New role of human ribosomal protein S3: Regulation of cell cycle via phosphorylation by cyclin-dependent kinase 2

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Abstract. Human ribosomal protein S3 (hRpS3) is a component of the 40S ribosomal subunit that associated in protein synthesis. hRpS3 has additional ribosomal functions such as DNA repair, transcription, metastasis, and apoptosis via interaction with numerous signaling molecules and has different modifications. Cdk-dependent kinases (CDKs) are heterodimeric serine/threonine protein kinases that regulate cell cycle progression. Among its members, the Cdk1-cyclin B complex is known to control cell progression in the G2/M phase, while Cdk2-cyclin E/A complexes function in G1/S and S/G2 transition. In our previous study, we observed interaction between hRpS3 and Cdk1. The present study investigated the interaction between hRpS3 and Cdk2. Cdk2 phosphorylated hRpS3 at amino acid residues S6 and T221 during the S-phase. Furthermore, hRpS3 knockdown delayed cell cycle progression by modulating the expression of cell cycle-related proteins, including cyclin B1 and cyclin E1. These findings suggest that hRpS3 is involved in Cdk2-mediated cell cycle regulation.

Introduction
Cyclin-dependent kinase (Cdk) 2 regulates the initiation and progression of the S phase of the cell cycle, and the regulation of Cdk2 involves cyclin binding and phosphorylation (1). Monomeric Cdk2 is inactive, and its activation requires binding to cyclin. Cdk2 associates with cyclin E to drive G1/S phase transition, and it associates with A-type cyclins to mediate DNA replication during S phase (2-5). The Cdk2/cyclin complex is recognized by multiple kinases, which phosphorylate the T14, Y15, and T160 amino acid residues in the complex. Phosphorylation of T14 and Y15 induces inactivation of Cdk2; activation of Cdk2 requires de-phosphorylation of both T14 and Y15 by Cdc25 and phosphorylation of T160 by CDK activating kinase (CAK) (6,7). Cdk2 also mediates the cell cycle inhibitory and tumor-suppressing activities of P21 Cip1 and P27 Kip1, respectively (2,8-10).

Human ribosomal protein S3 (hRpS3) is a component of the small ribosomal subunit, which is involved in protein synthesis (11-13). The functions of hRpS3 are not limited to protein translation, and it performs multiple extra-ribosomal activities, such as in DNA repair, cell death, inflammation, and tumorigenesis (13-19). These extra-ribosomal functions of hRpS3 are induced by its interactions many signaling molecules, which result in different post-translation modifications that confer different abilities to hRpS3 (13-19). hRpS3 is phosphorylated by IKKβ on Serine 209, which is important for nuclear localization (7). In response to DNA damage, hRpS3, phosphorylated by protein kinase C-δ (PKCδ) at serine 6 (S6) and Threonine 221 (T221) residues, translocates into the nucleus for repair (11,12,20). Moreover, hRpS3 is also phosphorylated by extracellular signal-regulated kinase 1 (ERK1), which is a MAP kinase that plays an important role in the regulation of cell growth (11,21,22). ERK1 phosphorylates hRpS3 on Threonine 42 (T42) (11,22); moreover, this phosphorylation is necessary for the nuclear translocation of hRpS3 in response to DNA damage (21). Threonine 70 (T70) of hRpS3 is phosphorylated by Akt, which promotes its nuclear translocation, thereby preventing hRpS3-induced apoptosis by inhibiting its interaction with E2F1 (13,20).

In this study, we showed that Cdk2 also interacts with hRpS3. Moreover, using bioinformatics tools (NetPhos2.0, www.cds.dtu.dk/services/NetPhos; KinasePhos, kinasephos.mbc.nctu.edu.tw), we found eight sites for putative Cdk2-mediated phosphorylation in hRpS3. We also found that Cdk2 phosphorylates S6 and T221 of hRpS3. Cdk2 is active in S phase, and our in vitro kinase assay revealed that
hRpS3 phosphorylation in cells arrested in the S-phase was more than two folds that in asynchronous control cells. Cdk2 phosphorylates the non-ribosomal form of hRpS3, and this phosphorylation is important for nuclear transport of hRpS3. We also observed that knockdown of hRpS3 delays cell cycle progression. Our findings demonstrate that hRpS3 is involved in cell cycle regulation and that this function may be linked to DNA damage repair.

