AD_____

Award Number: DAMD17-97-1-7147

TITLE: Germline Mutations of the Ataxia-Telangiectasia Gene, ATM, as a Risk Factor for Radiation-Associated Breast Cancer

PRINCIPAL INVESTIGATOR: Kenneth Offit, M.D.

CONTRACTING ORGANIZATION: Sloan-Kettering Institute for Cancer Research
New York, New York 10021

REPORT DATE: July 1999

TYPE OF REPORT: Annual

PREPARED FOR: U.S. Army Medical Research and Materiel Command
Fort Detrick, Maryland 21702-5012

DISTRIBUTION STATEMENT: Approved for Public Release;
Distribution Unlimited

The views, opinions and/or findings contained in this report are those of the author(s) and should not be construed as an official Department of the Army position, policy or decision unless so designated by other documentation.

20000308 026

DTIC QUALITY INSPECTED 3

REPORT DOCUMENTATION PAGE

Form Approved
OMB No. 0704-0188

Public reporting burden for this collection of information is estimated to average 1 hour per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden, to Washington Headquarters Services, Directorate for Information Operations and Reports, 1215 Jefferson Davis Highway, Suite 1204, Arlington, VA 22202-4302, and to the Office of Management and Budget, Paperwork Reduction Project (0704-0188), Washington, DC 20503.

1. AGENCY USE ONLY *(Leave blank)*	2. REPORT DATE	3. REPORT TYPE AND DATES COVERED
	July 1999	Annual (1 July 98 - 30 Jun 99)

4. TITLE AND SUBTITLE

Germline Mutations of the Ataxia-Telangiectasia Gene, ATM, as a Risk Factor for Radiation-Associated Breast Cancer

5. FUNDING NUMBERS

DAMD17-97-1-7147

6. AUTHOR(S)

Kenneth Offit, M.D.

7. PERFORMING ORGANIZATION NAME(S) AND ADDRESS(ES)

Sloan-Kettering Institute for Cancer Research
New York, New York 10021

8. PERFORMING ORGANIZATION REPORT NUMBER

9. SPONSORING / MONITORING AGENCY NAME(S) AND ADDRESS(ES)

U.S. Army Medical Research And Materiel Command
ATTN: MCMR-RMI-S
504 Scott Street
Fort Detrick, Maryland 21702-5012

10. SPONSORING / MONITORING AGENCY REPORT NUMBER

11. SUPPLEMENTARY NOTES

12a. DISTRIBUTION / AVAILABILITY STATEMENT

Approved for public release; distribution unlimited

12b. DISTRIBUTION CODE

13. ABSTRACT *(Maximum 200 words)*

Ataxia-telangiectasia (AT) is an autosomal recessive disorder characterized by neurologic and other abnormalities as well as an approximately 100-fold increase in cancer incidence. A single gene, *ATM*, appears to be mutated in the majority of AT families. *ATM* functions as a central component of the cellular response to DNA damage. It has been postulated that heterozygotes for *ATM* mutations have an increased susceptibility to radiation-induced breast cancer. To test this hypothesis we performed a study to test the frequency of *ATM* mutations in woman with breast cancer after therapeutic radiation therapy for Hodgkin's disease. Cases were compared to matched controls treated for Hodgkin's disease with no breast cancer. We observed no truncating mutations of *ATM* in either group, and no difference in the rate of missense mutations (37% controls vs 21% cases). Some missense mutations were observed at polymorphic (>1%) frequencies. Further studies analyzing the frequencies of these mutations in the general population, as well as functional studies, will be necessary to resolve their potential role as susceptibility alleles for lymphoid cancers.

14. SUBJECT TERMS
Breast Cancer

15. NUMBER OF PAGES
19

16. PRICE CODE

17. SECURITY CLASSIFICATION OF REPORT	18. SECURITY CLASSIFICATION OF THIS PAGE	19. SECURITY CLASSIFICATION OF ABSTRACT	20. LIMITATION OF ABSTRACT
Unclassified	Unclassified	Unclassified	Unlimited

FOREWORD

Opinions, interpretations, conclusions and recommendations are those of the author and are not necessarily endorsed by the U.S. Army.

____ Where copyrighted material is quoted, permission has been obtained to use such material.

____ Where material from documents designated for limited distribution is quoted, permission has been obtained to use the material.

____ Citations of commercial organizations and trade names in this report do not constitute an official Department of Army endorsement or approval of the products or services of these organizations.

____ In conducting research using animals, the investigator(s) adhered to the "Guide for the Care and Use of Laboratory Animals," prepared by the Committee on Care and use of Laboratory Animals of the Institute of Laboratory Resources, national Research Council (NIH Publication No. 86-23, Revised 1985).

__/__ For the protection of human subjects, the investigator(s) adhered to policies of applicable Federal Law 45 CFR 46.

____ In conducting research utilizing recombinant DNA technology, the investigator(s) adhered to current guidelines promulgated by the National Institutes of Health.

