Merkel cell polyomavirus T-antigens regulate *DICER1* mRNA stability and translation through HSC70

**Highlights**

- MCPyV T-antigen and HSC70 interaction regulates *DICER1* expression
- HSC70 directly binds to ARE in the 3′UTR of *DICER1* for expression regulation
- An unknown motif in *DICER1* CDS is also required for its expression regulation by LT
- The LT-HSC70 interaction can regulate other ARE-containing mRNAs
Merkel cell polyomavirus T-antigens regulate DICER1 mRNA stability and translation through HSC70

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SUMMARY
Merkel cell carcinoma is an aggressive skin malignancy, mostly caused by Merkel cell polyomavirus (MCPyV). MCPyV T-antigens can induce mature microRNA expressions through the DnaJ domain, but its underlying mechanism is still unknown. Here, we report that the T-antigens induce protein expression and mRNA stability of DICER1, a key factor in microRNA biogenesis, through heat shock cognate 70 (HSC70). HSC70 directly interacts with the AU-rich elements (ARE) of DICER1 mRNA in both coding and 3’ untranslated region in the presence of MCPyV T-antigen. The T-antigen/HSC70 interaction could induce luciferase activity of synthetic ARE-containing reporter, as well as the stability of ARE-containing mRNAs, suggesting a broader role of MCPyV T-antigens in regulating multiple mRNAs via HSC70. These findings highlight a new role for the interaction of HSC70 and MCPyV T-antigens in mRNA regulation and an undescribed regulatory mechanism of DICER1 mRNA stability and translation through its direct interaction with HSC70.

INTRODUCTION
Merkel cell carcinoma (MCC) is a neuroendocrine skin cancer with poor prognosis, of which 80% is caused by Merkel cell polyomavirus (MCPyV) (Feng et al., 2008; Shuda et al., 2008). In MCC, the MCPyV is found integrated into tumor DNA with a mutation in the large T-antigen (LT) gene, expressing truncated large T-antigen (trLT) and small T (sT), which both preserve the DnaJ domain at their N-terminals (Shuda et al., 2008). This domain is conserved among polyomaviruses, which recruits heat shock cognate 70 (HSC70) chaperone (Sullivan and Pipas, 2002). In Simian virus 40 (SV40), this interaction is required for viral replication (Campbell et al., 1997), neoplastic transformation (Srinivasan et al., 1997; Sullivan et al., 2000), and promoting cell-cycle progression (Kim et al., 2001; Sullivan et al., 2000; Zalvide et al., 1998). In MCPyV, a point mutation D44N within this motif disrupts the interaction between HSC70 and MCPyV T-antigen (Kwun et al., 2009), abolishing the effect of the LT on promoting viral replication (Kwun et al., 2009) and cell proliferation (Houben et al., 2015).

HSC70 is a member of the heat shock protein 70 (HSP70) family, which is constitutively expressed and functions as a chaperone that binds nascent polypeptides and facilitates their correct folding for preventing protein aggregation, as well as targets misfolded proteins for degradation (Kim et al., 2013). Besides its protein chaperone functions, HSC70 and various homologs in the HSP70 family can directly bind to selective mRNAs, regulating their stability and translation (Castello et al., 2012; Kishor et al., 2013, 2017; Matsui et al., 2007; Zimmer et al., 2001). RNA binding of HSP70 family members is affected by both ATP binding and co-chaperones (Henics et al., 1999; Zimmer et al., 2001), suggesting that the RNA substrate recognition resembles the protein-binding property of chaperone activity, which may assist in proper folding of its targeted RNA during mRNA decay and/or translation. The underlying mechanism of HSP70 family in mRNA metabolism remains to be uncovered.

HSP70 family proteins preferentially bind to AU-rich elements (ARE), which are cis-regulatory elements consisting of AUUUA motif(s) mostly present in the 3’untranslated region (3’UTR) of several mRNAs that typically modulate mRNA decay (Kishor et al., 2013, 2017; Matsui et al., 2007; Zimmer et al., 2001). These chaperone proteins have been suggested to contribute to ARE-mediated decay through two different mechanisms. First, HSP70 proteins may alter the composition of key protein complexes that affect mRNA decay. For example, AU-rich element RNA-binding protein 1 (AUF1), a key ARE-binding protein, binds to poly(A)-binding protein (PABP) and dissociate it from the poly(A) tail, leading to mRNA decay.
HSP70 proteins may disrupt the AUF1-PABP interaction through direct HSP70-AUF1 binding that sequesters AUF1 from ARE-mediated mRNA decay; thus PABP remains associated with poly(A) tail and protects the mRNA from ribonucleases (Muhlrad and Parker, 1992). Second, HSP70 may directly bind to ARE and inhibit mRNA decay. For example, Hsc70 binds to ARE in the 3'UTR of Bim mRNA and stabilizes the transcript during steady state conditions (Matsui et al., 2007). Upon treatment with interleukin-3, Bim mRNA is destabilized (probably through its association with RNA destabilizing factor) with concurrent loss of RNA-binding potential of HSC70 and its interaction with co-chaperones (Matsui et al., 2007). The molecular mechanism of how mRNA stability is regulated by the direct HSC70-RNA interaction is still unclear.

Recently, we observed that T-antigens increased expression of mature microRNAs (miRNA), but not their primary transcripts, through the DnaJ domain of viral T-antigens (Kumar et al., 2020). However, the underlying mechanism of how MCPyV T-antigens regulate miRNA expression remains unknown. One possible mechanism could be the regulation of miRNA biogenesis factors that affect miRNA processing. miRNAs undergo multiple processing steps to produce the functional mature miRNAs (Ha and Kim, 2014). For canonical miRNAs, the primary transcripts are processed by the RNase III enzyme DROSHA together with DiGeorge critical region 8 (DGCR8) in the nucleus, to yield ~70-nt stem-loop precursor miRNAs. These precursor miRNAs are then exported to the cytoplasm via the Exportin-5/Ran-GTP pathway, for further processing into ~22-nt miRNA duplexes by another RNase III enzyme DICER1 and its partner transactivation responsive RNA-binding protein 2 (TARBP2). These key proteins are frequently deregulated and associated with disease progression or poor outcomes in multiple tumor types (Merritt et al., 2008; Zhang et al., 2014) and can also contribute to tumorigenesis in vivo (Kumar et al., 2007, 2009; Ravi et al., 2012; Zhang et al., 2014). Notably, some viruses have been demonstrated to modulate the expression of miRNA processing factors. For example, human papillomavirus (HPV) oncoproteins can increase DROSHA and DICER1 expressions (Harden and Munger, 2017), whereas Kaposi’s sarcoma-associated herpesvirus (KSHV) can upregulate DICER1 and TARBP2 levels (Happel et al., 2016). These data prompted us to investigate whether MCPyV T-antigens regulate miRNA processing factors.

Here, we report that MCPyV T-antigens induce DICER1 protein expression and mRNA stability through the DnaJ domain and that the interaction between HSC70 and T-antigen is required for DICER1 regulation. In addition, we show that HSC70 directly binds to ARE in the coding region and 3'UTR of DICER1 mRNA and stabilizes the transcript. These findings reveal a new role for HSC70-binding site of MCPyV T-antigens and the involvement of LT-HSC70 interaction in regulating DICER1 mRNA stability and protein expression.

**RESULTS**

**MCPyV T-antigens induce DICER1 protein expression and stabilize DICER1 mRNA via DnaJ domain**

Our previous study has shown that MCPyV T-antigens can induce mature miRNA expressions through DnaJ domain, whereas no changes were observed on corresponding primary miRNAs (Kumar et al., 2020), suggesting that T-antigens can regulate miRNA processing. This finding led us to investigate whether MCPyV T-antigens can regulate key miRNA processing factors. We transfected codon-optimized LT (LTco), tumor-derived truncated LT (LT339), and codon-optimized sT (sTco) into MCPyV– MCC cell lines and evaluated the effect on expression levels of DROSHA, DGCR8, DICER1, and TARBP2 using western blotting (Figures 1A and S1). As compared with DROSHA, DGCR8, and TARBP2, protein expression of DICER1 was markedly increased upon transfection with sTco, LTco, or LT339 (Figure S1). Increased protein expression of DICER1 by MCPyV T-antigens was consistently observed in multiple cell lines (Figures 1B and 1C).

Given that the DnaJ domain of MCPyV T-antigens is required for regulation of miRNA expression (Kumar et al., 2020), we determined whether the HSC70-binding site in the DnaJ domain of MCPyV T-antigens is required for DICER1 expression regulation. To address this question, we constructed two types of HSC70-binding defective mutants, LTcoD44N and LTcoΔ. The LTcoD44N mutant has a substitution of aspartic acid with asparagine at residue 44, whereas the LTcoΔ contains a deletion of the HPDKGG (aa 42-47) core motif of DnaJ domain (Figure 1A). These mutants exhibited loss of interaction between HSC70 and LT, as evaluated by co-immunoprecipitation (co-IP) (Figures S2A and S2B). Indeed, we observed that both mutants abolished the effect of LT on increased DICER1 protein expression (Figure 1D).

