

TP53 Binding to BRCA1 and RAD51 in MCF7 and MDA-MB-468 Breast Cancer Cell Lines *In vivo* and *In vitro*

Mozhgan Rasti ^{*}, and Tayebeh Azimi

Recombinant Lab, Department of Biochemistry, Faculty of Medicine, Shiraz University of Medical Sciences, Shiraz, Iran

Abstract

Background: Tumour suppressor genes such as TP53, BRCA1 and RAD51 are involved in DNA repair and their malfunctions result in genomic instability and cancer. Wild type (WT) TP53 binds to BRCA1 and RAD51 *in vivo* and *in vitro*. However, mutated TP53 in tumours can interfere with WT TP53 function. We studied how mutation of TP53 in MDA-MB-468 cell line could affect its binding capacity and interfere with WT TP53 interaction with these DNA repair proteins.

Methods: Binding capacity of mutated TP53 in MDA-MB-468 breast cancer cell line to BRCA1 and RAD51 proteins in comparison to WT TP53 in MCF7 cell line was studied by Immunoprecipitation. *In vitro* studies were performed by GST-WT p53 pull-down assays in these cell lines to assess the interaction of GST-WT p53 with BRCA1 and RAD51 proteins.

Results: The results showed that mutated TP53 in MDA-MB-468 cells interacted with BRCA1 protein *in vivo* and did not effect WT TP53 binding to this protein *in vitro*. The Immunoprecipitation assays revealed that the mutated TP53 did not bind to RAD51 in comparison to WT TP53. However, this mutated protein could not interfere with binding of RAD51 to GST-WT p53 in MDA-MB-468 cell line by *in vitro* experiment.

Conclusion: It was found that WT TP53 interactions with BRCA1 and RAD51 did not interfere with mutated TP53 in MDA-MB-468 cell line. In addition, RAD51 did not bind to TP53 with R273C mutation *in vivo*.

Avicenna J Med Biotech 2015; 7(2): 76-79

Keywords: BRCA1 protein, Immunoprecipitation, RAD51 protein, TP53

Introduction

Breast cancer is a world-wide cancer in women. In Iranian women especially the age of its diagnosis has decreased recently ¹. Breast cancer genome analyses have shown that there are a few tumour suppressor genes that are frequently mutated such as breast cancer 1, early onset (BRCA1) and tumour protein p53 (TP53) genes ². TP53 is mutated in the majority of sporadic breast cancers. Most of these cancer related mutations in TP53 are in their DNA binding domain that reveal a gain-of-function for TP53 ³. In such cases TP53 mutated levels are increased in breast tumours and can interfere with repair of DNA double strand breaks ^{4,5}. In normal tissue only a small amount of TP53 is present that binds to DNA to induce cell cycle check point activation, cellular apoptosis, autophagy and senescence ⁶. BRCA1 mutation is associated with hereditary breast and ovarian cancers ². It binds to Mre11-Rad50-NBS1 (MRN) complex which is a DNA repair complex for DNA double stranded break and ZBRK1, a zinc finger transcriptional repressor ⁷. In addition, it regulates cell cycle check points ⁸. On the other hand,

BRCA1 interacts with RAD51 recombinase (RAD51) during homologous recombination mediated DNA double-strand break repair forming a nucleoprotein filament on DNA and its mutation can increase the risk of breast cancer ⁹⁻¹¹.

It has been shown that Wild Type (WT) TP53 is able to bind both to BRCA1 and RAD51 to conduct genomic stability ^{12,13}. However, mutated TP53 in tumours not only interferes with WT TP53 function but also impair chemotherapy responses ¹⁴. In the present study, our aims are to study how a mutation of TP53 protein in its DNA binding domain can affect its binding capacity and interfere with WT TP53 binding to these DNA repair proteins. Therefore, we evaluated the binding capacity of WT TP53 to these two proteins in MCF7 and MDA-MB-468 cell lines in comparison to mutated TP53 in MDA-MB-468 breast cancer cell line *in vivo* and *in vitro*. Such studies can elucidate the molecular mechanism of mutated TP53 in comparison to wild type one in order to provide novel diagnostic and therapeutic insights.