____ In the conduct of research utilizing recombinant DNA, the investigator(s) adhered to the NIH Guidelines for Research Involving Recombinant DNA Molecules.

____ In the conduct of research involving hazardous organisms, the investigator(s) adhered to the CDC-NIH Guide for Biosafety in Microbiological and Biomedical Laboratories.

Kenneth Ogata 28 July '99

PI - Signature Date

TABLE OF CONTENTS

I. INTRODUCTION

Ataxia-telangiectasia (AT) is an autosomal recessive disorder characterized by neurologic abnormalities (cerebellar degeneration, ataxia, and progressive mental retardation), dilation of blood vessels (oculocutaneous telangiectasia), immune deficiencies, and premature aging.[1] AT homozygotes also suffer from an approximately 100-fold increase in cancer incidence, with leukemias and lymphomas being particularly common.[2-4] A single gene, *ATM*, appears to be mutated in the majority of AT families studied to date.[5] The exact function of the *ATM* gene product is unknown, but the coding sequence bears similarity to yeast genes that serve cell cycle checkpoint and DNA repair functions.[6-9] Evidence suggests that *ATM* functions as a central component of the cellular response to DNA damage.[10]

AT heterozygotes do not manifest the multisystem abnormalities characteristic of the homozygous state. However, these individuals may also be prone to develop malignancies. In a large prospective follow-up study of 161 AT families, the risk of all cancers in female heterozygotes was 3.5 times that of non-carriers.[11] In particular, heterozygous females appeared to have at least a 5.1 times excess risk of breast cancer. Smaller, retrospective studies have also indicated that these individuals may suffer from an increased risk of cancer in general and female breast cancer in particular.[4,12,13] The issue of AT heterozygote cancer susceptibility is not trivial. Although AT is a rare disease, the population frequency of AT heterozygosity is estimated to be 1.4%, and Swift has speculated that up to 8% of all breast cancers in the United States may occur in women who are carriers of an abnormal AT allele.[12,14] The true prevalence of *ATM* mutation among unselected women with breast cancer is probably considerably lower than this figure. Using an exon-scanning PCR single-strand conformation polymorphism (PCR-SSCP) assay, Vorechovsky *et al* were unable to detect any *ATM* mutations in 38 unselected primary breast cancer cases.[15] A recent analysis of 401 women with early-onset breast cancer demonstrated a mutation frequency of only 0.5%.[35] Even accepting the limitations of this method and the possible existence of intronic or regulatory mutations, it is unlikely *ATM* mutations are present in a significant fraction of unselected women with breast cancer. However, because of design limitations, the available studies are unable to exclude a significant relative risk associated with the inheritance of an *ATM* mutation, particularly if a second insult is required to unveil the inherited susceptibility.

It remains possible that women who carry a mutant *ATM* allele are predisposed to breast cancer after exposure to a particular environmental factor, specifically radiation. Swift and his coworkers have suggested that diagnostic, therapeutic, or occupational exposure to radiation may predispose heterozygotes to the development of cancer.[11] This observation is biologically consistent with reports that cultured fibroblasts from AT heterozygotes suffer from an *in vitro* defect in the cellular response to radiation damage.[16,17] Unfortunately, the available studies suffer from significant methodological limitations and the conclusions are by no means universally accepted.[18] A creative approach is required to prove or refute Swift's hypothesis.

If women carrying a single mutant *ATM* allele are indeed susceptible to the genotoxic effects of radiation, one would expect to see an excess number of AT heterozygotes among women with radiation-induced breast cancer. The identification of such women is generally problematic due to difficulties in the quantitation of radiation exposure and in the establishment of a cause-effect relationship between that exposure and the subsequent development of malignancy. However, one well-defined group that does appear to be prone to develop radiation-associated breast cancer is women receiving therapeutic irradiation for Hodgkin's disease (HD). Several large studies have determined that women receiving radiotherapy for HD have a relative breast cancer risk of 1.3-2.2 when compared to controls.[19-26] The latency period is quite long, and the risk appears to be most significant after 15 years of follow-up.[20,21,24] Women who receive their radiotherapy before the age of 30 appear to be more prone to develop radiation-associated breast cancer, and those treated during adolescence have the greatest risk of all.[20,21,27] In a series collected at Stanford University, woman receiving radiotherapy between the ages of 10 and 19 years were 39 times more likely than controls to develop breast cancer in their third decade.[21] Interestingly, this relative risk correlates with an absolute risk of approximately 1.6%,[28] which is similar to the projected frequency of heterozygosity for *ATM* mutation in the general population.

Yahalom *et al* had previously identified 37 patients treated at MSKCC for breast cancer occurring after radiotherapy for HD.[29] We propose to study this cohort for the presence of *ATM* mutations and to perform a case-control to define the risk of breast cancer associated with such mutations. This novel approach will test the hypothesis that *ATM* mutations predispose women to breast cancer after radiation exposure. The approach is particularly innovative because it utilizes a cohort with defined radiation-associated cancer, and is not hindered by the difficulties inherent in studies that attempt to retrospectively attribute cancer to radiation exposure.

