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A genetic screen identifies the Triple T complex required for DNA damage signaling and ATM and ATR stability

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In response to DNA damage, cells activate a complex signal transduction network called the DNA damage response (DDR). To enhance our current understanding of the DDR network, we performed a genome-wide RNAi screen to identify genes required for resistance to ionizing radiation (IR). Along with a number of known DDR genes, we discovered a large set of novel genes whose depletion leads to cellular sensitivity to IR. Here we describe TTI1 (Tel two-interacting protein 1) and TTI2 as highly conserved regulators of the DDR in mammals. TTI1 and TTI2 protect cells from spontaneous DNA damage, and are required for the establishment of the intra-S and G2/M checkpoints. TTI1 and TTI2 exist in multiple complexes, including a 2-MDa complex with TEL2 (telomere maintenance 2), called the Triple T complex, and phosphoinositide-3-kinase-related protein kinases (PIKKs) such as ataxia telangiectasia-mutated (ATM). The components of the TTT complex are mutually dependent on each other, and act as critical regulators of PIKK abundance and checkpoint signaling.

[Keywords: TTI1; TEL2; TTI2; PIKK; TTT complex; IR sensitivity]

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Organisms experience significant amounts of DNA damage, which threatens their survival on a daily basis. To address this damage and optimize survival, cells evolved the DNA damage response (DDR), a complex signal transduction network that activates transcription and DNA repair, and coordinates cell cycle transitions in order to maintain genomic stability. The central regulators of the DDR are ataxia telangiectasia-mutated (ATM) and ataxia-telangiectasia and Rad3-related (ATR), two members of the mammalian phosphoinositide-3-kinase-related protein kinase (PIKK) family. ATM is critical for the cellular response to DNA double-strand breaks (DSBs), and mutations in ATM lead to the neurodegenerative and cancer predisposition disease ataxia telangiectasia [Lavin 2008]. ATR is activated by replication stress, and is an essential gene in budding yeast and mice. In addition, a hypomorphic mutation in ATR results in Seckel syndrome, a developmental disorder characterized by growth retardation and microcephaly [Cimprich and Cortez 2008]. In mammals, there are four additional members of the PIKK family that have diverse functions. They include DNA-PKcs (the DNA-dependent protein kinase catalytic subunit), which is crucial for DNA repair of DSBs by nonhomologous end-joining, and SMG1 (suppressor with morphological effect on genitalia 1), which regulates nonsense-mediated decay of aberrant mRNA. In addition, this family includes mTOR (mammalian target of rapamycin), which controls cell growth in response to environmental cues, mitogenic signals, and nutrient availability, and TRRAP (the catalytically inactive transformation/transcription domain-associated protein), a regulator of gene expression via its association with histone acetyltransferase complexes.

Recently, a common interacting protein between all six of the mammalian PIKKs that affects their stabilities was identified as TEL2 (telomere maintenance 2) [Takai et al. 2007]. TEL2 was originally identified in Saccharomyces cerevisiae in a screen for mutants that altered telomere length [Lustig and Petes 1986]. It was subsequently identified in Caenorhabditis elegans in a screen for mutants that altered telomere length [Lustig and Petes 1986]. It was subsequently identified in Caenorhabditis elegans in a screen for mutants that altered telomere length [Lustig and Petes 1986]. It was subsequently identified in Caenorhabditis elegans in a screen for mutants that altered telomere length [Lustig and Petes 1986]. It was subsequently identified in Caenorhabditis elegans in a screen for mutants that altered telomere length [Lustig and Petes 1986]. It was subsequently identified in Caenorhabditis elegans in a screen for mutants that altered telomere length [Lustig and Petes 1986]. It was subsequently identified in Caenorhabditis elegans in a screen for mutants that altered telomere length [Lustig and Petes 1986]. It was subsequently identified in Caenorhabditis elegans in a screen for mutants that altered telomere length [Lustig and Petes 1986]. It was subsequently identified in Caenorhabditis elegans in a screen for mutants that altered telomere length [Lustig and Petes 1986]. It was subsequently identified in Caenorhabditis elegans in a screen for mutants that altered telomere length [Lustig and Petes 1986]. It was subsequently identified in Caenorhabditis elegans in a screen for mutants that altered telomere length [Lustig and Petes 1986]. It was subsequently identified in Caenorhabditis elegans in a screen for mutants that altered telomere length [Lustig and Petes 1986]. It was subsequently identified in Caenorhabditis elegans in a screen for mutants that altered telomere length [Lustig and Petes 1986]. It was subsequently identified in Caenorhabditis elegans in a screen for mutants that altered telomere length [Lustig and Petes 1986]. It was subsequently identified in Caenorhabditis elegans in a screen for mutants that altered telomere length [Lustig and Petes 1986]. It was subsequently identified in Caenorhabditis elegans in a screen for mutants that altered telomere length [Lustig and Petes 1986].
heavily implicated in the ATR signaling pathway and the S-phase checkpoint (Collis et al. 2007, 2008; Shikata et al. 2007; Danielsen et al. 2009). The connection between TEL2 and the PIKK family helps explain the pleiotrophic nature of the phenotypes of TEL2 mutants in different organisms. However, it is not known whether TEL2 works alone or is a part of a larger machine involved in PIKK regulation. The determination of the precise composition of the TEL2 complex will be critical to the elucidation of its mechanism of action in promoting PIKK activity.