Materials and methods

Cell lines. Human embryonic kidney cells (HEK293) were grown in Dulbecco’s Modified Eagle’s medium (DMEM; Invitrogen Life Technologies, Carlsbad, CA, USA) containing 10% fetal bovine serum (FBS, Invitrogen Life Technologies) and 1% penicillin-streptomycin solution (Sigma-Aldrich, St. Louis, MO, USA) at 37°C in a 5% CO₂ incubator.

Plasmid construction. Gene fragments corresponding to cDNA coding regions of human RpS3 (Accession no. NM_001005) and Cdk2 (accession no. NM_001798) were amplified using PCR. A Cdk2 fragment was inserted into the EcoRI and SalI sites of pCMV Tag2C (Stratagene, La Jolla, CA, USA), and hRpS3 fragments were inserted into the BamHI and Xhol sites of pCMV Tag3A (Stratagene). All constructs were confirmed by restriction enzyme mapping and DNA sequencing.

pDNA transfection. HEK293 cells were seeded in 6-well plates at a density of 1x10⁴ cells per well and incubated for 24 h before transfection. Recombinant plasmid DNAs were transiently transfected into 80% confluent HEK293 cells using Lipofectamine® 2000 reagent (Invitrogen Life Technologies) according to the manufacturer's protocol. After 24 h of incubation with the plasmids, the cells were collected and lysed in lysis buffer (50 mM Tris-HCl, pH 8.0; 100 mM NaCl; 5 mM EDTA; 1 mM NaN₃; 1 mM NaVO₃; 1% Nonidet P-40; 10 µg/ml of PMSF; and protease inhibitor cocktail) for 40 min at 4°C. After centrifugation, supernatants were collected and used in immunoblot analysis and immunoprecipitation (IP).

GST pull-down assay. For GST pull-down assays, GST-Cdk2 was adsorbed onto glutathione-Sepharose 4B beads (GE Healthcare, Bucks, UK). Histone-purified hRpS3 protein was then incubated with GST or GST fusion protein in binding buffer (50 mM Tris-HCl, pH 7.5; 150 mM NaCl; 1 mM EDTA; 0.3 mM DTT; 0.1% NP-40; and protease inhibitor cocktail). The binding reaction was performed for 3 h at 4°C; subsequently, the beads were washed and bound proteins were subjected to SDS-PAGE analysis. Proteins were detected by immunoblot analysis.

Co-IP. HEK293 cells were seeded 5x10⁴ cells/60-mm² dish and cultured 80-90% confluency before transfection. Lysates were collected from cells transiently transfected with different sets of plasmid DNAs and incubated for 3 h with described antibodies at 4°C with gentle rotation. Antibodies and bound proteins were incubated with protein A/G-Sepharose beads (Santa Cruz Biotechnology, Inc., Santa Cruz, CA, USA) at 4°C for 12 h. Samples were precipitated with centrifugation, washed with PBS, and mixed with gel loading buffer.

Immunoprecipitated samples were resolved on SDS-PAGE gel and subjected to immunoblot analysis.

Immunoblot analysis. Proteins resolved on 8-12% SDS-PAGE gel were transferred onto polyvinylidene difluoride (PVDF) membranes (Pall Corporation, East Hill, NY, USA). The membranes were blocked with 5% non-fat dried milk in TBS-T (TBS with 0.05% Tween-20) for 20 min at room temperature. The membranes were incubated with primary antibodies against anti-hRpS3, p-Ser hRpS3, p-Thr hRpS3 (Cell Signaling Technology Inc., MA, USA), anti-Cdk2, Cdk1, cyclin B1, cyclin E1 (Santa Cruz Biotechnology). After appropriate washing, the membranes were incubated with horseradish peroxidase (HRP)-conjugated secondary antibodies (Santa Cruz Biotechnology). Protein bands were detected using ECL Western blotting detection reagents (Thermo Fisher Scientific, Waltham, MA, USA).

Cell cycle synchronization. To arrest cells the in the S-phase, cells transfected for 24 h were treated with 3 mM thymidine (an inhibitor of DNA synthesis) for 18 h (23,24). Then, the cells were washed with medium and incubated in fresh medium for 2 h. To collect G2/M phase cells, the cells were treated with 0.1 µg/ml nocodazole (a mitotic inhibitor) for 12 h (23,24). An aliquot of cells was stained with propidium iodide (PI; Sigma-Aldrich) and analyzed by fluorescence-activated cell sorting (FACS).