II. BODY

A. Protocol Submission and IRB Approval

After DAMD17-97-1-7147 was awarded, the full protocol was submitted to the Memorial Sloan-Kettering Institutional Review Board for approval. Formal approval for the study was obtained on June 24, 1997 and the study was assigned local protocol number 97-81. Patient ascertainment and recruitment began immediately.

B. Subject Ascertainment/Recruitment

At the time of preparation of this report, 34 patients with histories of both breast cancer and Hodgkin's disease identified in the original cohort of Yahalom et al. had undergone genetic counseling, provided informed consent, had donated a blood sample, and had preliminary genotyping information available. 27 Control subjects had genetic results available.

All participants have undergone complete pretest genetic counseling. In the context of that counseling, information has been gathered regarding the clinical features of the breast cancer and Hodgkin's disease diagnoses, as well as the personal risk factors for breast cancer. Individuals with an apparent inherited predisposition to breast cancer have been identified and counseled as to their testing options (e.g. *BRCA1*or *BRCA2*), although such testing has not and will not be provided as part of this study.

In addition, the parents of a child with ataxia-telangiectasia have donated samples to serve as positive controls for the mutation detection techniques being developed.

C. Laboratory Methods

1. Creation of Lymphoblastoid Cell Lines

To facilitate mutation detection, lymphoblastoid cell lines have been established from peripheral blood lymphocytes for 30 cases (4 in progress) by Epstein-Barr-virus transformation.[36]

2. Protein Truncation Testing (PTT) Analysis

PTT presents several advantages when compared to other mutation screening methods. Large gene segments of 2-3Kb can be analyzed at the same time in one reaction and mutations that lead to truncated protein products are efficiently detected.[37,38] Since more than 70% of the mutations in the ATM gene lead to truncated proteins, we chose PTT for initial screening of patient-derived cDNAs.

cDNA Preparation.

Total RNA was then isolated from the lymphoblastoid cell lines using the Ultraspec RNA isolation system (Biotecx Laboratories, INC., Houston, TX) according to manufacturer's instructions. RNA pellets were resuspended in 100µl of DEPC treated water. The total RNA was reverse transcribed with random hexamers using the Superscript cDNA Preamplification Kit (Life Technologies) to generate cDNA. Reverse transcriptions were carried out in a volume of 20µl containing 1-3 µg of RNA, 100 ng Random Hexamers, 20 mM Tris-HCl (pH 8.4), 50 mM KCl, 0.5 mM dNTP, 10 mM DTT, and 200U Superscript II Reverse Transcriptase. The reaction tubes were incubated at 42°C for 60 min and then 70°C for 15 min to terminate the reaction. The product was treated with 2U E.coli Rnase H and incubated at 37°C for 20 min. The cDNA was then diluted with 20µl sterile water and stored at -20°C. A 2.5µl aliquot of cDNA was used as a subsequent PCR template. The entire

coding region of the ATM transcript, composed of 63 exons covering 9.2kb, was divided into 7 overlapping regions. RT-PCR was used to produce transcription templates for PTT. Total RNA was isolated from lymphoblastoid cell lines using the Ultraspec RNA isolation system (Biotecx Laboratories, INC., Houston, TX) according to manufacturer's instructions. RNA pellets were resuspended in 100µl of DEPC treated water. The total RNA was reverse transcribed with random hexamers using the Superscript cDNA Preamplification Kit (Life Technologies) to generate cDNA. Reverse transcriptions were carried out in a volume of 20ul containing 1-3 µg of RNA, 100 ng Random Hexamers, 20 mM Tris-HCl (pH 8.4), 50 mM KCl, 0.5 mM dNTP, 10 mM DTT, and 200U Superscript II Reverse Transcriptase. The reaction tubes were incubated at 42°C for 60 min and then 70°C for 15 min to terminate the reaction. The product was treated with 2u E.coli Rnase H and incubated at 37°C for 20 min. The cDNA was then diluted with 20µl sterile water and stored at -20°C. A 2.5ul aliquot of cDNA was used as a subsequent PCR template.

Primers.

The entire coding region of the 9.2 Kb of ATM transcript and the adjacent untranslated region was divided into 7 overlapping regions (e, f, g, a, b, c, and d) and each segment was analyzed separately. Forward primers were designed to include a T7 promoter sequence for the initiation of transcription by T7 RNA polymerase and an ATG sequence for initiation of translation Primers being used for the PTT analysis are shown in Table 1. along with the optimal annealing temperatures.

RT-PCR.

PCR of each region was carried out in a volume of 50µl containing 1.5 mM $MgCl_2$, 20 mM Tris-HCl (pH 8.0), 50 mM KCl, 200 µM dNTP mix, forward and reverse primers at 10 pM, and 3.75U of Platinum Taq DNA Polymerase (Life Technologies). PCR of each region included of an initial denaturation of 3 min at 94°C, 35-40 cycles of 1 min at 94°C, 1 min at 50-56°C, 3 min at 72°C and a final extension step for 10 min at 72°C.