In this study, we report the results of a genome-wide RNAi screen for genes required for IR resistance in which we identify a novel regulator of the DDR, TTI1 (Tel two-interacting protein 1), which plays a critical role in DNA damage resistance and the establishment of cell cycle checkpoints. We find that TTI1 and its associated protein, TTI2 (C8ORF41), are required for DDR signaling through control of the protein levels of ATM, ATR, and a group of related PIKKs, suggesting a connection to TEL2. TTI1 and TTI2 associate physically with TEL2 to form a conserved trimeric complex called the Triple T complex. Interestingly, TTI1 and TTI2 are also required for maintaining the protein levels of TEL2. However, expression of exogenous TEL2 cannot restore DNA damage resistance in TTI1-depleted cells, indicating a specific role for TTI1 in the DDR. We propose that the TTT complex is a master regulator of PIKK abundance and DDR signaling.

Results
A genome-wide RNAi screen for genes required for IR resistance

We recently developed retroviral barcoded, microRNA-based shRNA libraries targeting the entire human genome that can be used in complex pools of shRNAs that can be deconvoluted by microarray technology (Paddison et al. 2004; Silva et al. 2005, 2008; Schlabach et al. 2008). In this study, we performed a genome-wide loss-of-function screen to identify genes whose inhibition leads to sensitivity to ionizing radiation (IR) in U2OS cells (Fig. 1A). We screened U2OS cells with 74,905 retroviral shRNAs targeting 32,293 unique human transcripts (including 19,542 RefSeq transcripts) in six pools of 13,000 shRNAs per pool and a “focused” sublibrary consisting

Figure 1. A genome-wide RNAi screen to search for novel DDR genes. (A) Schematic of the pool-based shRNA screen for regulators of the DNA DSB response in mammalian cells. shRNA pools were deconvoluted using microarray hybridization. (B) The screen behavior of a representative shRNA pool of 12,965 shRNAs. shRNAs are graphed based on their mean-normalized log2 Cy5/Cy3 ratio, with a ratio of less than −1 indicating a decrease in the relative abundance of the shRNA greater than twofold, and a ratio of >1 representing an increase in the relative abundance of the shRNA greater than twofold. The population of shRNAs causing IR sensitivity is shaded in red, and IR resistance is shaded in green. (C, D) Gene ontology analysis of the candidate genes required for IR resistance. (C) Significance refers to the −log(P-value) determined by the right-tailed Fisher’s exact test. Threshold is at 1.25 = −log (P = 0.05). The analysis includes 760 genes from Supplemental Table S1 whose gene symbols were identified by Ingenuity’s Knowledge Base. (D) The candidate genes required for IR resistance were classified into several biological processes by the PANTHER program. The analysis includes 760 genes from Supplemental Table S1 whose Entrez Gene IDs were identified by PANTHER.
of 8000 shRNAs against kinases, phosphatases, ubiquitin ligases, and genes implicated in cancer. The shRNA pools in MSCV-PM were introduced into U2OS cells at a multiplicity of infection [MOI] of 1 following a protocol described previously [Schlabach et al. 2008; Luo et al. 2009; Wang et al. 2009]. The cells were puromycin-selected, treated with 5 Gy IR [or left untreated], and propagated for ~2 wk, and the end samples were collected for processing for pool deconvolution. Using microarray hybridization, we monitored the change in relative abundance of each shRNA from the untreated cells to the IR-treated cells. We identified both a set of shRNAs that dropped out of the IR-treated cells and a set that was enriched in the IR-treated cells [Fig. 1B]. Actual screen data are plotted as a normalized log2 [Cy5/Cy3] ratio. Our screen identified 850 shRNAs corresponding to 813 genes that are required for IR resistance in U2OS cells [Supplemental Table S1; see the Materials and Methods for the criteria used to select hits]. The candidate IR-enriched shRNAs are listed in Supplemental Table S2.

To better understand our IR sensitivity data set, we performed a bioinformatics analysis of the genes targeted by the shRNAs. We used Ingenuity Pathway Analysis [Ingenuity Systems, http://www.ingenuity.com] to determine the significance of certain functional categories in our list by determining their enrichment relative to the total number of genes in their respective categories [Fig. 1C]. The DNA replication, recombination, and repair category and the cell cycle category showed a highly significant enrichment in our data set. Enrichment was also found for genes involved in cancer, protein degradation, post-translational modifications, and cell signaling [Fig. 1C]. Unexpectedly, the category that showed the most enrichment was RNA post-transcriptional modification, which includes genes involved in spliceosome assembly, RNA splicing, polyadenylation of mRNA, and processing of mRNA and RNA. This functional group was also highly enriched in the ATM/ATR substrates identified in our proteomics screen [Matsuoka et al. 2007]. We also annotated the list of candidate IR sensitivity genes in the gene ontology format. Using PANTHER, we found many biological processes to be enriched, including nucleic acid metabolism, protein modification, signal transduction, and cell cycle [Fig. 1D, Thomas et al. 2003; Mi et al. 2007]. Importantly, we found a significant number of genes that have remained unclassified, indicating the potential of this screen to uncover novel DDR genes.

