Genotoxic stress/p53-induced DNAJB9 inhibits the pro-apoptotic function of p53

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DNAJB9 is a recently isolated member of the molecular chaperone gene family, whose precise function is largely unknown. In the present study, we have identified **DNAJB9** as an inducible gene of the tumor suppressor p53. **DNAJB9** expression was induced by p53 or genotoxic stress in a p53-dependent manner, which was mediated by the Ras/Raf/ERK pathway. In addition, depletion of **DNAJB9** by using siRNAs greatly increased genotoxic stress/p53-induced apoptosis, suggesting that **DNAJB9** inhibits the pro-apoptotic function of p53. We also found that **DNAJB9** physically interacts with p53 through its J domain, through which it inhibits the pro-apoptotic function of p53. Moreover, **DNAJB9** colocalized with p53 in both cytoplasm and nucleus under genotoxic conditions. Together, these results demonstrate that **DNAJB9** is a downstream target of p53 that belongs to the group of negative feedback regulators of p53.

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The tumor suppressor p53 is activated upon DNA damage and inhibits cell proliferation by inducing cell cycle arrest, senescence or apoptosis. p53 exerts these actions principally via binding to the consensus binding motifs in the genome, thereby activating the transcription of its target genes. It has been known that, among the target genes, *p21^{Cip1/WAF1}* mediates cell cycle arrest while apoptosis is mediated by *Bax*, *DR5*, *IGF-BP3*, *NOXA*, *PERP*, *Pidd* or *PUMA*.^{1–7}

Intriguingly, another group of target genes have been reported to inhibit p53 function and to act as negative feedback regulators of p53, which include *COP1*, cyclooxygenase-2 (*COX-2*), *DDR1*, *HB-EGF*, *Mdm2* and *Pirh2*. The *COP1*, *Mdm2* and *Pirh2*, direct transcriptional targets of p53, promote degradation of p53 via their E3 ubiquitin ligase activity, thereby inhibiting p53 transcriptional activity and p53-dependent cell cycle arrest and apoptosis.^{8–10} *DDR1* is another direct transcriptional target of p53 and is able to protect mouse embryonic fibroblasts (MEFs) and HCT116 cells from apoptosis caused by γ -radiation and doxorubicin.¹¹ *COX-2* is indirectly induced by p53-mediated activation of the Ras/Raf/ERK pathway and inhibits apoptosis caused by genotoxic stress in normal human epithelial cells, fibroblasts and endothelial cells.^{12,13} *HB-EGF* is induced by DNA damage in a p53-dependent manner and inhibits H₂O₂- and mitomycin C-induced apoptosis in MCF-7 cells.^{14,15} These negative feedback regulators of p53 have been thought to protect cells from fatal stresses resulting from p53 activation.¹²

Molecular chaperones are specialized proteins that bind to substrate proteins and facilitate their proper folding, unfolding, translocation or degradation. In most organisms including humans, heat shock protein 70s (Hsp70s) are major chaperones, whose activity is regulated by co-chaperones

such as DNAJ proteins. DNAJ proteins bind directly to Hsp70s to stimulate their ATPase activity, leading to the stabilization of interactions between Hsp70s and substrate proteins.¹⁶ At least 41 DNAJ proteins have been identified in the human genome, all of which have a conserved 70-amino acid (aa) sequence, the J domain, which is essential for binding to Hsp70s and stimulation of their ATPase activity.¹⁷

DNAJB9, also known as Mdg-1 or ERdj4, is a member of the DNAJ protein family.^{17,18} It is located in the endoplasmic reticulum (ER), where it acts as a co-chaperone for BiP, the ER member of the Hsp70s.¹⁹ **DNAJB9** has been shown to be induced by external stimuli including heat, methanol, ethanol, sodium chloride, tunicamycin and thapsigargin.^{19–22} In addition, it has been reported that **DNAJB9** inhibits tunicamycin-induced cell death in SK-N-SH cells.²¹ The cellular function of this DNAJ protein, however, remains largely unknown.

In the present study, we show that **DNAJB9** is an inducible gene of p53 under genotoxic conditions. **DNAJB9** expression is induced by p53 in response to DNA damage, which is mediated by the Ras/Raf/ERK pathway. In addition, we present evidence that **DNAJB9** inhibits the pro-apoptotic function of p53 through a physical interaction with p53. These results suggest that **DNAJB9** is a downstream target of p53 and acts as a negative feedback regulator of p53 under genotoxic conditions.

