REPORT

An ataxia-telangiectasia-mutated (ATM) kinase mediated response to DNA damage down-regulates the mRNA-binding potential of THOC5

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ABSTRACT

In response to DNA damage, transcription is blocked by inhibition of RNA polymerase II activity. The regulation of a preexisting pool of mRNAs, therefore, plays a key role in DNA repair, cell cycle arrest, or inhibition of differentiation. THOC5 is a member of the THO complex and plays a role in the export of a subset of mRNA, which plays an important role in hematopoiesis and maintaining primitive cells. Since three serine residues in the PEST domain of THOC5 have been shown to be directly phosphorylated by ataxia-telangiectasia-mutated (ATM) kinase, we examined the THOC5-dependent mRNA export under DNA damage. We show here that DNA damage drastically decreased the cytoplasmic pool of a set of THOC5-dependent mRNAs and impaired the THOC5/mRNA complex formation. The mRNP complex formed with nonphosphorylation mutant (S307/312/314A) THOC5, but not with a C-terminal deletion mutant after DNA damage, suggesting that the C-terminal domain of THOC5, but not its phosphorylation in the PEST domain, is necessary for the regulation of the mRNA-binding potency of THOC5. The cytoplasmic THOC5-dependent mRNAs were recovered by treatment with ATM kinase-specific or p53-specific siRNA, as well as by treatment with ATM kinase inhibitor, KU55933, under DNA damage conditions, suggesting that the ATM-kinase-p53 pathway is involved in this response to the DNA damage. Furthermore, the treatment with KU55933 blocked DNA damage-induced THOC5mRNP complex dissociation, indicating that activation of ATM kinase suppresses the ability of THOC5 to bind to its target mRNAs.

Keywords: FMIP/THOC5; THO complex; mRNA export complex; DNA damage; ATM kinase

INTRODUCTION

DNA damage response (DDR) involves multiple highly conserved checkpoint pathways that become activated on genotoxic stress. Checkpoint activation triggers a cascade of events that ultimately lead to cell cycle arrest or apoptosis. Key components of this response are the ataxia-telangiectasia-mutated (ATM) and the ATM- and RAD3-related (ATR) kinases that typically function as sensors of DNA damage. In metazoans, the integrity of these pathways is critical to protect cells from agents that cause uncontrolled cellular proliferation and differentiation (Motoyama and Naka 2004). A

consequence of ATM kinase loss is a variety of effects, including extinction of hematopoietic stem cells (Ito et al. 2004). Recent analysis of substrates for ATM and ATR kinases revealed that 23 substrates out of 700 belong to a group responsible for “pre-mRNA processing,” and this included members of TREX (transcript/export), which is a protein-complex coupling transcription and nuclear mRNA export. These include THOC4/Aly and FMIP/THOC5/chr22orf19 (Matsuoka et al. 2007), which play roles in mRNA export, suggesting a potential connection between DNA damage and RNA export machinery. In the yeast system, it has been previously shown that Sub2-Yra1 and Thp1-Sac3, the two main mRNA export complexes, are required for efficient transcription-coupled repair (Gaillard et al. 2007). However, the exact role of the TREX complex in mammalian systems under DNA damage conditions is not clear. The THO complex, which is a submember of TREX, was originally

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identified in *Saccharomyces cerevisiae* as a 4-protein complex (THO2p, Hpr1p, Mft1p, and Thp2p) (Chavez and Aguilera 1997; Piruat and Aguilera 1998; Chavez et al. 2001; Jimeno et al. 2002; Strasser et al. 2002; Reed and Cheng 2005) that plays a role in transcriptional elongation, nuclear RNA export, and genome stability. In higher eukaryotes such as *Drosophila melanogaster* (Rehwinkel et al. 2004) or humans (Masuda et al. 2005), three proteins (THOC1/hHpr1/p84, THOC2/hRlr1, and THOC3) and three additional unique proteins were identified, namely, THOC5/Fms-interacting protein (FMIP) (Tamura et al. 1999), THOC6, and THOC7, as members of the THO complex. We have previously shown that depletion of THOC5 by siRNA or ectopic expression causes abnormal hematopoiesis and abnormal muscle differentiation in mouse myeloid progenitor or mesenchymal progenitor cell lines, indicating that the THO complex is essential for the differentiation process in mouse (Tamura et al. 1999; Mancini et al. 2007; Carney et al. 2009). Furthermore, we generated interferon-inducible *THOC5* knockout mice to demonstrate that THOC5 is essential for maintaining primitive cells, such as embryonic or hematopoietic stem cells, but not terminally differentiated cells (Mancini et al. 2010), similar to ATM kinase (Ito et al. 2004). THOC5 is phosphorylated by several tyrosine kinases (Tamura et al. 1999; Pierce et al. 2008), protein kinase C (Mancini et al. 2004), and ATM kinase (Matsuoka et al. 2007), suggesting that extracellular stimulation regulates the function of THOC5. To obtain further insight into the role of THOC5 at the molecular level, we performed a microarray analysis using mouse embryonic fibroblasts (MEF) in the presence or absence of THOC5. Surprisingly, only 71 functionally known genes were down-regulated more than threefold by depletion of THOC5/FMIP. Moreover, >43% of these genes are involved in differentiation and development, such as *HoxB3* or *polycomb-CBX2*. (Guria et al. 2011), confirming that the THO complex plays a key role in the differentiation and development system. In this work, we have studied the mechanism of THO complex response to DNA damage signaling. We show that THOC5-dependent mRNAs were spliced, but not detected in the cytoplasm after DNA damage, suggesting that THOC5 is a mediator between DNA damage and the mRNA export of a subset of genes. Since approximately half of the identified THOC5-dependent genes are involved in the differentiation process (Guria et al. 2011), THOC5 may play a role in protecting cells from agents that cause uncontrolled differentiation.

