SPIN1 promotes tumorigenesis by blocking the uL18 (universal large ribosomal subunit protein 18)-MDM2-p53 pathway in human cancer

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Abstract

Ribosomal proteins (RPs) play important roles in modulating the MDM2-p53 pathway. However, less is known about the upstream regulators of the RPs. Here we identify SPIN1 (Spindlin 1) as a novel binding partner of human RPL5/uL18 that is important for this pathway. SPIN1 ablation activates p53, suppresses cell growth, reduces clonogenic ability, and induces apoptosis of human cancer cells. Mechanistically, SPIN1 sequesters uL18 in the nucleolus, preventing it from interacting with MDM2, and thereby alleviating uL18-mediated inhibition of MDM2 ubiquitin ligase activity towards p53. SPIN1 deficiency increases ribosome-free uL18 and uL5 (human RPL11), which are required for SPIN1 depletion-induced p53 activation. Analysis of cancer genomic databases suggests that SPIN1 is highly expressed in several human cancers, and its overexpression is positively correlated with poor prognosis in cancer patients. Altogether, our findings reveal that the oncogenic property of SPIN1 may be attributed to its negative regulation of uL18, leading to p53 inactivation.

Key words: SPIN1; uL18 (human RPL5); MDM2-p53 pathway; cell proliferation; apoptosis; tumorigenesis.
Introduction

The well-documented tumor suppressor p53, referred as “the guardian of the genome”, is activated upon exposure to a myriad of cellular stresses. While loss of wild type p53 causes fatal damages to the genome, it is not surprising that the TP53 gene is mutated in more than 50% human cancers, and the functions of p53 are often impeded through various mechanisms in the remainder. One predominant negative regulator of p53 is the E3 ubiquitin ligase MDM2, an oncoprotein that conceals the N-terminal transcriptional activation (TA) domain of p53 and deactivates this protein by either abrogating its transcriptional activity, or inducing its nuclear export and ubiquitination. A plethora of cellular stress could stabilize p53 by blocking the MDM2-p53 feedback loop. For example, p19ARF inhibits MDM2-mediated p53 ubiquitination and proteolysis by associating with MDM2. Another pathway is the so-called ribosomal proteins (RPs)-MDM2-p53 pathway. Accumulating evidence has continuously verified this pathway as an emerging mechanism for boosting p53 activation in response to ribosomal stress or nucleolar stress over the past decade. Ribosomal stress is often triggered by aberrant ribosome biogenesis caused by nutrient deprivation, inhibition of rRNA synthesis, or malfunction of ribosomal proteins and/or nucleolar proteins. Earlier studies showed that disruption of ribosomal biogenesis induces translocation of a series of ribosomal proteins, including uL18 (human RPL5), uL5 (human RPL11), uL14 (human RPL23), eS7 (human S7) and uS11 (human S14) from the nucleolus to the nucleoplasm and bind to MDM2, blocking its ability to ubiquitinate p53 and consequently stabilizing p53 to maintain cellular homeostasis. Although there are a few proteins that have been identified to regulate this RPs-MDM2-p53 pathway, such as PICT-1 inhibition of uL5 and SRSF1 activation of uL18, it still remains to determine if there are more proteins that can regulate the RPs-MDM2-p53 pathway. In this present study, we identified SPIN1 as a new uL18 inhibitory regulator.

SPIN1, a new member of the SPIN/SSTY family, was originally identified as a highly expressed protein in ovarian cancer. The oncogenic potential of SPIN1 was later supported by the observation that overexpression of SPIN1 increases transformation and tumor growth ability of NIH3T3 cells. Signaling pathways responsible for SPIN1 functions include PI3K/Akt, Wnt and RET that are highly relevant to tumorigenesis. In addition, SPIN1 acts as a reader of
histone H3K4me3 and stimulates the transcription of ribosomal RNA-encoding genes \(^{32-34}\), suggesting its role in rRNA synthesis.

In screening uL18-associated protein complexes using co-immunoprecipitation followed by mass spectrometry, we identified SPIN1 as one of the potential uL18 binding proteins. We confirmed the specific interaction of SPIN1 with uL18, but not with uL5 or uL14, and also found out that by binding to uL18, SPIN1 prevents the inhibition of MDM2 by uL18 and promotes MDM2-mediated p53 ubiquitination and degradation. Also, SPIN1 knockdown induced ribosomal stress by facilitating the release of ribosome-free uL18 or uL5, accompanying p53 activation. Furthermore, SPIN1 knockdown inhibited cell proliferation and induced apoptosis in a predominantly p53-dependent manner in vitro and in vivo, consequently suppressing tumor growth in a xenograft model. Therefore, these results for the first time demonstrate that SPIN1 can regulate the RP-MDM2-p53 pathway by directly interacting with uL18, and suggest SPIN1 as a potential molecule target in this pathway for developing anti-cancer therapy in the future.

RESULTS

SPIN1 interacts with uL18.

Our and others’ studies previously demonstrated that uL18 can stabilize p53 by binding to MDM2 and inhibiting its E3 ligase activity toward p53 \(^{19,35}\). In order to identify potential upstream regulators that may modulate the uL18-MDM2-p53 circuit, we performed co-immunoprecipitation (co-IP) using HEK293 cells that stably expressed Flag-uL18 with the anti-Flag antibody, and the co-immunoprecipitated proteins were cut out for mass spectrometric (MS) analysis (Fig. 1A). The MS results not only revealed several previously described p53 regulatory proteins, such as MYBBP1A, PRMT5 and SRSF1, as uL18 binding proteins (Table 1), but also identified SPIN1 as a novel uL18-binding protein candidate that was previously shown to play a role in tumorigenesis and rDNA transcription \(^{30,34}\).

Next, we confirmed the interaction between SPIN1 and uL18 by performing a series of reciprocal co-IP assays. As expected, ectopic SPIN1 was specifically pulled down by ectopic uL18 and vice versa in HCT116\(^{p53-/-}\) cells (Fig. 1B and 1C). Their interaction was also verified in HEK293 cells (Figure 1-figure supplement 1). Also, we validated the interaction between endogenous SPIN1 and uL18 in HEK293 cells using anti-SPIN1 antibody (Fig. 1D). Interestingly, only uL18, but not uL5, was co-immunoprecipitated with SPIN1. In line with this
result, when comparing ectopic Flag-uL18 with Flag-uL5 and Flag-uL14, we found that only uL18, but not the other RPs, could pull down Myc-SPIN1 (Fig. 1E), further bolstering the specific interaction between uL18 and SPIN1. Taken together, these results demonstrate that SPIN1 specifically binds to uL18, but not uL5 or uL14, in cancer cells.

**SPIN1 knockdown inhibits proliferation and induces apoptosis of cancer cells by activating p53.**

Previous and recent studies showed that SPIN1 is a potential oncogene \(^{29,30,36}\), and uL18 can stabilize p53 by binding to MDM2 \(^{19}\). We therefore wondered if the interaction between SPIN1 and uL18 could confer any role to SPIN1 in regulation of the p53 pathway. First, we determined if depletion of SPIN1 might affect p53-dependent cellular outcomes. Interestingly, we found that knockdown of SPIN1 dramatically elevates p53 protein level in several wild type p53-containing cells, including U2OS, H460 and HCT116\(^{p53+/+}\) cells (Fig. 2A), without affecting TP53 mRNA expression (Fig. 2B). Consistently, protein and mRNA levels of p53 target genes, such as p21 and PUMA, were also increased in response to SPIN1 knockdown (Fig. 2A and 2B). Moreover, the effects of SPIN1 siRNA on p53 activity were dose-dependent (Figure 2-figure supplement 1A & 1B). The effect was unlikely due to off-target effects of siRNA, as ectopic expression of FLAG-SPIN1 reversed p53 activation by siRNA-mediated knockdown of SPIN1 in HCT116\(^{p53+/+}\) (Figure 2-figure supplement 1C). Conversely, overexpression of SPIN1 in HCT116\(^{p53+/+}\) decreased the protein levels of p53 and its targets, such as p21 and PUMA, and the mRNA levels of these target genes, without affecting TP53 mRNA level (Fig. 2C and 2D).

