | POLE        | 3                                             | G1/S                 | ✓            | ✓              | ✓             |
| FEN1        | 3                                             | G1/S                 | ✓            | ✓              | ✓             | POLE        | 3                                             | G1/S                 | ✓            | ✓              | ✓             |
| FIGN1       | 3                                             | G1/S                 | ✓            | ×              | ✓             | RAGAP1      | 3                                             | G2/M                 | ✓            | ×               | ✓             |
| GINS1       | 3                                             | G1/S                 | ✓            | ×              | ✓             | RB1         | 3                                             | G1/S                 | ✓            | ✓              | ✓             |
| GMNN         | 3                                            | G1/S                 | ✓            | ×              | ✓             | RFC4        | 3                                             | G1/S                 | ✓            | ✓              | ✓             |
| GSTE1       | 3                                             | G2/M                 | ✓            | ✓              | ✓             | RFC5        | 3                                             | G1/S                 | ✓            | ✓              | ✓             |
| H2AFX       | 3                                             | UNKN                 | ✓            | ✓              | ✓             | RMI1        | 3                                             | G1/S                 | ✓            | ×               | ×             |
| HAT1        | 3                                             | UNKN                 | ✓            | ×              | ✓             | RINGHE2A    | G1/S                                              | ✓                     | ✓            | ×               | ×             |
| HELLS       | 3                                             | G1/S                 | ✓            | ×              | ✓             | SAS6        | UNKN                                              | ✓                     | ×            | ×               | ×             |
| HMMR        | 3                                             | G2/M                 | ✓            | ×              | ✓             | SMC2        | UNKN                                              | ✓                     | ×            | ×               | ×             |
| ITGB3BP     | 3                                             | UNKN                 | ✓            | ×              | ✓             | TICRR       | 3                                             | G2/M                 | ✓            | ✓              | ✓             |
| KIAA0101    | 3                                             | G1/S                 | ✓            | ×              | ✓             | TIMELESS    | UNKN                                              | ✓                     | ×            | ×               | ✓             |
| KIF11       | 3                                             | G2/M                 | ✓            | ✓              | ✓             | TPX2        | G2/M                                              | ✓                     | ✓            | ×               | ×             |
| KIF15       | 3                                             | G2/M                 | ✓            | ✓              | ✓             | TMS         | 3                                             | G2/M                 | ✓            | ×               | ×             |
| KIF20B      | 3                                             | G2/M                 | ✓            | ×              | ✓             | UBE2C       | 3                                             | G2/M                 | ✓            | ×               | ×             |
| KIF4A       | 3                                             | G2/M                 | ✓            | ✓              | ✓             | UHRF1       | 3                                             | G1/S                 | ✓            | ×               | ✓             |

Table 3. Genes upregulated after HPV E7 expression with an identification-overlap of three in six datasets. 92 genes were identified in 3 of the 6 datasets as being deregulated by HPV E7 (extracted from Table S1). Information whether the gene is a cell cycle gene, including the phase of peak expression, and whether it is a DREAM, MMBOFOX1 or pRB-E2F target were extracted from Fischer et al.\(^5\). UNKN, timing of peak expression of the cell cycle gene is unknown; X, cell cycle-dependent expression was not reported in the datasets.

CDKN2A was previously reported to be upregulated by HPV E7 through a different mechanism, namely epigenetic derepression\(^6\).

To be considered a high confidence HPV E7-deregulated gene, we employed a threshold of at least three datasets that identify the gene as upregulated by E7. Remarkably, 139 of 141 genes (98.6%) that passed these criteria
are predicted cell cycle genes, and 134 (95.0%) are DREAM targets (Tables 2 and 3). The cell cycle genes represent genes with peak expression during G1/S or G2/M phases. Although pRB is the best known target protein of E7, only 87 (61.7%) of the high confidence E7-deregulated genes are predicted pRB-E2F targets. It is important to note that pRB-E2F targets largely represent the G1/S subgroup of DREAM-targeted cell cycle genes. The finding that most HPV E7-deregulated genes are DREAM targets is consistent with the previous finding that disruption of the DREAM complex is critical to prevent cell cycle arrest in HPV-infected cells. Together, our findings indicate that DREAM target genes are generally deregulated by HPV E7 expression.

