

AD _____

Award Number: DAMD17-02-1-0460

TITLE: Prevention of Radiation-Induced Breast Cancer
By Amifostine

PRINCIPAL INVESTIGATOR: Michael M. Weil, Ph.D.

CONTRACTING ORGANIZATION: The University of Texas M.D. Anderson
Cancer Center
Houston, Texas 77030

REPORT DATE: June 2003

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.

20031204 086

REPORT DOCUMENTATION PAGEForm Approved
OMB No. 074-0188

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 this 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 June 2003	3. REPORT TYPE AND DATES COVERED Annual (1 Jun 2002 - 31 May 2003)
--	------------------------------------	--

4. TITLE AND SUBTITLE Prevention of Radiation-Induced Breast Cancer By Amifostine	5. FUNDING NUMBERS DAMD17-02-1-0460
--	---

6. AUTHOR(S) Michael M. Weil, Ph.D.

7. PERFORMING ORGANIZATION NAME(S) AND ADDRESS(ES) The University of Texas M.D. Anderson Cancer Center Houston, Texas 77030 E-Mail: Sa64001@odin.mdacc.tmc.edu	8. PERFORMING ORGANIZATION REPORT NUMBER
--	---

9. SPONSORING / MONITORING AGENCY NAME(S) AND ADDRESS(ES) U.S. Army Medical Research and Materiel Command 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)

Exposure to ionizing radiation at doses used in cancer radiotherapy and diagnostic radiology can increase breast cancer risk in women less than 45 years of age. Amifostine is a currently used cytoprotective agent. In addition to its cytoprotective effect, amifostine has been reported to inhibit radiation-induced mutagenesis in cultured cells and to protect against radiation-induced tumorigenesis in rodent model systems. This project is a pre-clinical study designed to determine if amifostine might be effective in preventing breast cancer initiation by medical exposures to ionizing radiation. The experiments will determine if amifostine is protective in a murine model of breast cancer initiation and, if so, determine the optimum dose, route and timing for its administration. The first year's objective was to test a high dose of amifostine administered I.P. prior to irradiation for reduction of ductal dysplasia in an outgrowth assay. In the first year 412 mammary fat pads were transplanted with mammary epithelial cells from irradiated donors. Of these, 306 have been harvested and examined as whole mounts. No dysplasias have been seen, but due to the low outgrowth frequency no conclusions can be drawn yet on the effectiveness of amifostine.

14. SUBJECT TERMS Prevention of diagnostic and treatment induced long-term morbidity, chemoprevention, radiation-induced breast cancer	15. NUMBER OF PAGES 7
	16. PRICE CODE

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

Table of Contents

Cover.....	1
SF 298.....	2
Table of Contents.....	3
Introduction.....	4
Body.....	4
Key Research Accomplishments.....	6
Reportable Outcomes.....	6
Conclusions.....	7
References.....	7
Appendices.....	7

INTRODUCTION:

Historical medical exposures to radiation have been found to increase breast cancer risk. Some currently used diagnostic radiology and cancer radiotherapy procedures deliver doses to mammary tissue in susceptible girls and women that is comparable to the historical exposures. The research supported through DAMD17-02-1-0460 is designed to determine if a chemoprotective agent, amifostine, can prevent radiation-induced ductal dysplasias that progress to mammary tumors. The research design takes advantage of an inbred mouse strain that is susceptible to radiation-induced mammary tumors.

BODY:

The approved statement of work is:

Task 1. To determine if the frequency of radiation-induced ductal dysplasia in BALB/cByJ mice can be reduced by 400 mg/kg amifostine (Months 1-12):

- a. Mice required: 20 BALB/cByJ donors and 400 CB6F1 recipients. The recipients are F1 hybrids of BALB/c and C57BL/6J that are a bit more robust than BALB/c and give lower background levels in the dysplasia assay.

DONORS	TREATMENT	RECIPIENTS
10 BALB/c	saline	200 CB6F1
10 BALB/c	amifostine	200 CB6F1

- b. Methods: Assays on donor mice are set up weekly. In each assay a donor mouse will receive either saline or 400 mg/kg amifostine by intraperitoneal injection. Half an hour later the mouse will be irradiated to the whole body with 1 Gy in a ¹³⁷Cs small animal irradiator. After six weeks the donor will be sacrificed and the mammary epithelial cells will be harvested. The mammary epithelial cells will be transplanted into cleared mammary fat pads in 20 recipients (two fat pads per recipient). Ten weeks later the fat pads will be harvested from the recipients and processed as whole mounts. The whole mounts will be examined for ductal dysplasias and any suspected dysplasias will be confirmed by histological examination.
- c. The results of these experiments will indicate if a high dose of amifostine can reduce the frequency of radiation-induced ductal dysplasias in a susceptible mouse strain.
- d. Since amifostine has been shown to be active against radiation-induced tumorigenesis in other rodent models we anticipate that it will be effective in reducing radiation-induced ductal dysplasia in this system. However, if it is not we will screen other sensitive mouse strains (Weil in press) and other chemopreventive agents, including N-acetylcysteine and captopril .