Results

DNAJB9 is induced by p53 under genotoxic conditions. We have previously observed that the transcript for **DNAJB9** is increased in response to p53 expression in DNA microarray analyses using EJ-p53, which is a stable cell line

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Abbreviations: Ad, adenovirus; COX-2, cyclooxygenase-2; ER, endoplasmic reticulum; Hsp, heat shock protein; MEF, mouse embryonic fibroblast

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that expresses p53 under the control of a tetracycline-regulated promoter.^{12,23} To investigate whether *DNAJB9* is indeed an inducible gene of p53, we first monitored *DNAJB9* expression in EJ-p53 cells by Northern and Western blot analysis. Removal of tetracycline from EJ-p53 cells resulted in an increase of p53 levels as well as p21, a known p53 target, as expected. In this state, the amount of *DNAJB9* mRNA and protein was found to increase substantially (Figure 1a), indicating that *DNAJB9* is induced by exogenous p53.

p53 is activated in response to DNA damage.²⁴ Therefore, to examine whether *DNAJB9* is induced by endogenous p53, we monitored *DNAJB9* expression upon DNA damage in SK-N-SH and U2OS cells harboring wild-type p53. When we administered doxorubicin, a genotoxic drug, to these cells, it was observed that both mRNA and protein levels of *DNAJB9* were increased substantially (Figures 1b and c). The amount of p53 and p21 was also increased by doxorubicin treatment, confirming that DNA damage occurred and p53 was activated by doxorubicin treatment in these cells (Figures 1b and c). Moreover, the quantity of *DNAJB9* was also increased by other genotoxic drugs including actinomycin D, etoposide and mitomycin C in SK-N-SH cells (Figures 1d–g). These results

suggest that *DNAJB9* is induced by endogenous p53 under genotoxic conditions.

To verify the involvement of p53 in the induction of *DNAJB9* under genotoxic conditions, we analyzed *DNAJB9* expression in *p53*(+/+) and *p53*(–/–) MEFs. The amount of *DNAJB9* mRNA and protein was prominently increased by doxorubicin treatment in *p53*(+/+) MEFs, which was not observed in *p53*(–/–) MEFs (Figure 1h). The p53 and p21 levels were elevated in response to doxorubicin treatment in *p53*(+/+) MEFs but not in *p53*(–/–) MEFs, confirming that p53 was activated only in *p53*(+/+) MEFs (Figure 1h). These results demonstrate that the induction of *DNAJB9* is dependent on p53 activation, verifying that *DNAJB9* is induced by p53 under genotoxic conditions.

Together, these data suggest that *DNAJB9* is indeed an inducible gene of p53 under genotoxic conditions.

***DNAJB9* is indirectly induced by p53 through the Ras/Raf/ERK pathway.** We next investigated the mechanism by which *DNAJB9* is induced by p53. For this purpose, we first tested a possibility that *DNAJB9* is a direct transcriptional target of p53. According to the PubMed database, the gene encoding human *DNAJB9* is located on the chromosome 7

Figure 1 *DNAJB9* is induced by p53 under genotoxic conditions. (a) EJ-p53 cells were cultured in the presence of 1 μ g/ml of tetracycline (Tet). To induce p53 expression, tetracycline was removed and cells were cultured for the indicated times. Total RNA and protein samples were prepared and subjected to Northern and Western blot analysis, respectively. GAPDH and β -actin were used as loading controls. SK-N-SH (b) and U2OS (c) cells were treated with 1 μ M of doxorubicin (Dox) for 8 h. Northern and Western blot analyses were performed. Doxorubicin (1 μ M) (d), actinomycin D (10 ng/ml) (e), etoposide (20 μ M) (f) or mitomycin C (20 μ M) (g) was administered to SK-N-SH cells. Cells were harvested at the indicated times and subjected to Western blot analysis. (h) *p53*(+/+) and *p53*(–/–) MEFs were treated with 1 μ M of doxorubicin for 8 h. Total RNA and protein samples were prepared and subjected to Northern and Western blot analysis

ranging from the nucleotide number 108210189 to 108215294. To find the p53-binding motifs in the promoter or intron regions of *DNAJB9*, we analyzed the 3 kb upstream region (108207188–108210188) and the gene itself by using the AliBaba2 program. The results showed that there are no p53-binding motifs in the 3 kb upstream region as well as in the *DNAJB9* gene (Supplementary Figures 1a and b), suggesting that *DNAJB9* is not a direct transcriptional target of p53.

It has been previously reported that the Ras/Raf/ERK pathway is activated by p53 under genotoxic conditions.¹⁴ In addition, we have previously shown that the Ras/Raf/ERK pathway mediates the induction of COX-2 by p53.¹² Therefore, we tested the possibility that the MAPK pathway is involved in the p53-induced *DNAJB9* expression in genotoxic conditions. The amount of *DNAJB9* and phosphorylated ERK1/2 (p-ERK1/2) were increased dramatically by doxorubicin treatment in SK-N-SH and U2OS cells, which was remarkably attenuated by the pretreatment of a MEK1/2 inhibitor, U0126 (Figures 2a and d). On the contrary, the pretreatment of SB203580 (a p38-MAPK inhibitor) or SP600125 (a JNK inhibitor) did not affect the *DNAJB9* expression in the presence of doxorubicin (Figures 2b and c), suggesting that the p38 and JNK pathway are not involved in the p53-induced *DNAJB9* expression. These results suggest that the MEK/ERK pathway is involved in the induction of *DNAJB9* by p53 under genotoxic conditions.