To further support our findings, we also generated the HPDKGG deletion mutant in LT339 (LT339Δ; Figures 1A and S2C) and a point substitution in the LxCxE motif (LT339E216K). The E216K mutation has been
**A**

| Cell Line | DICER1 | LT | trLT | sT | GAPDH |
|-----------|--------|----|------|---|-------|
| MCC13     | 🟢     | 🟠 | 🟠   | 🟠 | 🟠     |
| LTco      | 🟠     | 🟠 | 🟠   | 🟠 | 🟠     |
| LTcoD44N  | 🟠     | 🟠 | 🟠   | 🟠 | 🟠     |
| LTcoE216K | 🟠     | 🟠 | 🟠   | 🟠 | 🟠     |

**B**

| Cell Line | DnaJ | MUR | OBDMURLTcoD44N |
|-----------|------|-----|----------------|
| MCC13     | 🟢   | 🟠  | 🟦             |
| LTco      | 🟠   | 🟠  | 🟦             |
| LTcoD44N  | 🟠   | 🟠  | 🟦             |
| LTcoE216K | 🟠   | 🟠  | 🟦             |

**C**

| Act. D treatment (hours) | DICER1/GAPDH |
|--------------------------|--------------|
| 0                         | 3.0          |
| 0.5                       | 2.5          |
| 1.0                       | 2.0          |
| 1.5                       | 1.5          |
| 2.0                       | 1.0          |

**D**

| Cell Line | DnaJ | MUR | OBDMURLTcoΔ |
|-----------|------|-----|-------------|
| MCC26     | 🟢   | 🟠  | 🟦          |
| LTco      | 🟠   | 🟠  | 🟦          |
| LTcoD44N  | 🟠   | 🟠  | 🟦          |
| LTcoE216K | 🟠   | 🟠  | 🟦          |

**E**

| Act. D treatment (hours) | DICER1/GAPDH |
|--------------------------|--------------|
| 0                         | 3.0          |
| 0.5                       | 2.5          |
| 1.0                       | 2.0          |
| 1.5                       | 1.5          |
| 2.0                       | 1.0          |

**F**

| Act. D treatment (hours) | DICER1 mRNA stability |
|--------------------------|------------------------|
| 0                        | 1.0                    |
| 1                        | 0.5                    |
| 2                        | 0.2                    |
| 3                        | 0.1                    |
| 4                        | 0.0                    |
| 5                        | 0.0                    |

**G**

| Act. D treatment (hours) | DICER1 mRNA stability |
|--------------------------|------------------------|
| 0                        | 1.0                    |
| 1                        | 0.5                    |
| 2                        | 0.2                    |
| 3                        | 0.1                    |
| 4                        | 0.0                    |
| 5                        | 0.0                    |

**Legend**

- 🟢: Act. D treatment (hours) 0 or 5
- 🟠: Act. D treatment (hours) 0.5
- 🟦: Act. D treatment (hours) 1.0
- ns: not significant
Figure 1. MCPyV T-antigens induce DICER1 protein expression and mRNA stability

(A) Illustration of plasmids expressing different MCPyV T-antigens (LTco, full-length LT; LT339, truncated LT (trLT) and sT; sTco, sT only), HSC70-binding defective (LTcoD44N, LTcoA, and LT339A), and RB-binding defective (LT339D216K) LT mutants. LTcoD44N has a substitution of aspartic acid with asparagine at residue 44, whereas both LTcoA and LT339A contain a deletion of the HPDKGG core motif (aa 42-47). LT339D216K has a substitution of glutamic acid with lysine in the LxCxE RB-binding motif.

(B) Western blot analysis of the effect of MCPyV T-antigens or vector control (pCTR) on DICER1 expression after 48 h of transfection. trLT, truncated LT.

(C) Quantification of DICER1 protein expression in Figure 1B. GAPDH was used for normalization. *p < 0.05, **p < 0.01 by paired t test with Benjamini–Hochberg correction.

(D) Western blot analysis of MCPyV– MCC and HEK293 cell lines transfected with LTco, LTcoD44N, LTcoA, or pCTR after 48 h.

(E) Western blots showing the effect of cells transfected with pCTR, LT339, LT339A, or LT339D216K on DICER1 expression.

(F and G) DICER1 mRNA stability in (F) MCC14/2 cells after transfection with various T-antigens or (G) WaGa cells transfected with shTA, shsTA, or shCTR. Both cell lines were treated with actinomycin D (Act. D, 2 μg/ml) after 48 h. Cells were harvested at 0, 2, 4, and 6 h after Act. D treatment for RT-qPCR analyses. DICER1 expression was normalized to GAPDH, and the treatment at 0 h was set as 1. t1/2 refers to the time needed to reach half amount of decaying mRNA. p values were calculated by two-way ANOVA.

(B, D, and E) The sizes of the molecular weight markers are given to the right of the western blots. The numbers below the western blots refer to the ratio of DICER1/GAPDH relative to their respective pCTR.

(C, F, and G) Each biological replicate is depicted by a circle, or the number of replicates is indicated by n. Error bars represent SEM.

reported to abolish RB binding to LT (Shuda et al., 2008), and the defective RB binding to LT339 was verified by co-IP (Figure 52D). Unlike LTco, LT339 expressed both trLT and sT (Figure 1E). Concordant with LTco, the LT339A DnaJ mutant failed to induce DICER1 protein expression, whereas increased DICER1 expression was still observed in the RB-binding defective LT339D216K mutant (Figure 1E).

MCPyV T-antigen is required for HSC70-mediated DICER1 regulation

To further investigate whether HSC70 is required for regulating DICER1 expression, we silenced HSC70 using two different siRNAs (siHSC70#1 and siHSC70#3) in MCPyV+ WaGa and MKL-1 cell lines. Indeed, silencing of HSC70 resulted in decreased expression of DICER1 (Figures 2A and 2B). However, no significant change of DICER1 protein expression was observed in MCPyV– MCC cell lines upon silencing of HSC70 (Figures 2C and 2D), indicating that HSC70 only regulates DICER1 expression in the presence of viral T-antigen. To support this notion, we compared the effect of T-antigens on DICER1 with and without silencing of HSC70. DICER1 expression was increased (2.5–3.3 folds) upon ectopic expressions of T-antigen (sTco, LTco or LT339) compared with pCTR; however, HSC70 knockdown partially diminished the effect (Figures 2E and 2F).

To further strengthen the involvement of HSC70 in MCPyV T-antigen-mediated DICER1 regulation, we assessed whether ectopically expressed siRNA-resistant His-tagged HSC70 (His-HSC70) could rescue the silencing effect of HSC70 in MCPyV+ WaGa cells. The results showed that expression of His-HSC70 reversed the effect of HSC70 knockdown on DICER1 expression (Figures 2G and 2H). In addition, inhibition of HSC70 in the MCPyV+ WaGa cells reduced DICER1 mRNA stability (Figure 2I). Together, our results indicate that MCPyV T-antigens regulate DICER1 protein expression and mRNA stability through HSC70 and that the interaction between MCPyV T-antigen and HSC70 is crucial for DICER1 regulation.

HSC70 directly binds to 3’UTR and CDS of DICER1

Given the involvement of LT-HSC70 interaction in regulation of DICER1 mRNA stability and protein expression, we assessed whether this interaction binds directly to DICER1 mRNA. We first pulled down RNAs bound to LT using RNA immunoprecipitation (RIP) and quantified DICER1 mRNA by RT-qPCR. The results showed an enrichment of DICER1, but not GAPDH, mRNA in LT immunoprecipitates compared with IgG control (Figure 3A), suggesting that DICER1 mRNA is a part of LT-HSC70 ribonucleoprotein complex. To further determine the importance of LT-HSC70 interaction on DICER1 mRNA binding, we performed RIP on HEK293 cells transfected with pCTR, LTco, or LTcoD44N and found that DICER1 mRNA was significantly enriched in LTco-, but not LTcoD44N, transfected cells compared with pCTR (Figure 3B). We also made..
**Figure A**

Western blots showing expression levels of DICER1, HSC70, and GAPDH in WaGa and MKL-1 cells treated with siCTR, siHSC70#1, and siHSC70#3.

**Figure B**

Graphs displaying the ratio of DICER1 to GAPDH in WaGa and MKL-1 cells treated with different siRNA sequences.

**Figure C**

Western blots showing expression levels of DICER1, HSC70, and GAPDH in MCC13, MCC14/2, and MCC26 cells treated with siCTR, siHSC70#1, and siHSC70#3.

**Figure D**

Graphs displaying the ratio of HSC70 to GAPDH in MCC13, MCC14/2, and MCC26 cells treated with different siRNA sequences.

**Figure E**

Western blot of MCC26 cells treated with pCTR, sTco, LTco, and LT339.

**Figure F**

Graph showing the ratio of DICER1 to GAPDH in pCTR, sTco, LTco, and LT339 treated cells.

**Figure G**

Western blots showing expression levels of DICER1, His-HSC70, HSC70, and GAPDH in WaGa cells treated with siCTR and siHSC70#1.

**Figure H**

Graphs displaying the ratio of DICER1 to GAPDH in WaGa cells treated with different siRNA sequences.

**Figure I**

Graphs showing changes in DICER1 mRNA stability in WaGa cells treated with siCTR and siHSC70#1 over time following Act. D treatment.