PTT Analysis

A 25µl reaction mix containing 2-5µl of RT-PCR product, 12.5µl of TnT rabbit reticulocyte lysate (Promega),3.5µl of master mix containing 1.0µl of [^{35}S] methionine (1000 Ci/mmol, NEN Life Science), 0. 5µl of RNAsin (40u/µl), 0.5µl of amino acid mix, 0.5µl of RNA polymerase and 1µl of TnT buffer was be incubated at 30°C for 1.5 hr. Following the PTT, 5µl of product was electrophoresed on 12.5 % SDS-polyacrylamide gels.

Following the electrophoresis, the gel was soaked in Amplify solution (Amersham Life Science) for half an hour, dried and subjected to autoradiography. Figure (1) shows a representative picture of the expected size protein fragments in regions B and D in four of the subjects.

Methodology for detection of the missense mutation 5557 G/A transition in exon 39 in the control population.

In the course of this study, a particular missense mutation (5557G/A) was detected at increased frequency in both cases and controls (*vide infra*). To determine the frequency of this mutation in an unaffected control population a 184 bp segment of the exon 39 was amplified using the following primers:

Forward primer 5' TTTAATATGTGTCAACGGGGCA 3'

Reverse Primer 5' CAGATTTCTCCATGATTCATTTAGA 3'

The reverse primer was designed such that it created a unique Bgl II restriction site (A'GATC) in the normal sequence, whereas this site was abolished in the mutant sequence in which the G/A transition occurred. A 25 µl PCR reaction containing 50 ng of DNA in PCR buffer containing 60mM Tris–HCl, 15 mM Ammonium sulphate,2 mM $MgCl_2$ 0.25 mM dntps, 0.2µ M of the primers and 1U of Taq. polymerase. Amplifications were performed using a DNA thermal cycler 9600(Perkin Elmer Cetus) for approximately 40 cycles at 55°C annealing temperature, followed by running 10 µl of the product on a 3% agarose gel to check for amplification. A second 10 µlaliqout of the PCR product was digested with 10U of BglII (New England Biolabs) at 37°C for 3-4h. The digestion products were run on 3% nusieve GTG agarose (FMC products) . The restriction digestion yielded a 157 and 27 bp products for a normal genotype and a 184 bp product in case of the mutant. (Fig.3). This analysis resulted 26 mutants (20.5%) out of the 127 controls screened.

Mutant DNA samples were amplified by PCR as described above and purified using a Qiaquick PCR purification kit by Qiagen. Purified PCR product was used as a template for sequencing using the dideoxynucleotide chain termination method using the fmol DNA sequencing system by Promega. The reaction products were analyzed on a 8% polyacrylamide /7M urea/.1M Trisborate/2.5mm EDTA gels. After electrophoresis, the gel was transfererd to blotting paper, dried and subjected to autoradiography. Fig (2). Shows some representative mutants with a G/A transition, along with the negative and the positive controls.

3. Restriction Endonuclease fingerprinting (REF)/Sequencing Approach

The ability of this method to detect deletions and insertions as well as base substitutions in homozygotes and heterozygotes which are otherwise missed by PTT has been documented.[31-33,39] While methodology for PTT was put in place and utilized for this proposal (see Appendix A), it was desirable to utilize the most sensitive method of mutation detection on the samples ascertained. In order to have the greatest sensitivity to detect missense mutations, in collaboration with colleagues in the laboratory of Dr. Y. Shiloh (who originally identified the ATM gene), sequence analysis was conducted. DNA was extracted by the ProteinaseK/SDS method. cDNA was prepared from RNA by reverse-transcription reaction with Superscript reverse-transcriptase (Gibco-BRL). cDNA was amplified with Expand Long Template PCR system (Boheringer Mannheim) with primer ATMF and ATMR. The amplified cDNA was divided into two sub fragments (ATM RA and ATM RB) by the following primers (ATMin,AR and ATMBF, ATMout, respectively). The two cDNAs were then directly sequenced. Sequencing was preformed on a ABI377 sequencer with BigDyes modified dNTPs.Sequence chromatograms were analyzed with Sequencher software (Gene codes corporation). Whenever a mutation was found in the cDNA, the corresponding genomic DNA region was amplified with the appropriate primers and sequenced for confirmation.

D. Results

Aim 1. Identification of cases of women who had developed breast cancer after exposure to therapeutic chest irradiation for Hodgkin's disease, and identify appropriate controls.

As projected in the initial design, we have been successful in recruiting 34 patients with breast cancer following therapeutic radiation. An additional two cases were ascertained from the University of Pennsylvania. Twenty nine matched controls have been identified from MSKCC, and one from the University of Pennsylvania. Controls are matched for date of birth (within 3 years), date of diagnosis of Hodgkin's disease (within 5 years, age at time of radiation therapy (within 4 years), radiation dose (within 40 Gy).

Aim 2: Identify women in the cases and controls who are heterozygous for germline ATM mutations

Results of PTT analysis

No protein truncating mutations of ATM were identified in cases or controls.