* Corresponding author:
Mozhgan Rasti, Ph.D.,
Recombinant Lab, Department
of Biochemistry, Faculty of
Medicine, Shiraz University of
Medical Sciences, Shiraz, Iran
Tel: +98 711 32303029
Fax: +98 711 32359847
E-mail:
rasti31@yahoo.com
Received: 30 Jul 2014
Accepted: 1 Nov 2014

Materials and Methods

Cell culture

Both MDA-MB-468 cells which possess a mutation in codon 273 of the TP53 gene and MCF7 cells with WT TP53 gene were used in this study (Pasture Institute, Iran). These cancer cell lines were grown in RPMI 1640 (Biosera, UK) supplemented with L-glutamine to 2 mM, and 10% fetal calf serum (Cinagen, Iran) and sub cultured 2-3 times in a week.

In vitro experiment by GST pull-down assay

The p53-Glutathione S-transferase (GST) fusion protein was expressed and purified as described previously¹⁵. The MCF7 and MDA-MB-468 cell lysates were solubilized in a lysis buffer containing 10 mM Tris-HCl pH=7.4, 0.825 M NaCl and 1% NP-40 and then sonicated and cleared by centrifugation. All protein concentrations were measured by Bradford protein assay and concentrations of samples were calculated based on standard curve¹⁶. For GST pull-down assay¹⁷, 50 µg of GST- WT p53 or GST proteins was mixed with 5 mg of either the MCF7 or MDA-MB-468 cell lysates for 2 hr at 4°C. The pull-down complexes with GST and GST-WT p53 proteins were eluted from the glutathione-agarose by 25 mM reduced glutathione in 50 mM Tris-HCl solution pH=8.0. The eluted samples were resolved by Sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) and then subjected to Western blot analysis with appropriate antibody.

In vivo experiment by immunoprecipitation of proteins

The studied cancer cells were lysed in buffer and protein concentrations were measured as described above. Immunocomplexes were precipitated from 5 mg of protein lysate¹⁷, using 5 µg of primary antibodies of anti-BRCA1 (SC-641; Santa Cruz, USA) or anti-TP53 serum¹⁵. IgG Ab was used as negative control. After 2 hr mixing at 4°C on a rotator, 30 µl of packed protein G agarose beads (Sigma, UK) was added into the protein-antibody complex and mixed for a further 1 hr. Immunocomplexes bound to the beads were then spun and washed, prior to re-suspending in the SDS sample buffer for SDS-PAGE analysis.

Western blot analyses

GST pull-down or immunoprecipitated samples with 50 µg proteins of appropriate lysates were electrophoresed on 8 and 12% SDS-PAGE gels for detection of BRCA1 and RAD51, TP53 proteins, respectively. GST pull-down complexes were immunoblotted with 1:100 dilution of anti-BRCA1 (SC-641; Santa Cruz, USA) and 1:1000 dilution of anti-RAD51 (ab63801; Abcam, USA) antibodies. In addition, anti-TP53 serum¹⁵ and anti-RAD51 (ab63801; Abcam, USA) primary antibodies were diluted 1:1000 for analyzing of immunoprecipitated proteins in blots. A secondary antibody of horseradish peroxidase-conjugated anti-rabbit (Sigma, USA) with dilution of 1:1000 and chemilumi-

nescence substrates (ECL; Amersham Bioscience AB) were used to detect the immuno-labeled bands.

Results

TP53 mutation in MDA-MB-468 cell line did not affect its binding capacity to BRCA1

In order to understand the binding capacity of mutated TP53 in MDA-MB-468 cell line in comparison to wild type one in MCF7 cells, cell lysates of these cells were immunoprecipitated with antibody against BRCA1 and blotted for TP53. The results (Figure 1A) showed that both mutated and WT p53 proteins were able to bind with this repair protein. GST pull-down assays were performed with GST-WT p53 fusion protein synthesized in *Escherichia coli* (*E. coli*) and these lysates to analyze by BRCA1 immunoblotting. *In vitro* results (Figure 1B) revealed that GST-WT p53 bound to BRCA1 and mutated TP53 in MDA-MB-468 cell line did not interfere with binding of the wild type fusion protein to BRCA1. BRCA1 did not bind to GST alone showing there was no binding affinity between GST alone and BRCA1. Figure 1B shows that the expression of BRCA1 in MDA-MB-468 cell line was more than MCF7 cell line that provided more proteins for binding to GST-WT p53 in MDA-MB-468 cell line rather than MCF7 cells. Therefore, co-precipitation of GST-WT p53 with BRCA1 in MCF7 cell line was underrepresented as judged from autoradiograph.