**DDR and cell cycle genes uncovered in the screen**

Many genes with a known link to the DDR, as well as genes that function in DNA replication and cell division, were identified in the screen. We found a subset of genes for which multiple independent shRNAs were depleted from the IR-treated population greater than twofold that included several known DDR regulators as well as genes not previously linked to the DDR [Fig. 2A, Supplemental Table S1 [all multiple shRNA hits are highlighted in pink]; see below]. Our screen scored four shRNAs against the ATM kinase, which is central to the entire DDR. In addition, four shRNAs scored against DNA-PKcs, a kinase involved in nonhomologous end-joining DNA repair. Also, UBC13 and NBN1/MERIT40 both scored with multiple shRNAs. UBC13 and NBN1 are both required for recruitment of BRCA1 to sites of DNA damage. NBN1 was identified in a preliminary version of this screen [Wang et al. 2009]. We used the Ingenuity pathway analysis software to create a network of genes known to be involved in the DDR, DNA replication, or cell cycle progression [Fig. 2B]. In Figure 2B, the genes marked in red indicate that at least one shRNA against this gene scored in the screen [Supplemental Table S3; see the Materials and Methods for a description of how this list was generated]. To validate shRNAs using an independent assay, shRNAs were tested using the multicolor competition assay [MCA] [Smogorzewska et al. 2007]. In brief, uncolored U2OS cells expressing a control shRNA were mixed in a 50:50 ratio with U2OS cells expressing dsRed and the shRNA of interest, and were either left untreated or treated with IR. Eight days after IR, the percentage of dsRed-expressing cells was measured by FACS, and IR-treated cells were normalized to untreated cells to correct for proliferation differences. We used this viability assay to validate several genes that have been linked previously to DNA repair, DNA replication, or cell cycle [Fig. 2C]. A full list of validated shRNAs using MCA is shown in Supplemental Table S4, and, overall, 115 out of 292 tested shRNAs validated for a rate of ~40%. As mentioned above, two of the strongest hits were ATM and DNA-PK, and both validated with multiple shRNAs in our independent viability assay. In addition, the DDR kinase ATR, UBC13, and NBN1 validated using MCA. We also identified C9orf80, which was characterized recently using proteomics as a member of a trimeric complex containing SSB1/2 and INTS3 that is important for the DDR [Huang et al. 2009; Li et al. 2009; Skaar et al. 2009]. Other validated hits include Topoisomerase 1 [TOP1], the replication licensing factor MCM6, and NIPBL, a gene involved in sister chromatid cohesion and mutated in Cornelia de Lange syndrome [Tonkin et al. 2004]. We also found genes whose yeast orthologs have been shown to be required for DNA damage resistance, including RBBP4/CAF-1 p48 subunit/RRAP48 [S. cerevisiae] MS1/CAC3 and RNF20 [S. cerevisiae] BRE1A [Kaufman et al. 1997; Giannattasio et al. 2005; Linger and Tyler 2005].

**TTI1 is required for DNA damage resistance**

Our screen identified a previously uncharacterized ORF, KIAA0406 [Entrez GeneID: 9675; accession no.: NM_014657], which we will refer to as TTI1 for reasons described below. TTI1 is highly conserved throughout evolution, with orthologs in mice, chickens, flies, frogs, fish, plants, and yeast [Supplemental Fig. S1]. We identified one shRNA targeting TTI1 that scored greater than twofold reduction in two independent microarray hybridizations [Fig. 3A; see the Materials and Methods for details on hybridizations]. To validate that TTI1 depletion leads to increased IR sensitivity, we used the MCA described above with multiple shRNAs and siRNAs [Fig.
3B, D). The TTI1 shRNA that scored in the screen (#3) depleted TTI1 mRNA [Fig. 3C] and led to decreased cell viability in response to IR, mitomycin C (MMC), and camptothecin (CPT) [Fig. 3B,F]. We further tested two additional shRNAs and four siRNAs that target distinct regions of the TTI1 sequence [Fig. 3B,D]. Both shRNAs and three of four siRNAs gave DNA damage sensitivity to IR, ultraviolet (UV), or both, which correlated with the degree of TTI1 protein depletion [Fig. 3B–E]. An additional shRNA, TTI1 #4, also led to strong TTI1 depletion and DNA damage sensitivity [Fig. 7D,E, below]. The effect of ATM depletion is shown in each case for comparison [Fig. 3B,D, Supplemental Fig. S4]. Furthermore, when we reintroduced a TTI1 cDNA lacking the shRNA #3 target site, which exists in the 3’ untranslated region [UTR] of the endogenous gene, into cells treated with TTI1 shRNA #3, the shRNA-resistant cDNA reversed the DNA damage sensitivity of the TTI1 shRNA [Fig. 3F]. To further confirm our results using the MCA assay, we performed colony survival assays and found that TTI1 depletion led to significant IR sensitivity at multiple doses of damage [Supplemental Fig. S3]. Thus, TTI1 is required for resistance to DNA damage.

**TTI1 controls the G2/M and intra-S-phase DNA damage checkpoints**

To further explore the role of TTI1 in the DDR, we monitored the ability of cells depleted of TTI1 to arrest the cell cycle in S and G2 phases in response to IR. To test G2 arrest, TTI1-depleted cells were treated with 10 Gy IR, and, after 1 h, were treated with nocodazole for 17 h to allow accumulation in mitosis, measured by phospho-S10 histone H3 staining. TTI1-depleted cells were unable...
CPT was monitored. Error bars represent the standard deviation of three replicates. For TTI1 shRNA #3, and, after 2 d, were mixed with control cells for MCA. Viability in response to 3 Gy of IR, 75 nM MMC, or 5 nM CPT was monitored. Error bars represent the standard deviation of three replicates. ( Supplemental Fig. S6). This finding helps to explain the IR sensitivity phenotype and checkpoint defects seen in TTI1-depleted cells.

TTI1 depletion leads to defective DDR signaling

Since TTI1 shRNA-treated cells displayed defective DNA damage-induced cell cycle arrest, we examined DDR signaling itself. We found that IR-induced phosphorylation of several targets of ATM—including CHK2, NBS1, SMC1, and KAP1—was reduced in TTI1-deleted cells (Fig. 5A). In addition, we found that phosphorylation of CHK1 on Ser 317 was very low in TTI1-depleted cells treated with IR or UV compared with a control shRNA (Supplemental Fig. S6). These defects correlate with a strong reduction in ATM and ATR protein levels (Fig. 5A; Supplemental Fig. S6). This finding helps to explain the IR sensitivity phenotype and checkpoint defects seen in TTI1-depleted cells.