We next generated both HCT116\(^{p53+/+}\) and HCT116\(^{p53-/-}\) cell lines that express scramble shRNA or SPIN1 shRNA to evaluate biological outcomes of SPIN1 knockdown. As illustrated in Fig. 2E, the expression of p53 and some of its target genes were markedly induced when SPIN1 was knocked down by its specific shRNA in HCT116\(^{p53+/+}\) cells, but not in HCT116\(^{p53-/-}\) cells. Using these cell lines for cell viability assays, we observed that SPIN1 ablation more dramatically represses the cell viability of HCT116\(^{p53+/+}\) than that of HCT116\(^{p53-/-}\) cells (Fig. 2F). In line with this observation, SPIN1 depletion also led to more predominant reduction of HCT116\(^{p53+/+}\) colonies than that of HCT116\(^{p53-/-}\) colonies, though both of the reductions were statistically significant (Fig. 2G). Furthermore, the percentage of cells undergoing apoptosis caused by SPIN1 shRNAs was much higher in HCT116\(^{p53+/+}\) cells than in HCT116\(^{p53-/-}\) cells, as measured
by sub-G1 population (Fig. 2H). Consistently, induction of apoptosis by SPIN1 knockdown was also evidenced in Annexin V assay in U2OS cells (Fig. 2I). Collectively, these data suggest that SPIN1 plays an oncogenic role at least partially by inactivating the p53 pathway, although SPIN1 may also possess a p53-independent role in cancer cell growth and survival.

**SPIN1 promotes p53 degradation by enhancing MDM2-mediated ubiquitination.**

Since SPIN1 knockdown affected only the protein, but not the mRNA, levels of p53 (Fig. 2A-2D), we next sought to determine the underlying mechanism. We first performed a cycloheximide-chase experiment using HCT116<sup>p53+/+</sup> cells. As shown in Figures 3A and 3B, knockdown of SPIN1 markedly prolonged p53’s half-life from 35 mins to 56 mins, as compared to scramble siRNA. Inversely, ectopic SPIN1 greatly shortened p53’s half-life, from 39 mins to ~22 mins (Fig. 3C and 3D). To further evaluate the effect of SPIN1 on MDM2-mediated p53 ubiquitination, which is the main mechanism responsible for p53 turnover<sup>12,19,22,37</sup>, we then performed an in vivo ubiquitination assay by transfecting HCT116<sup>p53-/-</sup> cells with plasmids indicated in Figure 3E. The results clearly showed that ectopic SPIN1 enhances MDM2-mediated p53 ubiquitination in a dose-dependent manner. Consistently, co-transfection of SPIN1 with MDM2 led to a stronger reduction of p53 protein levels, which was abrogated by proteasome inhibitor MG132 (Fig. 3F). Interestingly, the induction of p53 degradation by SPIN1 was MDM2-dependent, as overexpression of SPIN1 failed to repress ectopic p53 protein expression in <i>Trp53</i> and <i>MDM2</i> double knockout MEF cells (Fig. 3G). Together, these results demonstrate that SPIN1 reduces p53 stability by enhancing MDM2-mediated ubiquitination and degradation.

**SPIN1 prevents uL18 from MDM2 binding by sequestering it in the nucleolus**

Besides its role as a component of ribosome, uL18 has some well-established extra-ribosomal functions, acting as a bridge in connecting p53 activation to cellular stress response machinery<sup>8,9</sup>. Upon ribosomal stress, uL18 can translocate from the nucleolus to the nucleoplasm of a cell, where it binds to MDM2<sup>19,38</sup>, leading to stabilization of p53 and consequently p53-dependent cell growth arrest, apoptosis or senescence. We then investigated if SPIN1 might regulate this function of uL18, since SPIN1 could bind to uL18 (Figure 1), knockdown of SPIN1 led to p53 activation (Fig. 2), and SPIN1 stimulated MDM2-mediated p53 ubiquitination (Fig. 3). First, as
overexpression of uL18 induced the protein levels of p53 and its targets, such as p21 and MDM2, in wild type p53-containing U2OS cells (Fig. 4A). This induction of the p53 pathway by uL18 was markedly reduced by co-transfected SPIN1 (Fig. 4A). Since the effect of uL18 on p53 is through uL18’s interaction with MDM2 and consequent inhibition of its E3 ligase activity towards p53\(^9\), we tested if SPIN1 may affect uL18-MDM2 interaction. Interestingly, our co-immunoprecipitation result showed that ectopic Myc-SPIN1 dramatically reduces the amount of Flag-uL18 co-immunoprecipitated with HA-MDM2 in a dose-dependent manner, although Myc-SPIN1 itself did not co-immunoprecipitate with HA-MDM2 (Fig. 4B and Figure 4-figure supplement 1A). This effect was specific to the uL18-MDM2 interaction, as Myc-SPIN1 overexpression did not alter the interactions between uL5 and MDM2 (Fig. 4C). Our immunofluorescence result (Fig. 4D and Figure 4-figure supplement 1B) showed that SPIN1 and uL18 are clearly co-localized in the nucleolus, suggesting that SPIN1 might sequester uL18 in the nucleolus and thus prevent it from binding and inactivating MDM2 in the nucleoplasm. Taken together, these results demonstrate that SPIN1 is a regulator of the uL18-MDM2-p53 pathway, acting by preventing uL18 from interaction with MDM2.

**SPIN1 depletion also causes ribosomal stress, activating p53.**

Previous studies showed that SPIN1 could recognize H3K4 methylation and stimulate rRNA gene expression, unveiling its role in rRNA synthesis\(^{32,34}\). Disruption of rRNA synthesis leads to disassembly of ribosomal precursors and release of ribosome-free ribosomal proteins from the nucleolus\(^{16,19,22}\). Based on these lines of information, we speculated that dysregulation of SPIN1 itself might also impact ribosome biogenesis, resulting in accumulating ribosome-free ribosomal proteins to activate p53. To test this speculation, we first carried out a sucrose gradient fractionation assay using scramble- and SPIN1-shRNA transfected HCT116\(^{p53++}\) cells. The collected fractions were subjected to Western blot (WB) analysis. As anticipated, the levels of uL18 and uL5 in the soluble and ribosome-unbound fractions were markedly increased in SPIN1-depletion cells, accompanying with elevated p53 and MDM2 protein levels (Fig. 5A). Interestingly, the binding between endogenous uL18/uL5 and MDM2 increased upon SPIN1 knockdown, resembling ribosomal stress (Fig. 5B). Indeed, as expected, knockdown of SPIN1 reduced the expression of pre-rRNA and rRNA (Figure 5-figure supplement 1A).
Moreover, as clearly illustrated in Figure 5C, overexpression of SPIN1 compromised p53 activation induced by actinomycin D or 5-Fu treatment, which was reported to trigger ribosomal stress that in turn triggers the formation of RPs-MDM2 complex\textsuperscript{10,19,23,39}. In addition, the Y170A mutant of SPIN1, which loses the ability to interact with trimethylated K4\textsuperscript{33,34}, was still able to suppress p53 activity (Figure 5-figure supplement 1B), suggesting that SPIN1 regulation of p53 is independent of the activity of SPIN1 in regulating rRNA expression. Interestingly, our mapping results showed that the critical amino acids Y170, F141 and Y177 for trimethylated K4 interaction\textsuperscript{33,34} are all located in the SPIN1 Tudor 2 domain that is responsible for uL18 binding (Figure 5-figure supplement 2A & 2B). Tudor 2 domain truncated mutant of SPIN1 failed to suppress p53 or increase rRNA expression (data not shown). Interestingly, both the N- and C-termini of uL18 were found to bind to SPIN1 (Figure 5-figure supplement 2C and 2D), and these two fragments were required for uL18-MDM2 binding as well (Figure 5-figure supplement 2E & 2F), further supporting our observation that SPIN1 could compete with MDM2 for uL18 binding (Fig. 4B).