Mutated TP53 in MDA-MB-468 cell line was not able to bind RAD51

Immunoprecipitation of TP53 in MDA-MB-468 and MCF7 cell lysates and blotting for RAD51 revealed that mutated TP53 in MDA-MB-468 cell line was not able to bind RAD51. However, this *in vivo* experiment for MCF7 cell line showed that the precipitated immunocomplex included WT TP53 and RAD51 (Figure 2A). Interaction of WT p53 to RAD51 in MCF7 cell

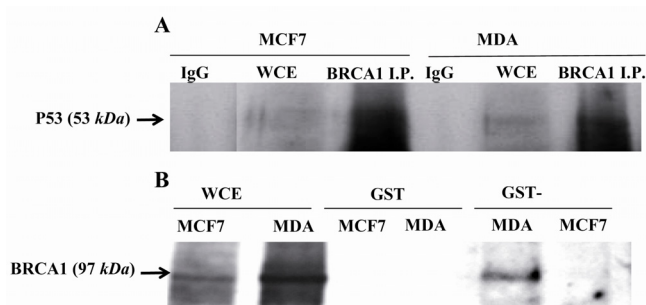


Figure 1. TP53 bound to BRCA1 *in vivo* and *in vitro*. A) *In vivo* binding of wild type (WT) TP53 in MCF7 cell line and mutated TP53 in MDA-MB-468 (MDA) to BRCA1. Cell lysates were immunoprecipitated with Ab against BRCA1 and IgG and then the immunocomplexes were Western blotted for anti-p53. B) Mutated TP53 did not interfere with *in vitro* binding of WT TP53 to BRCA1. BRCA1 was pulled down in MDA-MB-468 (MDA) and MCF7 cell lysates by GST-WT p53 that was a fusion protein synthesized in prokaryotes cells and analyzed by Western blotting with anti-BRCA1. WCE whole-cell extract, I.P. Immunoprecipitate.

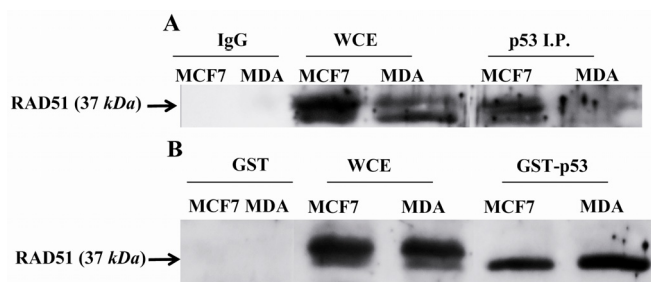


Figure 2. RAD51 only bound to wild type (WT) TP53 *in vivo* and *in vitro*. A) RAD51 bound to WT TP53 in MCF7 cell line but not to the mutated one in MDA-MB-468 (MDA) cells. Cell lysates were Immunoprecipitated with Ab against TP53 and IgG then analyzed by Western blotting with anti-RAD51. B) Mutated TP53 did not interfere with *in vitro* binding of WT TP53 to RAD51. RAD51 was pulled down in MDA-MB-468 (MDA) and MCF7 cell lysates by GST-WT p53 that was a fusion protein synthesized in prokaryotes cells to assay the binding capacity of RAD51 to synthesized TP53. Western blot analysis was performed with anti-RAD51. WCE whole-cell extract, I.P. Immunoprecipitate.

lysate was confirmed by pulling down of GST- WT p53 fusion protein with RAD51 and blotting for RAD51 in two studied cell lines. Figure 2B shows that GST-WT p53 pulled down the RAD51 proteins in these two cell lines. However, mutated TP53 was not able to bind the RAD51 in MDA-MB-468 cells, the mutated TP53 did not interfere with binding of GST- WT P53 to RAD51 in this cell line. RAD51 did not bind to GST alone that showed there was not binding affinity between GST alone and RAD51.

Discussion

WT TP53 is stabilized and accumulated in normal cells due to the cellular stress and DNA damage. However, mutant alleles of TP53 are highly expressed in one third of breast tumors¹⁴. In addition, it has been shown that mutant p53 proteins are able to inhibit WT TP53 through dominant negative gain of function. Different mutated p53 proteins may inhibit DNA damage response, cell cycle control and apoptosis process induced by the WT one. Therefore, mutated forms can drive tumor formation, invasion and metastasis that impair chemotherapy responses which are based on the WT TP53 function¹⁴. MDA-MB-468 cell line is hemizygous for a mutated p53 gene which makes a R273C mutant and ectopic expression of WT TP53 in these cells cannot overcome the gain of function of mutant form¹⁸. This mutant TP53 protein is reported to cooperate with oncogenes for transforming cells and inhibits p73 protein¹⁴.