To confirm the involvement of the Ras/Raf/ERK pathway in the induction of *DNAJB9* by p53, we analyzed the effect of a dominant-negative (DN) mutant form of Ras (H-RasN17) or Raf-1 (K375M) on p53-induced *DNAJB9* expression.

In EJ-p53 cells, overexpression of p53 by tetracycline removal increased the amount of *DNAJB9* and p-ERK1/2, which was almost completely abolished by a transient transfection of DN-Ras or DN-Raf-1 (Figure 2e). These results confirm that the Ras/Raf pathway is involved in the induction of *DNAJB9* by p53.

All these results suggest that the Ras/Raf/ERK pathway mediates the induction of *DNAJB9* by p53 under genotoxic conditions.

DNAJB9 inhibits the pro-apoptotic function of p53 under genotoxic conditions. We next investigated the physiological role of *DNAJB9* that has been induced by p53 under genotoxic conditions. For this purpose, we generated three siRNAs specific for *DNAJB9* (si-JB9 #1, #2 and #3) and a scrambled siRNA (si-control) as a control, and confirmed that all three siRNAs are effective in preventing *DNAJB9* expression (Supplementary Figure 2a).

We transfected si-JB9 #1 and #2 as well as si-control into SK-N-SH and U2OS cells, and then treated cells with doxorubicin. After 24 h, the sub-G₁ apoptotic cell population was measured by flow cytometry. The data showed that the doxorubicin treatment increased sub-G₁ cell population in si-control-transfected cells as compared with DMSO (the vehicle) treatment (from 3.1 to 7.5% in SK-N-SH; from 1.4 to 12.2% in U2OS), which was dramatically enhanced in si-JB9-transfected SK-N-SH cells (23.1% for #1 and 21.2% for #2) and U2OS cells (29.1% for #1 and 29.0% for #2; Figures 3a and b). These results suggest that *DNAJB9* acts as a survival factor under genotoxic conditions.

Figure 2 *DNAJB9* is induced by p53-mediated activation of the Ras/Raf/ERK pathway. SK-N-SH cells were pretreated with U0126 (a), SB203580 (b) or SP600125 (c) at the concentrations of 0 (DMSO only), 10 or 20 μM for 1 h. Cells were then treated with 1 μM of doxorubicin (Dox) for 8 h. Protein samples were prepared and analyzed by Western blotting. β-actin was used as a loading control. (d) U2OS cells were pretreated with U0126 for 1 h, and then treated with 1 μM of doxorubicin for 8 h. Protein samples were analyzed by Western blotting. (e) EJ-p53 cells grown in the medium containing 1 μg/ml of tetracycline (Tet) were transfected with DN-Ras, DN-Raf-1 or an empty vector (pcDNA3). After tetracycline removal, cells were further cultured for 24 h and subjected to Western blot analysis

On the other hand, the amounts of p53, bax and cleaved caspase-3 were increased by doxorubicin treatment in si-control-transfected SK-N-SH and U2OS cells as compared with DMSO (Figures 3c and d), showing that the p53-dependent and caspase-3-mediated apoptotic pathway was activated by doxorubicin treatment in these cells. In addition, si-JB9 #1 and #2 elevated the amount of bax and cleaved caspase-3 as compared with si-control in the presence of doxorubicin (Figures 3c and d), suggesting that DNAJB9 inhibits the p53-dependent and caspase-3-mediated apoptosis under genotoxic conditions.

To verify that DNAJB9 inhibits the p53-dependent apoptosis under genotoxic conditions, we evaluated effects of the siRNAs on doxorubicin-induced apoptosis in *p53*(+/+) and *p53*(-/-) MEFs. Doxorubicin treatment increased apoptotic rates in both si-control-transfected *p53*(+/+) MEFs (from 1.4 to 6.0%) and *p53*(-/-) MEFs (from 2.0 to 10.4%) as compared with DMSO treatment (Figures 3e and f). The si-JB9 #1 and #2 enhanced the apoptotic rates in *p53*(+/+) MEFs as compared with si-control in the presence of doxorubicin (25.2% for #1; 26.1% for #2; Figure 3e), while

the apoptotic rates were not enhanced by the si-JB9s in *p53*(-/-) MEFs (10.4% for #1; 10.3% for #2; Figure 3f). We also observed that the si-JB9s did not enhance apoptotic rates in the presence of doxorubicin in another *p53*-null cell line, H1299 (Figure 3g and Supplementary Figure 2c). These results verify that DNAJB9 inhibits the p53-dependent apoptosis under genotoxic conditions.