RESULTS

DNA damage drastically decreased the cytoplasmic pool of a set of THOC5-dependent mRNAs

Three THOC5 serine residues, 307, 312, and 314, within the PEST-like domain, were previously identified as phosphor-

ylation sites by ATM kinase using irradiated HEK293 cell extracts (Matsuoka et al. 2007). We confirmed THOC5 phosphorylation after DNA damage by ATM kinase using an ATM kinase substrate antibody. HEK293 cells were treated with etoposide (0.2, 2.0, or 20 μ M) for 2 h, then cells were lysed and anti-THOC5 antibody used for immunoprecipitation, followed by ATM substrate (pS/T-Q)-specific immunoblot. At each concentration of etoposide used, THOC5 was detected by ATM substrate-specific antibody. To further confirm whether ATM kinase directly phosphorylates THOC5 in vitro, we performed in vitro kinase assay using *flag*-tagged *ATM kinase* cDNA (Kim et al. 1999) and GST-THOC5 (289–378) as substrate. GST-THOC5 (289–378) was phosphorylated by ATM kinase and the mutation of the three serine residues abolished phosphorylation (Supplemental Fig. 1). Thus, after confirming phosphorylation of THOC5 upon etoposide treatment, we then considered the effects of DNA damage and ATM kinase on THOC5-mediated mRNA export. We assessed the protein level of several members of TREX, such as THOC1, THOC5, THOC7, or Aly after etoposide treatment (2 h) in mouse embryonic fibroblasts (MEF). As shown in Figure 1A, the level of these proteins did not change. Furthermore, phosphorylation of p53 was observed only after treatment with etoposide, indicating that a DNA damage signal is induced under this condition in MEF cells (Fig. 1A). In addition, the level of THOC5 mRNA was not changed, suggesting that etoposide treatment did not influence this or THOC5 protein. In addition, we examined the ability of THOC5 to interact with other components of the TREX complex, such as THOC1 or Aly after DNA damage; we did not observe any differences in the THOC1/THOC5 interaction or the THOC5/Aly interaction before and after etoposide treatment (data not shown). We then examined the mRNA export function of THOC5 after etoposide treatment. We had previously generated a MEF cell line from THOC5 *lox/lox* mice (Mancini et al. 2010). Infection of THOC5/FMIP *lox/lox* MEF with Adenovirus carrying *cre-recombinase* and *GFP* genes (*Ad-GFP-Cre*) resulted in depletion of THOC5/FMIP expression in these cells within 4 d after infection (Mancini et al. 2010; Guria et al. 2011). Using this system, we previously identified 10 THOC5-dependent genes, such as *Grb10 interacting GYF protein 2* (*Gigyf2* or *Tnrc15*), *protein kinase C binding protein 1* (*Prkcbp1*), *twisted gastrulation homolog 1* (*Twsg1*), *enhancer of yellow 2 homolog* (*Eny2*), *solute carrier family 11 (proton-coupled divalent metal ion transporters)*, *member 2* (*Slc11a2*), *DHHC domain containing 2* (*Zdhhc2*) *zinc finger, inhibitor of DNA binding 2* (*ID2*), or *gap junction membrane channel protein α 1* (*Gja1*) (Guria et al. 2011).

To examine whether THOC5-dependent mRNAs are exported into the cytoplasm after DNA damage, we treated MEF cells with etoposide (10 μ M), and 2 h later isolated nuclear and cytoplasmic RNA. To validate the quality of the nuclear and cytoplasmic fractions, the marker pro-

FIGURE 1. DNA damage drastically decreased the cytoplasmic pool of a set of THOC5-dependent mRNAs. (A) MEF cells were treated with DMSO alone (DMSO) or with 10 μ M of etoposide (etoposide) for 2 h, and cell extracts were then subjected to THOC5, THOC1, THOC7, Aly, pp53, p53, and GAPDH-specific immunoblot (immunoblot). In addition, RNA was isolated from the sister culture of A and supplied for THOC5 and actin-specific RT-PCR (RT-PCR). (B) A total of 5000 MEF cells were treated as described above, and then nuclear (Nuc) and cytoplasmic (Cyt) RNAs were isolated and applied for *Zdhhc2*, *Prkcbp1*, *Ggylf2*, *Twsg1*, *Pdrgr1*, *Gadd45a*, *Myc*, and *actin*-specific RT-PCR using primers as shown in Table 1. We performed three to four independent experiments, and we show one example of representative data. As control for fractionation, aliquots of protein extracts from each sample were supplied for THOC5, Histone H3 (nuclear fraction), and GAPDH (cytoplasmic fraction)-specific immunoblot (immunoblot). (C) Signal intensity from *Zdhhc2*, *Prkcbp1*, *Ggylf2*, *Gadd45a*, *Myc*, and *actin*-specific RT-PCR products from cytoplasmic fraction were quantified using TINA 2.0 software. The reduced percent signal intensity from the treated with DMSO alone (etoposide $-$; as 100%) of the treated with etoposide (etoposide $+$) were shown. Mean values $\pm$ SD from three independent experiments. (D) Samples were prepared as described in B, but cDNA from all samples were standardized by expression levels of *actin* mRNA in both fractions, then applied for *Zdhhc2*, *Prkcbp1*, *Slc11a2*, *Ggylf2*, *Eny2*, *Twsg1*, *Gja1* (THOC5 dependent), *Pdrgr1*, *Gadd45a*, *Myc*, *Dusp10*, and *actin* (THOC5 independent)-specific RT-PCR. (E) Signal intensity from *Zdhhc2*, *Prkcbp1*, *Slc11a2*, *Ggylf2*, *Twsg1*, *Gadd45a*, *Myc*, and *actin*-specific RT-PCR products was quantified using TINA 2.0 software. The percent signal intensity from the nuclear (Nuc) or the cytoplasmic (Cyt) fraction of total intensity (Nuc + Cyt) in MEF treated with DMSO alone (etoposide $-$) or with etoposide ($+$) for 2 h were shown. Mean values $\pm$ SD from three independent experiments.

