

Release Timing of Thyroid's Carcinoma Patients from the Hospital after Radiiodine (I-131) Therapy based on Dose rate Measurements

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Abstract: The therapeutic treatment of thyroid cancer generally comprises total or hemisphere thyroidectomy and oral administration of radioactive iodine (^{131}I). One consideration in this therapy is the radiation exposure dangers, which have been leading to the Regulatory Authority to guide line criteria on releasing patients from hospitals when the therapy procedure involves ^{131}I activities as greater than 0.91GBq (30mCi). This study was administered to assess the time dependent rate following the administration of ^{131}I activities and estimate the radioiodine effective half-life (Teff) inside the patient's body, which would be useful for releasing patients from hospitals and guide radiation protection recommendations also . Patients were divided into two groups the ablation group (A) and the follow up group (FU). Furthermore, Dose rate, was measured at 1.0 meter. Per administrated Activities was 3700MBq , (100mCi). Patients were measured post 2h after ^{131}I administration and at 24hour intervals up to 96hours. The results demonstrated a bi-exponential radioiodine clearance pattern up to five days with effective half-life Teff values of 15.9 and 25.3hours within the two phase, respectively. Also, majority of the patients follow up could be released from the hospital after 72 hours post-therapy administration as they reach the dose rate limit recommended by the regulatory authority by this time.

Keywords: thyroid cancer, patient release, effective half-life, radioactive iodine-131.

1 Introduction

The greatest potential radiation hazard encountered in the practice of nuclear medicine arises from the administration of radioiodine. Precautions are advised to limit the radiation exposure of member of the public and staff, with whom a treated patient may come into contact. However, recommendations are usually based on measurement of iodine retention or instantaneous dose rate treatment at home has been advocated but the majority of patients who receive high dose of radioiodine are isolated in hospital for a period of time for radiation protection purpose. Iodine-131 has been used for radionuclide therapy for cancer thyroid since 1942. Iodine-131 emits both beta rays with maximum energy of 606KeV , which travel a maximum distance of 3mm in tumor tissue and thereby ensure local treatment of the thyroid tissue. It also emits 364KeV gamma rays used for imaging. Its physical half-life of 8.04 days enables long-term irradiation of the target tissue includes guidelines for patients and all persons that may receive radiation exposures from patients under treatment, under all circumstances. Implementation of radiation protection

guidelines to protect the surroundings of the patient is necessary. The International Commission on Radiological Protection (ICRP) and the National Council on Radiation Protection and Measurements (NCRP), adopted this; these recommend exposure doses to the public of 1mSv/y . i.e., ICRP (1991) and NCRP (1993). Assessing this radiation exposure or the dose rate permits estimation of radioactive iodine inside a patient's body as a function of time post-therapy administration this helps the licensee to calculate the potential of radiation dose to others by means of ^{131}I effective half-time (Teff) inside a patient's whole body. The objective of the present study was to assess the time dependent dose rate following ^{131}I administration in thyroid cancer treatment to estimate the Teff and establish recommendations regarding the time necessary for the patients to be hospitalized in isolated wards. **2**

2.1 Materials and Methods

This study included the selection of 25 patients with cancer thyroid Patient's ages were between 27-57 years. Diagnosis told of papillary and follicular carcinoma. Patients were