TTI1 is required to maintain PIKK protein abundance

Since we observed a profound effect on ATM and ATR protein levels in cells expressing TTI1 shRNAs, we investigated the possibility that TTI1 might also control other PIKKs. In U2OS cells stably expressing a TTI1 shRNA, the levels of the other PIKK family members—including DNA-PKcs, mTOR, SMG1, and TRRAP—were also reduced (Fig. 5B). We did not see an effect on the levels of PI3K subunits p110α or p85α (data not shown). Two independent shRNAs targeting TTI1 were used for this analysis, with shRNA #2 producing greater depletion of TTI1 and a correspondingly stronger effect on PIKK levels. These data suggest that TTI1 may function in other cellular processes in addition to the DDR. Consistent with this, we found that signaling downstream from mTORC2, one of the mTOR-containing complexes, was defective in TTI1-depleted cells [Supplemental Fig. S8; Hresko and Mueckler 2005; Sarbassov et al. 2005].

To investigate the mechanism by which TTI1 controls the levels of the PIKKs, we monitored the effect of TTI1 depletion on PIKK transcripts. TTI1 depletion does not have a significant effect on PIKK mRNA levels at early time points, where we already see strong reduction in PIKK protein [Supplemental Fig. S7A,B]. However, TTI1-depleted cell lines carried under puromycin selection for several days did display a decrease in transcript levels.
These results suggest that the initial decrease in PIKK protein levels occurs post-transcriptionally in the absence of TTI1.

TTI1 interacts with TEL2 and other PIKKs in mammals

The phenotype of TTI1-depleted cells suggests that TTI1 might play a direct role in PIKK stability by associating with PIKKs, reminiscent of the role proposed for TEL2 (Collis et al. 2007; Takai et al. 2007). Immunoprecipitation of Flag-HA-tagged TTI1 revealed the presence of the PIKKs ATM, ATR, DNA-PKcs, SMG1, and mTOR (Fig. 5C). Endogenous ATM and ATR were also present in immunoprecipitates of endogenous TTI1 (Fig. 5D,E). In addition, to further confirm the association of TTI1 with PIKKs, reciprocal coimmunoprecipitations were performed, and both endogenous ATM and Flag-tagged ATM were able to immunoprecipitate endogenous TTI1 (Fig. 5F; Supplemental Fig. S9). Interestingly, it appears that, following IR treatment, several PIKKs show a small reduction in their interaction with TTI1—in particular, those that are activated by DNA damage, ATM, ATR, and DNA-PKcs (Fig. 5C,E).

The effect on PIKK abundance and the physical association with PIKKs strongly suggested a role in the TEL2 pathway. Interestingly, a physical link between PIKKs and fungal TTI1 had been detected previously through mass spectrometry analysis of Tor1/2 complexes. *S. pombe* SPCC622.13c was found to associate with Tor1/Tor2 and, subsequently, Tel2/SPAC458.03 proteins, and was named Tti1 (Hayashi et al. 2007). In addition, a second protein, SPBC1604.17c/Tti2, was also identified as a Tel2-associated protein, and is conserved in humans (C8ORF41; Entrez GeneID: 80185) and other species (Supplemental Fig. S2). Although *S. pombe* Tti1 and Tti2 were found in Tor1/2 and Tel2 immunoprecipitates, their functional relationship to Tel2 and DNA damage...
signaling was not explored, and it is not known whether this interaction is conserved in humans. To examine the latter, Flag-HA-tagged TTI1 expressed in U2OS cells was immunoprecipitated and found to coimmunoprecipitate endogenous human TEL2 [Fig. 6A]. In addition, tagged TEL2 (C-terminal Flag-HA tag) and Myc-tagged TTI1 reciprocally coimmunoprecipitated each other, and this interaction was not significantly altered in response to DNA damage [Fig. 6C; Supplemental Fig. S10]. To examine endogenous protein association, we generated an antibody against a C-terminal peptide in TTI1. This antibody was capable of immunoprecipitating endogenous TTI1 and TEL2 [Fig. 6B]. Immunoprecipitation of human TTI2 also coimmunoprecipitated TEL2 and TTI1 [Fig. 6D]. In addition, we identified TTI1 by mass spectrometry in HA-TTI2 immunoprecipitates [data not shown].

To further confirm that TTI1, TEL2, and TTI2 all interact together as a complex, we examined whether these proteins cofractionate on a sizing column. We found that all three proteins cofractionate on a Superose 6 column in a large complex of molecular weight 2000 KDa [Fig. 6E]. In addition, TTI1 and TEL2 peaked in a smaller complex (>450 kD, fractions 12 and 13), while TTI2 was also present in these fractions. Protein molecular weight standards and where they fractionate on the Superose 6 column are listed in Supplemental Table S5. The fractionation of TTI1, TEL2, and TTI2 did not significantly change if the cells were treated with IR [data not shown]. To determine whether the TTT complex was intact in the peak fractions, we used fractions 2, 12, 13, 15, and 18 as input material for immunoprecipitation with HA antibody-conjugated beads, and found that TEL2 and TTI1 interacted with HA-TTI2 in fraction 2 [Fig. 6F]. Despite significantly less TTT in fraction 2, HA-TTI2 brought down more TEL2 in fraction 2 than in fraction 12. This suggests that TEL2 in fraction 12 is bound much more weakly than in fraction 2, but the TTT1–TTT2 interaction remains strong in fraction 12. Importantly, ATM coimmunoprecipitated only with the TTT complex in fraction 2, suggesting that this fraction contains the active complex, and that, perhaps, PIKKs help it assemble. Altogether, these data suggest that the entire TTT complex is conserved throughout eukaryotic evolution.