Results of Full Sequence Analysis

Of 27 control cases analyzed thus far, the missense change 5557 G/A, resulting in an asp->asn change was noted in 8 (29.6%) of samples. The same mutation was noted in 6 of 34 (17.6%) of cases. A nearby mutation 5558 A/T causing an asp->val change was noted in 1 of 27 controls and 2 of 34 cases. The following missense mutations were observed in 2 of 27 (7.4%) of cases: 1810C/T (pro->ser); 2119 T/C (ser->pro), 2289 T/A (phe->leu), 3161C/G pro->arg, 4388 T/G phe->cys, and 6919C/T leu->phe. The 2119 T/C, and 3161 C/G were also observed in 1 of 34 (2.9%) of cases. The 4258 T/G phe->cys was observed in a single control and two of the cases.

In all a total of 14 missense mutations were detected in control and case specimens. Not including the 5557 G/A polymorphism, 10 of 27 (37%) of controls and 7 of 34 cases (21%) demonstrated missense changes

Aim 3: Determine relative risk of radiation associated breast cancer associated with the presence of a germline ATM mutation

The results of the PTT analysis and the preliminary results of the sequence analysis show no significant difference of the prevalence of mutations in cases or controls.

Based on preliminary power calculations, assuming a heterozygote frequency of 1.4% in the general population, and based on a sample size of 30 cases and controls, and the absence of detection of truncating mutations of ATM in the Hodgkin's breast cancer group rules out a large effect of the ATM mutations (relative risk of 22).

E. Discussion; Unexpected findings, and Additional studies in progress.

The finding of the relatively high frequency of missense mutations in both cases and controls was unanticipated. Some of the missense mutations observed have been observed previously. Thus, the mutation 2572 T/C, 3161 C/G, and 6235 G/A had previously been observed in breast tumors, and in B cell cancers.[17] Looking at know structural motifs of ATM, including the three exposed loops, the leucine zipper, the HEAT repeat and the helix turn helix region near the COOH terminus, only one of the reported missense mutations are in these critical regions. Another mutation, 3161 C/G occurs in an exon between an exposed loop and a HEAT repeat. The functional sugnificance of these mutations remains unclear.

The findings of this project argue against a substantial predisposition to radiation-induced breast cancer resulting from ATM mutations (fulfilling the primary specific aim of this proposal). However, the finding of substantial numbers of missense mutations among women with Hodgkin's Disease (with and without breast cancer) raised the possibility that such mutations may confer a susceptibility to Hodgkin's disease. In order to

investigate this hypothesis, a control series of 127 unmatched samples from normal donors was obtained in collaboration with the New York Blood Center. These samples were available as part of an unrelated study as part of an ongoing protocol that allowed genotyping of anonymized DNA samples. Dr P. Kolachna utilizing reagents supported by this project, devised a method of mutation detection for the most common polymorphism 5557G/A (see Methods section). This analysis confirmed the presence of this mutation in 26/127 (20.5%) of cases studied. These data confirm that this mutation is a common polymorphism of ATM, and unlikely to be related to lymphoma risk. However, the incidence of the other mutations detected in the Hodgkin's cases needs to be assessed in a more appropriate series of controls. This determination of the frequency of the other missense changes in a panel of controls matched for ethnic status will be critical to test the possibility that these polymorphisms may confer an increased risk for lymphoid cancer. These experiments are outside the scope of work of the initial proposal but emerge as a result of scientific discovery made possible by the project.

III. KEY RESEARCH ACCOMPLISHMENTS

This study has:

- Assembled a cohort of women and DNA repository of subjects with breast cancers occurring after radiation therapy for Hodgkin's disease

- Shown that there is no increase in truncating mutations of ATM in this cohort of women, suggesting that the increase in breast cancer risk attributable to a germline ATM mutation, if extant, is not very large.

- Demonstrated that there is a significant proportion of missense mutations of unknown biological significance in both the cases and controls (i.e. patients with Hodgkin's Disease). This finding merits additional experiments—both epidemiological (case control) and molecular genetic (functional studies of mutations).

IV. REPORTABLE OUTCOMES

This study has established a DNA resource for a unique group of human subjects with putative radiation induced tumors. This will be available for other studies (e.g. of BRCA mutations and radiation sensitivity). The project provided a research opportunity for the clinical and laboratory post-docs involved in the study (Drs. Robson and Kolachna). The results of this analysis have not yet been communicated but will be prepared for submission.

V. CONCLUSIONS

Within the power constraints defined in the initial proposal, a search for truncating mutations of the ATM gene, as well as a separate full-sequencing analysis performed in collaboration with the investigator who cloned the ATM gene, did not reveal any difference in incidence of ATM mutations in women exposed to ionizing radiation who developed breast cancer after Hodgkin's disease, and matched controls with the same exposure who did not develop Hodgkin's disease. These data argue against a large radiation-associated risk conferred by inheritance of a germline ATM mutation. This project did reveal several missense changes of the ATM gene, some at polymorphic frequencies in Hodgkin's disease cases. The potential significance of the mutations remains to be established and will require additional studies.