To further confirm that DNAJB9 inhibits the pro-apoptotic function of p53, we evaluated the effect of siRNAs on apoptosis triggered by p53 overexpression. SK-N-SH cells were transfected with siRNAs and then infected by adenoviruses expressing p53 (Ad-p53) or green fluorescence protein (Ad-GFP) as a control. After 24 h, cells were harvested and the apoptotic rates were measured by flow cytometry. The data showed that Ad-p53 infection triggered apoptosis in si-control-transfected cells as compared with Ad-GFP (from 1.4 to 6.8%), which was remarkably enhanced in si-JB9-transfected cells (20.3% for #1 and 18.2% for #2; Figure 3h). In addition, si-JB9 #1 and #2 elevated the amount of bax and cleaved caspase-3 as compared with si-control in the presence of Ad-p53

Figure 3 DNAJB9 inhibits the pro-apoptotic function of p53. SK-N-SH (a), U2OS (b), *p53*(+/+) MEFs (e), *p53*(-/-) MEFs (f) and H1299 (g) cells were transfected with a scrambled siRNA (si-control), si-DNAJB9 #1 (si-JB9 #1) or si-DNAJB9 #2 (si-JB9 #2). Cells were then exposed to 1 μ M of doxorubicin (Dox) or DMSO (a vehicle) for 24 h and the sub-G₁ cell population was measured by flow cytometry. SK-N-SH (c) and U2OS (d) cells were treated as in a and b, respectively. Cell lysates were prepared and subjected to Western blot analysis. (h) SK-N-SH cells were transfected with si-control, si-JB9 #1 or si-JB9 #2, and then infected by Ad-GFP or Ad-p53. After 30 h, the sub-G₁ cell population was measured by flow cytometry. (i) SK-N-SH cells were treated as in h and subjected to Western blot analysis. Data represent averages $\pm$ S.D. (* P < 0.05 versus si-control)

(Figure 3i). These results confirm that DNAJB9 inhibits the pro-apoptotic function of p53.

All these data suggest that DNAJB9 inhibits the pro-apoptotic function of p53 under genotoxic conditions.

DNAJB9 physically interacts with p53 through the J domain. DNAJB9 is a molecular chaperone whose entity is to bind to other proteins. This led us to hypothesize that DNAJB9 might inhibit the pro-apoptotic function of p53 through a physical interaction. To test this hypothesis, we performed immunoprecipitation assays in SK-N-SH and U2OS cells under the treatment of doxorubicin. The results showed that DNAJB9 was co-precipitated with p53 by an anti-p53 antibody (Figures 4a and c). Consistent with this finding, p53 was co-precipitated with DNAJB9 by an anti-DNAJB9 antibody (Figures 4b and d). These results suggest that DNAJB9 physically interacts with p53 under genotoxic conditions.

DNAJB9 consists of 223 aa, which contains an amino-terminal hydrophobic sequence (aa 7–23), a J domain (aa 24–93) and a glycine-rich domain.²¹ On the basis of this

structural information, we generated (myc)₆-tagged DNAJB9 and its deletion mutants designated as FL (aa 1–223), NJ (aa 1–93), N (aa 1–23), J (aa 24–93) and ΔJ (truncated, aa 30–93; Figure 4e). SK-N-SH cells were transfected with these myc-DNAJB9 plasmids and then treated with doxorubicin to enrich for endogenous p53. The subsequent immunoprecipitation assays using an anti-myc antibody showed that p53 was co-precipitated with myc-DNAJB9-FL, –NJ and –J but not with myc-DNAJB9-N and –ΔJ (Figure 4f). Reverse immunoprecipitation assays using an anti-p53 antibody also showed that myc-DNAJB9-FL, –NJ and –J were co-precipitated with p53, but myc-DNAJB9-N and –ΔJ were not (Figure 4g). These results suggest that DNAJB9 interacts with p53 through the J domain.

DNAJB9 colocalizes and interacts with p53 in both cytoplasm and nucleus under genotoxic conditions. It has been known that the primary location of DNAJB9 is the ER whereas p53 shuttles between the cytosol and the nucleus.^{19,21} This raises us a question of how DNAJB9 and p53 existing in different compartments can physically

Figure 4 DNAJB9 interacts with p53 through the J domain. SK-N-SH (a) and U2OS (c) cells were treated with 1 μ M of doxorubicin (Dox) for 8 h. Cell lysates were prepared and immunoprecipitation experiments were performed with an anti-p53 antibody (FL-393). The precipitates were subjected to Western blotting using antibodies against DNAJB9 and p53. SK-N-SH (b) and U2OS (d) cell lysates were immunoprecipitated by the anti-DNAJB9 antibody, and the precipitates were subjected to Western blotting using antibodies against p53 and DNAJB9. (e) Schematic representation of the structure of human DNAJB9 and its deletion mutants. (f and g) SK-N-SH cells were transfected with expression vectors for myc-DNAJB9 deletion mutants or an empty vector (Vt) as indicated, and then treated with 1 μ M of doxorubicin for 8 h. In f, immunoprecipitation was carried out using an anti-myc antibody, and the samples were then probed with anti-p53 or anti-myc antibodies. In g, immunoprecipitation was carried out using an anti-p53 antibody (FL-393), and the samples were then probed with anti-myc and anti-p53 antibodies