teins GAPDH (cytoplasmic fraction), Histone H3 (nuclear fraction) were used, plus THOC5-specific assay immunoblotting. GAPDH was detected primarily in the cytoplasmic fraction, while Histone H3 was detected only in the nuclear fraction (Fig. 1B). In addition, THOC5 were detected only in the nuclear fraction after treatment with DMSO alone or with 10 μ M of etoposide (Fig. 1B), indicating that drug treatment does not influence THOC5 nuclear location. RNA samples of both fractions obtained from 5000 cells were subjected to RT-PCR. Critically, only spliced forms of mRNA can be detected, since we chose the primer pair for RT-PCR, which is located on two different exons (Table 1). In all cases, most mRNA were detected in the cytosol, while mRNAs in the nuclear fraction were effectively undetectable, except for *actin* mRNA (Fig. 1B). Significantly, by quantifying the signal intensity, a reduction of $>85\%$ of THOC5-dependent mRNAs, such as *Zdhhc2*, *Prkcbp1*, and *Ggylf2* were observed in the cytoplasmic fraction after etoposide treatment (Fig. 1C). As a control, the THOC5-independent genes, *myc* and *actin* (Guria et al. 2011), were examined. The expression pattern of these genes was not altered by etoposide treatment (Fig. 1B,C; Table 1). In order to further validate these observations, we examined the *growth arrest and DNA-damage-inducible*, α (*Gadd45a*) and *p53* and *DNA damage-regulated gene 1* (*PDRG1*) mRNAs that play a role in DNA repair and are well known to be stabilized after DNA damage (Luo et al. 2003; Zhang et al. 2006; Zumbun et al. 2009; Reinhardt et al. 2010). As expected, both mRNAs were exported to the cytosol (Fig. 1B,C).

We next examined whether the THOC5-dependent mRNAs were transcribed in the nucleus. The amount of RNA was standardized by using equal expression levels of *actin* mRNA in both fractions (Fig. 1D,E). Eight THOC5-dependent mRNAs were then analyzed by RT-PCR. Seven out of eight THOC5-dependent mRNAs were barely detectable in the cytoplasmic fraction after cells had been treated with etoposide for 2 h,

TABLE 1. Comparison of THOC5-dependent mRNAs and mRNAs that are down-regulated by DNA damage