Additional notes:
- **DICER1 mRNA stability** in WaGa cells: t1/2 = 2.2 h for siCTR and t1/2 = 1.2 h for siHSC70#1.
- **p-value** for mRNA stability: p = 0.0007, ***
- **n = 3** for each treatment condition.
Figure 2. HSC70 is required for MCPyV T-antigen-mediated regulation of DICER1 protein expression and mRNA stability

(A) MCPyV+ cell lines were transfected with two different siRNAs targeting HSC70 (siHSC70#1 and siHSC70#3) or siRNA control (siCTR) for 48 h, followed by evaluation of the effect on DICER1 expression using western blotting.
(B) Quantification of DICER1 and HSC70 protein expressions in Figure 2A.
(C) Western blot analysis of MCPyV– MCC cell lines transfected with siHSC70 or siCTR after 48 h of transfection.
(D) Quantification of DICER1 and HSC70 in Figure 2C.
(E) MCC26 cells were co-transfected with various T-antigens and siHSC70#1 (+) or siCTR (−). The numbers below the western blots refer to the DICER1/GAPDH ratios relative to cells transfected with pCTR and siCTR.
(F) Quantification of DICER1/GAPDH in Figure 2E. The paired data are connected by gray lines. Error bars, SEM.
(G) Western blot analysis of the effect of ectopically expressed siRNA-resistant His-tagged HSC70 (His-HSC70) on DICER1 expression in WaGa cells transfected with siHSC70 or siCTR.
(H) Quantification of DICER1/GAPDH in Figure 2G.
(I) DICER1 mRNA stability in WaGa cells transfected with siCTR or siHSC70#1. Actinomycin D (Act. D, 2 μg/mL) was added after 48 h of transfection, followed by RT-qPCR. GAPDH was used as endogenous control. ***p < 0.001 by two-way ANOVA.
(B, D, and H) GAPDH was used as loading control. Expression levels were normalized to siCTR. Error bars, SEM. *p < 0.05, **p < 0.01, ***p < 0.001 and ns, not significant by paired t test with Benjamini–Hochberg correction. Each biological replicate is depicted by a circle.

several attempts to pull down endogenous HSC70 in MCPyV+ MCC cell lines using different HSC70 antibodies (sc-7298 from Santa Cruz Biotechnology, NBP2-67335 from Novus Biologicals and ab51052 from Abcam). Unfortunately, none of these antibodies could pull down the HSC70 complexes (data not shown), which is likely due to lack of accessible epitopes for the antibodies. Instead, we ectopically expressed His-HSC70 in HEK293 cells and pulled down exogenous HSC70 using His-tagged antibody (Figure 3C). The results showed an enrichment of DICER1 mRNA in cells co-transfected with His-HSC70 together with LTco but not with LTcoD44N (Figure 3C).

We next investigated the interaction between LT-HSC70 and DICER1 mRNA using biotin pull-down assays with biotinylated RNA probes from full-length 5'UTR (SU), CDS (CDS FL), and 3' UTR (3U FL) of DICER1 (Figures 3D and 3E). We detected HSC70 in the ribonucleoprotein complex using probes spanning DICER1 CDS and 3'UTR but not its 5' UTR or negative control (FlucCTR, biotinylated RNA probe from firefly luciferase gene). However, LT was not detected in the complex, suggesting that DICER1 is directly associated with HSC70 but not with LT.

To further map specific regions of DICER1 that binds to HSC70, we generated 8 probes spanning the CDS and 3 probes spanning the 3'UTR of DICER1 (Figure 3D). Among the CDS probes, CDS_H (nt 5693–6061) showed the strongest association with HSC70 (Figures 3E and S3A). Furthermore, the 3U_2 (nt 7253–8770) in 3' UTR of DICER1 also showed strong interaction with HSC70 (Figures 3E and S3A). Interestingly, both regions contain nonamer motifs of ARE (WWAUUUAWW; Figure 3D), suggesting that HSC70 may preferentially bind to these motifs of DICER1.

To further define the role of T-antigen in HSC70-DICER1 mRNA interaction, we repeated the biotin pull-down assays using the CDS_H and 3U_2 probes in MCPyV+ (WaGa and MKL-1) and the MCPyV–MCC14/2 cell lines. Consistently, HSC70 was detected in the ribonucleoprotein complex from both MCPyV+ cell lines, but barely detectable in the MCPyV–MCC14/2 cell line (Figure S3B), indicating that T-antigen is required for the HSC70-DICER1 mRNA interaction. Overall, our results indicate that HSC70 directly binds to the CDS and 3'UTR of DICER1 mRNA in MCPyV+ MCC cell lines.

The ARE motif in the DICER1 3'UTR is important for MCPyV T-antigen-mediated expression regulation

To determine the role of DICER1 3'UTR in expression regulation, we co-transfected Renilla luciferase reporter containing the wild-type sequence of DICER1 3'UTR and various MCPyV T-antigens into HEK293 cells (Figures 4A and 4B). Our results showed that all T-antigens enhanced luciferase activity compared with vector control (pCTR). On the other hand, LTcoD44N or silencing of HSC70 abolished the effect mediated by LTco (Figures 4B and 4C).

To dissect the role of ARE in this regulation, we selected an ARE (5'-AAUAUUUAUAA-3'; nt 8130–8140; Figure S4), located within the 3U_2 region, which showed strong association with HSC70 by biotin pull-down assay (Figure 3E). We generated two mutant constructs, where either three U > G substitutions were created (Mut_uG) or the whole ARE motif was deleted (Mut_A) (Figure 4A). Consistently, LTco enhanced...
Figure 3. DICER1 mRNA is present in the LT-HSC70 complex and directly associated with HSC70

(A–C) Evaluation of DICER1 mRNA in LT or His-HSC70 immunoprecipitation (IP). RNA immunoprecipitation was performed on (A) WaGa cells or (B) HEK293 cells transfected with LTco, LTcoD44N or pCTR, using LT antibody, as well as on (C) HEK293 cells co-transfected with His-HSC70 together with LTco or LTcoD44N.
Figure 3. Continued

LTcoΔD44N using His-tagged antibody. Upper: IP efficiency was validated by western blotting. IgG IP or pCTR was used as a negative control. WCL, whole-cell lysate. Lower: quantification of DICER1 and GAPDH mRNA in LT or His IP using RT-qPCR. Enrichment of mRNA was calculated by dividing the mRNA in LT IP to input and compared with their expression levels in IgG IP control (A and C) or pCTR (B). Error bars, SEM. *p < 0.05 by paired t test. Each biological replicate is depicted as a circle. (D) Illustration of DICER1 mRNA and various RNA probes synthesized for biotin pull-down assays. Probes spanning the full length of 5'UTR, CDS, and 3'UTR of DICER1 are indicated by SU (orange), CDS FL (red), and 3U FL (green), respectively. Fragment probes spanning the CDS are presented as (A) to (H) (red). whereas the probes for 3'UTR fragments are shown as 3U_1, 3U_2, and 3U_3 (green). The numbers next to the probe name refer to the nucleotide positions in the transcript (ENST00000526495.5). Arrowheads indicate various ARE motifs predicted by AREsite (Gruber et al., 2010). (E) Biotinylated RNA probes (20 pmol) were incubated with protein lysates from WaGa cells for 5–6 h at 4°C, and then pulled down by streptavidin beads. The proteins bound to DICER1 RNA probes were detected by western blotting. Biotinylated RNA probes from firefly luciferase gene (FlucCTR) were included in each experiment as a negative control. WCL, whole-cell lysate; blank, empty lane. The same blots with high and low exposure for HSC70 detection are shown in Figure S3A.

Luciferase activity of DICER1 3'UTR WT, but not the luciferase activity of DICER1 3'UTR Mutsub and MutΔ (Figure 4D), suggesting that this ARE in DICER1 3'UTR is important for T-antigen regulation of DICER1 expression. In addition, RNAfold (http://rna.tbi.univie.ac.at/cgi-bin/RNAWebSuite/RNAfold.cgi) predicted a linear secondary structure of this ARE region, making it accessible for HSC70 binding (Figure S4).

To determine whether MCPyV T-antigen functions in general ARE-mediated gene regulation, we cloned an artificial ARE sequence (AREWT) or mutant ARE (AREMUT) into Renilla luciferase vector and evaluated the effect of LTco or LTcoΔD44N on these reporter activities (Figure 4E). Similar to the observations for DICER1 3'UTR reporter, LTco, but not LTcoΔD44N, induced luciferase activity of the AREWT, and no enhanced luciferase activity was observed in the AREMUT (Figure 4E). These findings suggest that LT-HSC70 interaction can regulate multiple ARE-containing mRNAs. To address this question, we quantified mRNA half-life of two genes, BCL2L11 and MYC, which have cis-regulatory ARE within their mRNA (Brewer and Ross, 1988; Matsu et al., 2007). The mRNA half-life of BCL2L11 was increased by LTco, whereas LTcoΔD44N showed no effect on its mRNA stability (Figure 4F). Concordantly, the BCL2L11 protein level was also induced by LTco but not by the HSC70-binding defective LT mutants (Figure 4G). Although LTco had a subtle effect on MYC mRNA stability (Figure 4F), its protein level was markedly induced in cells transfected with LTco compared with pCTR or the HSC70-binding defective LT mutants (Figure 4G). These data suggest that MCPyV T-antigens can regulate a number of mRNAs containing ARE via HSC70.