interact with each other. To solve this problem, we performed Western blot analysis using cytoplasmic and nuclear fractions from SK-N-SH and U2OS cells. The majority of DNAJB9 was detected in the cytoplasmic fraction in resting states. Upon doxorubicin treatment, however, the amount of DNAJB9 increased in both

fractions. As for p53, it was detected mainly in the nuclear fraction in resting states and its amount was increased in both fractions by doxorubicin treatment (Figures 5a and b). These results demonstrate that DNAJB9 and p53 colocalize in both cytoplasmic and nuclear fractions under genotoxic conditions.

Figure 5 DNAJB9 colocalizes and interacts with p53 in both cytoplasm and nucleus under genotoxic conditions. SK-N-SH (a) and U2OS (b) cells were treated with doxorubicin (1 μ M) for 8 h. Cytoplasmic (Cy) and nuclear fractions (Nu) were prepared and subjected to Western blot analysis using antibodies for DNAJB9 and p53. Tubulin and PARP were used as loading controls for the cytoplasmic and nuclear protein, respectively. (c and d) SK-N-SH cells were exposed to doxorubicin (1 μ M) for 8 h. Cells were then stained with antibodies for DNAJB9 (c) or p53 (d) as described in the Materials and Methods. DAPI was used to stain the nuclei. Images were obtained with a confocal microscope. (e and f) SK-N-SH cells were treated with doxorubicin for 8 h and then cytoplasmic and nuclear fractions were prepared. In e, immunoprecipitation was carried out using an anti-p53 antibody (FL-393), and the samples were probed with anti-DNAJB9 or anti-p53 antibodies. In f, immunoprecipitation was carried out using an anti-DNAJB9 antibody, and the samples were probed with anti-p53 and anti-DNAJB9 antibodies

In confocal microscopy, DNAJB9 was detected mainly in the perinuclear region in resting states, which was consistent with the previous report.²¹ Intriguingly, however, DNAJB9 was detected in the cytoplasm dispersedly as well as in the nucleus under doxorubicin treatment (Figure 5c). As for p53, it was mainly detected in the nucleus while small amount of p53 was detected in the cytoplasm in resting states. The nuclear localization of p53 became more prominent under the treatment of doxorubicin (Figure 5d). These results demonstrate that DNAJB9 and p53 colocalize in both cytoplasm and nucleus under genotoxic conditions.

We then performed immunoprecipitation assays using cytoplasmic and nuclear fractions from SK-N-SH cells treated with doxorubicin. The results showed that DNAJB9 was co-precipitated with p53 by an anti-p53 antibody in both fractions (Figure 5e). Conversely, p53 was co-precipitated with DNAJB9 by an anti-DNAJB9 antibody in both fractions (Figure 5f). These results suggest that DNAJB9 interacts with p53 in both cytoplasm and nucleus under genotoxic conditions.

Together, these data suggest that DNAJB9 colocalizes and interacts with p53 in both cytoplasm and nucleus under genotoxic conditions.

DNAJB9 inhibits the pro-apoptotic function of p53 through a physical interaction. We next asked whether DNAJB9 could inhibit the pro-apoptotic function of p53 through a physical interaction. For this purpose, we transfected SK-N-SH cells with myc-DNAJB9 plasmids or an empty vector and then treated cells with doxorubicin. After 48 h, the sub-G₁ apoptotic cell population was measured by flow cytometry. The data showed that doxorubicin treatment increased sub-G₁ cell population from 1.5 to 47.9% in vector-transfected cells. In contrast, myc-DNAJB9-FL, -NJ and -J reduced the apoptotic rates remarkably to 15.2%, 14.4% and 13.8%, respectively. Myc-DNAJB9-N and -ΔJ had no significant effect on the apoptotic rates as compared with the vector (-N, 46.4% and -ΔJ, 52.6%; Figures 6a and b). These results suggest that the J domain of DNAJB9 is critical

Figure 6 DNAJB9 inhibits the pro-apoptotic function of p53 through a physical interaction. (a) The J domain of DNAJB9 inhibits genotoxic stress-induced apoptosis. SK-N-SH cells were transfected with expression vectors for myc-DNAJB9 deletion mutants or an empty vector (Vt), and then treated with 1 μ M of doxorubicin (Dox) for 48 h. The sub-G₁ cell population was measured by flow cytometry. (b) Representative flow cytometry data of a are shown. (c) The J domain of DNAJB9 inhibits p53-induced apoptosis. SK-N-SH cells were transfected with expression vectors for myc-DNAJB9 deletion mutants or an empty vector (Vt), and then infected by Ad-GFP or Ad-p53 for 48 h. The sub-G₁ cell population was measured by flow cytometry. (d) Representative flow cytometry data of c are shown. Data represent averages $\pm$ S.D. (* P < 0.05)

to inhibit the p53-dependent apoptosis under genotoxic conditions.