Symbol	Gene ID	Primers for RT-PCR (product size) ^a	Cytoplasmic mRNA in THOC5 ^{-/-} cells (Guria et al. 2011) ^b	Cytoplasmic mRNA in Etoside-treated cells (2 h) ^c
Grib10 interacting GYF protein 2 (Gigylf2)	BB782031	Forward: 5'-CGCCGACTGGAAGAGAAC-3'; reverse: 5'-TTGCTGTGTAGACTGTGAC-3' (235 bp)	More than fivefold reduction	More than fivefold reduction
Protein kinase C binding protein 1 (Prkcbp1)	BM239405	Forward: 5'-AAACCCGACAGACTTCCAC-3'; reverse: 5'-CGGCTGTATAAACATCTCTGAG-3' (245 bp)	More than fivefold reduction	More than fivefold reduction
Gap junction membrane channel protein α 1 (GJA1)	M63801	Forward: 5'-CAGCCTCCAGGAGTTCAC-3'; reverse: 5'-GAGAGATGGGGAAGACTTGT-3' (292 bp)	More than fivefold reduction	More than fivefold reduction
Twisted gastrulation homolog 1 (TWSG1)	AF292033	Forward: 5'-ACTGTGTCGGTATGTCAAC-3'; reverse: 5'-GGAGACGATGTTCCAGTTCAG-3' (146 bp)	More than threefold reduction	More than fivefold reduction
Enhancer of yellow 2 homolog (ENY2)	AV033253	Forward: 5'-ATGGTGTAGCAAGATGAACA-3'; reverse: 5'-ATGCTTGTAGCTGATCTTCCA-3' (144 bp)	More than threefold reduction	More than fivefold reduction
Solute carrier family 11 (proton-coupled divalent metal ion transporters) member 2 (Slc11a2)	BB771579	Forward: 5'-CAATGTCTTGTCTGTCCT-3'; reverse: 5'-GCGACCATTTAGGTTTCAAGAT-3' (293 bp)	More than threefold reduction	More than fivefold reduction
DHHC domain containing 2 (Zdhhc2)	BB342242	Forward: 5'-CTACTACGCTACGCCATCC-3'; reverse: 5'-TCAGCAATCTTCTCTGCA-3' (189 bp)	More than fivefold reduction	More than fivefold reduction
Zinc finger, inhibitor of DNA binding 2 (ID2)	NM_010496	Forward: 5'-CAGGCTCCAGGACGCCCT-3'; reverse: 5'-CCGGAGAACGACACCTGGGC-3' (310 bp)	No change	No change
p53 and DNA damage-regulated gene 1 (PDRG1)	NM_178939	Forward: 5'-GGTCAAGTAGAGGAGTTGGC-3'; reverse: 5'-CTTAGGGTGAGGCAATTTGATGA-3' (178 bp)	Twofold up-regulation	Twofold up-regulation
Growth arrest and DNA-damage-inducible, α (GADD45a)	NM_007836	Forward: 5'-CCCACTGATCATGTAGCG-3' (440 bp); reverse: GCTGGATTCCTTTGGCCCT-3';	Twofold up-regulation	1.2-fold up-regulation
Myc	NM_001177352	reverse: 5'-CGAACATAGGATGGAGCA (274 bp)	No change	No change
Dual specificity phosphatase 10 (DUSP10)	NM_022019	Forward: GGCAGTACTTTGAAGAGGCC-3'; reverse: 5'-GTTGAGGTCGGGGAAATAATTG (200bp)	No change	No change
Actin	NM_007393	Forward: 5'-AACACCCAGCCATGTACGTAG-3'; reverse: 5'-GTGTGGCATAGAGGTCTTACCG-3' (509 bp)	No change	No change

^aAll mRNA are detected in the nucleus as a spliced form.^bCompared with THOC5^{+/+} cells.^cCompared with untreated cells.

even in the presence of THOC5 (Fig. 1D; Table 1); however, in the nuclear fraction, spliced mRNAs were detected. The effect was not observed with THOC5-independent mRNAs, suggesting that THOC5-dependent mRNA export may be impaired after etoposide treatment. We obtained similar data using hydroxyurea (20 mM)-treated MEF (data not shown), suggesting that DNA damage may cause the loss of function of THOC5. However, one THOC5-dependent mRNA, *ID2*, was still exported into the cytoplasm after etoposide treatment (data not shown). In addition, we examined the THOC5-independent genes, *myc*, *dual specificity phosphatase 10 (DUSP10)*, *Gadd45a*, and *PDRG1* (Guria et al. 2011). The level of these mRNAs was not significantly altered (Fig. 1D). We then examined the ratio between nuclear and cytoplasmic spliced mRNA by quantifying the signal intensity. All seven THOC5-dependent mRNAs in the cytoplasmic fraction were drastically reduced (Fig. 1E), while etoposide treatment caused only slight changes of the ratio between cytoplasmic and nuclear THOC5-independent mRNAs. Although the half-lives of THOC5-dependent mRNAs, *Zdhhc2*, *Prkcbp1*, and *Gigyf2* are longer than 2 h in the presence or absence of etoposide, these mRNAs disappear from the cytoplasmic fraction within 2 h after etoposide treatment (Supplemental Fig. 2), suggesting that the decrease of cytoplasmic THOC5-dependent mRNAs may be in part due to faster cytoplasmic degradation.

THOC5 does not interact with mRNA after DNA damage

To determine whether the response to the DNA damage is directly dependent on THOC5, we analyzed the mRNP complex (containing THOC5) before and after DNA damage. We have previously shown that THOC5-dependent genes, such as *Gigyf2* or *Prkcbp1*, were detected in the mRNP complex using THOC5 as bait (Guria et al. 2011). Two-sister culture of 4×10^6 MEF cells were treated with DMSO alone or with 10 μ M of etoposide for 2 h. After centrifugation, cell extracts from 2×10^6 MEF cells were precipitated by THOC5 antibody or rabbit control IgG with protein G Sepharose. RNA were isolated from immunoprecipitates and supplied for RT-PCR using *Gigyf2*, *Prkcbp1*, *Zdhhc2*, and *actin*-specific primers. The endogenous *Gigyf2*, *Prkcbp1*, and *Zdhhc2*, but not *actin* mRNA, were detected in the THOC5-specific immunoprecipitates obtained from the DMSO-treated MEF. However, none of these mRNA species bound to THOC5 after etoposide treatment (Fig. 2A). No mRNAs were detected in precipitates with protein G Sepharose using control rabbit IgG or no IgG (Fig. 2A). Similar data were obtained using exogenous TAP-THOC5-Myc. MEF cells were transfected with TAP-THOC5-Myc (Fig. 2B), and were then treated with DMSO alone or with 10 μ M of etoposide for 2 h. After isolation of the mRNP complex using streptavidin-conjugated beads, RNA were isolated

and supplied for RT-PCR using *Gigyf2*, *Prkcbp1*, *Zdhhc2*, and *actin* specific primers. The endogenous *Gigyf2*, *Prkcbp1*, and *Zdhhc2* mRNAs, but not *actin* mRNA, were detected in the TAP-THOC5-Myc complex; however, after DNA damage, neither of the THOC5-dependent mRNAs were detected in the complex (Fig. 2C). Furthermore, a THOC5-mRNAs complex was isolated exclusively from the nuclear fraction, but not from the cytoplasmic fraction (data not shown), and THOC5 stayed in the nucleus after DNA damage (Fig. 1B), strongly suggesting that DNA damage causes dissociation of THOC5 with mRNAs in the nucleus.