To further confirm the role of these free forms of ribosomal proteins in SPIN1 ablation-induced p53 activation, we knocked down uL18 or uL5 using siRNA with or without SPIN1 depletion in U2OS cells. Strikingly, the reduction of either uL18 or uL5 abrogated SPIN1 knockdown-induced p53 levels, as well as its target p21, as compared to scramble siRNA-transfected cells (Fig. 5D and 5E). Collectively, these data indicate that knockdown of SPIN1 could also lead to ribosomal stress, releasing ribosome-free uL18 and uL5, which are required for p53 activation induced by SPIN1 depletion.

**SPIN1 depletion impedes xenograft tumor growth.**

To translate the above-described cellular functions of SPIN1 into more biological significance, we established a xenograft tumor model by inoculating the aforementioned HCT116 (both p53\textsuperscript{+/+} and p53\textsuperscript{-/-}) cell lines that expressed scramble shRNA or SPIN1 shRNA into NOD/SCID mice, and monitored tumor size for 18 days. As illustrated in Fig. 6A and 6B, SPIN1 knockdown more markedly slowed down the growth of xenograft tumors derived from HCT116\textsuperscript{p53+/+} cells than that from HCT116\textsuperscript{p53-/} cells. Notably, SPIN1 depletion also reduced the growth of tumors derived from HCT116\textsuperscript{p53-/} cells, suggesting that SPIN1 might possess a p53-independent function required for cancer cell growth. In line with the tumor growth curve, the reduction of
tumor mass and weight by SPIN1 knockdown was more profound in HCT116\(^{p53+/+}\) groups (~60% reduction in weight) than that in HCT116\(^{p53-/-}\) groups (~30% reduction in weight) (Fig. 6C and 6D). To confirm our cell-based findings, we performed qRT-PCR and WB analysis using the xenograft tumors. As expected, the mRNA levels of p21 and PUMA were significantly increased upon SPIN1 knockdown in HCT116\(^{p53+/+}\), but not in HCT116\(^{p53-/-}\) tumors (Fig. 6E and 6F). Consistently, the protein levels of p53 and its target PUMA were elevated in HCT116\(^{p53+/+}\) groups, but not in HCT116\(^{p53-/-}\) groups (Fig. 6G and Figure 6-figure supplement 1). Taken together, these results demonstrate that SPIN1 depletion retards tumor growth by mainly activating p53, though SPIN1 might also possess p53-independent functions in regulation of cell growth and survival.

The data presented above suggest that SPIN1 plays an important role in tumorigenesis. Therefore, we further searched some available genomic and gene expression database for SPIN1 expression in cancers. Interestingly, our analysis of TCGA genome database \(^{40,41}\) indicated that the SPIN1 gene is markedly amplified in a panel of cancers, including prostate, sarcoma, lung, stomach, breast, head and neck, pancreas and colorectal cancers (Figure 6-figure supplement 2A). Consistent with this observation, the analysis of Oncomine database \(^{42}\) also showed that SPIN1 mRNA expression is extensively upregulated in melanoma tissues when compared with normal skin tissues (~2.367 folds upregulation, Figure 6-figure supplement 2B). Moreover, using databases \(^{40,43,44}\) that contain gene expression profiles of clinical cancer samples combined with patient outcomes, we found that overexpression of SPIN1 is correlated with poorer prognosis in patients with breast cancer, colorectal cancer and gastric cancer (Figure 6-figure supplement 2C-2F). Elevated protein expression was also observed in a panel of human colon tumor samples compared with normal tissues (Figure 6-figure supplement 3). These data further support that SPIN1 may play an oncogenic role in human cancer progression.

Based on these data and the aforementioned results, we proposed a model for the role of SPIN1 in regulation of p53 (Fig. 7). Under the condition of low SPIN1 level, nucleolar uL18 escapes from the nucleolus into the nucleoplasm, and works together with uL5 to bind MDM2 and to inhibit its E3 ubiquitin ligase activity towards p53, consequently leading to p53 activation and p53-dependent cell growth arrest and apoptosis, suppressing cancer cell survival (Fig. 7A). But when SPIN1 levels are high or abnormally elevated in cancer cells, SPIN1 retains uL18 in the nucleolus, thereby preventing uL18 from suppression of MDM2 activity and resulting in p53
degradation, favoring tumor cell growth (Fig. 7B). This conjecture is further supported by the aforementioned xenograft experiment.

**Discussion**

The tumor suppressor p53 provides a critical brake on cancer development in response to ribosomal stress, as impairing this ribosomal stress-uL18/uL5-p53 pathway could accelerate tumorigenesis in c-Myc transgenic lymphoma mice. However, it remains largely elusive whether this pathway is subjected to the regulation by other yet unknown proteins. In our attempt to understand molecular insights into this possible regulation, we identified SPIN1, the nucleolar protein important for rRNA synthesis, as a novel regulator of the uL18-MDM2-p53 pathway through interplay with uL18 (Fig. 6E). Our studies as presented here provide the first line of evidence for that SPIN1 acts as an upstream regulator of uL18’s accessibility to MDM2 for p53 activation.

Using IP-MS analysis, we identified SPIN1 as a new uL18-associated protein (Fig. 1A). Our biochemical and cellular experiments using co-IP and GST pull down assays further validated the direct association of SPIN1 with uL18 (Fig. 1B-1D; Figure 5-figure supplement 2A-2D). Moreover, we found that SPIN1 and uL18 co-localized in the nucleolus by immunofluorescence assay (Fig. 4D and Figure 4-figure supplement 1B). Remarkably, SPIN1 specifically binds to uL18, but not uL5 and uL14, as no binding was detected between ectopic SPIN1 and uL5 or uL14 by co-IP (Fig. 1E). Interestingly, SPIN1 does not appear to bind to MDM2, as it was not co-immunoprecipitated with MDM2 either (Fig. 4B). Although our previous reports described a complex of uL18/uL5/uL14-MDM2, our present findings indicate that SPIN1 may work with uL18 in a separate complex that is different from reported RPs-MDM2 complexes. Also our results suggest that SPIN1 may retain uL18 in the nucleolus so that the latter is unable to shuttle to the nucleoplasm and to inhibit MDM2 activity towards p53 (Fig. 7).