To verify the effect of the J domain on the p53-dependent apoptosis, we analyzed the effect of the J domain on apoptosis triggered by p53 overexpression. SK-N-SH cells were transfected with myc-DNAJB9 plasmids or an empty vector and then infected by Ad-GFP or Ad-p53. After 48 h, cells were harvested and the apoptotic rates were measured by flow cytometry. The data showed that Ad-p53 infection triggered apoptotic response in vector-transfected cells as compared with Ad-GFP (sub-G₁, from 2.6 to 27.0%), which was remarkably inhibited by myc-DNAJB9-FL, -NJ and -J (10.2%, 9.1% and 8.9%, respectively). Myc-DNAJB9-N and -ΔJ had no significant effect on the apoptotic rates (31.2% and 24.1%, respectively; Figures 6b and d). These results confirm that the J domain of DNAJB9 is critical to inhibit the p53-dependent apoptosis.

All these results demonstrate that the J domain of DNAJB9 is critical to inhibit the pro-apoptotic function of p53, suggesting that DNAJB9 inhibits the pro-apoptotic function of p53 through a physical interaction.

Discussion

p53 is an important tumor suppressor whose mutation is found in half of human cancers. In response to DNA damage, p53 is activated to induce cell cycle arrest, senescence or apoptosis, through which it prevents proliferation of cells harboring damaged DNA.²⁴ Since p53 exerts these actions mainly via inducing the expression of its downstream target genes, it is indispensable to identify its downstream target genes and to characterize them for understanding the action mechanism of p53. In the present study, we have identified *DNAJB9* as a novel inducible gene of p53.

Most known target genes of p53 including *Bax* and *p21* have been shown to mediate p53 function such as cell cycle arrest or apoptosis under genotoxic conditions. Accumulating evidence, however, indicates that another group of target genes do not mediate but inhibit p53 function, thereby acting as negative feedback regulators of p53.²⁵ For example, *COP1*, *COX-2*, *DDR1*, *HB-EGF*, *Mdm2* and *Pirh2* are induced by p53 but inhibit p53-dependent cell cycle arrest or apoptosis under genotoxic conditions.^{8–15} In the present study, we have identified *DNAJB9* as a novel inducible gene of p53 that belongs to the group of negative feedback regulators of p53.

We have shown here that DNAJB9 is induced by p53 through the Ras/Raf/ERK pathway. There is still a possibility that other signaling pathways are also involved in the DNAJB9 induction by p53. Given the results, however, that the DNAJB9 induction by p53 was almost completely abolished by DN-Ras, DN-Raf-1 and a MEK1/2 inhibitor (Figure 2), we reason that the Ras/Raf/ERK cascade is the main pathway to induce the DNAJB9 expression by p53. It has been previously reported that p53 activates the Ras/Raf/ERK pathway under genotoxic conditions¹⁴ and that COX-2, another downstream target of p53, is induced by p53/Ras/Raf/ERK cascade,¹² supporting our conclusion that the Ras/Raf/ERK pathway mediates the induction of DNAJB9 by p53. Further studies are required to understand the complete signaling networks by which p53 induces DNAJB9 expression.

Human *DNAJB9* was first cloned in SK-N-SH cells as one of the genes induced by ER stressors such as tunicamycin and thapsigargin.²¹ The primary location of DNAJB9 is the ER, where it binds to BiP/GRP78, the major Hsp70 in the ER, and stimulates the ATPase activity of BiP.¹⁹ Although it has been recently proposed that DNAJB9 is involved in the ER-associated degradation of misfolded proteins,²⁶ the cellular function of DNAJB9 is largely unknown. We here found a novel function of DNAJB9 under genotoxic conditions: DNAJB9 is induced by p53 and inhibits the pro-apoptotic function of p53 (Figures 1–3). A previous report that DNAJB9 inhibits tunicamycin-induced cell death in SK-N-SH cells supports our conclusion that DNAJB9 is a survival factor.²¹

We here found that DNAJB9 inhibits the pro-apoptotic function of p53 through a physical interaction and the physical interaction occurs in both cytoplasm and nucleus under genotoxic conditions (Figures 4–6). These findings suggest a possibility that the cytoplasmic interaction between DNAJB9 and p53 might promote the degradation of p53. However, our data showed that depletion of DNAJB9 by siRNAs rather decreased p53 levels under genotoxic conditions (Figures 3c, d and i) and DNAJB9 overexpression increased p53 levels (Supplementary Figure 2b), suggesting that DNAJB9 does not downregulate p53 levels.