Deletion of nonamer ARE in the CDS of DICER1 partly reverses the effect of LT

Our biotin pull-down assays revealed that HSC70 interacted with the CDS of DICER1 and fragment H (nt 5693–6061) had the strongest interaction (Figure 3E). We determined whether DICER1 CDS alone (without 3'UTR) is sufficient to regulate DICER1 expression, by co-transfecting a plasmid expressing the full-length CDS of DICER1 together with T-antigen or pCTR in DICER1 knockout cell line (NoDice 2-20) (Figures 5A and S5A). Our results showed that all T-antigens (sTco, LTco and LT339) increased expression of exogenous DICER1 (Figure S5A). However, the HSC70-binding defective LT mutant (LTcoΔD44N) or HSC70 knockout (siHSC70#1) abolished the effect of T-antigen-induced DICER1 expression (Figures 5B and S5B).

To assess whether the nonmeric ARE in the fragment H of DICER1 CDS can regulate DICER1 expression, we constructed an in-frame deletion mutant, which has a deletion of the nonmeric ARE in the CDS of DICER1 (CDSΔ; Figure 5A). By comparing the effect of LT in NoDice 2-20 cells, we observed 4-fold increased expression of DICER1 in the CDSΔ, and < 2-fold in CDSΔ (Figure 5C), suggesting that the nonmeric ARE in the DICER1 CDS partly contributes to expression regulation by the viral T-antigens.

AUF1 does not contribute to DICER1 expression regulation in MCC

To our knowledge, AUF1 is the only RNA-binding protein that binds to DICER1 mRNA and destabilize the transcripts (Abdelmohsen et al., 2012). To determine whether AUF1 plays a role in DICER1 regulation in MCC, we silenced AUF1 using two different siRNAs (siAUF1#1 and siAUF1#2) in MCPyV+ (WaGa and MKL-1) and MCPyV− (MCC26) cell lines and evaluated the effect on DICER1 protein expression. Silencing of AUF1 did not alter DICER1 protein expression in any of the three cell lines (Figures 6A and 6B). Similarly, no changes of DICER1 mRNA stability were observed upon silencing of AUF1 in WaGa (Figure 6C). In addition, no AUF1 was detected in the DICER1 biotin pull-down assays (Figures S3A and S3B). Furthermore, protein fractionation experiments revealed that AUF1 was predominantly present in the nuclear fraction, whereas HSC70 and endogenous/
**A**

DICER1 3'UTR WT

DICER1 3'UTR Mut<sub>sub</sub>

DICER1 3'UTR Mut<sub>a</sub>

8130 8140

WT...CUUAAAUUUUAUAAAC...

Mut<sub>sub</sub>...CUUAAGAUGAGAAAC...

Mut<sub>a</sub>...CUUU----------AC...

**B**

Renilla/firefly luciferase activity (normalized to pCTR)

**C**

Renilla/firefly luciferase activity (normalized to their respective pCTR)

**D**

Renilla/firefly luciferase activity (normalized to their respective pCTR)

**E**

ARE<sub>WT</sub>

ARE<sub>MUT</sub>

**F**

BCL2L11 mRNA stability (normalized to GAPDH)

(t<sub>1/2</sub> = 1.36 h)

(t<sub>1/2</sub> = 2.12 h)

(t<sub>1/2</sub> = 1.33 h)

**G**

HEK293

MCC14/2
Figure 4. ARE motif in the 3'UTR contributes to gene expression regulation by MCPyV T-antigen

(A) Illustration of various Renilla luciferase reporters containing wild-type (WT) or mutations (Mut) of DICER1 3'UTR. Two types of mutants were generated: MutWT contains three nucleotide substitutions (underlined), and MutΔ has a deletion of the ARE motif (nt 8130–8140), which are highlighted in red. (B and C) Luciferase activity was assessed in HEK293 cells 48 h after co-transfection with DICER1 3'UTR WT luciferase reporter and (B) various MCPyV T-antigens or (C) pCTR or LTco together with siHSC70 or siCTR. (D) The effect of LTco on luciferase activity from reporters containing wild-type (WT) or mutations (MutWT) or MutΔ of DICER1 3'UTR was evaluated in HEK293 cells after 48 h of transfection. (E) Upper: Illustration of Renilla luciferase reporters containing synthetic ARE sequences (AREWT) or ARE mutant (AREMUT). In the AREMUT, three U > G substitutions were introduced in the middle U of the pentameric ARE motif. Lower: The graph shows the effect of LTco and LTcoD44N on AREWT luciferase activity, as well as the effect of LTco on AREMUT luciferase activity in HEK293 cells. (B–E) Error bars, SEM; *p < 0.05, ***p < 0.001 by paired t test with Benjamini--Hochberg correction for multiple comparisons. Each biological replicate is depicted as a circle. (F) RT-qPCR analysis of BCL2L11 and MYC in HEK293 cells transfected with pCTR, LTco, or LTcoD44N after addition of actinomycin D (Act. D) at the indicated time points. mRNA levels were normalized to GAPDH. p values were calculated by two-way ANOVA. (G) Western blot analysis of MYC and BCL2L11 in cells transfected with pCTR, LTco, and HSC70-binding defective LT mutants (LTcoD44N and LTcoΔ). MYC/GAPDH and BCL2L11/GAPDH ratios relative to pCTR are shown below the western blots.

exogenous LT were highly abundant in the cytoplasm (Figures 4D and 6E), suggesting that LT-HSC70 interaction binds to DICER1 mRNA in the cytoplasm, thus enhancing its mRNA stability and translation. Our results also indicate that AUF1 is not involved in DICER1 regulation in MCC.

DISCUSSION

This study reveals a novel role for LT-HSC70 interaction in RNA regulation. Particularly, we show that this interaction regulates DICER1 mRNA stability and protein expression through the ARE located in the CDS and 3'UTR of DICER1.

MCPyV T-antigens posttranscriptionally regulate DICER1 expression via HSC70

Here, we demonstrate that (1) ectopic expression of MCPyV T-antigen or silencing of endogenous T-antigens affect DICER1 mRNA stability; (2) DICER1 mRNA is enriched in the LT-HSC70 ribonucleoprotein complex but not in the complex of HSC70-binding defective LT mutant; (3) DICER1 mRNA directly interacts with HSC70. In addition, we also observed that ectopic expression of MCPyV T-antigens did not affect proteolysis of DICER1 (data not shown), excluding the possibility that the increased expression of DICER1 protein level was due to inhibition of protein degradation pathway.

It is noteworthy that both sT and LT can regulate DICER1 expression through HSC70. Although we did not generate HSC70-binding defective sTcoD44N, the HPDKGG deletion mutant of LT339 (LT339Δ) expressed both trLT and sT. Notably, both trLT and sT were detected in His IP from cells expressing LT339 but not LT339Δ (Figure 52C2). These results support that sT also binds to HSC70 and the deletion of the HPDKGG motif abolished its interaction with HSC70. In addition, ectopic expression of sTco also induced DICER1 protein expression, and silencing of HSC70 reversed the effect of sTco (Figure 2E). Collectively, these data support that both LT and sT regulate DICER1 expression via HSC70.

The identification of DICER1 mRNA as a direct target of HSC70 in the present study also expands the involvement of HSC70 in regulation of mRNA stability in a broader context. Prior to this study, Hsc70 was only known to bind to Bim mRNA in a cytokine-dependent manner (Matsui et al., 2007). It is noted that cochaperones, Bag-4, CHIP, Hsp40, and Hip, are required for Hsc70 binding to Bim mRNA (Matsui et al., 2007). Similarly, MCPyV T-antigen is required for HSC70-DICER1 mRNA interaction and DICER1 expression regulation; thus, the DnaJ domain of MCPyV T-antigen can act as co-chaperone for regulating the RNA-binding potential of HSC70. Similar to Bim (Matsui et al., 2007), we show that AUF1 is not involved in regulating mRNA stability of DICER1. These findings suggest that HSC70-mediated mRNA stabilization of Bim and DICER1 is unlikely due to disruption of AUF1–PABP interaction, leading to sequestration of AUF1 from its mRNA decay activity. Alternatively, both data support direct interaction between HSC70 and mRNAs that contributes to mRNA stability via an unknown mechanism.

ARE in the 3'UTR of DICER1 contributes to expression regulation by the LT-HSC70 interaction

Here, we demonstrate that HSC70 directly binds to CDS and 3'UTR of DICER1, and both regions of DICER1 can contribute to expression regulation by the LT-HSC70 interaction. Similarly, AUF1 has also been shown to interact with both CDS and 3'UTR of DICER1 mRNA (Abdelmohsen et al., 2012). These findings suggest...
**Figure 5. The nonameric ARE in the CDS partly contributes to LT-induced DICER1 expression**

(A) Illustration of plasmids containing the full-length CDS of DICER1 (CDS\textsubscript{WT}) or an in-frame deletion of aa1889-1891, which corresponds to the nonameric ARE (CDS\textsubscript{Δ}).

(B) Western blot analysis of NoDice 2-20 cells co-transfected with DICER1 CDS\textsubscript{WT} and LT\textsubscript{co}, LT\textsubscript{co D44N}, or pCTR.