SPIN1 expression in cells is tightly controlled, as several studies have shown that SPIN1 expression could be negatively monitored by some tumor suppressive non-coding RNAs, such as miR-489 and miR-219-5p. Moreover, elevated expression of SPIN1 was strongly correlated with advanced histological stage, chemoresistance and metastasis in patients with breast cancer. Consistent with the aforementioned oncogenic role of SPIN1, our study as presented here showed that SPIN1 depletion by its specific shRNA leads to the augment of the p53-dependent cancer cell growth arrest and apoptosis. This is at least partly because SPIN1 can...
promote MDM2-dependent ubiquitination and degradation of p53 (Fig. 3), which is highly likely attributable to its capability to prevent uL18 from binding to MDM2 through retaining uL18 in the nucleolus (Fig. 4).

Also, knockdown of SPIN1 led to the increase of ribosome-free uL18 and uL5 levels, of the uL18/uL5-MDM2 complex, and of p53 level and activity (Fig. 5A and 5B). The activation of p53 by knocking down SPIN1 is due to the ribosomal stress caused by the depletion of this nucleolar protein, as SPIN1 is required for rRNA synthesis by RNA polymerase I. Also, consistent with these observations, overexpression of SPIN1 reduced the activation of p53 by Actinomycin D treatment (Fig. 5C), whereas knockdown of uL18 or of uL5 impaired the activation of p53 by SPIN1 knockdown (Fig. 5D and 5E). Several genes have been implicated to modulate the RPs-MDM2-p53 pathway through interplay with ribosomal proteins. In particular, SRSF1 was identified as a component of the RP-MDM2-p53 complex, and could stabilize p53 via uL18. Different from their studies, SPIN1 specifically forms an independent complex with uL18, but not MDM2 or other ribosomal proteins, such as uL5 or uL14, and acts as a negative regulator of p53. Therefore, our present findings unveil a novel mechanism for suppression of the uL18-MDM2-p53 pathway by SPIN1, whose depletion consequently leads to p53-dependent cell growth inhibition and apoptosis.

Consistent with its oncogenic activity, SPIN1 is often amplified in a panel of cancer types with less or no p53 mutation based on our analysis of human samples available in TCGA database (Figure 6-figure supplement 2A-2B). In addition, elevated SPIN1 expression correlates with poor prognosis in breast, colorectal and gastric cancer patients (Figure 6-figure supplement 2C-2F). Western blot analysis of a panel of human colon samples revealed that SPIN1 is expressed at significantly higher levels in tumors than in normal tissues (Figure 6-figure supplement 3). These findings further indicate that SPIN1 acts as a potential oncogene. In line with these observations, we found that overexpression of SPIN1 promotes cancer cell survival, while knockdown of SPIN1 leads to cancer cell death as well as the suppression of cancer cell growth and colony formation predominantly in wild type p53-containing cancer cells (Fig. 2). Remarkably, knockdown of SPIN1 inhibited xenograft tumorigenesis derived from human colon cancer cells, which was much more significantly in HCT116p53+/+ cells than in HCT116p53−/− cells (Fig. 6). Consistently, bio-informatic analysis on gene expression data of 644 colorectal tumors downloaded from Genomic Data Commons showed that SPIN1
gene expression was correlated with genes (22 genes) enriched in p53 signaling pathway (Figure 6-figure supplement 4). These results demonstrate that SPIN1 can promote tumor growth and survival by inactivating p53 and its pathway (Fig. 7).

Intriguingly, we also found that SPIN1 ablation had a moderate inhibitory effect on cell growth in p53-null HCT116 cells in vitro and in vivo as mentioned above (Fig. 2 and 6). These findings suggest that SPIN1 must also possess p53-independent oncogenic effects, which might be explained by two possible mechanisms. First, SPIN1 has been reported to execute its oncogenic potentials by activating Wnt and PI3K/Akt pathways, both of which are closely correlated with cancer progression. Second, since our previous study has demonstrated that uL18 and uL5 could activate TAp73 through association with MDM2, it is possible that the SPIN1-uL18 interaction might impose suppression on TAp73 activity as well, ultimately leading to cell growth arrest and apoptosis.

Recent studies have demonstrated the role of SPIN1 in rRNA transcription, which provides a clue that dysregulation of SPIN1 may perturb ribosome biogenesis. In fact, in our current study, we observed that SPIN1 depletion per se increases the levels of ribosome-free uL18 and uL5, accompanying elevated p53 protein levels (Figure 5), which recapitulates the effects of ribosomal stress. Our observation that p53 induction caused by SPIN1 depletion could be abrogated by knockdown of either uL18 or uL5 further supports this hypothesis. Therefore, while it is conceivable that SPIN1 counteracts p53 by blocking the interaction between uL18 and MDM2 as discussed above, the mechanism by which disruption of SPIN1 causes ribosomal stress may be also responsible for p53 activation.

In summary, our findings unveil SPIN1 as another novel and important regulator of the MDM2-p53 pathway by predominantly inhibiting the association of uL18 with MDM2 to modulate p53 activity (Fig. 7) and provide more molecular insights into the fine regulation of this pathway.

Materials and Methods
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Cell culture and transient transfection.

U2OS, H1299, HEK293 and H460 cells were purchased from American Type Culture Collection (ATCC). HCT116<sup>p53+/+</sup> and HCT116<sup>p53-/-</sup> cells were generous gifts from Dr. Bert Vogelstein at the John Hopkins Medical institutes. MEF<sup>p53-/-; Mdm2-/-</sup> cells were generous gifts from Dr. Guillermina Lozano from MD Anderson Cancer Center, the University of Texas. STR profiling was performed to ensure cell identity. No mycoplasma contamination was found. All cells were cultured in Dulbecco’s modified Eagle’s medium (DMEM) supplemented with 10% fetal bovine serum, 50 U/ml penicillin and 0.1 mg/ml streptomycin and were maintained at 37°C in a 5% CO₂.
humidified atmosphere. Cells were seeded on the plate the day before transfection and then transfected with plasmids as indicated in figure legends using TurboFect transfection reagent according to the manufacturer’s protocol (Thermo Scientific, R0532). Cells were harvested at 30-48 h post-transfection for future experiments.

**Plasmids and antibodies.**

The Myc-tagged SPIN1 plasmid was generated by inserting the full-length cDNA amplified by PCR into the pcDNA3.1/Myc-His vector at EcoR I and Bam HI, using the following primers, forward-CGGAATTCatgaagaccccattcggaaag; reverse-CGGGATCCggatgttttcaccaaatcgtag. Flag-SPIN1 was generated by inserting SPIN1 cDNA into 2Flag-pcDNA3 at BamHI and XhoI sites. The primers used for PCR amplifying reverse transcribed mRNA were: forward-CGGGATCCaagaccccattcggaaagaca; reverse-CCGCTCGAGctaggatgttttcaccaaatcgtag. The GST-tagged SPIN1 fragments, His-tagged SPIN1, GFP-tagged SPIN1 and FLAG-tagged SPIN1-Y170A plasmids were generous gifts from Drs. Bing Zhu from Institute of Biophysics, Chinese Academy of Sciences, and Haitao Li from Tsinghua University, Beijing, China. The plasmids SPIN1 shRNA-1 and -2 were purchased from Sigma-Aldrich (St Louis, MO, USA). The plasmids encoding HA-MDM2, Flag-uL18, Flag-uL5, Flag-uL14, GFP-uL18, p53, His-Ub, GST-MDM2, His-uL18 and GST-uL18 fragments were described previously. Anti-Flag (Sigma-Aldrich, catalogue no. F1804, diluted 1:3000), anti-Myc (9E10, Santa Cruz Technology, catalogue no. sc-40, diluted 1:1000), anti-GFP (B-2, Santa Cruz Technology, catalogue no.sc-9996, diluted 1:1000), anti-SPIN1 (Proteintech, catalogue no. 12105-1-AP), anti-p53 (DO-1, Santa Cruz Technology, catalogue no. sc-126, diluted 1:1000), anti-p21 (CP74, Neomarkers, Fremont, catalogue no. MS-891-P0, diluted 1:1000), anti-PUMA (Proteintech, catalogue no. 55120-1-AP), anti-β-actin (C4, Santa Cruz Technology, catalogue no.sc-47778, diluted 1:5000), anti-GAPDH (Proteintech, catalogue no. 10494-1-AP), were commercially purchased. Antibodies against MDM2 (2A9 and 4B11), uL18 and uL5 were described previously.

