

# Alteration of Lipid Peroxidation and Total Antioxidant Capacity in Patients With Head and Neck Cancers Following Radiotherapy

Soheila Manifar,<sup>1</sup> Farid Abbassi,<sup>2</sup> Iraj Mirzaii Dizgah,<sup>3</sup> Roya Khatami,<sup>2,\*</sup> Mostafa Esmseil,<sup>2</sup> and Afshin Almasi<sup>4</sup>

<sup>1</sup>Department of Oral and Dental Disease, School of Dentistry, Tehran University of Medical Sciences, Tehran, IR Iran

<sup>2</sup>Department of Oral and Dental Disease, School of Dentistry, Shahed University, Tehran, IR Iran

<sup>3</sup>Department of Physiology, School of Medicine, AJA University of Medical Sciences, Tehran, IR Iran

<sup>4</sup>Clinical Research Development Center, Department of Biostatistics and Epidemiology, School of Public Health, Kermanshah University of Medical Sciences, Kermanshah, IR Iran

\*Corresponding author: Roya Khatami, Department of Oral and Dental Disease, School of Dentistry, Shahed University, Tehran, IR Iran. Tel/Fax: +98-2188956227, E-mail: Royakhatami74@yahoo.com

Received 2015 June 4; Revised 2015 July 14; Accepted 2015 July 14.

## Abstract

**Background:** Head and neck cancers are one of the main causes of cancer related death in the world. The common approach in the treatment of head and neck cancers is radiotherapy either alone or in combination with other therapeutic strategies.

**Objectives:** The aim of this study was to evaluate the effects of radiotherapy on lipid peroxidation and total antioxidant levels as biomarkers of oxidative stress in patients with head and neck cancers.

**Patients and Methods:** Head and neck cancers were documented by two blinded expert pathologists. Whole saliva samples were collected from 30 patients with head and neck cancers before and after radiotherapy and 30 healthy individuals. The unstimulated whole saliva malondialdehyde (MDA), as an indicator of lipid peroxidation, and total antioxidant capacity (TAC) were assayed by thiobarbituric acid and decolorization of ABTS radical cation method, respectively. Data was analyzed by ANOVA followed by Tukey's post hoc test.

**Results:** The levels of antioxidants capacity were significantly increased before radiotherapy in patients in comparison to healthy control ( $P < 0.05$ ). Moreover, antioxidants capacity levels were significantly enhanced after radiotherapy in patients. Furthermore, lipid oxidation levels slightly decreased after radiotherapy in non-significant manner.

**Conclusions:** Lipid peroxidation slightly reduced in patients with head and neck cancers after radiotherapy. However, antioxidant status improved in these patients after radiotherapy.

**Keywords:** Antioxidant, Head and Neck Neoplasms, Malondialdehyde, Oxidative Stress, Radiotherapy

## 1. Background

Head and neck cancer is one of the most common causes of cancer related death and oral cavity with 2% leading reason of death worldwide (1). Head and neck cancers are a heterogeneous groups of tumors originated from different anatomic sites. Prevalence of head and neck cancer related to diverse factors including age, race, sex and geography are differentially changed (2). Incidence of head and neck cancers is largely associated to smoking and alcohol drinking (3).

Free radicals as inevitably product in metabolic process in normal cells play important roles in cancer pathogenesis. Elevated reactive oxygen species (ROS) induces DNA damage, genomic instability and tumor suppressor genes reduction as well as increased expression of proto oncogenes (4). Increased levels of lipid peroxidation, shown by malondialdehyde (MDA) have been reported in different types of cancer. Conversely, the levels of total antioxidant capacity and related enzymes such as glutathione peroxidase, superoxide dismutase and catalase decrease in patients with head and neck cancers (5, 6).

Radiotherapy is the common approach in the treatment of head and neck cancers. Radiotherapy induces DNA damage in exposed cells producing high levels of ROS in site of tumor that ultimately cause cell death. Furthermore, radiotherapy changes signal transduction pathways leading to increased apoptosis rates in cancer cells (7, 8). Nonetheless, effects of radiotherapy on oxidative stress are minimally investigated with conflicting results. Some studies reported that levels of oxidative stress decreased following radiotherapy and conversely, enhanced levels of oxidative stress have been reported by some other investigations (9).