(C) The effect of LT\textsubscript{co} on exogenous DICER1 expression from DICER1 CDS\textsubscript{WT} and CDS\textsubscript{Δ} was evaluated in NoDice 2-20 cells using western blotting.
that sequence elements residing within these regions can affect DICER1 expression. At the posttranscriptional level, the 3’UTR is the most common site for expression regulation. This region usually contains regulatory elements for mRNA decay, such as ARE- and miRNA-binding sites (Mayr, 2017). For DICER1, multiple miRNAs, e.g. miR-103/107 family (Martello et al., 2010), miR-192 (Feinberg-Gorenstein et al., 2013), and miR-630 (Rupaimoole et al., 2016), have been demonstrated to target its 3’UTR. Besides miRNAs, to our knowledge, only two ARE-binding proteins, AUF1 and HSC70, are known to interact with the 3’UTR of DICER1. Notably, the HSC70 interaction sites of DICER1 3’UTR are similar to AUF1, in which only the first half of the 3’UTR interacts with both HSC70 and AUF1. Although it is unclear whether HSC70 and AUF1 bind to the same ARE site(s) of DICER1 3’UTR, we could not detect AUF1 in our biotin pull-down assays in both MCPyV+ and MCPyV– MCC cell lines. Furthermore, silencing of AUF1 did not affect DICER1 protein expression and mRNA stability. These findings suggest that HSC70 is unlikely competing with AUF1 for the same ARE sites, releasing DICER1 from AUF1-mediated mRNA decay. Whether the direct interaction between HSC70 and mRNAs affect mRNA stability through inhibition of deadenylation has yet to be uncovered. Alternatively, the binding of HSC70 to DICER1 3’UTR may inhibit miRNAs from associating with their target sites, thereby releasing DICER1 from miRNA-mediated repression. Similar mechanism has also been observed for other RNA-binding proteins, such as Dnd1 (Kedde et al., 2007) and RBM38 (Léveillé et al., 2011), in which both proteins bind to uridine-rich regions near miRNA target sites and block miRNA accessibility.

**A coding region determinant in DICER1 mRNA is required for LT-mediated DICER1 expression**

Although ARE are mostly described within the 3’UTR of mRNAs, mapping of ARE-binding proteins’ binding sites using high-throughput method, e.g. PAR-CLIP, has identified binding motifs beyond 3’UTR, including CDS (e.g. LARP4B [Küspert et al., 2015; Van Nostrand et al., 2020]) and intronic regions (e.g. HuR [Lébedeva et al., 2011; Mukherjee et al., 2011]). We therefore investigated fragment H of the DICER1 CDS (nt 5693–6061), which showed the strongest interaction with HSC70 and has a putative nonameric ARE. Although deletion of the ARE significantly reduced the effect of T-antigen compared with DICER1 CDSWT, LT still increased DICER1 expression with CDS, but to a lesser extent. These results suggest that other sequences in fragment H or other regions within the DICER1 CDS may also contribute to DICER1 expression regulation. Nevertheless, this motif still partly contributes to increased DICER1 expression by T-antigen.

HSP70 family proteins, including HSC70, are associated with polysomes, and these chaperones have been demonstrated to enhance translation elongation by regulating nascent peptide folding, leading to continued elongation (Liu et al., 2013; Shalgi et al., 2013). In addition, HSP70 proteins can also enhance translation initiation by maintaining interaction with Pab1, the yeast homolog of human PABP (Horton et al., 2001; Zhong and Arndt, 1993), or solubility of eIF4G (Cuesta et al., 2000), which strengthens the translation initiation complexes. In both scenarios, HSP70/HSC70 is not expected to bind directly to mRNAs. However, the LT-HSC70 interaction could still recruit translational complexes for mRNA translations.

Alternatively, DICER1 mRNA may contain a translation-dependent coding region determinant (CRD) that regulates its expression. It is known that some mRNAs contain CRD that can affect their stability through endonucleolytic cleavage or deadenylation during translation (Bernstein et al., 1992; Grosset et al., 2000; Mattijssen et al., 2017; Noubissi et al., 2006; Wisdom and Lee, 1991). There are several examples, such as c-myc (Bernstein et al., 1992; Wisdom and Lee, 1991), c-fos (Grosset et al., 2000), β-TrCP (Noubissi et al., 2006), and LARP4 (Mattijssen et al., 2017). For c-myc, the CRD is located in the last 249 nucleotides of the CDS, which can trigger rapid mRNA turnover (Wisdom and Lee, 1991). This region is associated with an RNA-binding protein, called CRD-BP (also known as IGF2BP1), which protects the CRD from degradation by polysome-associated endonuclease(s) during translational pausing (Lemm and Ross, 2002). Given that translational pausing is linked to heat shock (Shalgi et al., 2013) and proteotoxic stress (Liu et al., 2013), it
is tempting to speculate that the interaction between HSC70 and DICER1 CDS may protect the mRNA from endonucleolytic attack, deadenylation, and/or enhance mRNA translation.

**LT-HSC70-mediated RNA regulation and potential clinical implications**

In addition to DICER1, we also show that the LT-HSC70 interaction can induce luciferase activity of synthetic ARE-containing reporter, as well as mRNA stability or protein expression of endogenous ARE-containing mRNAs, BCL2L11, and MYC. These results suggest that the LT-HSC70 interaction can regulate multiple mRNAs. Further investigations are warranted to map the LT-mediated HSC70-mRNA interactions on a genome-wide scale and investigate the functional roles of these interactions in MCC.
Although the clinical implications of our findings are presently unknown, HSC70 is the most abundantly expressed HSP70 family member in MCC cell lines (Adam et al., 2014). Importantly, MAL3-101, an inhibitor of HSP70 ATPase activity, has been demonstrated to induce apoptosis in MCC cell lines and inhibit tumor growth in MCC xenograft model (Adam et al., 2014), suggesting that inhibition of HSP70/HSC70 activity can be a potential therapeutic strategy for MCC treatment.

Based on our findings, we propose a model that the DnaJ domain of T-antigen recruits HSC70 that binds to DICER1 mRNA in both CDS and 3'UTR, leading to mRNA stabilization and enhanced protein expression. In HSC70-binding defective mutant of T-antigen or without MCPyV T-antigen, HSC70 can no longer bind to DICER1 mRNA and regulate its expression.

Limitations of study
This study demonstrates that MCPyV T-antigens, including both LT and sT, regulate DICER1 mRNA stability and protein expression through HSC70. Although the enrichment of DICER1 mRNA in His-HSC70 immunoprecipitates of cells expressing LT and detection of HSC70 using DICER1 mRNA probes in the biotin pull-down assays provide strong evidence of direct HSC70-DICER1 mRNA interaction, we were unable to pull down endogenous HSC70 using several commercial HSC70 antibodies. The evidence of direct interaction with the endogenous HSC70 will further establish the significance of our findings.

Another limitation of this study is the underlying mechanism of how the HSC70-DICER1 mRNA interaction contributes to DICER1 mRNA stability and translation. Further experiments are warranted to investigate whether distinct mechanisms are involved for HSC70-mediated regulation of DICER1 3'UTR and CDS. Although the functional roles of MCPyV T-antigen-mediated DICER1 regulation have not been investigated in this study, we previously demonstrated that the DnaJ domain of MCPyV LT is required for miRNA regulation (Kumar et al., 2020). However, it is still unclear how the viral T-antigens contribute to selective miRNA processing. One possible explanation could be attributed to the binding of RNA-binding proteins, e.g. KSRP (Trabucchi et al., 2009) and lin28 (Heo et al., 2009; Viswanathan et al., 2008), which can promote or prevent processing of their targeted precursor miRNAs.

Although our current data indicate that the LT-HSC70 interaction can regulate multiple ARE-containing miRNAs, other LT-HSC70-regulated miRNAs have yet to be identified using high-throughput sequencing approach and functionally characterized. The results of these studies will shed new insights into the mechanisms and functions of MCPyV T-antigens in viral infection and tumorigenesis.

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Supplemental information can be found online at https://doi.org/10.1016/j.isci.2021.103264.
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### STAR METHODS

#### KEY RESOURCES TABLE

| REAGENT or RESOURCE | SOURCE | IDENTIFIER |
|---------------------|--------|------------|
| **Antibodies**      |        |            |
| Rabbit polyclonal anti-AUF1 (HNRNPD) antibody | Novus Biologicals | NBP1-88915 RRID: AB_11037375 |
| Mouse monoclonal BCL2L11 antibody (H-5) | Santa Cruz Biotechnology | sc-374358 RRID: AB_10987853 |
| Rabbit monoclonal anti-DGCR8 antibody (EPR18757) | Abcam | ab191875 RRID: AB_2892625 |
| Rabbit monoclonal anti-DICER1 antibody (D38 × 10^3) | Cell Signaling Technology | 5342 RRID: AB_10692484 |
| Mouse monoclonal anti-DROSHA antibody (C-7) | Santa Cruz Biotechnology | sc-393591 RRID: AB_2732793 |
| Rabbit monoclonal anti-GADPH antibody (14C10) | Cell Signaling Technology | 5174 RRID: AB_10622025 |
| Mouse monoclonal anti-His-Tag antibody (H-3) | Santa Cruz Biotechnology | sc-8036 RRID: AB_627727 |
| Mouse monoclonal anti-Histone H3 antibody (1G1) | Santa Cruz Biotechnology | sc-517576 RRID: AB_2848194 |
| Mouse monoclonal anti-HSC70 antibody (8-6) | Santa Cruz Biotechnology | sc-7298 RRID: AB_627761 |
| Rabbit monoclonal anti-HSC70 antibody (SR39-04) | Novus Biologicals | NBP2-67335 RRID: NBP2-67335 |
| Mouse monoclonal anti-MCPyV large T-antigen antibody (CM2B4) | Santa Cruz Biotechnology | sc-136172 RRID: AB_2013156 |
| Mouse monoclonal anti-MCPyV T-antigen antibody (clone 2t2) | Merck KGaA | MABF2316 RRID: MABF2316 |
| Mouse monoclonal anti-MYC antibody (9E10) | Santa Cruz Biotechnology | sc-40 RRID: AB_2857941 |
| Rabbit polyclonal anti-RB1 antibody (C-15) | Santa Cruz Biotechnology | sc-50 RRID: AB_632339 |
| Mouse monoclonal anti-TARBP2 (TRBP2) antibody (D-5) | Santa Cruz Biotechnology | sc-51424 RRID: AB_2732792 |
| Mouse monoclonal anti-α-TUBULIN antibody (B-7) | Santa Cruz Biotechnology | sc-5286 RRID: AB_628411 |
| IRDye 680RD Goat anti-Rabbit IgG Secondary Antibody | LI-COR Biosciences | 926-68071 RRID: AB_10956166 |
| IRDye 800CW Goat anti-Mouse IgG Secondary Antibody | LI-COR Biosciences | 926-32210 RRID: AB_621842 |