**GST fusion protein-protein interaction assay.**

GST-tagged SPIN1 or GST-tagged uL18 fragments were expressed in *E. coli* and conjugated with glutathione-Sepharose 4B beads (Sigma-Aldrich). His-tagged SPIN1 and His-tagged uL18 were purified using a Ni-NTA (QIAGEN, Valencia, CA, USA) column, and eluted with 0.5 M
imidazole. Protein-protein interaction assays were conducted as described previously. Briefly, for Fig. 6A, 500 ng of purified His-tagged uL18 protein were incubated and gently rotated with the glutathione-Sepharose 4B beads containing 300ng of GST-SPIN1 fragments or GST only at 4°C for 4h. For Fig. 6C, 300 ng of purified His-tagged SPIN1 protein were incubated and gently shaked with the glutathione-Sepharose 4B beads containing 200ng of GST-uL18 fragments or GST only at 4°C for 1h. The mixtures were washed three times with GST lysis buffer (50 mM Tris/HCT pH 8.0, 0.5% NP-40, 1 mM EDTA, 150 mM NaCl, 10% glycerol). Bound proteins were analyzed by IB with the antibodies as indicated in the figure legends.

**Reverse transcription (RT) and quantitative RT-PCR analysis.**

Total RNA was isolated from cells or tissues using Trizol (Invitrogen, Carlsbad, CA, USA) following the manufacturer’s protocol. Total RNAs of 0.5 or 1.0 μg were used as template for reverse transcription using poly-(T)20 primers and M-MLV reverse transcriptase (Promega, Madison, WI, USA). Quantitative RT-PCR (RT-PCR) was performed using SYBR Green Mix following the manufacturer’s protocol (BioRad, Hercules, CA, USA). The primers for SPIN1, p53, p21, PUMA and GAPDH cDNA are as follows: SPIN1, 5’-CAGAGCTGATGCAGGCCAT-3’ and 5’-ACTGGGTAACAGGGCCATTG-3’, p53, 5’-CCCAAGCAATGGATGATTTGA-3’ and 5’-GCCATTCTGGGAGCTTCATCT-3’; p21, 5’-CTGGACTGTTTTCTCTCGGCTC-3’ and 5’-TGTATATTCAGCATTGTGGGAGGA-3’; PUMA, 5’-ACAGTACGACGGCCGAGACAA-3’ and 5’-GGCGGGTGCGACGCACCTAATT-3’; pre-rRNA, 5’-GCTCTACCTTACCTACCTGG-3’ and 5’-TGAGCCATTCCGAGTTTCAC-3’; 18S rRNA, 5’-GCTTCACCTGGACTCAACACGGGC-3’ and 5’-AGCTATCAATCTGTCAATCCTG-3’; rRNA, 5’-TGAGAAGACGGTCGAACTTG-3’ and 5’-TCCGGGCTCCGTTAATGATC-3’; GAPDH, 5’-GATTCCACCCATGGCAAATT-3’ and 5’-AGCATCGCCACCACCTTGATT-3’.

**Flow cytometry analysis.**

Cell transfected with scramble shRNA or SPIN1 shRNAs as indicated in the figure legends were fixed with 70% ethanol overnight and stained in 500 μl of propidium iodide (PI, Sigma-Aldrich) stain buffer (50 μg/ml PI, 200 μg/ml RNase A, 0.1% Triton X-100 in phosphate-buffered saline) at 37°C for 30 min. The cells were then analyzed for DNA content using a BD Biosciences
FACScan flow cytometer (BD Biosciences, San Jose, CA, USA). Data were analyzed using the CellQuest (BD Biosciences) and Modfit (Verity, Topsham, ME, USA) software programs.

**Annexin V assay.**
Cells transfected with scramble shRNA or SPIN1 shRNA were split into 96-well plate and IncuCyte® Annexin V Green Reagent for apoptosis was added to each well at the time of seeding. Cell apoptosis was monitored using IncuCyte® S3 live-cell imaging system.

**Cell viability assay.**
To assess the long-term cell survival, the Cell Counting Kit-8 (CCK-8) (Dojindo Molecular Technologies, Rockville, MD, USA) was used according to the manufacturer’s instructions. Cell suspensions were seeded at 2000 cells per well in 96-well culture plates at 12 h post-transfection. Cell viability was determined by adding WST-8 at a final concentration of 10% to each well, and the absorbance of these samples was measured at 450 nm using a Microplate Reader (Molecular Device, Spectrmax M5e, Sunnyvale, CA, USA) every 24 h for 5 days.

**Colony formation assay.**
Cells were trypsinized and seeded at equal number of cells on 60-mm plates. Media were changed every 4 days until the colonies were visible. Puromycin was added into the media for selection at a concentration of 2 μg/ml. Cells were fixed with methanol and stained with crystal violet solution at RT for 30 min. ImageJ was used for quantification of the colonies.

**Western blot analysis.**
Cells were harvested and lysed in lysis buffer consisting of 50 mM Tris/HCl (pH 7.5), 0.5% Nonidet P-40 (NP-40), 1 mM EDTA, 150 mM NaCl, 1mM dithiothreitol (DTT), 0.2 mM phenylmethylsulfonyl fluoride (PMSF), 10 μM pepstatin A and 1 mM leupeptin. Equal amounts of clear cell lysates (20-80 μg) were used for WB analysis as described previously. Human samples originally obtained from Indiana University Simon Cancer Center Solid Tissues Bank were ground and lysed in lysis buffer before Western blot analysis.

**In vivo ubiquitination assay.**
HCT116\(^{p53-/}$ cells were transfected with plasmids encoding p53, HA-MDM2, His-Ub or Myc-SPIN1 as indicated in the figure legends. At 48 h after transfection, cells were harvested and split into two aliquots, one for WB analysis and the other for ubiquitination assay. Briefly, cell pellets were lysed in buffer I (6 M guanidinium-HCl, 0.1 M Na\(_2\)HPO\(_4$/NaH\(_2\)PO\(_4\), 10 mM Tris-HCl (pH 8.0), 10 mM β-mercaptoethanol) and incubated with Ni-NTA beads (QIAGEN) at room temperature for 4 h. Beads were washed once with buffer I, buffer II (8 M urea, 0.1 M Na\(_2\)HPO\(_4$/NaH\(_2\)PO\(_4\), 10 mM Tris-HCl (pH 8.0), 10 mM β-mercaptoethanol), and buffer III (8 M urea, 0.1 M Na\(_2\)HPO\(_4$/NaH\(_2\)PO\(_4\), 10 mM Tris-HCl (pH 6.3), 10 mM β-mercaptoethanol). Proteins were eluted from beads in buffer IV (200 mM imidazole, 0.15 M Tris-HCl (pH 6.7), 30% glycerol, 0.72 M β-mercaptoethanol, and 5% SDS). Eluted proteins were analyzed by WB with indicated antibodies as previously reported \(^56\).