In the recent years, there has been an increasing interest in saliva-based analyses, because saliva collection methods are simple and noninvasive. Oral fluid sampling is safe for both the operator and patient and has easy and low-cost storage. Since saliva was put forth as a potential diagnostic tool, its use for surveillance of disease and general health has become a highly desirable goal in healthcare and medical research (10-13).

Increasing attempts to establish saliva as a diagnostic

matrix have compelling reasons behind. In this regard, it clearly offers an inexpensive, noninvasive and easy-to-use screening method. In addition, it has several advantages over serum and urine for collection, storage, shipping and voluminous sampling. Moreover, handling of oral fluid during laboratory procedures is far easier than blood, because it does not clot, thus reducing the number of required manipulations. Furthermore, noninvasive nature of saliva collection approach could dramatically reduce anxiety and discomfort and thereby increase patients' willingness to continue health-related examinations over time (14-16).

## 2. Objectives

Here, the objective of the present study was to evaluate oxidative stress status in patients with head and neck cancers before and after radiotherapy using measurements of lipid peroxidation and total antioxidant capacity levels.

## 3. Patients and Methods

### 3.1. Patients and Clinical Conditions

Whole intact saliva samples were obtained from 47 patients with head and neck cancers referred to Imam Khomeini Hospital (Tehran, Iran) for radiotherapy from January to September 2014, comprising 23 males and 24 females in pre- and post-radiation. Seventeen patients were excluded because of xerostomia and finally 9 males and 21 females evaluated in pre- and post-radiation. The mean age of patients was 46.8 years (ranged 17 - 62 years) and they were compared with a control group of 30 healthy individuals including 16 females and 14 males with a mean age of 27.9 years without any history of smoking and alcohol consumption. The conditions of all patients concerning antibiotic prophylaxis, nutrition and preoperative therapy were controlled. In this study, unstimulated saliva was used and saliva induction was prevented by avoiding patients and healthy controls for drinking, eating, smoking and all kinds of actions that may induce saliva secretions two hours before samples collection. Collected samples were kept at -70°C until use.

Head and neck tumors were documented by two blinded expert pathologists. Patients with other diseases such as connective tissue pathologies, ulcerative colitis and vasculitis were excluded. The study was performed according to Declaration of Helsinki (2000) of the World Medical Association and all patients filled an informed written consent. Moreover, the study was approved by the Ethical Committee of Shahed University.

### 3.2. Determination of Lipid Peroxidation Using Malondialdehyde (MDA) and Total Antioxidant Capacity (TAC)

TAC of saliva was determined by measuring its ability to decolorization of ABTS radical cation according to pre-

vious fully described methods and the assay calibrated with Trolox (17).

Saliva MDA levels were determined by a method based on reaction with thiobarbituric acid (TBA) at 90 - 100°C (18). MDA and TBA react together in the TBA test reaction, to produce a pink pigment having an absorption maximum at 532 nm. The reaction was performed at pH 2 - 3 at 90°C for 15 minutes. The sample was mixed with two volumes of cold 10% (w/v) trichloroacetic acid to precipitate protein. The precipitate was palliated by centrifugation and an aliquot of supernatant was reacted with an equal volume of 0.67% (w/v) TBA in a boiling water bath for 10 min. After cooling, the absorbance was read at 532 nm. The results were expressed as pmol/mL according to a standard curve, which was prepared with a serial dilution of standard 1, 1, 3, 3-tetramethoxypropane.