**Figure 6.** Human TTI1 and TTI2 associate physically with TEL2. (A) Epitope-tagged TTI1 can coimmunoprecipitate endogenous TEL2. Flag-HA-TTI1 was transiently transfected into 293T cells, and proteins immunoprecipitated with HA antibody-conjugated beads were separated by SDS-PAGE, transferred to PVDF, and immunoblotted with anti-TEL2 antisera or anti-Flag-HRP. (B) An antibody generated against the C terminus of TTI1 coimmunoprecipitated endogenous TTI1 and TEL2. Normal rabbit IgG was used for a control immunoprecipitation. (C) TEL2 can coimmunoprecipitate TTI1 independently of IR treatment. MYC-TTI1 (N-terminal MYC tag) and/or TEL2-FHA (C-terminal Flag-HA tag) were transiently transfected into 293T cells, and proteins were immunoprecipitated with HA antibodies and immunoblotted with anti-MYC or anti-HA antibodies. Thirty-six hours after transfection, cells were either left untreated or treated with 10 Gy of IR for 4 h prior to harvesting. (D) TTI2 interacts with both TTI1 and TEL2. FHA-TTI1 (N-terminal Flag-HA tag) was transiently transfected into 293T cells, and HA-immunoprecipitated proteins were immunoblotted with anti-TEL2 and anti-TTI1 antibodies. (E) TTI1, TEL2, and TTI2 cofractionate in a large-molecular-weight complex on Superose 6. 293 TRex cells containing a doxycycline-inducible Flag-HA-TTI2 were treated with 2 μg/mL doxycycline for 24 h prior to preparation of whole-cell extract. The molecular weight standards and the corresponding fractions they eluted off of the Superose 6 column are listed in Supplemental Table S5. (F) Fractions from extracts run on a Superose 6 column (shown in E) were used as input for immunoprecipitation with HA antibody-conjugated beads.

**All three TTT complex components are required for DNA damage resistance**

To investigate more completely the involvement of the individual TTT complex subunits in the DDR, we assayed viability in response to IR and MMC in cells stably expressing TTI1, TEL2, or TTI2 shRNAs. All shRNAs led to efficient depletion of these components, as monitored by immunoblotting [TTI1 and TEL2] or RT–PCR [TTI2] [Fig. 7A; Supplemental Fig. S14]. Three shRNAs against TEL2 led to decreased survival in response to IR and MMC, with TEL2 #1 giving the strongest phenotype [and most efficient depletion] [Fig. 7D, left panel]. Also, three shRNAs against TTI2 led to decreased survival in response to IR and MMC, with TTI2 #2 having the strongest effect [Fig. 7D, right panel]. The phenotypes of ATM, ATR, and TTI1 shRNAs are shown for comparison. Furthermore, TEL2 and TTI2 shRNAs also led to decreased levels of ATM and ATR, consistent with the decreased viability results [Fig. 7E]. MCA assays done with cells depleted of TTT complex subunits using siRNAs are provided for comparison [Supplemental Fig. S13B,C]. Thus, overall, our data support a critical role for the entire TTT complex in the DDR.

**TTT complex components depend on each other for their stability**

While all three members of the TTT complex associate together, the extent to which each subunit was required was unclear. To address this, we examined the relationship between TTI1 and TEL2. U2OS cells expressing shRNAs against TTI1 showed reduced levels of TEL2...
protein (Fig. 7A) but not TEL2 mRNA (Supplemental Fig. S11), suggesting an effect on protein stability. The effect on TEL2 abundance was also observed in an independent cell line (293T) (data not shown). Expression of an shRNA-resistant TTI1 cDNA restored TEL2 and PIKK protein levels. U2OS cells stably expressing a control (FF) or TTI1 shRNA #3 [targeting the 3′ UTR of endogenous TTI1] were infected with an empty vector or Flag-HA-TTI1 for 2 d. Whole-cell extracts were immunoblotted with the indicated antisera. (C) Overexpression of TEL2 does not restore IR resistance to TTI1-depleted cells. U2OS cells stably expressing TTI1 shRNA #3 were infected with retroviruses expressing MSCV-HA-Flag-empty, MSCV-HA-Flag-TTI1, or MSCV-TEL2-HA-Flag, and, after 2 d, were mixed with control cells for MCA. Viability in response to 3 Gy of IR or 75 nM MMC was monitored. (D) Each subunit of the TTT complex is required for DNA damage resistance. MCA was performed using U2OS cells expressing dsRed, and the individual shRNAs are indicated as in Figure 2C. Viability in response to 3 Gy of IR or 75 nM MMC was monitored. shRNA sequences are listed in Supplemental Table S6. (E) Knockdown of TTT subunits affects ATM and ATR levels and levels of the other TTT subunits. Immunoblots corresponding to the viability assays shown in D. The lysates from cells expressing FF, ATM, ATR, TTI1 #1, and TTI1 #2 shRNAs are the same in both panels to allow for comparison between the effects of TEL2 and TTI2 depletion. (F) TTI1 is critical for interaction of the TTT complex with ATM. U2OS cells were transfected with siRNA duplexes and incubated for 72 h prior to harvesting for preparation of whole-cell extracts. Immunoprecipitations were performed using control, TTI1, or TEL2 [ProteinTech Group] antisera. TTI1, TEL2, and TTI2 pools of an equimolar mix of four siRNAs from Dharmacon were used [See Supplemental Table S7].