**Immunoprecipitation.**

Immunoprecipitation (IP) was conducted using antibodies as indicated in the figure legends. Briefly, ~500-1000 μg of proteins were incubated with the indicated antibody at 4 °C for 4 h or overnight. Protein A or G beads (Santa Cruz Biotechnology) were then added, and the mixture was incubated at 4°C for additional 1 to 2 h. Beads were wash at least three times with lysis buffer. Bound proteins were detected by WB analysis with antibodies as indicated in the figure legends.

**RNA interference.**

SiRNAs against SPIN1, uL18 and uL5 were commercially purchased from Ambion. SiRNAs (20-40 nm) were introduced into cells using TurboFect transfection reagent following the manufacturer’s instruction. Cells were harvested 48-72 h post-transfection for WB or RT-PCR.

**Immunofluorescence staining**

Cells were fixed in 4% paraformaldehyde (PFA) for 25 min, followed by permeabilization in 0.3% Triton X-100 for 20 min. The fixed cells were blocked with 5% bovine serum albumin for 30 min, and then the cells were incubated with indicated antibodies at 4°C overnight. Cells were then washed and incubated with the corresponding secondary antibody and 4′-6-diamidino-2-
phenylindole (DAPI) for nuclear staining. The cellular localization of SPIN1 or uL18 was examined under a confocal microscope (Nikon, ECLIPSE Ti2).

**Sucrose gradient fractionation and ribosome profiling.**

This assay was performed following the protocol previously described. Briefly, cells were harvested at 70-80% confluence after halting translation by 100 μg/ml cycloheximide incubation for 10 min. Cells were lysed in lysis buffer (10 mM Tris-HCl (pH 7.4), 5 mM MgCl₂, 100 mM KCl, 1% Triton X-100) and gently sheared with a 26-gauge needle for 4 times. Lysates were subjected to 10-50% sucrose gradient centrifugation and the fractions were collected through BR-188 Density Gradient Fractionation System (Brandel, Gaithersburg, MD, USA).

**Generating stable cell lines**

Briefly, scramble shRNA or SPIN1 shRNAs purchased from Sigma were transfected into HCT116⁰⁵³⁺/⁺ and HCT116⁰⁵³⁻/- cells using TurboFect reagent. The cells were maintained at 37°C in a 5% CO₂ humidified atmosphere for 48 h and were split to two aliquots, one for WB analysis and the other for selection using final concentration of 2 μg/ml puromycin in growth medium.

**Mouse xenograft experiments.**

Seven-week-old female NOD/SCID mice were purchased from Jackson Laboratories. Mice were randomized into two groups (6 mice in each) and subcutaneously inoculated with 5×10⁶ HCT116 cells that stably expressing scramble shRNA or SPIN1 shRNA in the right and left flanks, respectively. Tumor growth was monitored every other day with electronic digital calipers (Thermo Scientific) in two dimensions. Tumor volume was calculated with the formula: tumor volume (mm³) = (length×width²)/2. Mice were sacrificed by euthanasia, and tumors were harvested and weighed. To detect p53 activation and apoptosis in vivo, the RNAs and proteins were disrupted from tumors via homogenization in Trizol or RIPA buffer, and then subjected to RT-qPCR and WB analysis. The experiment was not blind and was handled according to approved institutional animal care and use committee (IACUC) protocol (#4275R) of Tulane University School of Medicine. The maximum tumor volume per tumor allowed the IACUC committee is 1.5 cm diameter or 300 mm³ per tumor.
TCGA data analysis

From Genomic Data Commons (https://portal.gdc.cancer.gov/), we downloaded the digital gene expression data of 644 colorectal cancer tumors, which was generated using a RNA-seq platform by the Cancer Genome Atlas (TCGA). In the data set, gene expression levels were measured with FPKM (Fragments Per Kilobase of transcript per Million mapped reads) and normalized using the Upper Quantile method.

We condensed the data by excluding the genes that were not expressed in over 75% samples. Logarithm transformation was applied to the expression levels of the remaining ~28700 Ensembl genes. The transcriptional correlations between the SPIN1 gene and the others genes were evaluated using Pearson correlation coefficient (r). The corresponding p-values were estimated by t-tests. On the cutoffs, including the absolute value of r being larger than 0.3 and Bonferroni adjusted p-value being less than 0.01, ~4500 significant genes were selected. The functional enrichment test of the selected genes was performed using the DAVID tool (https://david.ncifcrf.gov/). The heatmap was generated using R function, heatmap.2().

Statistical testing

All in vitro experiments were performed in biological triplicate and reproduced at least twice. The Student’s two-tailed t-test was used to determine mean difference among groups. P<0.05 was considered statistically significant, asterisks represent significance in the following way: *, p<0.05; **, p<0.01. The term “n.s” indicates that no significant difference was found. All the data are presented as mean ± SEM.

ACKNOWLEDGEMENTS

We thank Drs. Bert Vogelstein and Guillermina Lozano for offering HCT116 cells and MEF cells, respectively, Drs. Bing Zhu and Haitao Li for offering plasmids, the Lu lab members and Dr. Hee-Won Park for active discussion and suggestions. Hua Lu and Shelya X Zeng were supported in part by NIH-NCI grants R01CA095441, R01CA172468, R01CA127724, R21CA190775, and R21 CA201889. Kun Zhang was supported in part by NIH grant 2G12MD007595.

COMPETING INTERSTS
The authors declare that they have no conflict of interest.

REFERENCE


**Figure Legend**

**Figure 1. SPIN1 binds to uL18, but not uL5, or uL14.** (A) Identification of SPIN1 as a candidate of uL18 binding protein by immunopurification and mass spectrometric analysis. Lysates from HEK 293 cells were immunoprecipitated with the anti-Flag antibody. Bound proteins were visualized on a coomassie staining SDS-PAGE gel. Several bands were excised and subjected to mass spectrometry. One of them was identified as SPIN1 (Spindlin 1). The polypeptides identified from these bands are listed in Table 1. (B) and (C) SPIN1 interacts with uL18. (B) HCT116<sup>p53/-</sup> cells were transfected with plasmids encoding Myc-SPIN1 and Flag-uL18, and 48 h later cell lysates were collected for immunoprecipitation (IP) analysis using the anti-Flag antibody. (C) HCT116<sup>p53/-</sup> cells were transfected with plasmids encoding Flag-SPIN1 and GFP-uL18 for 48 h and harvested for WB analysis with indicated antibodies. (D) The interaction between endogenous SPIN1 and uL18. The HEK 293 cell lysates were immunoprecipitated with anti-SPIN1 or control immunoglobulin G (IgG), followed by WB analysis with anti-SPIN1, anti-uL18 and anti-uL5. (E) SPIN1 was specifically immunoprecipitated by uL18, but not uL5 or uL14. H1299 cells were co-transfected with Myc-SPIN1 and Flag-uL18, Flag-uL5 or Flag-uL14 as indicated and subjected to IP with the anti-Flag antibody, followed by WB analysis with indicated antibodies.