Figure 1. HPV E7 abrogates p53-mediated downregulation of DREAM target genes. (A) HCT116 wild-type and (B) HCT116 HPV E7-expressing cells were treated with Nutlin-3a or doxorubicin for 24 h. Untreated cells served as control. Semiquantitative RT-PCR was performed. The log 2-fold change of mRNA expression of treated compared to untreated cells is displayed. GAPDH served as a negative control for the p53 response, while p21 (CDKN1A) and PLK4 were tested as positive controls. (C) HCT116 wild-type and E7-expressing cells were treated with Nutlin-3a for 24 h or left untreated. Protein levels were analyzed through immunoblotting and β-actin levels served as loading control. Cropped blot images are displayed; full images are included in Supplementary Figure S1.
High risk HPV E7 abrogates p53-p21-DREAM-mediated repression of cell cycle genes. Given that p53-p21-dependent downregulation of the DREAM target gene PLK4 was disturbed by HPV E7, we asked whether disruption of the p53-p21-DREAM pathway was a general phenomenon upon HPV E7 expression. The p53-p21-DREAM pathway is best characterized in the HCT116 colon carcinoma cell line, and thus, we employed HCT116 cells stably transfected with HPV-16 E7 expression plasmids. We treated wild-type and HPV E7-expressing HCT116 cells with p53-stabilizing Nutlin-3a or the DNA intercalator doxorubicin and compared changes in mRNA levels to untreated control cells (Fig. 1). Consistent with earlier findings, the mRNA levels of the well-established DREAM target genes B-MYB (MYBL2), E2F1, CDC25C, Survivin (BIRC5), KIF23, ORC1 and RAD51 were downregulated in HCT116 wild-type cells treated with Nutlin-3a or doxorubicin compared to untreated cells (Fig. 1A). Most importantly, downregulation of these genes was abrogated upon HPV E7 expression (Fig. 1B). With B-MYB (MYBL2), E2F1, KIF23 and CDC25C serving as examples, western blot analyses showed that protein levels followed decreased mRNA levels. Nutlin-3a treatment led to reduced B-MYB, E2F1, KIF23 and CDC25C protein levels in HCT116 wild-type cells but not in E7-expression HCT116 cells (Fig. 1C). In contrast to the abrogated repression of cell cycle genes, p21 (CDKN1A) was still induced in response to p53 activation even when HPV E7 is present (Fig. 1B). Notably, HCT116 cells that express HPV E7 displayed increased expression of DREAM target genes upon treatment with doxorubicin, but not in the presence of Nutlin-3a, when compared to untreated cells. Doxorubicin can induce G1/S and G2/M cell cycle arrest, while Nutlin-3a mainly induces G1/S arrest. A doxorubicin-induced increase in G1/M cell cycle population leads to increased mRNA levels of late cell cycle genes when the p53 pathway is not active or blocked, which has been observed previously. To explore whether findings from HCT116 cancer cells are also observed in primary cells, we tested for mRNA expression changes following Nutlin-3a treatment in wild-type mouse keratinocytes compared to cells that were expressing HPV E7. Similar to the results from HCT116 cells, wild-type but not E7-expressing mouse keratinocytes displayed decreased mRNA levels of DREAM target genes upon Nutlin-3a treatment. Induction of Cdkn1a (p21), however, was not impaired by E7 expression (Fig. 2).