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divided into two groups the ablation group (A) and the follow up group (FU). The initial assessment for both groups included measurement of serum thyrotropin (TSH), thyroglobulin performed two weeks after surgery, and Imaging ran with iodine I-131 of thyroid remnants or metastasis. If no remnants are present, patients were not treated with radioactive iodine I-131 and were not started on L-thyroxin (T4) replacement therapy. For ablation of residual thyroid tissue, patients were treated with single dose corresponding to 3700MBq (100mCi) I-131. Patients are isolated in special wards for 4-5 days before discharge from hospital. The environmental background of the survey in the absence of patients was evaluated to be 0.214 ± 0.032 $\mu\text{Sv/h}$, gamma specific constant and correction for the conversion ratio of 0.869 between exposure in R and dose in rad= rem and then in μSv . Original values in Roentgen, corrected by $\times 0.87$ to get rad, and $\times 10$ to get μSv . Average dose rates were measured daily for 5 days for all patients at 0.1m, 0.5m and 1m from the midline abdomen anteriorly, Posterior. in addition, lateral direction of the back, at mid trunk and laterally axially line of patients. Averages of readings were calculated giving Mean, Standard deviation,. The instrument used was RDS-120 Universal Survey Meter (Turku, Finland). The detector is energy- compensated two halogen quenched Geiger Mueller tubes with a referred accuracy of $< \pm 5\%$. All the measurements were taken after emptying the bladder and background radiation was subtracted from the data. The average value for each data set (dose rates from all directions) was used in Teff and 131I activity calculation as this methodology seems less sensitive to iodine distribution inside a patient's body. The average dose rate ($\mu\text{Sv/h}$) measurements by the dosimeter were corrected for environmental background in-patient at distance 1.0 meters post 2 hours to discharge patients after 96 hours.

3 Results and Discussion

Twenty- five therapies selective patients were administrated during the study period, this study divided patients into two groups: Ablative group: included 10 patients (group A), and Follow up groups: included 15 patients group (FU). All patients aged between (27-57) years, mean \pm standard deviation(41.9 ± 9.2) furthermore dose rate, was measured at 1.0 meter. The mean 131I activity administered to the patients was 3.7GBq. Table 1 shows the mean dose rates measured at 1.0 m from the patients and at different times of post-131I administration. All the patients were released from the hospital on the 3rd day. One example of the variation of the dose rate with time is given in Figure 1. This figure indicates that the dose rate decays in a bi-exponential manner where a fast clearance phase is followed by a slower one. The mean Teff to all patients obtained by means of MATLAB software were $15.8 (\pm 3.1)$ and $25.5 (\pm 5.8)$ hours for the initial fast and the second slower clearance phase, respectively. The Teff values calculated for each patient are given in Figure 2. It is

possible to extract from this figure, that for the initial phase 58% of the patients had a Teff value between 11 and 15 hours, and 32% between 16 and 20 hours. For the slower phase these values are 43% between 21 and 25 hours and 42% between 16 and 20 hours, respectively. A comparison between the mean Teff for all the patients and previously reported values are given in Table 2.

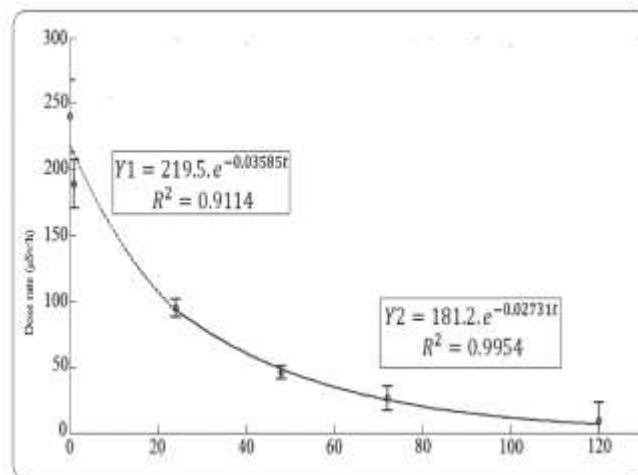


Fig. 1: Pattern of clearance of body burden with administered ^{131}I Post administration time (h).

Table 1: dose rate ($\mu\text{Sv h}^{-1}$) measured at 1.0 m from patients (n=25) as a function of time of post ^{131}I activity administration.

