

RAI is handled by the human body very much similar to stable iodine in the food. It is rapidly absorbed (>90%) from the duodenum into the blood by sodium iodide symporter (NIS) on the apical membrane of enterocytes. The RAI is taken up by the thyroid via iodide transporter (NIS) in the basal membrane of the thyroid follicular cell, in the same way as natural iodine, and is similarly processed through.

NIS is also detectable and active in some extrathyroidal tissues such as the salivary glands, gastric mucosa, and lactating mammary glands. Therefore, these tissues can take up RAI by the action of the NIS. However, contrary to thyroid follicular cells, extrathyroidal tissue shows no long-term retention of iodide. Therefore, in these tissues damage due to β particles of I-131 bombardment is generally of mild degree. Complications like xerostomia, ageusia, lacrimal gland dysfunction, thyroiditis, xerophthalmia, and epiphora have been reported in literature. These side effects can reduce the patient wellness and adversely affect the quality of life [3].

Radiation damage to the salivary glands is one of the common complications of radioiodine therapy as NIS is expressed in salivary glands, especially in the striated ducts of the gland. Primary saliva is produced in the acini of the salivary glands, which subsequently drains into intercalated, striated, and excretory ducts. During the transport in

the ductal system, composition of saliva is actively changed, e.g., sodium and chloride are reabsorbed, and potassium is excreted into the saliva. As radioiodine is mainly concentrated in the ductal system, beta radiation may generate luminal debris which may cause ducts narrowing. These processes can lead to obstruction of the ductal system, causing an inflammatory response in the secretory tissue (sialoadenitis), as well as glandular degeneration [4]. Reduced flow and increased transit of thickened saliva may lead to increased radiation dose delivery to ducts. Scarring after inflammation and collection of debris adds up on degree of obstruction. Consequently, persistent dry mouth adversely affects quality of life. There is reduced patient wellness and increased risk of dental carries.

Previous studies have reported that the salivary gland dysfunction is more frequent in patients receiving higher doses of 100 mCi in differentiated thyroid cancer compared with lower doses of less than 30 mCi [5]. Zanzonico estimated through the calculations based on the collection and counting of saliva samples from a ¹³¹I-treated thyroid cancer patient and a small-scale dosimetry analysis, indicating that the dose to the epithelial lining of the salivary gland ducts was 3- to 4-fold higher than the mean gland dose. This study was based on the speculation that a 1,000-rad-

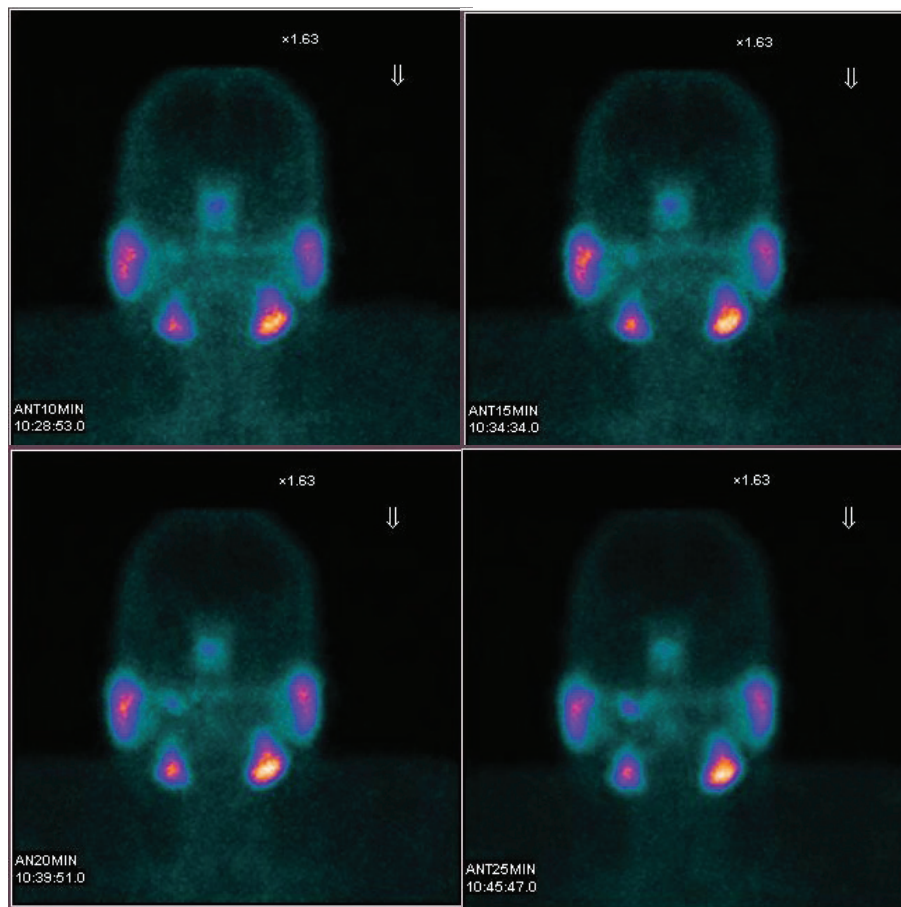


Figure 1. Scintigraphy of salivary gland done with Tc99m, in the patient treated with I-131, shows retention of radiotracer in the left submandibular gland with no clearance even with lemon stimulus, favoring salivary gland duct obstruction.

plus dose to the salivary duct lining is more consistent with sialadenitis than the mean salivary dose of only 300 rad [6].