Bacterial and virus strains

| **TOP10 Chemically Competent E. coli** | Thermo Fisher Scientific | C4004010 |

Chemicals, peptides, and recombinant proteins

| **TRizol Reagent** | Thermo Fisher Scientific | 15596026 |
| **Actinomycin D** | Sigma-Aldrich | A1410 |
| **Lipofectamine LTX Reagent with PLUS Reagent** | Thermo Fisher Scientific | 15338030 |

(Continued on next page)
### REAGENT or RESOURCE SOURCE IDENTIFIER

| REAGENT or RESOURCE | SOURCE | IDENTIFIER |
|---------------------|--------|------------|
| Lipofectamine™ 3000 Transfection Reagent | Thermo Fisher Scientific | L3000008 |
| Ingenio electroporation solution | Mirus Bio LLC | MIR50111 |
| Pierce BCA Protein Assay Kit | Thermo Fisher Scientific | 23225 |
| SuperSignal West Femto Maximum Sensitivity Substrate | Thermo Fisher Scientific | 34095 |
| Protein G Sepharose 4 Fast Flow beads | Sigma-Aldrich | GE17-0618-01 |
| Dynabeads M-280 Streptavidin | Thermo Fisher Scientific | 11205D |
| Biotinylated UTP (Bio-11-UTP) | Thermo Fisher Scientific | AM8450 |
| LDS loading buffer | Thermo Fisher Scientific | NP0008 |
| Sample Reducing Agent | Thermo Fisher Scientific | NP0009 |
| RIPA buffer | Thermo Fisher Scientific | 89900 |
| NP40 buffer | Thermo Fisher Scientific | FN0021 |
| RNaseOUT Recombinant Ribonuclease Inhibitor | Thermo Fisher Scientific | 10777019 |
| Phenylmethanesulfonyl fluoride | Sigma-Aldrich | 93482 |
| Dithiothreitol | Thermo Fisher Scientific | P2325 |
| Proteinase inhibitor | Sigma-Aldrich | P8340 |
| MultiScribe Reverse Transcriptase | Thermo Fisher Scientific | 4311235 |
| Platinum Taq DNA Polymerase | Thermo Fisher Scientific | 10966018 |

### Critical commercial assays

| TaqMan gene expression assay: DICER1 | Thermo Fisher Scientific | Hs00229023_m1 |
| TaqMan gene expression assay: GAPDH | Thermo Fisher Scientific | Hs99999905_m1 |
| TaqMan gene expression assay: BCL2L11 | Thermo Fisher Scientific | Hs01076940_m1 |
| TaqMan gene expression assay: MYC | Thermo Fisher Scientific | Hs00153408_m1 |
| Monarch PCR & DNA Cleanup Kit | New England Biolabs | T1030 |
| HiScribe T7 High Yield RNA Synthesis Kit | New England Biolabs | E20405 |
| MEGAtall Transcription Clean-Up Kit | Thermo Fisher Scientific | AM1908 |
| Dual-Glo Luciferase Assay | Promega Biotech | E2920 |
| Q5 Site-Directed Mutagenesis Kit | New England Biolabs | E0554 |
| Qproteome Cell Compartment Kit | Qiagen | 37502 |

### Experimental models: Cell lines

| Cell line | Source | RRID |
|-----------|--------|------|
| MCPyV- MCC13 | CellBank Australia | CBA-1338, RRID: CVCL_2583 |
| MCPyV- MCC14/2 | CellBank Australia | CBA-1340, RRID: CVCL_2584 |
| MCPyV- MCC26 | CellBank Australia | CBA-1341, RRID: CVCL_2585 |
| MCPyV+WaGa | Laboratory of Jurgen Becker | RRID: VCL_E998 |
| MCPyV+ MKL-1 | Laboratory of Nancy Krett | RRID: CVCL_2600 |
| HEK293 | ATCC | Cat# CRL-1573, RRID: CVCL_0045 |
| NoDice 2-20 | Bogerd et al. (2014) | N/A |

### Oligonucleotides

**In vitro transcription of DICER1 RNA probes (5’ to 3’)**

- **5’UTR forward:** This paper | N/A
- **5’UTR reverse:** This paper | N/A
- **3’UTR forward:** TAATACGACTCACTATAGGGGCGGAAGTGGGTGTTTGTA
- **3’UTR reverse:** TCATCCAGTTTCTTTGATTGC

(Continued on next page)
| REAGENT or RESOURCE | SOURCE | IDENTIFIER |
|---------------------|--------|------------|
| CDS_A forward:      | This paper | N/A        |
| TAATACGACTCCTATAGGGATGAAAGCCCTGCTTTGCA |
| CDS_A reverse:      | This paper | N/A        |
| TAAAGGCCACTTCTGTCAGT |
| CDS_B forward:      | This paper | N/A        |
| TAATACGACTCCTATAGGGATGAAAGCCCTGCTTTGCA |
| CDS_B reverse:      | This paper | N/A        |
| CCAAGTTGCACTTTTGTTATA |
| CDS_C forward:      | This paper | N/A        |
| TAATACGACTCCTATAGGGATGAAAGCCCTGCTTTGCA |
| CDS_C reverse:      | This paper | N/A        |
| TCTTGTGGTATCTTCAAGAG |
| CDS_D forward:      | This paper | N/A        |
| TAATACGACTCCTATAGGGATGAAAGCCCTGCTTTGCA |
| CDS_D reverse:      | This paper | N/A        |
| GCAGTCAAAGGCAGTGGAAG |
| CDS_E forward:      | This paper | N/A        |
| TAATACGACTCCTATAGGGATGAAAGCCCTGCTTTGCA |
| CDS_E reverse:      | This paper | N/A        |
| TGCAAAATAGATATGTTGTTG |
| CDS_F forward:      | This paper | N/A        |
| TAATACGACTCCTATAGGGATGAAAGCCCTGCTTTGCA |
| CDS_F reverse:      | This paper | N/A        |
| TTCCCGATCAGTGCTTTTTAA |
| CDS_G forward:      | This paper | N/A        |
| TAATACGACTCCTATAGGGATGAAAGCCCTGCTTTGCA |
| CDS_G reverse:      | This paper | N/A        |
| CTCTCTTTCTCTTATCTCTT |
| CDS_H forward:      | This paper | N/A        |
| TAATACGACTCCTATAGGGATGAAAGCCCTGCTTTGCA |
| CDS_H reverse:      | This paper | N/A        |
| TCAGCTATTGGGAACCTGAG |
| 3U_1 forward:       | This paper | N/A        |
| TAATACGACTCCTATAGGGATGAAAGCCCTGCTTTGCA |
| AGAACCAGC |
| 3U_1 reverse:       | This paper | N/A        |
| ACGGCAGTTTATCGAAACGT |
| 3U_2 forward:       | This paper | N/A        |
| TAATACGACTCCTATAGGGATGAAAGCCCTGCTTTGCA |
| CTGCGTG |
| 3U_2 reverse:       | This paper | N/A        |
| CAGTCTTTTACACACGTCGTCAG |
| 3U_3 forward:       | This paper | N/A        |
| TAATACGACTCCTATAGGGATGAAAGCCCTGCTTTGCA |
| TAAAGGACTG |
| 3U_3 reverse:       | This paper | N/A        |
| CCGCGAAGCAGTGATAACCTTAT |

Site-directed mutagenesis:

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| REAGENT or RESOURCE | SOURCE | IDENTIFIER |
|---------------------|--------|------------|
| LTco<sup>a</sup> sense: AACCCAGCTATTATGATG | This paper | N/A |
| LTco<sup>a</sup> antisense: GTGTTTCAAACAGGATCG | This paper | N/A |
| LT339<sup>a</sup> sense: AATCCTGTATAATGATGGAATTG | This paper | N/A |
| LT339<sup>a</sup> antisense: ATGCTTTAAGCACCTTTTTGG | This paper | N/A |
| LT339<sup>E216K</sup> sense: CTTCTGCGATAAATCACTTTCCTCCC | This paper | N/A |
| LT339<sup>E216K</sup> antisense: AGATCCTCCAGGTGC | This paper | N/A |
| DICER1 3′UTR Mut<sub>sub</sub> sense: TAGAAACTACCTGATGCGCAGGA | This paper | N/A |
| DICER1 3′UTR Mut<sub>sub</sub> antisense: CATCTTAAGAGATTTGTGATTAACTTGATC | This paper | N/A |
| DICER1 3′UTR Mut<sub>D</sub> sense: ACTACCTGTAGCCAGGAC | This paper | N/A |
| DICER1 3′UTR Mut<sub>D</sub> antisense: AAGAGATTTGTGATTAACTTGATCAG | This paper | N/A |
| DICER1 CDS<sub>D</sub> sense: AGGGGAAATTCAAAGGTGTTG | This paper | N/A |
| DICER1 CDS<sub>D</sub> antisense: TTCCTACTACTTCCACAG | This paper | N/A |
| Artificial ARE constructs | |
| ARE<sub>MUT</sub> sense: CTAGTTATGTATGATGATGATGTCGCC | This paper | N/A |
| ARE<sub>MUT</sub> antisense: GGCCGATACATACATACATAACATAA | This paper | N/A |
| ARE<sub>WT</sub> sense: CTAGTTATTTATTTATTTATTTATTTGC | This paper | N/A |
| ARE<sub>WT</sub> antisense: GGCCGATACATAAATAAAATAAAATAA | This paper | N/A |
| siRNA (uppercase, DNA; lowercase, RNA) | |
| siAUF1#2 sense: acgaggaggaagaagccaaTT | Raineri et al. (2004) | N/A |
| siAUF1#2 antisense: uggccuaauuccucucguuTT | Raineri et al. (2004) | N/A |
| siHSC70#1 | Qiagen | SI02661477 |
| siHSC70#3 | Qiagen | SI04236596 |
| siAUF1#1 | Qiagen | SI00300454 |
| Recombinant DNA | |
| LTco (pcDNA6 MCV LTco) | Shuda et al. (2011) | Addgene Cat#40200 RRID: Addgene_40200 |
| LTco<sup>a</sup> | This paper | N/A |
| LTco<sup>D44N</sup> | Kwun et al. (2009) | N/A |

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**REAGENT or RESOURCE** | **SOURCE** | **IDENTIFIER** 
--- | --- | --- 
`sTco (pcDNA6 MCV sTco)` | Shuda et al. (2011) | Addgene Cat #40201 RRID: Addgene_40201 
`LT339 (pcDNA6 MCV LT339 VS.4)` | Shuda et al. (2008) | Addgene Cat#28193 RRID: Addgene_28193 
`LT339Δ` | This paper | N/A 
`LT339E214K` | This paper | N/A 
`His-HSC70 (pcDNA6/FRT/TO HIS HSPA8)` | Hageman and Kampinga (2009) | Addgene Cat#19541 RRID: Addgene_19541 
`shTA (shRNA targeting LT and sT)` | Xie et al. (2014) | N/A 
`shsT (shRNA targeting sT)` | Xie et al. (2014) | N/A 
`DICER1 3′UTR (pIS1 DICER1 long UTR)` | Mayr and Bartel (2009) | Addgene Cat#21649 RRID: Addgene_21649 
`DICER1 3′UTR MutΔ` | This paper | N/A 
`DICER1 3′UTR MutΔ` | This paper | N/A 
`AREMut` | This paper | N/A 
`AREWT` | This paper | N/A 
`DICER1 CDSΔ` | Paulsson et al. (2020) | N/A 
`DICER1 CDSΔ` | This paper | N/A 

**Software and algorithms**
- R version 4.0.5 [https://www.R-project.org/](https://www.R-project.org/) N/A RRID:SCR_001905
- Rstudio [http://www.rstudio.com/](http://www.rstudio.com/) N/A RRID:SCR_000432
- Rstatix [https://github.com/kassambara/rstatix](https://github.com/kassambara/rstatix) N/A RRID:SCR_021240
- tidyverse [https://www.tidyverse.org/](https://www.tidyverse.org/) N/A RRID:SCR_019186
- Prism GraphPad 8.0 GraphPad Software, La Jolla California USA N/A RRID:SCR_002798

**RESOURCE AVAILABILITY**

**Lead contact**
Further information and requests for reagents should be directed to the lead contact, Weng-Onn Lui (weng-onn.lui@ki.se).

**Materials availability**
Plasmids generated in this study are available upon request.

**Data and code availability**
- All source data for Figures 1C, 1F, 1G, 2B, 2D, 2F, 2H, 2I, 3A–3C, 4B–4F, 5B, 5C, 6B, 6C, S1, and SSA, as well as uncropped Western blot images were deposited in Mendeley data [https://doi.org/10.17632/r3ghfxxs3b.1](https://doi.org/10.17632/r3ghfxxs3b.1).
- There is no original code reported in this work.
- Any additional information required is available from the lead contact upon reasonable request.

**EXPERIMENTAL MODEL AND SUBJECT DETAILS**
Cell lines used in this study include human MCC cell lines [WaGa (male), MKL-1 (male), MCC13 (female), MCC14/2 (male) and MCC26 (female)], human embryonic kidney cell line HEK293 (female) and DICER1 knockout cell line NoDICE 2-20 (derived from HEK293T). All MCC cell lines were verified by short tandem
repeat profiling as previously reported (Kumar et al., 2020). Loss of DICER1 expression was verified in NoDice 2-20 cells by Western blotting. Refer to the key resources table for information on the source of each cell line and to the method details below for culture conditions.

**METHOD DETAILS**

**Cell culture**
MCC cell lines were cultured in RPMI1640 medium supplemented with 10% (MCPyV+: WaGa and MKL-1) or 15% (MCPyV-: MCC13, MCC14/2 and MCC26) fetal bovine serum (FBS). HEK293 and NoDice 2-20 cell lines were cultured in DMEM media with 1 g/L glucose (HEK293) or 4.5 g/L glucose (NoDice 2-20) supplemented with 10% FBS.

**Plasmids and siRNAs**
MCPyV codon-optimized full-length LT (LTco; Addgene, #40200), truncated LT (LT339; Addgene, #28193), HSC70 expression vector (His-HSC70 contains the full-length coding sequence of HSPA8, NM_006597; Addgene, #19541), DICER1 3’UTR luciferase reporter (DICER1 3’UTR; Addgene, #21649) and firefly luciferase reporter (Addgene, #64784) plasmids were purchased from Addgene (Cambridge, MA, USA). Codon-optimized sT (sTco) and LTcoD44N were kindly provided by Drs. Yuan Chang and Patrick Moore (University of Pittsburgh, Pennsylvania, USA). The vector control for viral T-antigen (pCTR) was previously constructed by deleting the insert from the LT339 plasmid (Xie et al., 2014). The DICER1 CDSWT contains the full-length coding sequence (CDS) of DICER1 inserted between SalI and NotI restriction sites of pCMV6entry expression vector, which had been described in our previous study (Paulsson et al., 2020). MCPyV T-antigen mutant constructs (LTcoD, LT339D and LT339 E216K) and DICER1 3’UTR mutant constructs (Mutub, MutLa and CDSaLa) were generated with the Q5 Site-Directed Mutagenesis Kit (New England Biolabs, Ipswich, MA, USA; #E0554). Primers used for mutagenesis are listed in key resources table. For artificial ARE-containing luciferase reporter constructs, synthetic oligonucleotides containing wild-type (WT) or mutated (MUT) sequences of the ARE were phosphorylated, annealed and cloned into DICER1 3’UTR luciferase reporter backbone after digestion with SpeI and NotI (New England Biolabs, Ipswich, MA, USA). All mutant and ARE-containing vectors were confirmed by Sanger sequencing at the KIGene core facility.

Short hairpin RNA (shRNA) plasmids targeting either both MCPyV LT and sT (shTA), or only sT (shsT) were previously generated in our laboratory (Xie et al., 2014). Small interfering RNAs (siRNAs) silencing HSPA8/HSC70 (siHSC70#1, S302661477; siHSC70#3, S104236596), AUF1 (siAUF1#1, S100300454) and siRNA control (siCTR, #1027310) were purchased from Qiagen (Hilden, Germany). siAUF1#2, was designed in (Raineri et al., 2004) and oligos were synthesized by Integrated DNA Technologies (Leuven, Belgium). The oligos were phosphorylated using T4 polynucleotide kinase (New England Biolabs) and annealed in 1X annealing buffer (50 mM Tris, pH 7.5–8.0; 100 mM NaCl). All oligonucleotides used for plasmid constructions and siRNAs are detailed in Key resources table.

**Transfection**
MCPyV+ cell lines (WaGa and MKL-1) were transfected using Nucleofector electroporation system. Briefly, 3 million cells were pelleted and re-suspended in 100 µL Ingenio electroporation solution (MIR50111; Mirus Bio LLC, Madison, WI, USA) containing 2 µg plasmid DNA or 30 pmol siRNA. The mixture was transferred to a cuvette and electroporated using program D-24 (WaGa) or A-23 (MKL-1) of the Nucleofector™ 2b Device (Lonza, Basel, Switzerland). MCPyV- cell lines (MCC14/2, MCC26 and MCC13) were transfected with 2–10 µg plasmid DNA or 20 pmol siRNA using Lipofectamine LTX with PLUS (Thermo Fisher Scientific, #15338100). Similar procedures were applied to HEK293 and NoDice 2-20 cell lines, where Lipofectamine 3000 transfection reagent (Thermo Fisher Scientific, #L3000150) was used.