Time	4h	24h	48h	72h	96h
Dose rate	274 ± 34	137 ± 23	103 ± 20	78 ± 19	65 ± 14
Range	103-137	45-78	48-65	17-53	11-37

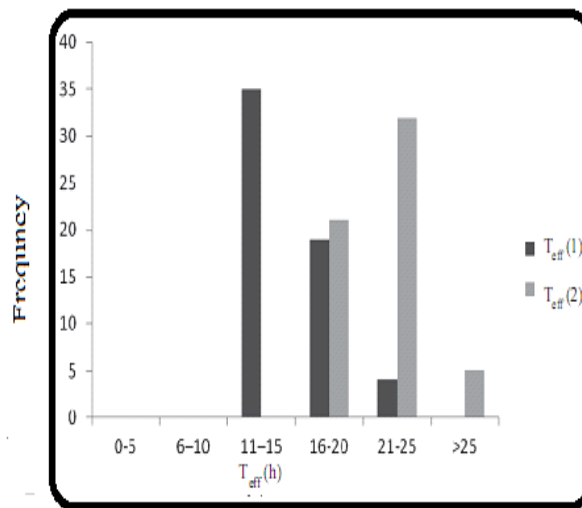


Fig. 2: distribution of effective half-life (T_{eff}) for two phases.

Table 2: comparison of T_{eff} from this study and from others studies.

	First phase T_{eff} (h)	Second phase T_{eff} (h)
Current study (n=25)	15.9	25.3
Ravichandran et al. (n=69)	14.4	22
Barrington et al. (n=86)	12.0	101.7
Tabei et al. (n=562)	11.76	38.64

n=number if subjects studied

4 Conclusions

Surgical resection of the thyroid gland followed by ^{131}I therapy has long been the standard treatment for differentiated thyroid cancer. The ^{131}I T_{eff} in total body of the patients with intact thyroid gland is generally estimated as 5.5d (132h), but in thyroidectomized patients this value is much shorter. The method used in this study estimation the T_{eff} was based on dose rate measurements performed during treatments of patients. This method is somewhat crude compared to bio kinetic studies where the metabolized and eliminated iodine is accounted. The mean dose rate measured at 1.0 m are presented with a range of values (Table 1) as a consequence of many variables involved such as the administered ^{131}I activity, different patients' gender, age and body weight. However, these dose rates constitute important information for radiation protection purposes because they are directly linked to the risk involved in the management of thyroid cancer patients after receiving ^{131}I therapy. Dose rate from patients decreased in a bi-exponential pattern, where there is a slow clearance of radioiodine in the first day post- ^{131}I administration and a slower clearance after this time. Also the rapid clearance in the first phase corresponds to the urinary excretion of radioiodine, whereas the second phase corresponds to the organified ^{131}I with a fast excretion, In a similar study by Barrington et al. (1996) with 86 patients, a bi-exponential pattern was found that the correspondent values for T_{eff} in the first and second phase were $0.50(\pm 0.09)$ and $4.28 (\pm 1.55)$ days, respectively. In past study by Ravichandran et al. (2010), the values were 14.4 and 22.0 hours, respectively. The results from this study, i.e. 15.9 and 25.3 hours, respectively (Table 2), are in approach with both these studies. Some researchers have revealed that a tri-exponential clearance pattern on differentiated thyroid cancer patients (Castronovo et al., 1983; Sasikala et al., 1996), but such a difference in the clearance model may be related to the data sets used for T_{eff} calculation, the methodology applied for dose rate measurement, population size, patients' characteristics and other factors. moreover, it is possible that the T_{eff} value not increase in

both phases according to the ^{131}I activity administered to patients as reported by Tabei et al. (2012) in a study with 562 patients and dosages corresponding to activities of ^{131}I ranging from 3.7GBq (100mCi) to 9.25GBq (250mCi). It was not possible to show this difference in our study due to the small sample size. According to Pacilio et al. (2005), about 80% of the administered ^{131}I activity is eliminated from a patient's body within 48 hours, while Thompson (2001) reported that between 35% and 75% is eliminated within the first 24 hours post administration in all most of patients. At the National Cancer Institute (NCI) of Egypt, all the patients are discharged from the hospital only when the dose rate measured at 1.0 m

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