**Figure 2. SPIN1 knockdown inhibits cell proliferation and induces apoptosis.** (A) SPIN1 knockdown induces protein levels of p53 and its target genes. U2OS, H460 and HCT116<sup>p53+/-</sup> cells were transfected with scramble siRNA (scr-siRNA) or SPIN1 siRNA and harvested 48 h
post-transfection for WB analysis with indicated antibodies. (B) SPIN1 knockdown induces mRNA levels of p53 target genes without effect on TP53 mRNA level. U2OS cells were transfected with scramble siRNA (scr-siRNA) or SPIN1 siRNA, and harvested 72 h post-transfection for RT-qPCR (mean ± SEM, n=2). (C) SPIN1 overexpression reduces protein levels of p53 and its target genes. HCT116\(^{p53+/-}\) cells were transfected with pcDNA or Flag-SPIN1 and harvested 48 h post-transfection for WB analysis with indicated antibodies. (D) SPIN1 overexpression reduces mRNA levels of p53 target genes without effect on TP53 mRNA levels. HCT116\(^{p53+/-}\) cells were transfected with pcDNA or Flag-SPIN1 and harvested 72 h post-transfection for RT-qPCR (mean ± SEM, n=2). (E) Knockdown of SPIN1 causes p53-dependent induction of p21 and PUMA. The protein levels of p53 and its targets in HCT116\(^{p53+/-}\) cells and HCT116\(^{p53-/-}\) cells that stably express scramble shRNA (scr-shRNA) or SPIN1 shRNAs were detected by WB analysis with indicated antibodies. (F) SPIN1 knockdown suppresses cell survival. HCT116\(^{p53+/-}\) and HCT116\(^{p53-/-}\) cells that stably expressed scramble or SPIN1 shRNAs were seeded in 96-well plate and cell viability was evaluated every 24 h by CCK-8 assays (mean ± SEM, n=2). (G) Knockdown of SPIN1 inhibits clonogenic ability of colorectal cancer cells, more significantly when the cells harbor wild type p53. HCT116\(^{p53+/-}\) cells and HCT116\(^{p53-/-}\) that stably expressed scramble or SPIN1 shRNAs were seeded on 60 mm plates. Puromycin selection was performed for 14 days. Colonies were fixed with methanol, and visualized by staining with crystal violet (mean ± SEM, n=3). (H) The effect of SPIN1 knockdown on apoptosis of HCT116\(^{p53+/-}\) cells and HCT116\(^{p53-/-}\) that stably expressed scramble or SPIN1 shRNAs (mean ± SEM, n=3). (I) U2OS cells were transfected with scramble or SPIN1 shRNA and incubated in IncuCyte® S3 chamber in the presence of IncuCyte® Annexin V Green Reagent for apoptosis. Positive stained cells were determined using IncuCyte® analysis software. *\(p<0.05\), **\(p<0.01\) by two-tailed t-test (C, D, G, H, I).

**Figure 3.** SPIN1 reduces p53 stability by enhancing MDM2-mediated ubiquitination. (A) and (B) p53-half-life is increased by SPIN1 knockdown. (A) HCT116\(^{p53+/-}\) cells transfected with scramble or SPIN1 siRNA for 48 h, treated with 100 μg/ml of cycloheximide (CHX), and harvested at different time points as indicated. The p53 protein was detected by WB analysis, quantified by densitometry and plotted against time to determine p53-half-lives (B). (C) and (D) SPIN1 overexpression shortens the half-life of p53. HCT116\(^{p53+/-}\) cells transfected with pcDNA
or Flag-SPIN1 for 48 h were treated with 100 μg/ml of cycloheximide and harvested at indicated time points for WB analysis with indicated antibodies (C). The intensity of each band was quantified, and normalized with β-actin and plotted (D). (E) SPIN1 promotes MDM2-induced p53 ubiquitination. HCT116^{p53−/−} cells were transfected with combinations of plasmids encoding HA-MDM2, p53, His-Ub or Myc-SPIN1, and treated with MG132 for 6h before being harvested for in vivo ubiquitination assay. Bound and input proteins were detected by WB analysis with indicated antibodies. (F) SPIN1 enhances MDM2-mediated p53 proteasomal degradation. HCT116^{p53+/+} cells were transfected with plasmids encoding HA-MDM2 and Flag-SPIN1, and treated with MG132 for 6h before harvested, followed by WB analysis with antibodies as indicated. (G) Ectopic SPIN1 does not change p53 protein level without MDM2. MEF^{p53−/−; Mdm2−/−} cells were transfected with combinations of plasmids encoding p53 with or without Flag-SPIN1, followed by WB analysis using antibodies as indicated.

**Figure 4. SPIN1 blocks uL18-MDM2 interaction by sequestering uL18 in the nucleolus.** (A) SPIN1 overexpression attenuates p53 activation induced by ectopic uL18. U2OS cells were co-transfected with plasmids encoding Flag-uL18 or Myc-SPIN1 for 36 h and harvested for WB analysis with indicated antibodies. (B) Overexpression of SPIN1 disrupts the uL18-MDM2 binding. Lysates were prepared from HCT116^{p53−/−} cells co-transfected with HA-MDM2, Flag-uL18, Myc-SPIN1 or the corresponding empty vectors for 48 h and analyzed by immunoprecipitated with the anti-HA antibody. Immunoprecipitates and 5% of inputs were immunoblotted with the indicated antibodies. (C) Overexpression of SPIN1 fails to disrupt the uL5-MDM2 interaction. Lysates were prepared from HCT116^{p53−/−} cells co-transfected with HA-MDM2, Flag-uL5 and Myc-SPIN1 for 48 h and analyzed by immunoprecipitated with the anti-HA antibody. Immunoprecipitates and 5% of inputs were immunoblotted with the indicated antibodies. (LC: light chain). (D) SPIN1 and uL18 co-localize in the nucleolus. H1299 cells were transfected with GFP-SPIN1 and Flag-uL18 for 36 h and then immunostained with the anti-Flag antibody (red), and counterstained with DAPI.

**Figure 5. SPIN1 depletion increases ribosome-free uL18 and uL5.** (A) Knockdown of SPIN1 releases free forms of uL18 and uL5. HCT116^{p53+/+} were transfected with scramble or SPIN1 shRNA for 36 h and subjected to sucrose gradient fractionation assay followed by WB analysis.
with indicated antibodies. (B) SPIN1 knockdown increases the endogenous uL18/uL5-MDM2 interaction. Cell lysates of HCT116<sup>p53<sup>+/+</sup></sup> cells transfected with scramble or SPIN1 shRNA were immunoprecipitated with MDM2 or control IgG, and analyzed by WB analysis with indicated antibodies. (C) SPIN1 overexpression counteracts p53 activation induced by ActD or 5-Fu. U2OS cells were transfected with pcDNA or Flag-SPIN1 for 48 h, and treated with ActD or 5-Fu for 12 h before harvested for WB analysis with indicated antibodies. (D) and (E) Knockdown of uL18 or uL5 compromises the induction of p53 by SPIN1 depletion. U2OS cells were transfected with scramble siRNA, SPIN1 siRNA, uL18 siRNA (D) or uL5 siRNA (E) as indicated for 48 h. Cell lysates were subjected to WB analysis with indicated antibodies.

**Figure 6. SPIN1 knockdown retards tumor growth more dramatically by inducing p53 activity.** (A) and (B) Growth curves of xenograft tumors derived from HCT116<sup>p53<sup>+/+</sup></sup> cells and HCT116<sup>p53<sup>-/-</sup></sup> cells that expressed scramble or SPIN1 shRNA. Data are represented as mean ± SEM, n=6. (C) The images of xenograft tumors that were harvested at the end of experiment. (D) Quantification of the average weights of collected tumors from the above experiments. (E) and (F) The mRNA levels of SPIN1, p53 and p53 target genes were detected in 6 tumors by RT-qPCR (mean ± SEM, n=6). (G) The protein levels of SPIN1, p53 and p53 targets were detected in 6 tumors samples by WB analysis with indicated antibodies. *p<0.05, **p<0.01 by two-tailed t-test (D, E, F, G).