FACS analysis. Cells were trypsinized and centrifuged for 10 min. The collected cells were fixed with 70% ethanol for 30 min, followed by centrifugation. After washing with ice-cold PBS, the cell pellets were resuspended in 0.5 ml PBS containing 50 µg/ml PI and 100 µg/ml RNase (Sigma-Aldrich). After 30 min of incubation in the dark, cell complex was estimated using a minimum of 10,000 cells per sample and was analyzed using a flow cytometer. Fluorescence emitted from the PI-DNA was analyzed using Cell Quest Alias software (BD Biosciences, Rockville, MD, USA).

In vitro Cdk2 kinase assay. After incubation of cell lysates with Cdk2-specific monoclonal antibody (5 µg), samples were incubated with 50 µl protein A/G-Sepharose beads at 4°C for 12 h. Immune complexes were washed with washing buffer (1:1 mixture of lysis buffer and PBS) and kinase reaction buffer (20 mM Tris-HCl, pH 7.4; 15 mM MgCl₂; and 1 mM DTT). After washing, samples were added to a mixture of 1 µg substrate (histone H1 or hRpS3), 200 µM ATP, and 5 µCi [³²P-γ] ATP in 30 µl of reaction buffer. Reactions were carried out for 25 min at 30°C and terminated by addition of SDS-PAGE sample buffer. Samples were subjected to SDS-PAGE, and phosphorylation was detected by autoradiography.

hRpS3 siRNA. hRpS3 siRNA oligonucleotides targeting the sequences 5'-GGGUCCUCUUGAUCUGCAATT-3' and 5'-UGCGAGUACUGGACCCTT-3' were purchased from Santa Cruz Biotechnology. Cells were transfected with hRpS3 siRNAs (5 nM each) using Lipofectamine® 2000 (Invitrogen Life Technologies). Following 24 h of transfection, the level of hRpS3 was determined using anti-hRpS3 antibody
A scrambled siRNA [green fluorescent protein (GFP) siRNA, 5'-GGGCACAAGCUGGAGUACAAC UAC A-3'] was used as the control.

**Statistical analysis.** All experiments were performed at least three times for statistical analysis. To determine the significance of the differences between samples, we performed the Student’s t-test using Microsoft Excel. Data were expressed as mean ± standard errors, and *P*<0.05 (paired two-tailed t-test, *P*<0.01, *P*<0.001) was considered to indicate statistical significance.

**Results**

**Cdk2 physically interacts with hRpS3.** To determine whether hRpS3 interacts with Cdk2, co-IP was conducted using HEK293 cells transfected with FLAG empty vector or with FLAG-tagged Cdk2 (FLAG-Cdk2) and c-Myc-tagged hRpS3. After 24 h incubation, cells were lysed and immunoprecipitated with anti-RpS3 antibody. The co-IP results indicated that Cdk2 interacted with ectopically expressed hRpS3 (Fig. 1A). A similar co-IP assay with antibodies against endogenous Cdk2 and hRpS3 also revealed that Cdk2 interacted with hRpS3 (Fig. 1B). Rabbit IgG was used as a control. To determine whether hRpS3 interacts directly with Cdk2, we conducted a GST pull-down assay (Fig. 1C). His-hRpS3 and GST-Cdk2 were generated and purified before conducting the assay. Purified His-hRpS3 interacted with GST-Cdk2 but not with GST alone, indicating that Cdk2 interacts directly with hRpS3.

Next, we investigated whether the interaction between Cdk2 and hRpS3 required Cdk2 activation. For this, we prepared FLAG-Cdk2 and a mutant of Cdk2 with modifications at T160 (T160A; Cdk2 phosphorylated at T160 is the active form), and transfected these into HEK293 cells (Fig. 1A; lanes 2 and 3). Co-IP revealed that substitution of threonine with alanine inhibited Cdk2 phosphorylation at the site. These results show that hRpS3 interacts only with the active form of Cdk2 (Fig. 1A; lanes 2 and 3).