Discussion
A cell uses several mechanisms to control proliferation. Hypo-phosphorylated forms of the pRB tumor suppressor block E2F-mediated induction of cell cycle genes required for the G1/S transition. In addition, activation of proliferation in cells with serious defects in replication leads to DNA damage and causes stabilization of p53,
which triggers cell cycle arrest or apoptosis. By employing E6 and E7 oncoproteins, human papilloma viruses have evolved two strategies to intercept the host's control of proliferation and response to infection. HPV E6 causes destruction of p53, and E7 forms a complex with pRB, thereby interfering with pRB's ability to form complexes with E2F transcription factors. Recently, a third mechanism based on E7 preventing DREAM complex formation was discovered. Here, we analyzed E7-mediated gene dysregulation using genome-wide data analysis and expression profiling of distinct cell cycle genes. Our analysis revealed that HPV E7 causes deregulation of a large number of cell cycle genes that are normally regulated by DREAM. Deregulation also affects p53 function through disruption of the p53-p21-DREAM pathway (Fig. 3). This mechanism is independent of HPV E6-mediated destruction of p53.

It is widely accepted that pRB controls the G1/S checkpoint and that it is required for G1/S transition. However, HPV-induced proliferation additionally requires deregulation of the G2/M checkpoint. The DREAM complex contributes to the G2/M checkpoint through downregulation of cell cycle genes in response to p53 activation. The HPV E7 oncoprotein deregulates target genes of the DREAM complex that comprise G1/S and G2/M cell cycle genes (Tables 1, 2 and 3 and Figs 1 and 2). Furthermore, the G1/S subgroup of DREAM target genes is also bound by pRB-E2F complexes. DREAM binds to these genes either through E2F promoter elements or through a combination of E2F and CHR-like elements. For clarity only genes with CHR or E2F sites are depicted as examples. When p130/p107 pocket proteins are sequestered and not available for DREAM repressor formation, protein complexes on E2F or CHR sites change their composition from repressor to activator complexes. CHR elements then bind MMB-FOXM1 and E2F sites bind activating E2F1-3-DP complexes, respectively. In conclusion, sequestration of p130 and p107 by HPV E7 abrogates p53-dependent repression of cell cycle genes and thus impairs cell cycle checkpoint control by p53.
were so stringent that the computational analysis rather missed candidates than to include false-positive genes. This suggests that the genes in Tables 2 and 3 are indeed high confidence targets deregulated by HPV E7, but that some additional target genes may have been missed. Taken together, our findings provide evidence that DREAM target genes are generally deregulated by HPV E7 expression.

It is important to note that pRB differs in its function from the pRB-like pocket proteins p107 and p130. While all pocket proteins pRB, p107 and p130 bind to LxCxE motifs, only p107 and p130 can be recruited to the MuvB core through an LxSxExL motif in LIN52 to form the DREAM complex57. HPV E7 possesses an LxCxE motif through which it binds pocket proteins4, and binding of E7 to p107 and p130 inhibits their interaction with the LxSxExL motif in LIN5257. Several other viral oncoproteins target the pocket proteins through LxCxE motifs, including adenovirus early-region 1A (E1A) and large T antigens of several polyomaviruses, such as SV40, JCV and BKV38. Consistent with this notion, also SV40 large T was reported to impair DREAM function39,40.

It is established that HPV destroys p53 function through marking it for degradation by the E6 oncoprotein40,41. This mechanism may be sufficient to block p53 activity completely. However, p21 is a central effector of the p53 response, and p21 can be activated independently of p53, for example through the MAPK and TGFβ pathways42. Also in the absence of HPV E6, we observe that the p53-p21-DREAM pathway is interrupted further downstream by E7 interfering with DREAM function and host cell cycle arrest (Tables 1, 2 and 3 and Figs 1 and 2). The data indicate that HPV employs several means to disrupt cell cycle checkpoints (Fig. 3).

In summary, the data reveal that deregulation of DREAM function by the HPV E7 oncoprotein may contribute substantially to the development of the many cancer types caused by HPV.

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**Author Contributions**

M.F. and K.E. conceived the study. M.F., S.U. and C.S. performed the experiments. M.F. performed the computational analysis. T.M.M. contributed reagents and advice. M.F. and K.E. interpreted the data and wrote the manuscript.