Another possibility is that the cytoplasmic DNAJB9–p53 interaction might prevent the nuclear translocation of p53. Actually, we observed that nuclear translocation of p53 was increased by DNAJB9 siRNAs but decreased by DNAJB9 overexpression (Supplementary Figures 3a and b). In addition, confocal data showed that DNAJB9 overexpression attenuates the nuclear translocation of p53 under doxorubicin treatment (Supplementary Figure 3c). These data suggest that the cytoplasmic DNAJB9–p53 interaction inhibits the nuclear translocation of p53, which could be one of the mechanisms by which DNAJB9 inhibits the pro-apoptotic function of p53.

Nonetheless, the confocal data also showed that DNAJB9 did not sequester p53 in the cytoplasm completely and the majority of p53 still translocated to the nucleus under doxorubicin treatment (Supplementary Figure 3c), suggesting that the cytoplasmic DNAJB9–p53 interaction is not the main mechanism by which DNAJB9 inhibits the pro-apoptotic function of p53. In addition, we observed that DNAJB9 abolished the p53 binding to the *Bax* promoter under doxorubicin treatment in chromatin immunoprecipitation assays (Supplementary Figure 3d). Therefore, we assume that the nuclear DNAJB9–p53 interaction might be a major mechanism by which DNAJB9 inhibits the pro-apoptotic function of p53. Further studies are necessary to test this possibility and to elucidate the function of DNAJB9 in the nucleus.

DNAJB9 is an ER-resident protein. In some conditions, however, DNAJB9 has been found in another compartment. For example, GFP-fused DNAJB9 was observed to translocate to the nucleus under the treatment of heat shock and methanol in COS-7 and HeLa cells.^{20,22} We report here that genotoxic stress is another cause to localize DNAJB9 in the nucleus (Figure 5). Further studies are required to elucidate the mechanism of how DNAJB9 enters the nucleus.

Figure 7 A biochemical link between DNAJB9 and p53 under genotoxic conditions

Collectively, we have established a biochemical link between DNAJB9 and p53 under genotoxic conditions. DNAJB9 is a downstream target of p53 and acts as a negative feedback regulator of p53 through a physical interaction (Figure 7). This study might contribute to understand action mechanisms of p53 and cellular functions of DNAJB9 under genotoxic conditions.

Materials and Methods

Plasmids, siRNAs and adenoviruses. Mammalian expression vectors for DN-Ras (H-RasN17)²⁷ and Raf-1 (K375M)²⁸ were generous gifts from Dr. Sam W Lee (Massachusetts General Hospital, Boston, MA, USA). The oligonucleotide sequences of siRNAs used in this study were: si-control, 3'-CCUACGCCACCAU UUCGU-5'; si-JB9 #1, 3'-ACACGUAAUUUGCCUAGU-5'; si-JB9 #2, 3'-CACUC UUAGGUCUUAGU-5'; si-JB9 #3, 3'-ACGAAGAGGAATATGGTT-5'. Adenoviruses expressing GFP and p53 were generated and amplified as described previously.^{29,30}

Generation of a polyclonal anti-DNAJB9 antibody. A rabbit polyclonal antibody was raised against a synthetic peptide, Ac-G-F-D-S-T-N-Q-H-T-V-Q-T-E-N-R-F-H-G-S-S-K-H-C, corresponding to residues 181–203 of human DNAJB9, as described previously.²¹ The antibody was purified using a protein A agarose affinity column (Upstate, Lake Placid, NY, USA).

Cell culture and transfection. SK-N-SH (human neuroblastoma), U2OS (human osteosarcoma), H1299 (human lung cancer), p53(+/+) and p53(-/-) MEFs were cultured in Dulbecco's modified Eagle's medium containing fetal bovine serum (10%) and penicillin/streptomycin (1%) in a 5% CO₂ incubator. EJ-p53 cells were cultured in the above medium in the presence of tetracycline (1 µg/ml) as described previously.¹² For transfection of plasmids and siRNAs, cells were seeded into 60-mm dishes and the transient transfection was performed using Lipofectamine2000 reagent (Invitrogen, Carlsbad, CA, USA).

FACS analysis. Cells were collected and resuspended in 1 ml cold PBS. Ethanol, pre-chilled at -20 °C, was added to cells drop wise with gentle vortexing. The resultant mixture was incubated at 4 °C overnight. The following day, cells were washed with 10 ml ice-cold PBS and permeabilized in a reagent consisting of 100 µg/ml RNase A and 50 µg/ml propidium iodide in PBS. Samples were incubated at 37 °C for 1 h and then analyzed by flow cytometry (FACScan, Becton Dickinson, Franklin Lakes, NJ, USA). The sub-G₁ cell population was measured using the CellQuest program (Becton Dickinson).¹²