We next examined whether phosphorylation of the PEST domain residues is involved in the down-regulation of the mRNP complex formation. We utilized the wild-type TAP-THOC5 and a mutant form generated that lacked ATM kinase phosphorylation sites, TAP-THOC5 (S307/312/314A). MEF cells were transfected with TAP-THOC5-Myc or TAP-THOC5 (S307/312/314A) (Fig. 2D) and Flag-tagged *Gigyf2* cDNA and were then treated with or without etoposide. After isolation of the mRNP complex using streptavidin-conjugated beads, RNA was isolated and subjected to RT-PCR using Flag and *Gigyf2*-specific primers. As expected, the exogenous flag-tagged *Gigyf2* mRNA was detected in the TAP-THOC5-Myc complex; however, after DNA damage, flag-tagged *Gigyf2* mRNA was not detected (Fig. 2D). Since the nonphosphorylation mutant also lost its RNA-binding potential after DNA damage (Fig. 2B,D), the phosphorylation itself is not required for the dissociation of *Gigyf2* mRNA from THOC5 after etoposide treatment. Given these data, we examined the domain that is essential for the mRNP complex dissociation using two further deletion mutants, THOC5 (1–499) (Fig. 2B) and THOC5 (1–289) (data not shown) (El Bounkari et al. 2009). Both mutants associated with *Gigyf2* mRNA before and after DNA damage equally well (Fig. 2D,E). These data suggest that the C-terminal domain (500–671), but not the PEST domain, is necessary for the regulation of THOC5 interaction, with its target mRNAs resulting from DNA damage. As a negative control, we cotransfected TAP-THOC5-Myc with β -globin cDNA, whose mRNA does not bind to THO complex (Merz et al. 2007). As expected, we did not detect β -globin mRNA from TAP-THOC5-Myc precipitates in any of the samples tested (Fig. 2D).

DNA damage impaired THOC5/mRNA complex formation via the ATM kinase pathway

In order to determine whether the effect of etoposide treatment on THOC5-dependent mRNAs is due to ATM-kinase activation, we used the ATM-kinase-specific siRNA, leading to a >10-fold down-regulation of ATM kinase within 48 h after transfection with ATM-kinase-specific siRNA (Fig. 3A). In cells depleted of ATM kinase, *Zdhhc2*, *Twsg1*, *Gigyf2*, or *Gja1* mRNAs were exported in the cytoplasmic fraction even 2 h after treatment with etoposide (Fig. 3B), suggesting

FIGURE 2. Etoposide treatment impairs the mRNA-binding potential of THOC5. (A) Two-sister culture of 4×10^6 MEF cells were treated with DMSO alone (DMSO) or with $10 \mu\text{M}$ of etoposide (etoposide) for 2 h. The cells were then extracted with lysis buffer. After centrifugation, cell extracts from 2×10^6 MEF cells were incubated with THOC5 antibody (anti THOC5), rabbit control IgG (CrIgG), and no IgG (—) for 1 h and were then precipitated with protein G Sepharose. RNAs were isolated from immunoprecipitates and supplied for RT-PCR using *Gigylf2*, *Prkcbp1*, *Zdhhc2*, and *actin*-specific primers. As control, immunoprecipitates were analyzed by THOC5-specific immunoblot (THOC5 [endo] Blot). Aliquots of cell extracts were supplied for RT-PCR (input). (B) Schematic representation of the pTAP (TAP-Vector, CBP: Calmodulin binding peptide; SBP: Streptavidin binding peptide) and pTAP carrying myc-tagged THOC5 (TAP-THOC5wt-Myc), and THOC5 mutants (TAP-THOC5 S307/312/314A-Myc and TAP-THOC5 [1-499]-Myc). (PEST) PEST-like domain; (LZ) putative leucine zipper. Numbers represent amino acid number. (C) pTAP (Cr) or TAP-THOC5wt-Myc were transfected into MEF cells. After 48 h, cells were incubated with DMSO alone (DMSO) or with $10 \mu\text{M}$ of etoposide (etoposide) for 2 h, and the RNP complex was then isolated using Streptavidin Sepharose. Bound fractions with Streptavidin Sepharose (bound) and cell extracts (input) were analyzed by immunoblotting using THOC5-specific antibody (Blot). Endogenous *Prkcbp1*, *Gigylf2*, *Zdhhc2*, and *actin* mRNAs in bound and input RNA samples were analyzed (RT-PCR). (D) pTAP, TAP-THOC5wt-Myc, TAP-THOC5 S307/312/314A-Myc, and TAP-THOC5 (1-499)-Myc (Blot) were cotransfected with *flag*-tagged *Gigylf2* cDNA or β *globin* gene cDNA into MEF cells. As control, we transfected MEF cells with pTAP empty vector (pTAP Cr). After 48 h, cells were treated as described above, and the RNP complex was then isolated using Streptavidin Sepharose. Bound fractions with Streptavidin Sepharose (bound) and cell extracts (input) were analyzed by immunoblotting using THOC5-specific antibody (Blot). *Gigylf2* mRNA in bound and input RNA samples were analyzed by *Flag* and *Gigylf2*-specific RT-PCR (Forward [*flag*]: 5'-ATGGATTACAAGGATGACGACGAT-3'; reverse [*Gigylf2*]: 5'-TTCATGCCGAAGTCTGGCAGC-3' [product size: 225bp]) or β *globin*-specific RT-PCR (Forward: 5'-AGCGGCCGCACACTTGCTTTTGA-3'; reverse: 5'-TCGAAGAACCTCTGGGTCCATGGG-3' [product size: 183 bp, spliced form]). (Mock) Cell extract from nontransfectants was applied for THOC5-specific immunoblotting and *flag*- and *Gigylf2*-specific RT-PCR. We have performed three independent experiments and we show one example of representative data. (E) Signal intensity from *Gigylf2*-specific RT-PCR products from the bound fraction using the full-length THOC5 and deletion mutant THOC5 (1-489) were quantified using TINA 2.0 software. The ratio of signal intensity before and after treatment with etoposide is presented from three independent experiments.