VI. REFERENCES

1. Zakian VA. ATM-Related Genes: What Do They Tell Us About Functions of the Human Gene? *Cell.* 1995;82(5):685-687.

2. Taylor AMR, Metcalfe JA, Thick J, Mak Y-F. Leukemia and Lymphoma in Ataxia Telangiectasia. *Blood.* 1996;87(2):423-438.

3. Morrell D, Cromartie E, Swift M. Mortality and Cancer Incidence in 263 Patients with Ataxia Telangiectasia. *Journal of the National Cancer Institute.* 1986;77(1):89-92.

4. Pippard EC, Hall AJ, Barker DJP, Bridges BA. Cancer in Homozygotes and Heterozygotes of Ataxia-Telangiectasia and Xeroderma Pigmentosum in Britain. *Cancer Research.* 1988;48:2929-2932.

5. Savitsky K, Bar-Shira A, Gilad S, Rotman G, Ziv Y, Vanagaite L, Tagle DA, Smith S, Uziel T, Sfez S, Ashkenazi M, Pecker I, Frydman M, Harnik R, Patanjali SR, Simmons A, Clines GA, Sartiel A, Gatti RA, Chessa L, Sanal O, Lavin MF, Jaspers NGJ, Taylor AMR, Arlett CF, Miki T, Weissman SM, Lovett M, Collins FS, Shiloh Y. A Single Ataxia Telangiectasia Gene with a Product Similar to PI-3 Kinase. *Science.* 1995;268:1749-1753.

6. Savitsky K, Sfez S, Tagle DA, Ziv Y, Sartiel A, Collins FS, Shiloh Y, Rotman G. The Complete Sequence of the Coding Region of the ATM Gene Reveals Similarity to Cell Cycle Regulators in Different Species. *Human Molecular Genetics.* 1995;4(11):2025-2032.

7. Sanchez Y, Desany BA, Jones WJ, Liu Q, Wang B, Elledge SJ. Regulation of RAD53 by the ATM-like Kinases MEC1 and TEL1 in Yeast Cell Cycle Checkpoint Pathways. *Science.* 1996;271:357-360.

8. Morrow DM, Tagle DA, Shiloh Y, Collins FS, Hieter P. TEL1, an S. cerevisiae Homolog of the Human Gene Mutated in Ataxia Telangiectasia, Is Functionally Related to the Yeast Checkpoint Gene MEC1. *Cell.* 1995;82(5):831-840.

9. Greenwell PW, Kronmal SL, Porter SE, Gassenhuber J, Obermaier B, Petes TD. TEL1, a Gene Involved in Controlling Telomere Length in S. cerevisiae, Is Homologous to the Human Ataxia Telangiectasia Gene. *Cell.* 1995;82(5):823-829.

10. Meyn MS. Ataxia-Telangiectasia and Cellular Responses to DNA Damage. *Cancer Research.* 1995;55:5991-6001.

11. Swift M, Morrell D, Massey R, Chase CL. Incidence of Cancer in 161 Families Affected by Ataxia-Telangiectasia. *New England Journal of Medicine.* 1991;325(26):1831-1836.

12. Swift M, Reitnauer PJ, Morrell D, Chase CL. Breast and Other Cancers in Families with Ataxia-Telangiectasia. *New England Journal of Medicine.* 1987;316(21):1289-1294.

13. Borresen A-L, Andersen TI, Tretli S, Heiberg A, Moller P. Breast Cancer and Other Cancers in Norwegian Families with Ataxia-Telangiectasia. *Genes, Chromosomes, and Cancer.* 1990;2:339-340.

14. Swift M, Morrell D, Cromartie E, Chamberlin AR, Skolnick MH, Bishop DT. The Incidence and Gene Frequency of Ataxia-Telangiectasia in the United States. *American Journal of Human Genetics.* 1986;39:573-583.

15. Vorechovský I, Rasio D, Luo L, Monaco C, Hammerström L, Webster ADB, Zaloudik J, Barbanti-Brodano G, James M, Russo G, Croce CM, Negrini M. The ATM Gene and Susceptibility to Breast Cancer: Analysis of 38 Breast Tumors Reveals No Evidence for Mutation. *Cancer Research.* 1996;56:2726-2732.

16. Paterson MC, naderson AK, Smith BP, Smith PJ. Enhanced Radiosensitivity of Cultured Fibroblasts from Ataxia Telangiectasia Heterozygotes Manifested by Defective Colony-Forming Ability and Reduced DNA Repair Replication after Hypoxic γ-Irradiation. *Cancer Research.* 1979;39:3725-3734.

17. Lavin MF, Le Poidevin P, Bates P. Enhanced Levels of Radiation-Induced G2 Phase Delay in Ataxia Telangiectasia Heterozygotes. *Cancer Genet Cytogenet.* 1992;60:183-187.