### 3.3. Statistical Analysis

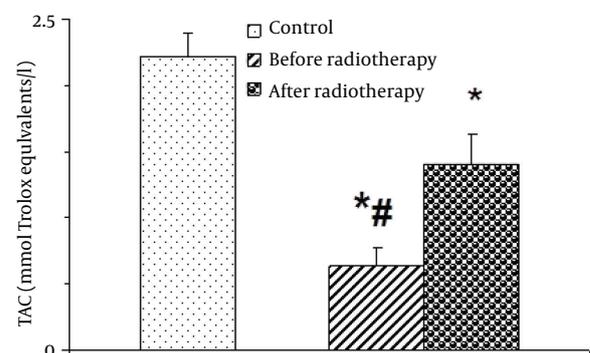
Differences of lipid peroxidation using Malondialdehyde (MDA) and total antioxidant capacity levels in saliva of patients and healthy control in various groups including pre-radiotherapy and post radiotherapy and comparison of these levels with control healthy group were analyzed by ANOVA and Tukey's as Post hoc using SPSS software version 16.00 (SPSS Inc. Chicago, IL, USA).

## 4. Results

### 4.1. Levels of Total Antioxidant Capacity

One-way ANOVA indicated that saliva total antioxidant capacity was altered by radiotherapy [ $F(2, 87) = 20.635$ ;  $P = 0.001$ ] (Figure 1). Post-hoc analysis showed that TAC was significantly low in head and neck cancer group before radiotherapy than after radiotherapy and control group. Interestingly, the level of TAC was remarkably enhanced following radiotherapy in patients compared with before radiotherapy ( $P < 0.05$ ).

**Figure 1.** Unstimulated whole saliva concentration of total antioxidant capacity (TAC) in patients with head and neck cancer before and after radiotherapy and control individuals

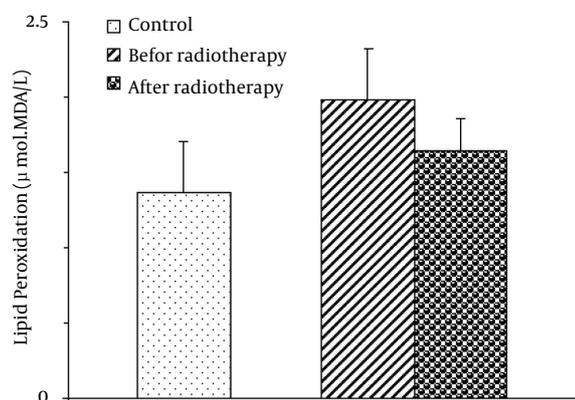


Data are expressed as mean  $\pm$  SEM. \*Different from control; # different from after radiotherapy,  $P < 0.05$ .

## 4.2. Levels of Lipid Peroxidation (MDA)

There was no significant difference in the lipid peroxidation as expressed by MDA levels among groups [F (2, 87) = 1.019; P = 0.365] (Figure 2).

**Figure 2.** Unstimulated whole Saliva Concentration of Malondialdehyde in Patients With Head and Neck Cancer Before and After Radiotherapy and Control Individuals



Data are expressed as mean  $\pm$  SEM.

## 5. Discussion

In the current study, total antioxidant capacity was low in patients with head and neck cancer before radiotherapy. After radiotherapy, the levels of total antioxidant capacity enhanced. Furthermore, lipid peroxidation was slightly high in patients with head and neck cancers, but the levels of MDA as marker of lipid peroxidation decreased following radiotherapy.

The oxidative stress induced lipid peroxidation to generate wide range of products including MDA. Previous investigations showed that MDA plays important roles in carcinogenesis through interaction with DNA and formation of DNA-MDA adduct. DNA-MDA adducts induce mutations in various genes such as tumor suppressor genes and oncogens as well as cell cycle alterations in several cancers (19).

Increased levels of lipid peroxidation alongside decreased levels of antioxidant capacity have been severally reported in patients with head and neck cancers (5, 20). Increased levels of lipid peroxidation and MDA are largely associated to damage red blood cell membranes enriched by polyunsaturated fatty acids. The levels of free radicals increase in head and neck cancers leading to enhanced lipid peroxidation. On the other hands, the levels of antioxidant capacity decreased following enhancement of ROS production in compensatory manner (21).