FIGURE 3. DNA damage impaired the THOC5/mRNA complex formation via the ATM kinase pathway. (A–E) MEF cells were transfected with siRNA against mouse *ATM kinase* (*ATMK*) (A,B,E) or *p53* (*p53*) (C–E) (Santa Cruz). (A,C) Forty-eight hours after transfection, cells were lysed and subjected to actin and ATM-kinase- (A) or p53- (C) specific immunoblot. (B,D) Sister cultures from B and D were treated with etoposide (10 μM) for 2 h, then nuclear (Nuc) and cytoplasmic (Cyt) RNAs were isolated, and applied for RT-PCR using *Zdhhc2*, *Twsg1*, *Gigyf2*, and *actin*-specific primers as described in Table 1. (E) Signal intensity from *Zdhhc2*, *Twsg1*, and *actin*-specific RT-PCR products was quantified using TINA 2.0 software. The percent signal intensity from the nuclear (Nuc) or the cytoplasmic (Cyt) fraction of total intensity (Nuc + Cyt) in MEF transfected with control siRNA (*siCr*), ATM-kinase-specific siRNA (*siATMK*), and p53-specific siRNA (*sip53*) were shown. Mean values ± SD from three independent experiments. (F) MEF cells were incubated with the potent inhibitor of ATM kinase KU55933 (5 μM) 1 h prior to treat with DMSO, or with etoposide for 2 h. Whole-cell extracts were subjected to THOC5, p53, pp53, and GAPDH-specific immunoblot. (G) The nuclear (Nuc) and cytoplasmic (Cyt) RNAs from the sister culture of (F: etoposide) were isolated and applied for RT-PCR using primers as described in Table 1. (H) TAP-THOC5wt-Myc was cotransfected with *flag*-tagged *Gigyf2* (*FlagGigyf2*) cDNA into MEF cells. After 48 h, cells were treated as described in F, and then RNP complexes were isolated using Streptavidin Sepharose. Bound fractions with Streptavidin Sepharose (bound) were analyzed by immunoblotting using THOC5-specific antibody (Blot) or RT-PCR using *Flag-Gigyf2*, *Prkcbp1*, *Zdhhc2*, and *actin*-specific primers (RT-PCR). (input) Input RNA samples were analyzed by RT-PCR; (endo) endogenous mRNA; (exo) exogenous mRNA.

that ATM kinase is required for the inhibition of THOC5-dependent mRNA export after DNA damage. To determine whether p53, a major target substrate of ATM kinase, influences the export of THOC5-dependent mRNAs, we down-modulated p53 using *p53*-specific siRNA. As shown in Figure 3C, p53 expression was abolished 48 h after siRNA transfection. In these cells, THOC5-dependent mRNA, such as *Zdhhc2*, *Twsg1*, *Gigyf2*, or *Gja1* mRNAs were still exported into the cytoplasm 2 h after etoposide treatment (Fig. 3D), indicating that the ATM-kinase/p53 pathway is required for this response to the DNA damage. The intensity of RT-PCR signals was also quantified (Fig. 3E): The export of THOC5-dependent mRNAs, such as *Zdhhc2* and *Twsg1*, were detected in the cytoplasm upon treatment with both ATM-specific (*siATMK*) and p53-specific (*sip53*) siRNAs, but not by the treatment with control siRNA (*siCr*) under DNA damage conditions.