**hRpS3 was phosphorylated by Cdk2 during the S-phase.** It is already known that the highest activity of Cdk2 is observed in the S-phase (1). Therefore, we investigated the association between hRpS3 phosphorylation and Cdk2 activity. Co-IP of asynchronous and synchronous cells arrested at the S and G2/M phases was conducted using anti-hRpS3 antibody (Fig. 2A). Consistent with the results in Fig. 1A, endogenous hRpS3 was found to interact with Cdk2. Moreover, Fig. 2A shows that this interaction was higher in the S-phase than in the G2/M phase. This result indicates that hRpS3 interacts with the active form of Cdk2 in the S-phase.

Cdk2 phosphorylates its substrate at the serine/threonine residues. Therefore, the possibility of hRpS3 being a substrate of Cdk2 was examined. We have previously demonstrated that Cdk1 interacts and phosphorylates hRpS3, and this phosphorylation increased in the G2/M phase (21). Thus, we conducted a

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**Figure 1. Cdk2 interacts with hRpS3 in vitro and in vivo.** (A) HEK293 cells were transfected with c-Myc-hRpS3 and empty vector, FLAG-Cdk2, or FLAG-Cdk2-T160A. Whole cell lysates were collected and immunoprecipitated with anti-FLAG antibody and protein A/G Sepharose beads. Immunoblot analysis was performed using anti-hRpS3 and anti-FLAG antibodies. (B) HEK293 cell lysates were immunoprecipitated with anti-Cdk2 antibody, and immunoblot analysis was performed with anti-Cdk2 and anti-hRpS3 antibodies. Rabbit immunoglobulin G (IgG) was used as a negative control. (C) Pull down of purified His-hRpS3 with GST-Cdk2 bound to glutathione sepharose beads. Lane 1 contains purified His-hRpS3 only (input). Lanes 2 and 3 are the pull-down pellets from GST and GST-Cdk2, respectively.
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Co-IP assay using asynchronous and synchronous cells (arrested at the S and G2/M phases). Whole cell lysates were subjected to immunoprecipitation with anti-hRpS3 antibody. Following immunoblot analysis using anti-hRpS3, hRpS3 was detected in both input and immunoprecipitation samples (Fig. 2B). Phospho-serine (p-Ser) levels in the immunoprecipitated sample were higher in the cells arrested at the S-phase than in the cells arrested at the G2/M phase. In contrast, an intense band of phospho-threonine (p-Thr) was detected in cells in the G2/M phase. Since Cdk2 is active in the S-phase, these results indicate that Cdk2 phosphorylates the serine of hRpS3 in the S-phase.

Fig. 1A shows that hRpS3 binds with the active form of Cdk2. To demonstrate the ability of Cdk2 to phosphorylate hRpS3, we conducted an in vitro kinase assay (Fig. 2C). Cells were synchronized at the G2/M- and S-phases using nocodazole and thymidine, respectively. Cell synchronization was confirmed by FACS analysis (Fig. 2C, upper histogram). The in vitro kinase assay was conducted using hRpS3 and histone H1. Cdk2 was isolated from the synchronized samples by immunoprecipitation with anti-Cdk2 antibody. histone H1, a known substrate of Cdk2, was used as a positive control. Autoradiography revealed phosphorylation of both histone H1 and hRpS3 by Cdk2. Moreover, phosphorylation of hRpS3 by Cdk2 increased in the S-phase.

Cdk1 and Cdk2 are closely related proteins with high similarity. Thus, there might be a cross-reaction among the antibodies used for co-IP. To exclude this possibility, we tested the specificity of anti-Cdk1 and anti-Cdk2 antibodies by immunoprecipitation. The first panel in Fig. 2D shows that the antibody against Cdk1 specifically recognized Cdk1 in input, but it did not detect Cdk1 or Cdk2 in the immunoprecipitation assay. The same result was obtained for the anti-Cdk2 antibody (Fig. 2D, second panel). Thus, the result for the in vitro Cdk1/2 kinase assay was specific for the indicated protein.