Figure 7. TTI1 and TTI2 are required for maintaining TEL2 protein levels, and are critical components of the TTT complex. (A) TTI1 is required to maintain TEL2 protein levels. Whole-cell extracts from U2OS cells stably expressing control, ATR, or TTI1 shRNAs were immunoblotted with TEL2, TTI1, or VINCULIN antisera. (B) Expression of an shRNA-resistant TTI1 cDNA can restore TEL2 and PIKK protein levels. U2OS cells stably expressing a control (FF) or TTI1 shRNA #3 [targeting the 3′ UTR of endogenous TTI1] were infected with an empty vector or Flag-HA-TTI1 for 2 d. Whole-cell extracts were immunoblotted with the indicated antisera. (C) Overexpression of TEL2 does not restore IR resistance to TTI1-depleted cells. U2OS cells stably expressing TTI1 shRNA #3 were infected with retroviruses expressing MSCV-HA-Flag-empty, MSCV-HA-Flag-TTI1, or MSCV-TEL2-HA-Flag, and, after 2 d, were mixed with control cells for MCA. Viability in response to 3 Gy of IR or 75 nM MMC was monitored. (D) Each subunit of the TTT complex is required for DNA damage resistance. MCA was performed using U2OS cells expressing dsRed, and the individual shRNAs are indicated as in Figure 2C. Viability in response to 3 Gy of IR or 75 nM MMC was monitored. shRNA sequences are listed in Supplemental Table S6. (E) Knockdown of TTT subunits affects ATM and ATR levels and levels of the other TTT subunits. Immunoblots corresponding to the viability assays shown in D. The lysates from cells expressing FF, ATM, ATR, TTI1 #1, and TTI1 #2 shRNAs are the same in both panels to allow for comparison between the effects of TEL2 and TTI2 depletion. (F) TTI1 is critical for interaction of the TTT complex with ATM. U2OS cells were transfected with siRNA duplexes and incubated for 72 h prior to harvesting for preparation of whole-cell extracts. Immunoprecipitations were performed using control, TTI1, or TEL2 [ProteinTech Group] antisera. TTI1, TEL2, and TTI2 pools of an equimolar mix of four siRNAs from Dharmacon were used [See Supplemental Table S7].
The Triple T complex controls PIKKs

still able to interact with ATM as well as in the control cells [Fig. 7F, left panel]. However, in cells depleted of TTI1, endogenous TEL2 is unable to coimmunoprecipitate ATM as efficiently as in control cells [Fig. 7F, right panel]. In cells depleted of TTI2, less ATM comes down with both TTI1 and TEL2 antisera, and this is most likely because TTI2 depletion has a stronger effect on TTI1 than TEL2 depletion does [Fig. 7E]. Thus, TTI1 is required for efficient binding of the TTT complex to the major DDR kinase, ATM, and therefore we argue that TTI1 plays a major role within the TTT complex and is a critical regulator of the DDR and likely other PIKK-regulated pathways in mammalian cells.

Discussion

We performed an unbiased genome-wide screen to find genes required to protect the cell from DNA DSBs. We identified a large number of DDR genes with known roles in DNA repair, replication, damage signaling, and nucleotide metabolism, as well as many genes not implicated previously in the DDR. Among these, we found a strong enrichment for genes in RNA metabolism. Genes involved in spliceosome assembly, RNA splicing, polyadenylation of mRNA, and processing of mRNA and rRNA were highly enriched. This is interesting due to the fact that, in previous screens for ATM and ATR substrates, a significant enrichment was also seen for these classes of proteins (Matsuoka et al. 2007; Stokes et al. 2007). Furthermore, in a screen to look for genes whose depletion causes genomic instability, Cimprich and colleagues (Paulsen et al. 2009) found a very strong enrichment for splicing factors and RNA-binding proteins, further underscoring a role for these factors in the DDR. It is possible that these proteins play roles in the DDR outside of an indirect role in splicing.

The strong enrichment observed for genes involved in known DDR processes suggests that many of the novel genes will also play a role in these processes. Among the novel genes, we found that the uncharacterized TTI1 gene is required for resistance to IR as well as other genotoxic agents, including UV light, MMC, and CPT. TTI1 is absolutely required for proper DDR signaling. Our results reveal that TTI1 interacts with TEL2 and a second protein, TTI2, to form the TTT complex required to stabilize the protein levels of the PIKKs. TTI1 is required to maintain the abundance of TEL2, as depletion of TTI1 results in a decrease in TEL2 protein levels, most likely by controlling its stability. TEL2 has been implicated in the DDR, in particular in response to agents that activate the ATR pathway (Jiang et al. 2003; Collis et al. 2007, 2008). However, the DNA damage sensitivity phenotype observed with TTI1 depletion is not simply due to the decrease in TEL2 levels, since restoration of TEL2 expression cannot restore DNA damage resistance. We argue that TTI1 is a critical component of the TTT complex that is likely involved in mediating the interaction of the complex with PIKKs, since TTT complex depleted of TTI1 can no longer interact efficiently with ATM.

We have evidence to suggest that the activated PIKKs do not bind as tightly to the TTT complex. First, although it is a small effect, we show that TTI1 interacts less well with ATM, DNA PK, ATR, and SMG1 once they are activated by IR [Fig. 5C, E]. Second, we observed that TTI1 does not localize to sites of DNA damage, indicating that TTT is unlikely to associate with activated ATM and ATR, which are localized to damage sites, further suggesting that ATM and ATR are released from TTI1 prior to their localization to DNA damage foci (data not shown). This is consistent with the observation of de Lange and colleagues (Takai et al. 2007) that TEL2 does not localize to sites of DNA damage. The simplest interpretation of these results is that TTT participates in proper folding of newly synthesized PIKKs. Once properly folded, TTT releases the PIKK, which is then capable of activation by the relevant signal transduction pathway. In the absence of TTT, PIKKs are poorly folded and fall subject to proteolysis, thereby reducing their levels. It is also possible that more complex relationships exist, including possible roles for TTT in recycling PIKKs from the active to the inactive state and back.