Free radicals, as mentioned above, play crucial roles in oxidative stress. Some studies showed that the levels of ROS products increased by radiotherapy resulting in

enhancement of oxidative stress (22, 23). Furthermore, some authors indicated that radiotherapy-mediated oxidative stress decreased using antioxidant such as alpha tocopherol (24). Conversely, some other studies indicated that lipid peroxidation decreased after radiotherapy alongside improvement of antioxidant capacity status (21, 25). Decreased levels of MDA may be related to death of tumor cells and tumor load decline as the main source of ROS.

In conclusions, lipid peroxidation slightly reduced in patients with head and neck cancers after radiotherapy. However, antioxidant status seems to be improved in these patients after radiotherapy.

## Acknowledgments

We would like to thank all patients and other volunteers for their contribution to this study.

## Footnote

**Authors' Contributions:**All authors contributed in the analysis and interpretation of data, drafting the manuscript and revising it, and gave final approval of the version to be published.

## References

1. Choong N, Vokes E. Expanding role of the medical oncologist in the management of head and neck cancer. *CA Cancer J Clin.* 2008;**58**(1):32-53. doi:10.3322/CA.2007.0004. [PubMed:18096865]
2. Simard EP, Torre LA, Jemal A. International trends in head and neck cancer incidence rates: Differences by country, sex and anatomic site. *Oral Oncol.* 2014;**50**(5):387-403. doi: 10.1016/j.oraloncology.2014.01.016. [PubMed: 24530208]
3. Freedman ND, Abnet CC, Leitzmann MF, Hollenbeck AR, Schatzkin A. Prospective investigation of the cigarette smoking-head and neck cancer association by sex. *Cancer.* 2007;**110**(7):1593-601. doi:10.1002/cncr.22957. [PubMed: 17724671]
4. Patel BP, Rawal UM, Rawal RM, Shukla SN, Patel PS. Tobacco, Antioxidant Enzymes, Oxidative Stress, and Genetic Susceptibility in Oral Cancer. *Am J Clin Oncol.* 2008;**31**(5):454-9. doi: 10.1097/COC.0b013e31816a61da. [PubMed: 18838881]
5. Rasheed MH, Beevi SS, Geetha A. Enhanced lipid peroxidation and nitric oxide products with deranged antioxidant status in patients with head and neck squamous cell carcinoma. *Oral Oncol.* 2007;**43**(4):333-8. doi: 10.1016/j.oraloncology.2006.02.013. [PubMed:16857409]
6. Beevi SS, Rasheed AM, Geetha A. Evaluation of oxidative stress and nitric oxide levels in patients with oral cavity cancer. *Jpn J Clin Oncol.* 2004;**34**(7):379-85. doi: 10.1093/jjco/hyh058. [PubMed: 15342664]
7. Lawrance JAL, Mais KL, Slevin NJ. Radiologically Inserted Gastrostomies: their use in Patients with Cancer of the Upper Aerodigestive Tract. *Clin Oncol.* 2003;**15**(3):87-91. doi: 10.1053/clon.2002.0199.
8. Sakhi AK, Russnes KM, Thoresen M, Bastani NE, Karlsen A, Smealand S, et al. Pre-radiotherapy plasma carotenoids and markers of oxidative stress are associated with survival in head and neck squamous cell carcinoma patients: a prospective study. *BMC Cancer.* 2009;**9**(1):458. doi:10.1186/1471-2407-9-458. [PubMed: 20025747]
9. Koskinen W. *Prognostic markers in head and neck carcinoma.* University of Helsinki, Faculty of Medicine, Institute of Clinical MedicineHaartman Institute, Department of Virology; 2006.
10. Agha-Hosseini F, Mirzaii-Dizgah I, Mahboobi N, Shirazian S, Harirchi I. Serum and Saliva MMP-3 in Patients with OLP and Oral