18. Boice JD, Miller RW. Risk of Breast Cancer in Ataxia Telangiectasia (Letter). *New England Journal of Medicine.* 1992;326(20):1358.

19. Boivin J-F, Hutchison GB, Zauber AG, Bernstein L, Davis FG, Michel RP, Zanke B, Tan CTC, Fuller LM, Mauch P, Ultmann JE. Incidence of Second Cancers in Patients Treated For Hodgkin's Disease. *Journal of the National Cancer Institute*. 1995;87(10):732-741.

20. van Leeuwen FE, Klokman WJ, Hagenbeek A, Noyon R, van den Belt-Dusebout AW, van Kerkhoff EHM, van Heerde P, Somers R. Second Cancer Risk Following Hodgkin's Disease: A 20-Year Follow-Up Study. *Journal of Clinical Oncology*. 1994;12(2):312-325.

21. Hancock SL, Tucker MA, Hoppe RT. Breast Cancer After Treatment of Hodgkin's Disease. *Journal of the National Cancer Institute*. 1993;85(1):25-31.

22. Swerdlow AJ, Douglas AJ, Hudson GV, Hudson BV, Bennett MH, MacLennan KA. Risk of Second Primary Cancers after Hodgkin's Disease by Type of Primary Treatment: Analysis of 2846 Patients in the British National Lymphoma Investigation. *British Medical Journal*. 1992;304:1137-1143.

23. Tucker MA, Coleman CN, Cox RS, Varghese A, Rosenberg SA. Risk of Second Cancers After Treatment for Hodgkin's Disease. *New England Journal of Medicine*. 1988;318(2):76-81.

24. Henry-Amar M. Second Cancer after the Treatment for Hodgkin's Disease: A Report from the International Database on Hodgkin's Disease. *Annals of Oncology*. 1992;3(Suppl. 4):S117-S128.

25. Curtis RE, Boice JD, Jr. Second Cancers After Radiotherapy for Hodgkin's Disease. *New England Journal of Medicine*. 1988;319(4):244-245.

26. Prior P, Pope DJ. Hodgkin's Disease: Subsequent Primary Cancers in Relation to Treatment. *British Journal of Cancer*. 1988;58:512-517.

27. Bhatia S, Robison LL, Oberlin O, Greenberg M, Bunin G, Fossati-Bellani F, Meadows AT. Breast Cancer and Other Second Neoplasms after Childhood Hodgkin's Disease. *New England Journal of Medicine*. 1996;334(12):745-51.

28. Shapiro CL, Mauch PM. Radiation-Associated Breast Cancer After Hodgkin's Disease: Risks and Screening in Perspective. *Journal of Clinical Oncology*. 1992;10(11):1662-1665.

29. Yahalom J, Petrek JA, Biddinger PW, Kessler S, Dershaw DD, McCormick B, Osborne MP, Kinne DA, Rosen PP. Breast Cancer in Patients Irradiated for Hodgkin's Disease: A Clinical and Pathologic Analysis of 45 Events in 37 Patients. *Journal of Clinical Oncology*. 1992;10(11):1674-1681.

30. Rassio D, Negrini M, Croce CM. Genomic Organization of the ATM Locus Involved in Ataxia-Telangiectasia. *Cancer Research.* 1995;55:6053-6057.

31. Liu Q, Sommer SS. Restriction Endonuclease Fingerprinting (REF): A Sensitive Method for Screening Mutations in Long, Contiguous Segments of DNA. *Biotechniques.* 1995; 18(3):470-477.

32. Gilad S, Khosravi R, Shkedy D, et. al. Predominance of Null Mutations in Ataxia-Telangiectasia. *Human Molecular Genetics* 1996;5(4): 433-439.

33. Gilad S, Bar-Shira A, Harnik R, et. al. Ataxia-Telangiectasia: Founder Effect Among North African Jews. *Human Molecular Genetics* 1996; 5(12):2033-2037.

34. Easton DF. Cancer Risks in A-T Heterozygotes. *International Journal of Radiation Biology.* 1994;66(6 Suppl):S177-182.

35. FitzGerald MG, Bean JM, Hegde, et. al. Heterozygous *ATM* Mutations do Not Contribute to Early Onset of Breast Cancer. *Nature Genetics* 1997;15:307-310.

36. Svedmyr EAJ, Leiobold W, Gatti RA. Possible use of established cell lines for MLR locus typing. *Tissue Antigens* 1975:5: 186-195.

37. Telatar M, Wang Z, Udar N, Liang T, Bernatowska-Matuszkiewicz E, lavin M, Shiloh Y, et al Ataxia-telagiectasia: Mutations in ATM cDNA detected by protein-truncating screening. *Am J Hum Genet* 1996:59:40-44.

38. Telatar M, Teroaka S, Wang Z,Chun HH, Liang T Sergi Castvelli-Bel, Udar N, et al . Ataxia-Telagiectasia : Identification and detection of founder-effect mutations in the ATM gene in ethnic populations. *Am J Hum Genet* 1998:62:86-97.