To confirm these data, we utilized a potent inhibitor of ATM kinase, KU55933 (Hickson et al. 2004; Adams et al. 2010). MEF cells were incubated with KU55933 1 h prior to the stimulation with etoposide. Under these conditions, the phosphorylation of p53 was drastically reduced (Fig. 3F). RNA was then isolated from the nuclear and cytosolic fractions. In agreement with data using an siRNA approach, the THOC5-dependent mRNAs, *Zdhhc2*, *Prkcbp1*, and *Gigyf2* were detected in the cytosolic fraction after treatment with ATM-kinase inhibitor (Fig. 3G). We next examined whether DNA damage impaired the THOC5/mRNA complex formation via the ATM-kinase pathway. TAP-THOC5wt-Myc was cotransfected with *flag*-tagged *Gigyf2* (*FlagGigyf2*) cDNA into MEF cells. After 24 h, cells were incubated with or without ATM-kinase inhibitor, KU55933, 1 h prior to the stimulation with DMSO alone or with etoposide for 2 h, and then RNP complexes were isolated using Streptavidin Sepharose. Bound fractions were analyzed by immunoblotting using THOC5-specific antibody. As shown in Figure 3H, the treatment with KU55933 did not change the level of TAP-THOC5-Myc. The *FlagGigyf2*, *Prkcbp1*,

Zdhhc2, and *actin* mRNAs-bound fraction were then analyzed by RT-PCR (Fig. 3H), and treatment with ATM-kinase inhibitor recovered the ability of THOC5 to bind to its target mRNAs, such as *FlagGiyf2*, *Prkcbp1*, and *Zdhhc2*, indicating that ATM kinase is required for DNA damage-induced dissociation of THOC5 with mRNA. The THOC5-independent mRNA, *actin* mRNA, was not detected in the TAP-THOC5-Myc complex under any conditions shown.

In summary, DNA damage impaired the complex formation of THOC5 with its target mRNAs via the ATM-kinase/p53 pathway in the nucleus.

DISCUSSION

DNA damage inhibits the activity of RNA polymerase II at a time when DDR proteins and DNA repair proteins are needed. One of the main post-transcriptional events after DNA damage is the altered stability of mRNAs. Because transcription is reduced after DNA damage, there is an increased need to regulate the production of proteins from a pre-existing pool of mRNAs (Mitchell and Tollervey 2000; Orphanides and Reinberg 2002). On the other hand, the production of proteins that play a role in proliferation or differentiation likely should be blocked, because these may cause uncontrolled cellular proliferation and inappropriate differentiation. mRNAs from a pre-existing pool are potentially influenced by RNA-binding proteins and non-coding RNAs, such as microRNA (Keene 2007; Bartel 2009). However, little is known about mRNA export under conditions that damage the DNA template and RNA itself. We have previously shown that depletion of the *THOC5* gene in MEF cells causes the down-regulation (more than three-fold) of 198 genes (Guria et al. 2011). Forty-three percent of these genes are involved in development and cell differentiation, suggesting that THOC5 plays a role in the export of only a subset of genes, but plays an important role in cell differentiation. Our data obtained from *THOC5* knockout mice revealed that primitive hematopoietic cells in vivo not only stopped growing, but became apoptotic after *THOC5* gene depletion (Mancini et al. 2010). In addition, conventional THOC1 or THOC5 knockout mice are embryonic lethal (Li et al. 2005; Mancini et al. 2010). On the other hand, we show in the MEF system that THOC5-deficient differentiated fibroblasts just cease growing by *THOC5* gene depletion (Guria et al. 2011). We show here that seven out of eight THOC5-dependent mRNAs that were examined were spliced, but were not detected in the cytoplasm within 2 h after DNA damage. Thus, the inhibition of mRNA-binding potential of THOC5 may be essential to block uncontrolled differentiation. Interestingly, it has been reported that the treatment of HeLa cells with DNA damage reagents, such as adriamycin or etoposide, suppressed *Glut3* (*Slc2a3*) mRNA expression (Watanabe et al. 2010). We have previously shown that knockout of

THOC5 down-regulated *Glut3* mRNA more than sevenfold (Guria et al. 2011), suggesting that the down-regulation of *Glut3* mRNA, which was observed under conditions of DNA damage, may also be due to the loss of mRNA-binding potential of THOC5.

In the yeast system, it has been shown that DNA damage impairs tRNA nuclear export via differential relocation of the karyopherin Los1, the principal tRNA export receptor, to the cytoplasm (Ghavidel et al. 2007). However, THOC5 does not change its subcellular localization nor the complex formation with another member of TREX such as THOC1 or Aly after DNA damage. Strikingly, THOC5-dependent mRNA did not bind to THOC5 after etoposide treatment. How does THOC5 lose its binding potential to mRNAs? Since an ATM kinase phosphorylation site-defective mutant of THOC5 also lost the mRNA-binding potential just as the wild-type does upon DNA damage, the phosphorylation of THOC5 by ATM kinase cannot play a role in the binding potential. Furthermore, since the N-terminal domain that binds to mRNAs (Katahira et al. 2009) contains several phosphorylation sites (Tamura et al. 1999; Pierce et al. 2008), a potential acetylation site, and a potential SUMOylation site, post-translational modification may play a role in the loss of the mRNA-binding potential of THOC5. Alternatively, THOC5-dependent mRNAs may be recruited to other unknown mRNA-binding proteins specifically after DNA damage, annexing them away from THOC5. It has been shown that DNA damage-inducing agents can activate the RNA-binding activity of specific proteins including nucleolin, nucleophosmin, and MCG10, that favors a double stem-loop RNA structure (Carrier et al. 1994; Zhu and Chen 2000; Yang et al. 2002). Notably, some of RNA-binding proteins, such as MCG10 or nucleolin are regulated by p53 (Zhu and Chen 2000; Daniely et al. 2002). On the other hand, the C-terminal deletion mutant still binds mRNAs after DNA damage, suggesting that DNA damage-induced dissociation of THOC5 with its target mRNAs is via the loss of binding potential of THOC5 itself. It could be that the modification of the C-terminal domain by DNA damage influences the THOC5 structure by binding with unknown proteins that are not members of TREX, or alters protein conformation. Interestingly, interactome analysis using THOC5 as bait in HEK293 cells reveals that one of the interacting partners is p53 (AD Whetton and T Tamura, unpubl.). Furthermore, exogenous expression of THOC5 in cancer cells causes accumulation and colocalization of p53 (S Ramachandran and T Tamura, unpubl.). However, we could not find a direct interaction between p53 and THOC5, or indeed, colocalization of the two proteins in MEF cells before or after DNA damage. We are currently studying the binding partners of the THOC5 C-terminal domain before and after DNA damage by interactome analysis, using the MEF cell system.