It will be important to determine whether a single subunit of the TTT complex contacts each of the structurally related PIKKs, or if different subunits primarily recognize different PIKKs. Baculovirus-produced Tel2 was reported to bind fragments of mTOR and ATM (Takai et al. 2007). It is possible that multiple components of the TTT complex can contact substrates, perhaps to create an extended folding template. A critical question to be resolved is the elucidation of the composition and assembly of the 2000-Da TTT complex. Are there additional subunits that contribute to this complex? Is this complex preformed, or is it assembly induced by the presence of newly translated PIKKs? Is the complex regulated in response to DNA damage? Now that the complex has been partially elucidated, reconstituting the entire TTT complex to allow its activity toward PIKKs to be explored biochemically becomes a possibility.

Is the TTT complex a chaperone? Currently, the evidence for this hypothesis is purely circumstantial. If the TTT complex acts as a chaperone to refold PIKKs, it might do so by recruiting a known chaperone/heat-shock protein, analogous to how Cdc37 recruits Hsp90 (Stepanova et al. 1996). Alternatively, it could perform chaperone duties itself, which would predict the need for energy consumption. Unfortunately, none of the TTT proteins contain conserved motifs that might give a clue to their molecular functions, and no ATPase motifs are discernable from the sequences of TTT proteins. Instead, both TTI1 and TTI2 appear to be largely composed of HEAT repeats, although they are highly divergent (K Hofmann, pers. comm.). HEAT repeats are generally thought to be protein–protein interaction regions, or may suggest some type of molecular scaffold function. Thus, identifying which non-PIKK proteins associate with TTT will be critical to the elucidation of its biochemical function. Other biochemical functions are possible for the TTT complex. For example, TTT could control a modification on PIKKs needed for their stability and activation. Alternatively, TTT could act to insert an essential cofactor needed for PIKK holoenzyme formation. The resolution of these issues will require a full...
elicitation of the TTT complex composition, and biochemical reconstitution of its role in PIKK regulation.

Materials and methods

A loss-of-function screen for radio-protective genes using shRNA libraries

The pool-based shRNA screen using half-hairpin barcode deconvolution was performed as described previously [Schlabach et al. 2008, Luo et al. 2009]. Briefly, U2OS cells were infected with pools of retroviral shRNAs at a representation of 1000 and an MOI of ∼1. The cells were treated with puromycin to remove cells that were not infected and were treated with IR or left untreated, and an initial sample (population doubling 0) was collected. The rest of the population was propagated in triplicate for ∼12 population doublings [14 d], and then end samples were harvested for genomic DNA isolation. At each passage, a minimum representation of 1000 was maintained. For each sample (INITIAL, UNTREATED-END, or IR-END), shRNA half-hairpin barcodes were PCR-recovered from genomic DNA and labeled with Cy5 or Cy3 dyes, and labeled amplicons were hybridized competitively to a microarray containing corresponding probes. Custom microarrays with half-hairpin probe sequences were from Roche Nimblegen or Agilent. Array hybridization and scanning protocols were based on the manufacturer’s instructions. The PCR primers used to amplify half-hairpin barcodes from genomic DNA samples were JH353F (5'-TGTGAAAGCCA CAGATGTA-3') and HHR2L (5'-ATGTATCAAAGAGATAGCA -9'). shRNAs with log2 ratios (average of three replicates) >1 or less than ∼1, which corresponds to a twofold change or greater, were considered enriched or depleted from the population, respectively. Two different competitive hybridizations were performed: untreated samples taken at the end of the screen (UNTREATED-END) versus IR-treated samples taken at the end of the screen (IR-END). The Focus library was used to screen a second cell line, the lung cancer cell line H460, using one dose of 2 Gy or 5 Gy of IR. In addition, the Focus library was also used to screen U2OS cells treated with multiple low doses of IR. Cells were treated with 2 Gy and then allowed to recover for 1 wk before being treated again with 2 Gy. A total of four 2 Gy doses of IR were given. Supplemental Table S1 reports the data from the genome-wide pools and the focus pool from U2OS cells treated with one dose of 5 Gy and competitive hybridization of UNTREATED-END versus IR-END. We used data collected from all of our IR screens for the network shown in Figure 2B, which is why some of these hits are not listed in Supplemental Table S1. Supplemental Table S3 lists all of the genes from Figure 2B that scored in our screens (those indicated in red), and what screen condition or hybridization they scored in.

Cell culture

U2OS osteosarcoma cells (American Type Culture Collection) were grown in McCoy’s 5A medium supplemented with 10% fetal bovine serum and antibiotics. 293T and 293 T Rex [Invitrogen] were cultured in DMEM supplemented with 10% fetal bovine serum and antibiotics.