- SCC. *J Contemp Dent Pract.* 2015;**16**(2):107-11. [PubMed: 25906800]
11. Mominzadeh M, Mirzaii-Dizgah I, Mirzaii-Dizgah MR, Mirzaii-Dizgah MH. Stimulated Saliva Aminotransaminase Alteration After Experiencing Acute Hypoxia Training. *Air Med J.* 2014;**33**(4):157-60. doi: 10.1016/j.amj.2014.03.004. [PubMed: 25049186]
  12. Mirzaii-Dizgah I, Riahi E. Salivary troponin I as an indicator of myocardial infarction. *Indian J Med Res.* 2013;**138**(6):861-5. [PubMed: 24521627]
  13. Agha-Hosseini F, Mirzaii-Dizgah I, Mirjalili N. Unstimulated whole saliva 25-hydroxycholecalciferol in patients with xerostomia in menopausal women. *Aging Clin Exp Res.* 2013;**25**(2):147-51. doi: 10.1007/s40520-013-0023-z. [PubMed: 23739899]
  14. Mirzaii-Dizgah I, Riahi E. Salivary high-sensitivity cardiac troponin T levels in patients with acute myocardial infarction. *Oral Dis.* 2013;**19**(2):180-4. doi: 10.1111/j.1601-0825.2012.01968.x. [PubMed: 22834943]
  15. Mirzaii-Dizgah I, Hejazi SF, Riahi E, Salehi MM. Saliva-based creatine kinase MB measurement as a potential point-of-care testing for detection of myocardial infarction. *Clin Oral Investigat.* 2011;**16**(3):775-9. doi: 10.1007/s00784-011-0578-z.
  16. Mirzaii-Dizgah I, Agha-Hosseini F. Unstimulated whole saliva parathyroid hormone in postmenopausal women with xerostomia. *J Contemp Dent Pract.* 2011;**12**(3):196-9. [PubMed: 22186816]
  17. Erel O. A novel automated direct measurement method for total antioxidant capacity using a new generation, more stable ABTS radical cation. *Clin Biochem.* 2004;**37**(4):277-85. doi: 10.1016/j.clinbiochem.2003.11.015. [PubMed: 15003729]
  18. Agha-Hosseini F, Mirzaii-Dizgah I, Farmanbar N, Abdollahi M. Oxidative stress status and DNA damage in saliva of human subjects with oral lichen planus and oral squamous cell carcinoma. *J Oral Pathol Med.* 2012;**41**(10):736-40. doi: 10.1111/j.1600-0714.2012.01172.x. [PubMed: 22582895]
  19. Blair IA. DNA adducts with lipid peroxidation products. *J Biol Chem.* 2008;**283**(23):15545-9. doi: 10.1074/jbc.R700051200. [PubMed: 18285329]
  20. Halliwell B. Oxidative stress and cancer: have we moved forward? *Biochem J.* 2007;**401**(1):1-11. doi: 10.1042/B]20061131. [PubMed: 17150040]
  21. Malathi M, Vijay M, Shivashankara AR. The role of oxidative stress and the effect of radiotherapy on the plasma oxidant-antioxidant status in head and neck cancer. *J Clin Diagnos Res.* 2011;**5**(2):249-51.
  22. Kasapović J, Pejić S, Stojiljković V, Todorović A, Radošević-Jelić L, Saičić ZS, et al. Antioxidant status and lipid peroxidation in the blood of breast cancer patients of different ages after chemotherapy with 5-fluorouracil, doxorubicin and cyclophosphamide. *Clin biochem.* 2010;**43**(16):1287-93. [PubMed: 20713039]
  23. Sabitha KE, Shyamaladevi CS. Oxidant and antioxidant activity changes in patients with oral cancer and treated with radiotherapy. *Oral Oncol.* 1999;**35**(3):273-7. doi: 10.1016/s1368-8375(98)00115-8. [PubMed: 10621847]
  24. Chitra S, Shyamala Devi CS. Effect of alpha-tocopherol on prooxidant and antioxidant enzyme status in radiation-treated oral squamous cell carcinoma. *Indian J Med Sci.* 2008;**62**(4):141-8. [PubMed: 18445980]
  25. Gupta A, Bhatt ML, Misra MK. Assessment of free radical-mediated damage in head and neck squamous cell carcinoma patients and after treatment with radiotherapy. *Indian J Biochem Biophys.* 2010;**47**(2):96-9. [PubMed: 20521622]