Confocal microscopy. Cells were seeded onto 18-mm coverslips and cultured overnight. The following day, cells were fixed in a 3.7% formaldehyde solution and permeabilized in a 0.2% Triton X-100 solution. The cells were incubated with the antibody for DNAJB9 or p53 (FL-393, Santa Cruz Biotechnology, Dallas, TX, USA) for 2 h and then with a FITC-conjugated anti-rabbit IgG (Sigma, St. Louis, MO, USA) for 1 h. After washing, nuclei were stained with DAPI (Sigma) for 5 min and images were taken with a confocal microscope (Olympus Fluoview, Olympus, Tokyo, Japan).²⁹

Northern blot analysis. Total RNA was extracted from cells with Trizol reagent (Life Technologies, Carlsbad, CA, USA), resolved on 1% formaldehyde-agarose gels

and subsequently transferred to zeta-probe GT blotting membranes (Bio-Rad, Berkeley, CA, USA). Hybridization was performed using ³²P-labeled probes, and the washed membranes were then subjected to autoradiography.²⁹

Western blot analysis. Cells were lysed in a lysis buffer (150 mM NaCl, 50 mM Tris-HCl, 1% NP-40, 0.25% sodium deoxycholate, and 0.1% SDS) with 1 mM PMSF, 1 µg/ml aprotinin, 1 µg/ml leupeptin and 1 µg/ml pepstatin. Proteins were resolved on 10 or 12% SDS-polyacrylamide gels and transferred to PVDF membranes, which were probed with specific antibodies. The immunoreactive protein complexes were detected by enhanced chemiluminescence (Amersham Bioscience, Buckinghamshire, UK).²⁹ The specific antibodies used in this study were: antibodies for p21, bax, cleaved caspase-3, ERK, p-ERK, p38, p-p38, JNK, p-JNK and PARP from Cell Signaling Technology (Danvers, MA, USA); an anti-p53 antibody (Ab-6) from Calbiochem (Billerica, MA, USA); anti-myc and anti-tubulin antibodies from Covance (Princeton, NJ, USA); and an anti-β-actin antibody from Sigma.

Subcellular fractionation. Cytoplasmic and nuclear fractions were prepared as described previously.^{29,31} Cells were washed with PBS and allowed to swell in buffer A (10 mM HEPES, pH 7.9, 10 mM KCl, 1 mM DTT, 1 mM PMSF, 1 µg/ml leupeptin, 1 µg/ml aprotinin and 1 µg/ml pepstatin). After 15 min, cells were lysed by adding NP-40 (final concentration, 0.5%) and the nuclei were pelleted by centrifugation at 1500 × g for 5 min. The supernatant was centrifuged at 15 000 × g for 5 min, and the cytoplasmic fraction was recovered. The nuclei were washed with buffer A and resuspended in buffer B (20 mM HEPES, pH 7.9, 0.4 M NaCl, 1 mM DTT, 1 mM PMSF, 1 µg/ml leupeptin, 1 µg/ml aprotinin and 1 µg/ml pepstatin). After incubation on ice for 15 min, the nuclear fraction was recovered by centrifugation at 18 000 × g for 5 min.

Immunoprecipitation. Cells were lysed in a lysis buffer (50 mM Tris-HCl, pH 7.4, 150 mM NaCl, 1% NP-40, 0.25% sodium deoxycholate, 0.1% SDS, 50 mM NaF, 1 mM sodium orthovanadate and a protease inhibitor cocktail) by passage through a 22G needle (Tae-Chang, Kongju, South Korea). The cell lysates were immunoprecipitated with an antibody against DNAJB9 or p53 (FL-393, Santa Cruz Biotechnology) at 4 °C, then mixed with Protein A/G PLUS-agarose beads (Santa Cruz Biotechnology) for 2 h. The beads were then washed three times with the lysis buffer, and bound proteins were subjected to SDS-PAGE and Western blot analysis.²⁹

Chromatin immunoprecipitation. Chromatin immunoprecipitation assays were performed according to the manufacturer's instruction (Upstate). Briefly, cells were fixed with 1% formaldehyde and the cell pellet was sonicated in a lysis buffer. After centrifugation, the supernatant was diluted, pre-cleared and mixed with 2 µg of an anti-p53 antibody (FL-393) or normal rabbit serum at 4 °C overnight. The next day, samples were mixed with protein A agarose/salmon sperm DNA for 1 h at 4 °C and the beads were washed several times. An elution buffer was added and the eluate was treated with 0.2 M NaCl for 4 h at 65 °C and then proteinase K for 1 h at 45 °C. DNA was recovered by phenol/chloroform extraction and amplified by polymerase chain reaction using primers specific for the human *Bax* promoter. Sequences were 5'-TAATCCAGCGCTTTGGAAG-3' (forward) and 5'-TGCAGA GACCTGGATCTAGC-3' (reverse) as described previously.³²

Statistical analysis. Comparisons between groups were evaluated by the student's *t*-test using the Excel program (Microsoft Co., Redmond, WA, USA). Differences between groups were considered statistically significant when the *P*-values were ≤ 0.05.

Conflict of Interest

The authors declare no conflict of interest.

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