Antibodies

Anti-human TTI1 antibody [YZ503-1] was affinity-purified from rabbit serum immobilized with KLH-conjugated hTTI peptide corresponding to amino acids #1071-1089 (SGQQNPYTTNVLQ LLKELQ-COOH) [Yenzym Antibodies, LLC]. Other antibodies use were TEL2B [a gift from T. de Lange], TEL2 [SC1997-1-AP, ProteinTech Group], ATM [M300-136A, Bethesda Laboratories], ATR [N-19] [M18-187, Santa Cruz Biotechnology], DNA-PK [Ab-2] [NA57, Calbiochem-EMD Biosciences], SMC1 [A300-394A, Bethesda Laboratories], TRRAP [A301-32A2, Reth Laboratories], mTOR [A2972, Cell Signaling Technology], HA-agarose [clone HA-7] [A2095, Sigma], C-MYC [9E10] [MMS-150, Covance], VINCLIN [V9131, Sigma], Flag M2-HRP [A8592, Sigma], EZview Red Flag-MAFII Antibody Gel [A2426, Sigma], P-H3 S10 [A06-570, Millipore], P-CHK1 S317 [A2344, Cell Signaling Technology], CHK1 [SC-8409, Santa Cruz Biotechnology], P-ATM S1981 [A200-301-400, Rockland], P-CHK2 T68 [A2661, Cell Signaling Technology], P-NBS1 S343 [A300-189A, Bethesda Laboratories], NBS1 [A300-187A, Bethesda Laboratories], P-KAP1 S424 [A300-767A, Bethesda], P-SMC1 S966 [A300-050A, SMC1 [A300-055A, Bethesda Laboratories], CHK2 [generated in our laboratory [Matsuoka et al. 1998]], P-AKT S473 [A4058, Cell Signaling Technology], AKT [A300-136A, Bethyl Laboratories], ATM [A300-136A, Bethyl Laboratories], P-TEL2B (a gift from T. de Lange), TELO2 [A300-767A, TEL2, TEL2 cDNAs are from the hORFeome V5.1 collection, and the Gateway recombination system [Invitrogen] was used to transfer the cDNAs from pDONR-223 to the following vectors: MSCV-IREs-PURO-N-HA-Flag, MSCV-PGK-PURO-C- HA-Flag, MSCV-PGK-PURO-N-MYC, MSCV-PGK-PURO-N- EGFP, MSCV-CMV-TO-N-HA-Flag PGK-Puro (inducible). The sequence of shRNAs and siRNAs used in this study are listed in Supplemental Tables S6 and S7, respectively. The shRNA library is in a MSCV-PURO-based vector (MSCV-PM, Open Biosystems nomenclature is pSMPI). The shRNAs with the “V3LHS” identifier were subcloned into the MSCV-PM vector from the pGIPZ vector. The shRNAs indicated “CUSTOM” were generated in our laboratory, and the 22-mer shRNA sequence is within the mir30 context and the MSCV-PM vector backbone (the same as the library).

Quantitative RT-PCR

Total RNA was prepared using the RNAeasy Plus kit (Qiagen), and reverse transcription to generate cDNA was performed using SuperScript III Reverse Transcriptase (#18080-044, Invitrogen). RT–PCR was done using Platinum Cybergreen Super Mix with

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Radio-resistant DNA synthesis assay

Radio-resistant DNA synthesis assays were done as described previously (Silverman et al. 2004). Briefly, U2OS cells were transfected with siRNA and, 24 h later, medium containing 10 nCi/mL [methyl-14C] thymidine (Amersham) was added, and the cells were incubated for 24 h. After another 24 h incubation with medium without label, the cells were irradiated with 15 Gy of IR. Following a 30-min incubation, the cells were pulse-labeled with 2.5 μCi/mL [methyl-3H] thymidine (Amersham) for 20 min, and then washed twice with medium containing 2.5 mM cold thymidine. Approximately 1012 cells were harvested by trypsinization, and TCA precipitation was performed on Whatman glass microfiber filters using a vacuum manifold. Following an ethanol wash, the filters were dried and counted using a liquid scintillation counter (Beckman LS6000). The ratio of 3H counts per minute to 14C counts per minute, corrected for those counts per minute that were the result of channel crossover, was a measure of DNA synthesis.

G2/M checkpoint assay

U2OS cells were transfected with 50 nM final concentration of siRNA using oligofectamine reagent (Invitrogen) following the manufacturer’s recommendations. After 4–6 d to allow for efficient gene knockdown, cells were treated with 10 Gy of IR and incubated for 1 h at 37°C. Nocodazole was added to the medium at a final concentration of 400 μg/mL, and cells were incubated for an additional 17 h at 37°C. Cells were fixed with ethanol, stained with phospho-S10 Histone H3 antisera followed by Alexa 488 secondary antibody and propidium iodide, and analyzed by FACS. The percentage of phospho-S10 Histone H3 cells in ATM, ATR, or TTF1 knockdown cells was normalized to FF control knockdown cells to give fold change mitotic index.

MCA

U2OS cells were retrovirally transduced with MSCV-dsRed [no selection marker] and FACS-sorted to obtain a more homogeneous population. The dsRed cells were then infected with retrovirus expressing individual shRNAs of interest, selected with puromycin, and then equally mixed with U2OS cells [uncolored] expressing a control shRNA targeting firefly luciferase. The mixed population was left untreated or treated with 5 Gy of γ-irradiation, 75–100 nM MMC, 5 nM CPT, or 5/fJ/m2 UV light; medium containing approximately 3 × 1012 cells were harvested for FACS analysis. For a graphical representation, the percent of dsRed-expressing cells in DNA damage-treated samples was normalized to that of untreated cells. We also normalized to the performance of dsRed cells carrying a control shRNA. For MMC- and CPT-treated cells, the medium containing the drug was removed after 12 h and replaced with fresh medium without drug. For siRNA experiments, U2OS cells were transfected with 50 nM siRNAs using oligofectamine, mixed for MCA, and treated with DNA damage at 72 h post-transfection.

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