Task 2. To determine if post-irradiation amifostine treatment can reduce the frequency of radiation-induced ductal dysplasia. If amifostine can be delivered post-irradiation it would be more acceptable as an agent to prevent radiotherapy-induced breast cancer, since it is less likely to provide tumor protection with this schedule of administration (Months 13-20):

a. Mice required: 15 donors and 300 recipients.

DONORS	AMIFOSTINE TREATMENT	RECIPIENTS
5 BALB/c	1/2 hour post-irradiation	100 CB6F1
5 BALB/c	2 hours post-irradiation	100 CB6F1
5 BALB/c	4 hours post-irradiation	100 CB6F1

- b. Methods: As described in Task 1 except amifostine will be injected ½, 2 and 4 hours following irradiation.
- c. The results of these experiments will determine if amifostine can prevent ductal dysplasia if it is delivered after irradiation and, if so, provide a first indication of the how much later.
- d. We anticipate a simple positive or negative result. However, if no protection is seen we may also assay a time as short as 5 minutes post-irradiation.

Task 3. To determine if lower doses of amifostine are effective in preventing radiation-induced ductal dysplasias. Reducing the dose would decrease side effects and alleviate concern about tumor protection if the drug is administered during tumor radiathery (Months 21-30):

a. Mice required: 15 BALB/c donors and 300 CB6F1 recipients.

DONORS	TREATMENT	RECIPIENTS
5 BALB/c	100 mg/kg amifostine	100 CB6F1
5 BALB/c	20 mg/kg amifostine	100 CB6F1
5 BALB/c	1 mg/kg amifostine	100 CB6F1

- b. Methods: As described in *Task 1*, except decreasing doses of amifostine will be used.
- c. The results of these experiments will determine the lowest dose of amifostine, down to 1 mg/kg, that will reduce the frequency of radiation-induced ductal dysplasia.
- d. We do not anticipate problems with this task.

Task 4. Test oral administration of amifostine. Oral administration is easier than intravenous delivery in patients that are not receiving other intravenous medications. (Months 31-36)

a. Mice required: 5 donors 100 recipients.

- b. Methods: As described in *Task 1*, except amifostine will be delivered by gavage.
- c. The results of these experiments will determine if amifostine administered p.o. will reduce the frequency of radiation-induced ductal dysplasia.
- d. If oral administration is ineffective we will examine other routes such as subcutaneous or intramuscular injection. We will also test compounds similar to amifostine but more suited to oral administration such as WR-3689, WR151327, N-acetylcysteine and captopril.

The task for the first year was to determine if high dose amifostine administered I.P. prevents radiation-induced ductal dysplasia. To accomplish this task a breeding colony was established, amifostine obtained and irradiations and transplants performed. The table below summarizes the assays that have been completed through the gross pathology review. Histological examination is being performed on several glands that had subtle, but not dysplastic, changes. An additional 106 glands have been transplanted but are still in different stages of analysis are not included in the table.

Treatment	Donors	Fat pads transplanted	Outgrowths	Dysplasias
Amifostine	6	148	39	0
none	6	158	34	0

The table reveals two shortcomings. The first is a technical matter, the low percentage of fat pads that have outgrowths. Ideally this number should be 70% but in these experiments it was only 24%. We have seen this before and other groups have also experienced similar reductions in outgrowth frequency. The decreased efficiencies have been transient and their cause(s) remain unknown. Although we had normal mammary epithelial cell yields from the donors and good cell viabilities we have, never the less, replaced all the reagents used in the assay. A second shortcoming is that only 412 transplants were made of the 800 proposed. Part of this was due to delays in setting up accounts and vendor shortages of BALB/cByJ mice.

KEY RESEARCH ACCOMPLISHMENTS:

- The ductal dysplasia assay has been set up in the laboratory
- 412 mammary fat pads have been transplanted with mammary epithelial cells from irradiated donors
- 306 glands have been harvested and examined as whole mounts

REPORTABLE OUTCOMES:

None:

CONCLUSIONS:

Additional outgrowths will have to be generated to determine if amifostine treatment prior to irradiation reduces the frequency of ductal dysplasia.

REFERENCES:

None.

APPENDICES:

None.