Cdk2 phosphorylates hRpS3 at specific residues. Based on the analysis of hRpS3 amino acid sequence using bioinformatics tools (NetPhos2.0, www.cds.dtu.dk/services/NetPhos; KinasePhos, kinasephos.mbc.nctu.edu.tw), eight possible sites for phosphorylation by Cdk2 were identified (Fig. 3A) (18). To observe which amino acid is phosphorylated by Cdk2, hRpS3 constructs containing a serine/threonine-to-alanine substitution were created and used as the substrate in the in vitro kinase assay. As shown in Fig. 3B, phosphorylation was significantly
lower in the hRpS3-S6A and -T221A mutants than in the wild type. This result suggests that Cdk2 phosphorylates the S6 and T221 residues of hRpS3. Coomassie blue staining in the lowest panel was used as a loading control.

hRpS3 knockdown induced phosphorylation of Cdk2 and P53. The physiological effect of interaction between Cdk2 and hRpS3 on the cell cycle was observed by knockdown of hRpS3. For hRpS3 knockdown, HEK293 cells were transfected with varying concentrations siRpS3. Cells transfected with siGFP were used as control. As shown in Fig. 4A, hRpS3 knockdown was successful in all transfected cells. Whole cell lysates were subjected to immunoblotting for the detection of phospho-Cdk2 (p-Cdk2) (T14, Y15; inactive form), phospho-P53 (p-P53), and P53. The expression of p-Cdk2 (T14, Y15) and p-P53 increased with increasing concentration of siRpS3. Phosphorylation of P53 induced production of p21, an inhibitor of Cdk2 activation (9). Therefore, increased phosphorylation of P53 and Cdk2 (T14, Y15) is indicative of delayed cell cycle progression. This result suggests that hRpS3 is involved in cell cycle regulation by inducing phosphorylation of P53.

Cell cycle delay was induced by knockdown of hRpS3. The effect of hRpS3 knockdown was examined in synchronous and asynchronous cells by FACs and immunoblotting. To synchronize cells at the late G1 phase, they were treated with aphidicolin (Aph). After 15 h, Aph was released into fresh medium, and cells were harvested at different release times such that they would synchronize and enter the S-phase in 2-4 h (22). Cells enter the G2-phase after 8 h of release.

We observed a slight difference in cell cycle progression in siRpS3-transfected cells and siGFP-transfected control cells (Fig. 4B). In the hRpS3 knockdown cells, the S and G2/M phases were delayed compared with that in siGFP-transfected control cells. siGFP-transfected cells entered the S-phase at 0-3 h and the G2/M phase at 5 h after release of Aph into fresh media. In contrast, the siRpS3-transfected cells were still in the G1-phase at 0-3 h, and entered the G2/M phase at 5 h after Aph release. Notably, at 5 h after Aph release, the number of siGFP-transfected cells that entered the G2/M phase was considerably higher than that of siRpS3-transfected cells. Thus, the hRpS3 knockdown cells showed delayed cell cycle progression in each phase compared with the siGFP-transfected control cells. These results suggest that hRpS3 is required for cell cycle progression.

Further, to elucidate the mechanism through which hRpS3 regulates cell cycle progression, we examined the expression of cell cycle-related proteins using immunoblotting (Fig. 4C). In the siGFP-transfected cells, the level of cyclin B1 (binds to Cdk1 and involved in regulating G2/M transition) significantly decreased after 10 h of Aph release (Fig. 4C). Additionally, in hRpS3 knockdown cells, the level of cyclin B1 significantly decreased after release for 12 h. These results indicate that hRpS3 knockdown induced delay of G2/M progression. The level of cyclin E1 (binds to Cdk2 and required for progression to the S-phase) in siGFP-transfected and siRpS3-transfected cells decreased after 8 h of release. The level of cyclin E1 in siGFP-transfected cells increased significantly after 12 h release, whereas it was not detected in siRpS3-transfected cells after 8 h of release. This expression pattern of cyclin E1 showed that hRpS3 knockdown delayed the progression to S-phase. These findings suggest that hRpS3 is regulates cell cycle progression modulating the expression of cell cycle-related proteins (cyclins B and E1).

Cell cycle progression is mainly regulated by CDKs. CDK activity increased or decreased as the cell cycle progressed. Among the CDKs, the Cdk1/cyclin B complex is known to control G2/M transition, while Cdk2/cyclin E1/A complexes regulate G1/S and S/G2 transitions (9,11,21,25). We found that the levels of Cdk2/cyclin E1 and Cdk1/cyclin B1 were decreased in hRpS3 knockdown cells. Thus, hRpS3 knockdown delayed cell cycle progression by downregulating the proteins involved in cell cycle regulation.