Genetic instability is another result of TP53 mutation that is because of inhibition of DNA repair pathway. Some of TP53 mutants can bind to MNR complex and impair recruiting taxia-telangiectasia-mutated (ATM) kinase to double strand break¹⁹. WT TP53 binds to BRCA1 and RAD51 to induce repair pathways in the case of any DNA damage^{12,13}. Many studies

showed that how BRCA1 is involved in DNA repair and cell cycle check points to maintain genome stability^{2,7,8,20}. BRCA1 binds to RAD51⁹ which possesses a main role in Homologous Recombination (HR) by repairing endogenous and exogenous DNA double-strand breaks¹¹. It happens during DNA replication and uses sister chromatid as a template to repair the DNA double-strand break²¹.

Our *in vitro* experiments using GST- WT p53 protein in studied cell lines confirmed previous studies¹³ that WT TP53 binds to RAD51 directly. We found that this interaction was not interfered by the mutated TP53 in MDA-MB-468 cell line. These data shows that RAD51 interacted with bacterial extracted GST- p53 protein and did not need post transcriptional modification of p53 for its binding. It has been reported that RAD51 physically interacts to GST-WT p53 and to a lesser extent a point mutant GST-273H-p53 protein. Mapping studies revealed that RAD51 binds two regions of amino acids 94-164 and 264-315 of TP53¹³. In addition, our *in vivo* result for RAD51 in MCF7 cell line is consistent with previous data in other cell lines²² and confirmed that it was able to bind WT TP53.

In the present study, for the first time it was shown that RAD51 protein failed to interact with mutated TP53 in MDA-MB-468 cell line *in vivo*. It reveals that despite the previous *in vitro* results for 273H-p53 point mutant with reduced amount of binding to RAD51¹³, R273C mutant diminished interaction activity completely *in vivo*. It implies a main role for this point mutant in TP53 for binding to RAD51. This finding can explain the genomic instability in MDA-MB-468 cell line that may result in the harsh metastasis property of these cells. It has been reported that BRCA1 binds to C-terminal domain of TP53 between amino acid residues 300-393¹².

Our *in vivo* experiments showed that BRCA1 was over expressed and precipitated with mutated TP53 in MDA-MB-468 cell line which indicated its point mutation in 273 codon did not effect its binding affinity to BRCA1. However, because low levels of BRCA1 were present in MCF7 cell line we could not show this interaction with WT TP53. This interaction has been revealed in other cell line previously¹². The *in vitro* results with GST-WT p53 confirmed previous finding for direct interaction of WT-p53 with BRCA1¹² and showed that mutated TP53 in MDA-MB-468 cell line could not interfere with its interaction to BRCA1. It has been shown that BRCA1 is a potent co-activator of TP53 in p21 and bax genes transcription¹². In MCF7 cells, TP53 is maintained at very low level by proteasome degradation when there is not any cellular stress. In MDA-MB-468 cell line, there is genomic instability that activates mainly ATM and related proteins²³. While ATM activation stabilizes TP53 and BRCA1 by phosphorylation²³, it may result in the increased expression levels of the mutated TP53 and BRCA1 that we observed in MDA-MB-468 cell line.

Conclusion

We found that BRCA1 protein but not RAD51 interacted to TP53 with R273C mutant *in vivo* and this point mutant in TP53 did not interfere with the binding capacity of WT TP53 to BRCA1 and RAD51.

Acknowledgement

This work was supported by grant No: 88-01-01-1395 from Shiraz University of Medical Sciences, Iran for Ms. Tayebe azimi's thesis. The authors would like to thank the School of Advanced Medical Sciences and Technologies for their support in performing this project.

Conflict of Interest

None of authors has any conflict of interest.