**Figure 7. A model for SPIN1 regulation of the uL18-MDM2-p53 pathway in cancer.** (see text in the Discussion for details).
Figure Supplement Legend

**Figure 1-figure supplement 1. SPIN1 interacts with uL18 in HEK293 cells.** HEK293 cells were transfected with plasmids encoding Myc-SPIN1 and Flag-uL18, and 48 h later cell lysates were collected for IP-WB analysis using anti-Flag antibody.

**Figure 2-figure supplement 1. SPIN1 knockdown increase p53 and its targets protein levels in a dose-dependent manner.** HCT116 p53+/+ (A) and U2OS (B) cells were transfected with a titrated concentration of SPIN1 siRNA (20 nM, 40 nM and 60 nM). And cells were harvested 48 h after transfection and followed by WB analysis with indicated antibodies. (C) U2OS cells were co-transfected with scramble siRNA (40 nM), SPIN1-siRNA (40 nM), with or without FLAG-SPIN1 as indicated. Cells were harvested 48 h after transfection for WB analysis.

**Figure 4-figure supplement 1. SPIN1 does not bind to MDM2, and SPIN1 and uL18 co-localize in the nucleolus.** (A) SPIN1 does not bind to MDM2. HCT116p53-/- cells were transfected with combination of plasmids encoding Myc-SPIN1 and HA-MDM2, followed by IP-WB analysis with indicated antibodies. (B) SPIN1 and uL18 co-localize in the nucleolus. HEK293 and H1299 SPIN1 stable cells were transfected with Flag-uL18 for 36 h and then immunostained with anti-Myc (red) and anti-Flag antibody (green), and counterstained with DAPI.

**Figure 5-figure supplement 1. SPIN1 knockdown reduces rRNA expression and SPIN1-Y170A mutant retains activity to repress p53.** (A). Scramble siRNA or SPIN1 siRNA was introduced into U2OS cells. RNA levels were analyzed using Q-PCR (*, p value < 0.05; **, p value < 0.01 by tailed t-test; n = 6). (B). SPIN1-Y170A inhibits p53. U2OS cells were transfected with control plasmids or FLAG-SPIN1-Y170A and the cells were collected for Western Blot analysis 48 hrs after transfection.

**Figure 5-figure supplement 2. Mapping of domains responsible for uL18-SPIN1 and uL18-MDM2 binding.** (A) uL18 interacts with the second Tudor like domain of SPIN1. Purified GST-tagged SPIN1 fragments, including aa 1-262(FL), aa 50-120, aa 121-262, aa 121-193, aa
194-262 and GST protein alone were incubated with purified His-uL18 protein for 3 hour at 4°C. Bound proteins were detected by WB analysis using anti-uL18 or coomassie staining. (B) A schematic diagram of uL18 binding regions on SPIN1 based on the result from (A). (C) SPIN1 interacts with both the N- and C-termini of uL18. Purified GST-tagged uL18 fragments, including aa1-297(FL), aa 1-50, aa 112-297, aa 39-253 or GST protein alone were rotated with purified His-SPIN1 protein for 1 hour at 4°C. Bound proteins were detected by WB analysis using anti-SPIN1 or coomassie staining. (D) A schematic diagram of SPIN1 binding regions on uL18 derived from the result from (C). (E) MDM2 interacts with both the N- and C-termini of uL18. Purified GST-tagged uL18 fragments, including aa1-297(FL), aa 1-50, aa 112-297, aa 39-253, aa 1-251 or GST protein alone were rotated with purified His-MDM2 protein for 4 hour at 4°C. Bound proteins were detected by WB analysis using anti-MDM2 (2A10) or coomassie staining. (F) A schematic diagram of uL18 binding regions on MDM2 based on the result from (E).

**Figure 6-figure supplement 1. Quantification of protein expression analyzed from xenograft tumors by Image J software.** All data were presented as mean ± SEM, n=6, *p<0.05, **p<0.01 by two-tailed t-test.

**Figure 6-figure supplement 2. High expression of SPIN1 is detected in multiple cancers and associated with poor prognosis in cancer patients.** (A) TCGA database was utilized, and the data were modified from the cBioPortal for Cancer Genomics (http://www.cbioportal.org/). (B) The expression profile of SPIN1 in cancers and normal tissues was searched in Oncomine Gene Browser (http://www.oncomine.org/). The results were from Talantov Melanoma database. 7 cases of normal skin and 45 cases of melanoma were analyzed in this figure. Correlation between SPIN1 upregulation and tumor stage, poorer prognosis or treatment resistance is not clear. (C and D) Overexpression of SPIN1 is correlated with overall survival and disease-free survival in breast cancer (http://www.cbioportal.org/), although the sample number of high SPIN1 patients is small and more samples are desired. (E) SPIN1 overexpression was associated with poor prognosis in colorectal cancer patients in an expression profile study from GSE17537 (http://www.PrognoScan.org/). (F) High expression of SPIN1 was correlated with poor overall survival in gastric cancer patients (http://www.kmplot.com).
Figure 6-figure supplement 3. Western blotting analyses of human colon cancer tissues (n=22) and normal colon tissue (n=20) and quantification of SPIN1 expression (Mean ± SEM, p<0.05)

Figure 6-figure supplement 4. Expression of genes involved in p53 pathway is correlated with SPIN1 expression. mRNA expression levels of 664 colorectal tumors were retrieved from Genomic Data Commons (https://portal.gdc.cancer.gov/). In the data set, gene expression levels were measured with FPKM (Fragments Per Kilobase of transcript per Million mapped reads) and normalized using the Upper Quantile method. All 644 tumor samples were sorted based on the expression level of SPIN1 from low to high.
Table 1. uL18-associated polypeptides identified from mass spectrometry analysis of proteins as shown in Figure 1A.

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</tbody>
</table>
Figure 6

A. HCT116 p53+/+
B. HCT116 p53-/-

Tumor Volume over Days after Inoculation

C. HCT116 p53+/+
D. HCT116 p53-/-

Tumor Weight over Days after Inoculation

E. Relative mRNA expression of SPIN1, p53, p21, PUMA in HCT116 p53+/+
F. Relative mRNA expression of SPIN1, p53, p21, PUMA in HCT116 p53-/-

G. Western Blot analysis of p53, SPIN1, PUMA, β-actin, GAPDH in HCT116 p53+/+ and HCT116 p53-/-.
Figure 1-figure supplement 1

Myc-SPIN1 + + (kDa)
Flag-uL18 +

Myc-SPIN1
-43
-34

Flag-uL18
-34

IP: anti-Flag

Input

Flag-uL18
-43

Myc-SPIN1
Figure 5-figure supplement 1

A

B

Flag-SPIN1-Y170A

MDM2

p53

p21

GAPDH
Figure 6-figure supplement 2

Panel A: Bar chart showing alteration frequency for different tumor types.

Panel B: Box plot comparing normal skin and melanoma with log_2 centered ratio.

Panel C: Kaplan-Meier curves for Breast Cancer patients with low and high SPIN1 expression.

Panel D: Kaplan-Meier curves for Breast Cancer patients comparing low SPIN1 to high SPIN1.

Panel E: Kaplan-Meier curves for Colorectal Cancer patients with low and high SPIN1 expression.

Panel F: Kaplan-Meier curves for Gastric Cancer patients comparing low and high SPIN1 expression.

Statistical significance: p<0.001 for Breast Cancer, p<0.01 for Gastric Cancer.
SPIN1 and 22 correlated p53 genes