39. Gilad S, Khosravi R, Harnik R, Ziv Y, Shkedy D,Galanty Y, Frydman M, Levi J, et al . Identification of mutations using extended RT-PCR and restriction endonuclease fingerprinting, and elucidation of the repertoire of A-T mutations in Israel. *Hum Mut* 1998:11:69-75.

Table 1. PTT[a] Primers being used for Mutation Screening of the ATM Gene

NAME OF THE PRIMER	NUCLEOTIDE SEQUENCE	TM[0] C[b]	FRAGMENT SIZE (bp)	REGION AMPLIFIED
ATMe	Forward(T7)[c]-GAAGTTGAGAAATTTAAGC Reverse AATGCAACTTCCGTAAGGC	50.0°	1316	76-1392
ATMf	Forward(T7)-GCAGATATCTGT Reverse GTAGGTTCTAGCGTGCTAGA	55.0°	1769	1048-2817
ATMg	Forward(T7)-AATGACATTGCAGATATTT Reverse TCAGTGCTCTGACTGGCACT	55.0°	1655	2437-4092
ATMa	Forward(T7)-ACGTTACATGAGCCAG Reverse TCCAAATGTCATGATTTTCAC	50.0°	1387	4048-5435
ATMb	Forward (T7)-CTGGCCTATCTACAGC Reverse CAACCTGCTAAGTGTGGGAT	55.0°	1247	5282-6529
ATMc	Forward (T7)-CAGTGGGACCATTGC Reverse TTCTGACCATCTGAGGTCTCC	55.0°	1534	6322-7856
ATMd	Forward (T7)-GATCACCCCATCACA Reverse TCACACCCAAGCTTTCCATC	55.0°	1521	7651-9172

a Protein Truncating Test

b Annealing Temperature

c Promoter sequence and Translation start codon were added to all the forward primers at their 5' end GGATCCTAATACGACTCACTATAGGAACAGACCATG.

Figure 1 : Mutation screening by Protein Truncation in regions B and D. Autoradiogram of ^{35}S-labeled protein fragment show the resolution of a 57.77 and 46.29 kD normal protein fragment in 4 of the subjects tested in the first 8 lanes., followed by a negative control and luciferase as a positive control.

Figure 2: PCR amplified exon 39 product of normal controls restriction digested with Bgl II and run on 3% gel. Figure shows. the negative control with a 157 bp band and the positive control with the 184 bp band in the right hand panel. The 27 bp band is not visible. The left hand panel shows the digestion products of some of the control population samples. #172, 177 and 181 are positive for the mutation, where as the remaining ones are negative.

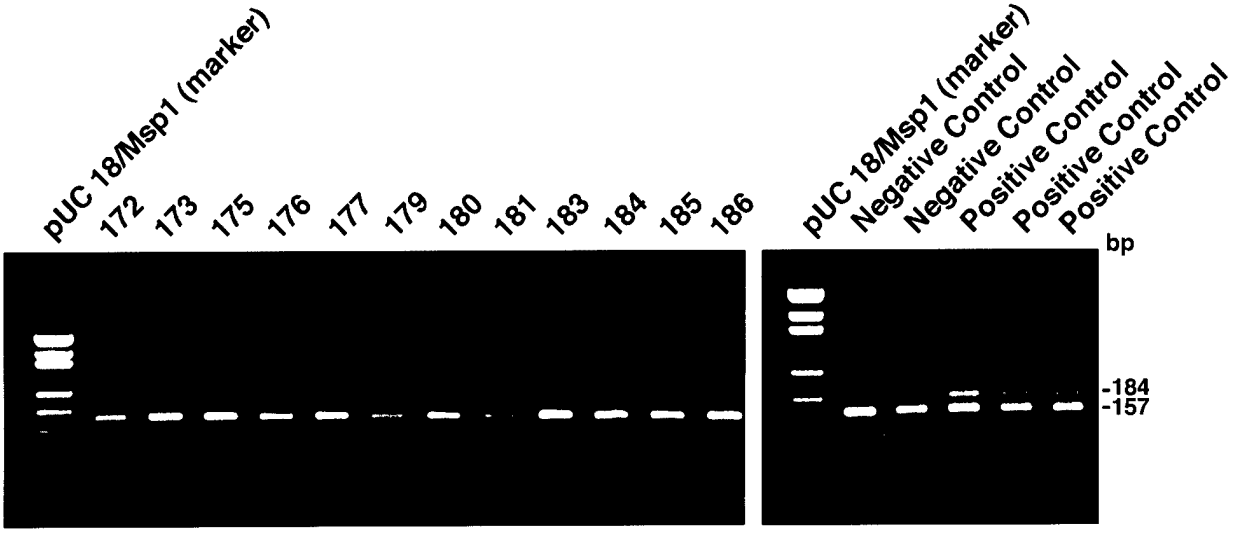

Figure 3: shows the G/A transition at position 458 leading to stretch of 4 A's in case of the positive controls and some of the mutants identified.(arrow), and the absence of A in the negative control.