**Western blot analysis**
Whole cell lysates were prepared using RIPA buffer (Thermo Fisher Scientific, #89900) supplemented with protease inhibitor (Sigma-Aldrich, #P2714) and 1% phenylmethylsulfonyl fluoride (Sigma-Aldrich, #93482). Protein concentrations were assessed using Pierce BCA Protein Assay Kit (Thermo Fisher Scientific, #23225). Protein lysates (30 µg) were separated in 4–12% NuPAGE Bis-Tris gels, and transferred to nitrocellulose membranes (Thermo Fisher Scientific, #LC2000). After blocking with 5% skim milk, the membranes were incubated with primary antibodies at 4°C overnight. After washing, the membranes were incubated with HRP- or fluorescence-conjugated antibodies for 1 h at room temperature. For HRP-conjugated
secondary antibodies, detection was performed using SuperSignal West Femto Maximum Sensitivity Substrate (Thermo Fisher Scientific, #34095). Images were acquired using Odyssey Fc Image system (LI-COR Biosciences, Lincoln, NE, USA) and analyzed in Image Studio Lite (LI-COR Biosciences, Lincoln, NE, USA). Antibodies used for Western blotting are detailed in key resources table.

RNA stability analysis
Actinomycin D (2 μg/mL, Sigma-Aldrich, #A1410) was added to the cells after 48 h of transfection, and cells were harvested at indicated time points after treatment. Cells were washed with cold PBS twice, and total RNA was isolated using Trizol reagent (Thermo Fisher Scientific, #15596018). RNA concentrations were measured using a Nanodrop Spectrophotometer (Thermo Fisher Scientific). Complementary DNA (cDNA) was synthesized with MultiScribe Reverse Transcriptase (Thermo Fisher Scientific, #4311235). TaqMan gene expression assays for DICER1 (Hs00229023_m1), GAPDH (Hs99999905_m1), BCL2L11 (Hs01076940_m1) and MYC (Hs00153408_m1) were purchased from Thermo Fisher Scientific (Waltham, MA). Relative mRNA expressions were first normalized to GAPDH using 2^ΔΔCT, and then compared to time point 0.

Immunoprecipitation (IP) and RNA-IP (RIP)
To prepare antibody-beads conjugation, 150 μL beads slurry (Protein G Sepharose 4 Fast Flow beads, GE17-0618-01, Sigma-Aldrich) was washed three times with 1 mL NT2 buffer (50 mM Tris-HCl, pH 7.4; 150 mM NaCl; 1 mM MgCl2; 0.5% NP-40) supplemented with 8 U/mL RNase inhibitor (RNaseOUT; Thermo Fisher Scientific; #10777019), followed by incubation with anti-His (Santa Cruz Biotechnology, sc-8036, 1:50 dilution), anti-HSC70 (Santa Cruz Biotechnology, sc-8036, 1:50 dilution), anti-HSC70 (Novus Biologicals, NB2-67335, 1:50 dilution), anti-HSC70 (Abcam, ab51052, 1:100 dilution) or anti-LT (Santa Cruz Biotechnology, sc-136172, 1:50 dilution) for 5–6 hours at 4°C. Unbound antibodies were washed away in NT2 buffer.

Forty-eight hours after transfection, cells were washed with cold PBS twice and lysed in NP40 buffer (Thermo Fisher Scientific, FNN0021), supplemented with 200 U/mL RNase inhibitor, 1% phenylmethylsulfonyl fluoride (Sigma-Aldrich, #93482), 1 mM Dithiothreitol (Thermo Fisher Scientific, P2325) and proteinase inhibitor (Sigma-Aldrich, P8340).

To pull down RNAs bound to target proteins, 3 mg of whole cell lysate was added to the antibody-beads slurry, supplemented with EDTA (1 M), RNase inhibitor (200 U/mL) and protease inhibitor. After incubation overnight at 4°C on a rotator, sepharose beads were pelleted and washed 4 × 10 minutes with 1 mL cold NT2 buffer, supplemented with RNase inhibitor 8 U/mL. For Western blot analysis, beads were eluted with LDS loading buffer (Thermo Fisher Scientific, NP0008) and Sample Reducing Agent (Thermo Fisher Scientific, NP0009), and the lysates were denatured and then subjected to 4-12% Bis-Tris gel electrophoresis. For RNA analysis, proteinase K (1 mg/mL) was added to beads slurry, and then incubated at 55°C for 30 minutes. Total RNA was isolated with Trizol Reagent and RT-qPCR was performed to quantify the mRNA levels of DICER1 and GAPDH. Enrichment of mRNA was calculated by dividing the amount of mRNAs in the IP samples to their input samples.

In vitro transcription and biotin pull-down assay
To prepare DNA templates for in vitro transcription (IVT), DICER1 fragments were amplified using Platinum Taq DNA Polymerase (Thermo Fisher Scientific, #10966018) with primers detailed in key resources table. DICER1 CDS and DICER1 3’UTR plasmids were used as templates for amplifying coding sequence and 3’UTR fragments, respectively. Total cDNA of WaGa cells was used as template for amplifying the 5’UTR of DICER1. PCR products were purified with the Monarch PCR & DNA Cleanup Kit (New England Biolabs, #T1030), and were then used for IVT with the HiScribe T7 High Yield RNA Synthesis Kit (New England Biolabs, #E2040S). Probes were labeled with biotinylated UTP (Bio-11-UTP, Thermo Fisher Scientific, #AM8450), cleaned by MEGAclear Transcription Clean-Up Kit (Thermo Fisher Scientific, #AM1908) and quantified using a Nanodrop Spectrophotometer (Thermo Fisher Scientific).

For pull-down assay, 20 pmol of biotinylated RNA probe was incubated with 300 μg of whole cell lysate for 5–6 hours at 4°C. Protein complexes were isolated with 40 μL of Dynabeads M-280 Streptavidin (Thermo Fisher Scientific, #11205D). After washing twice with 300 μL washing buffer (5 mM Tris-HCl, pH 7.5;
0.5 mM EDTA; 1 M NaCl) for 2 min, the pull-down mixture was analyzed by Western blotting. A biotinylated RNA probe from firefly luciferase gene (FlucCTR) was used as a control for non-specific binding.

**Luciferase reporter assay**

One µg of plasmid DNA encoding Renilla luciferase with various 3’UTR constructs (DICER1 3’UTR WT, DICER1 3’UTR Mutsub, DICER1 3’UTR Mutsub AREWT or AREMUT) together with 1 µg of firefly luciferase reporter plasmid and 1 µg of T-antigen plasmid (sTco, LT339, LTco, LTcoD44N or pCTR) or 20 pmol siHSC70#1/siCTR were co-transfected into 2x10⁵ HEK293 cells using Lipofactamine 3000. Luciferase activity was assessed using Dual-Glo Luciferase Assay (Promega Biotech, #E2920) and luminescence was detected by Spark 10M multimode microplate reader (Tecan Group Ltd, Männedorf, Switzerland). All experiments were performed in triplicates for each condition and repeated at least three times independently. The Renilla luciferase activity was divided by the firefly luciferase activity in each condition and normalized to the pCTR or siCTR.

**Subcellular fractionation**

MCPyV+ cells, WaGa and MKL-1 (~1x10⁷), were collected and washed with cold PBS twice. 200 µL subcellular fractionation buffer (20 mM HEPES, pH 7.4; 10 mM KCl; 2mM MgCl₂; 1mM EDTA; 2mM DTT; supplemented with protease inhibitor cocktail) were added to cell pellet and ground 10 times using a pestle. After 20-minute incubation on ice, the lysate was centrifuged at 700 g for 5 min and the supernatant was transferred to a new tube as cytoplasmic fraction. The remaining nuclear pellet was washed with subcellular fractionation buffer twice, and then resuspended in 200 µL RIPA buffer. After 20-minute incubation on ice, the nuclei were disrupted by sonication 3 x 15s in ice-cold water, followed by centrifugation at 16,000 g for 30 minutes to collect the supernatant as nuclear fraction. MCPyV-MCC26 cells were collected after 48 hours of transfection and fractionated using Qproteome Cell Compartment Kit (Qiagen, Cat#37502). Protein concentration was assayed using Pierce BCA Protein Assay Kit (Thermo Fisher Scientific, #23225), and ~50 µg of total protein was loaded for Western blot analysis. α-TUBULIN or GAPDH was used as a cytoplasm marker, while HISTONE3 or Lamin A/C was used as a nuclear marker.

**QUANTIFICATION AND STATISTICAL ANALYSIS**

Statistical tests were performed in R (version 4.0.5, Windows 10) using the package “rstatix”, unless specified otherwise. Two-tailed paired t-test was used to compare two groups in transfection assays. p-values for multiple tests were corrected with “Benjamini–Hochberg” method. RNA stability assay was analyzed in GraphPad Prism 8 (GraphPad Software, Inc, CA, USA), and one-phase decay model was applied to determine the half-life of mRNAs (t1/2). p-values for mRNA stability analysis were calculated by two-way ANOVA. Statistical significance was defined as a p < 0.05 and p values were indicated by asterisks as *p < 0.05, **p < 0.01, ***p < 0.001 and ns, not significant. Error bars stand for standard error of the mean (SEM), and numbers of observations or replicates (n) are indicated in figure legends.