References

- Mousavi SM, Montazeri A, Mohagheghi MA, Jarrahi AM, Harirchi I, Najafi M, et al. Breast cancer in Iran: an epidemiological review. *Breast J* 2007;13(4):383-391.
- Lee EY, Muller WJ. Oncogenes and tumor suppressor genes. *Cold Spring Harb Perspect Biol* 2010;2(10):a003236.
- Xu Y. Induction of genetic instability by gain-of-function p53 cancer mutants. *Oncogene* 2008;27(25):3501-3507.
- Song H, Hollstein M, Xu Y. p53 gain-of-function cancer mutants induce genetic instability by inactivating ATM. *Nat Cell Biol* 2007;9(5):573-580.
- Brosh R, Rotter V. When mutants gain new powers: news from the mutant p53 field. *Nat Rev Cancer* 2009;9(10):701-713.
- Menendez D, Inga A, Resnick MA. The expanding universe of p53 targets. *Nat Rev Cancer* 2009;9(10):724-737.
- Zheng L, Pan H, Li S, Flesken-Nikitin A, Chen PL, Boyer TG, et al. Sequence-specific transcriptional corepressor function for BRCA1 through a novel zinc finger protein, ZBRK1. *Mol Cell* 2000;6(4):757-768.
- Yu X, Chini CC, He M, Mer G, Chen J. The BRCT domain is a phospho-protein binding domain. *Science* 2003;302(5645):639-642.
- Davies AA, Masson JY, McIlwraith MJ, Stasiak AZ, Stasiak A, Venkitaraman AR, et al. Role of BRCA2 in control of the RAD51 recombination and DNA repair protein. *Mol Cell* 2001;7(2):273-282.
- Galkin VE, Wu Y, Zhang XP, Qian X, He Y, Yu X, et al. The Rad51/RadA N-terminal domain activates nucleosome protein filament ATPase activity. *Structure* 2006;14(6):983-992.
- Somyajit K, Subramanya S, Nagaraju G. RAD51C: a novel cancer susceptibility gene is linked to Fanconi anemia and breast cancer. *Carcinogenesis* 2010;31(12):2031-2038.
- Zhang H, Somasundaram K, Peng Y, Tian H, Zhang H, Bi D, et al. BRCA1 physically associates with p53 and stimulates its transcriptional activity. *Oncogene* 1998;16(13):1713-1721.
- Buchhop S, Gibson MK, Wang XW, Wagner P, Sturzbecher HW, Harris CC. Interaction of p53 with the human Rad51 protein. *Nucleic Acids Res* 1997;25(19):3868-3874.
- Goh AM, Coffill CR, Lane DP. The role of mutant p53 in human cancer. *J Pathol* 2011;223(2):116-126.
- Rasti M, Khatooni Z, Mostafavi-pour Z. Production and evaluation of polyclonal rabbit anti-human p53 antibody using bacterially expressed glutathione S-transferase-p53 fusion protein. *Iran J Biotechnol* 2012;10(4):240-248.
- Bradford MM. A rapid and sensitive method for the quantitation of microgram quantities of protein utilizing the principle of protein-dye binding. *Anal Biochem* 1976;72:248-254.
- Arabsolghar R, Azimi T, Rasti M. Mutant p53 binds to estrogen receptor negative promoter via DNMT1 and HDAC1 in MDA-MB-468 breast cancer cells. *Mol Biol Rep* 2013;40(3):2617-2625.
- Casey G, Lo-Hsueh M, Lopez ME, Vogelstein B, Stanbridge EJ. Growth suppression of human breast cancer cells by the introduction of a wild-type p53 gene. *Oncogene* 1991;6(10):1791-1797.
- Muller PA, Caswell PT, Doyle B, Iwanicki MP, Tan EH, Karim S, et al. Mutant p53 drives invasion by promoting integrin recycling. *Cell* 2009;139(7):1327-1341.
- Rodriguez MC, Songyang Z. BRCT domains: phosphopeptide binding and signaling modules. *Front Biosci* 2008;13:5905-5915.
- Johnson RD, Jasin M. Sister chromatid gene conversion is a prominent double-strand break repair pathway in mammalian cells. *EMBO J* 2000;19(13):3398-3407.
- Stürzbecher HW, Donzelmann B, Henning W, Knipschild U, Buchhop S. p53 is linked directly to homologous recombination processes via RAD51/RecA protein interaction. *EMBO J* 1996;15(8):1992-2002.
- Valentine JM, Kumar S, Moumen A. A p53-independent role for the MDM2 antagonist Nutlin-3 in DNA damage response initiation. *BMC Cancer* 2011;11:79.