Discussion

The cell cycle is regulated by the CDK family of proteins (26). CDKs are the catalytic subunits of a large family of heterodimeric serine/threonine protein kinases that regulate cell cycle progression (27). The catalytic activity of CDKs requires the binding of a regulatory subunit, cyclin, which is synthesized and degraded during each cycle (28). The catalytic activity of Cdk1 requires the binding of the cyclin B1 (27). When Cdk1/cyclin B1 activity is maximal during G2-M phase, this complex phosphorylates a number of proteins that regulate several cellular events (29). Cdk2 plays a role in G1/S transition and initiation of DNA synthesis in the S-phase by forming complexes with cyclins E and A (1,5,7).

DNA damage repair and cell cycle progression are tightly regulated. As a ribosomal protein, hRpS3, is also known to

Figure 3. S6 and T221 of hRpS3 were phosphorylated by Cdk1/2. (A) The amino acid sequence of hRpS3. hRpS3 has 8 possible phosphorylation sites (red), NLS, nuclear localization signal (yellow, amino acids 7-10); KH, K homologue domain (blue, amino acids 42-111). (B) Autoradiograph from kinase assays carried out using Cdk1/2 immunoprecipitated lysates of HEK293 cells expressing wild type or various point mutants of hRpS3. Equal amounts of samples were analyzed by coomassie staining.

Figure 4. C). In the siGFP -transfected cells, the level of
perform other extra-ribosomal functions, such as in DNA repair, via its N-glycosylase activity and by releasing 8-oxoG from a DNA lesion (30, 31). Furthermore, hRpS3 binds and stimulates the activity of uracil-DNA glycosylase (UNG), as well as the base excision repair (BER) enzymes, hOGG1 and APE/Ref-1 (32, 33). Previous reports have shown that hRpS3 can be translocated into the nucleus for regulation of cell cycle progression, similar to some other ribosomal proteins such as ribosomal protein L6 and ribosomal protein S13, which are involved in the regulation of cyclin E and the promotion of cell growth (34, 35). Therefore, different functions of hRpS3 are believed to result from the post-translational modifications that result from its interaction with different molecules.

Previously, we reported that hRpS3 physically interacts with and is phosphorylated by Cdk1, especially, in the G2/M phase. This phosphorylation occurs at T221 and is important for the nuclear translocation of hRpS3 (21). Here, we studied the interaction between hRpS3 and Cdk2 (Fig. 1). Interestingly, the interaction between hRpS3 and Cdk2 was increased in the S-phase. Previous observations prompted us to determine whether hRpS3 is a substrate of Cdk2. We conducted a Cdk2 kinase assay in asynchronous and synchronous cells (arrested in the S and G2/M phases) using hRpS3 and purified histone H1 as substrates (Fig. 2C) and observed that hRpS3 phosphorylation by Cdk2 occurs in the S-phase. These results revealed that hRpS3 interacts with Cdk2 and is phosphorylated at a serine residue by Cdk2. Subsequently, based on a previous report and the hRpS3 amino acid sequence analysis performed in this study, eight phosphorylation sites were identified (Fig. 3A). The serine/threonine residues in these putative phosphorylation sites were substituted by alanine and a Cdk1/2 kinase assay was performed (Fig. 3B). S6 and T221 of hRpS3 were phosphorylated by Cdk2, and T221 was phosphorylated by Cdk1.

Knockdown of hRpS3 induced phosphorylation of P53, which functions upstream of Cdk1/2, resulting in a delay in cell cycle progression and cell cycle arrest (Fig. 4A). Furthermore, we observed that knockdown of hRpS3 induced delay in cell cycle progression and downregulation of cyclins E1 and B1 (Fig. 4B and C). Our results show that the hRpS3 participates in cell cycle regulation by modulating the expression of cell cycle-related proteins.

In summary, we have shown that hRpS3 is involved in the Cdk2-mediated regulation of cell cycle progression. Our findings indicate a possible link between DNA damage repair and cell cycle progression. Further studies are required to determine other phosphorylation sites of hRpS3 that are linked to cell cycle progression and to elucidate the relationship between the mechanisms involved in DNA damage repair and cell cycle control.
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