This complication, although less common following low-dose treatment of RAI, warrants to take precautions to reduce their occurrence. Prevention of the I-131 induced sialadenitis can be done by parasympathomimetic drugs like pilocarpine and sialogogue agents, which accelerate salivary flow and hence radioactive clearance from the salivary glands [7]. However, not all studies advocate the efficacy of this approach [8]. Recently, amifostine is believed to be useful in the prevention of radiation damage through a cytoprotective effect [9]. The concurrent use of physiologic sialogogues like candy and gum, adequate hydration, good oral hygiene, and mouth-washes are the practical effective prevention methods [10].

Recently, a daily supplementation with 800 IU vitamin E for the duration of 5 weeks (1 week before to 4 weeks after RAI therapy) has been introduced to provide the protective effect against radiation-induced dysfunction in salivary glands [11].

Conclusion

We conclude that though the submandibular gland swelling due to submandibular duct obstruction following radioiodine therapy is rare. The occurrence of this complication even after low dose of radioiodine administration should be kept in mind. Maximum preventive actions may be taken before RAI treatment to reduce side effects including those on salivary glands.

Acknowledgement

None

List of Abbreviations

LEAP	Low energy all purpose
NIS	Sodium iodide symporters
RAI	Radioactive iodine

Consent for publication

Informed written consent was obtained from the patient to publish this case in a medical journal, anonymously.

Ethical approval

Ethical approval is not required at our institution for publishing a case report in a medical journal.

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References

- van Luijk P, Pringle S, Deasy JO, Moiseenko VV, Faber H, Hovan A, et al. Sparing the region of the salivary gland containing stem cells preserves saliva production after radiotherapy for head and neck cancer. *Sci Transl Med* 2015; 7:305ra147. <https://doi.org/10.1126/scitranslmed.aac4441>
- Malik M, Lim S, Khaw C, Amir S. Radioiodine I-131 for the therapy of Graves' disease. *Malays J Med Sci* 2009; 16:25–33.
- Mandel L, Liu F. Salivary gland injury resulting from exposure to radioactive iodine: case reports. *J Am Dent Assoc* 2007; 138:1582–7. <https://doi.org/10.14219/jada.archive.2007.0107>
- Van ND. Sialoadenitis secondary to 131I therapy for well-differentiated thyroid cancer. *Oral Dis* 2011; 17:154–61. <https://doi.org/10.1111/j.1601-0825.2010.01726.x>
- Armaghan F, Babak F, Karimi M, Davood B, Mohsen S, Alireza E, et al. Changes in salivary gland function following radioiodine therapy of thyroid diseases: a comparison of high-dose therapy for thyroid cancer and low-dose therapy for benign thyroid disease. *Iran J Nucl Med* 2015; 23:1–7.
- Zanzonico P. Radiation dose to patients and relatives incident to 131I therapy. *Thyroid* 1997; 7:199–204. <https://doi.org/10.1089/thy.1997.7.199>
- Aframian DJ, Helcer M, Livni D, Markitziu A. Pilocarpine for the treatment of salivary glands' impairment caused by radioiodine therapy for thyroid cancer. *Oral Dis* 2006; 12:297–300. <https://doi.org/10.1111/j.1601-0825.2005.01195.x>
- Silberstein EB. Reducing the incidence of 131I-induced sialadenitis: the role of pilocarpine. *J Nucl Med* 2008; 49:546–9. <https://doi.org/10.2967/jnumed.107.049411>
- Mandel SJ, Mandel L. Radioactive iodine and the salivary glands. *Thyroid* 2003; 13:265–71. <https://doi.org/10.1089/105072503321582060>
- Ma C, Xie J, Jiang Z, Wang G, Zuo S. Does amifostine have radioprotective effects on salivary glands in high-dose radioactive iodine-treated differentiated thyroid cancer. *Eur J Nucl Med Mol Imaging* 2010; 37:1778–85. <https://doi.org/10.1007/s00259-009-1368-6>
- Fallahi B, Beiki D, Abedi SM, Saghari M, Fard-Esfahani A, Akhzari F, et al. Does vitamin E protect salivary glands from I-131 radiation damage in patients with thyroid cancer? *Nucl Med Commun* 2013; 34:777–86. <https://doi.org/10.1097/MNM.0b013e328362b1f2>

Summary of the case

Patient (gender, age)	1	Male, 50 years old
Final Diagnosis	2	Salivary gland duct obstruction after radioiodine therapy
Symptoms	3	Swelling in the left submandibular region
Medications	4	N/A
Clinical Procedure	5	Salivary gland scintigraphy with 99mTc-pertechnetate
Specialty	6	Nuclear Medicine