It is presently unclear whether DNA damage causes inhibition of the export of THOC5-dependent mRNAs. A THOC5/mRNA complex was isolated exclusively from the

nuclear fraction, and THOC5 remained in the nucleus after DNA damage, strongly suggesting that DNA damage causes dissociation of THOC5 with mRNAs in the nucleus. These data also suggest that DNA damage may block THOC5-dependent mRNA export. On the other hand, although half-lives of THOC5-dependent mRNAs that were examined here are longer than 2 h, these mRNAs disappear from the cytoplasmic fraction within 2 h after etoposide treatment, suggesting that THOC5-dependent mRNAs are degraded rapidly in the cytoplasm. It may be that THOC5 or other proteins are bound to the mRNA under normal conditions and stabilize the RNA in the cytoplasm. Upon DNA damage, THOC5 (or another protein) no longer binds to the mRNAs, resulting in a shortened half-life for THOC5-dependent mRNAs. Both the inhibition of mRNA export and the faster cytoplasmic degradation may be involved in this response to the DNA damage.

We have previously shown that approximately half of identified THOC5-dependent genes are involved in the differentiation process (Guria et al. 2011), suggesting that THOC5 may play a role in protecting cells from agents that cause uncontrolled differentiation or alternatively disallow differentiation in inappropriate conditions.

MATERIALS AND METHODS

Cell culture, virus infection, plasmid construction, and DNA transfection

Mouse embryonic fibroblast (MEF) cells were grown in Dulbecco's modified Eagle's medium supplemented with 10% (v/v) FCS. Myc-tagged wild-type and mutant THOC5 were generated using pNTAP (Stratagene), as described previously (Tamura et al. 1999; El Bounkari et al. 2009). DNA transfection was performed with the Polyfect reagent as described by the manufacturer (Qiagen). *Gigv2* cDNA (IMAGE ID 5357184) was purchased by ImaGene. *β -globin* cDNA was a kind gift from H. Holtmann. *Flag*-tagged *ATM kinase* cDNA (Kim et al. 1999) was a kind gift from Michael Kastan. KU55933 was purchased by TOCRIS bioscience.

Antibodies

Mouse monoclonal antibodies against THOC5/FMIP (F6d) were described previously (Mancini et al. 2007). Rabbit polyclonal antibody against THOC5 was from Bethyl Laboratories, Inc. Mouse monoclonal antibodies against Aly and GAPDH, goat antibody against actin, and rabbit polyclonal against p53 were from Santa Cruz Biotechnology. Rabbit antibodies against phospho-p53 (Ser15) and Histone H3 were from Cell Signaling Technology, mouse monoclonal antibody against THOC1 (p84 N5) was from Gene Tex, Inc. Rabbit antibody against THOC7 was from Abgent.

Western blot procedures

Details of immunoblotting have been described previously (Koch et al. 2008). Results were documented on a LAS4000 imaging system (GE Healthcare Bio-Sciences).

RT-PCR analysis

Cytoplasmic RNA was isolated from MEF using the Qiagen RNeasy kit (Qiagen) according to the manufacturer's recommendations (Guria et al. 2011). Reverse transcription was carried out using oligo dT primers and Omniscript reverse transcriptase kit (Qiagen) following the instructions provided. Primer pairs for each PCR are shown in Table 1.

THOC5-mRNA complex isolation

Two-sister culture of 4×10^6 MEF cells were treated with DMSO alone or with etoposide for 2 h. After washing three times, cells were lysed with lysis buffer (10 mM Tris, 150 mM NaCl, 1 mM PMSF, 0.5% NP40, protease inhibitor cocktail, [Sigma-Aldrich] and RNase inhibitor), and were then frozen and thawed three times. After centrifugation, supernatants were incubated with rabbit polyclonal THOC5 antibody or rabbit control IgG with protein G Sepharose, then immunoprecipitates were washed three times. Alternatively, MEF cells were transfected with pNTAP vector (Stratagene) and the same vector carrying *myc*-tagged *THOC5*, and its mutants with or without *flag*-tagged *Gigv2* cDNA using the Polyfect reagent as described by the manufacturer (Qiagen), and the cells were allowed to grow for 48 h. Cells were then treated as detailed above. After centrifugation, supernatants were incubated with Streptavidin Sepharose (GE Healthcare Bioscience) in Streptavidin-binding buffer (lysis buffer with 10 mM β -mercaptoethanol and 0.5 mM EDTA), then washed three times. All steps were carried out at 4°C. Bound RNAs were used for RT-PCR analysis.

SUPPLEMENTAL MATERIAL

Supplemental material is available for this article.

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