Factors Associated with Clinical Outcomes of Differentiated Thyroid Cancer Following Radioiodine Therapy in Tanzania

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Abstract: Background: Thyroid cancer is the most common endocrine type of malignancy, accounting for 1-5% of all cancers worldwide. Most of the differentiated thyroid cancers are asymptomatic. Surgery is the mainstay of management to be followed by radioactive iodine (RAI). RAI accessibility is still a challenge in most developing countries including Tanzania. The aim of this study was to determine factors affecting the clinical outcome of patients with differentiated thyroid cancer (DTC) following RAI treatment in a resource limited setting. Methods: This was a prospective cohort study carried out from 2014 to 2018 at the Ocean Road Cancer Institute, in Tanzania. A total of 52 histologically proven differentiated thyroid cancer patients post- near or total thyroidectomy were recruited. All patients received RAI therapy until ablation was achieved, were maintained on thyroxine suppression dose, and were followed for two years. Results: A total of 52 differentiated thyroid cancer patients were recruited after surgery by convenience sampling. The median age of patients was 46 years (range 17-77), and 87% (n=45) were female. Distant metastases were detected in 60% of patients (n=20) at initial presentation. The most common clinical presentation was a neck mass without compression symptoms (85%). Analysis at the end of two years revealed that female gender, clinical-pathological presentation, and the absence of distant metastasis(es) at diagnosis and amount of RAI received, contributed significantly to improved outcome. Conclusion: In a limited resource setting, the outcome of DTC patients post RAI therapy can be improved by early diagnosis hence improving clinical outcome.

Keywords: Radioiodine Therapy, Thyroid Cancer, Tanzania

1. Introduction

Thyroid carcinoma is the most common type of malignant endocrine neoplasia, accounting for 1-5% of all cancers worldwide [1-2]. Follicular thyroid cancer (FTC) and papillary thyroid cancer (PTC) are the main subtypes and are derived from follicular epithelial cells [3]. The exact cause for their occurrence is not known, but dietary iodine deficiency has been implicated as a potential risk factor [4-5]. Most differentiated thyroid cancers (DTC) are asymptomatic and present as painless swelling in the anterior neck. Patients may also present with hoarseness, dysphagia, cough, and dyspnea, which are symptoms often associated with more advanced disease. These patients usually have cervical lymphadenopathy, or distant metastasis, commonly to the lungs or bones and rarely to the brain and liver [3, 6]. For patients with localized disease at presentation, the standard of care immediately after a diagnosis of a neck swelling is to perform surgery to remove the entire lesion and
the remaining normal thyroid tissue as well as the regional lymph nodes [7]. Within six to eight weeks of surgery, patients should receive a RAI ablation dose of 100mCi (3700 mBq) [8]. By this time a patient will also be hypothyroid due to removal of entire thyroid gland. Thyroxine hormone supplementation has to be instituted soon after [9]. Thereafter, patients are followed up after every six months to reassess if the patient has attained complete ablation before repeating surgery or RAI treatment [10].

According to American Thyroid Association (ATA) [11], the clinical outcome of RAI ablation dose should be assessed 6-8 months later. Neck ultrasound is useful in detecting any residual thyroid tissue in the thyroid bed and any presence of local and regional lymphadenopathy [12]. Further diagnostic ¹³¹I whole body scan (WBS) is used to assess the completeness of ablation by detecting any residual thyroid uptake on the thyroid bed or any focus of uptake anywhere in the body which signifies recurrent or distant disease. Thyroglobulin is a tumor marker for DTC and is also used to assess any DTC focus elsewhere in the body [11-12].

DTC represent approximately 1% of cases who receive care at Tanzania’s national cancer referral, Ocean Road Cancer Institute in Dar es Salaam. Currently, patients are diagnosed and undergo surgery at Muhimbili National Hospital and at other referral and regional hospitals and are subsequently referred to ORCI for RAI. We aimed to evaluate factors associated with clinical outcomes following RAI treatment in Tanzania.

2. Methodology

2.1. Study Design

This was a prospective cohort study of patients who received care for DTC at ORCI between 2014 and 2018. ORCI is the only specialized center for cancer treatment in Tanzania that offers RAI therapy and is currently home to an isolation facility for RAI treatment and two dedicated gamma cameras that are used for post RAI imaging as well as diagnostic whole body scan with ¹³¹I during follow up visits. Convenience sampling was used to recruit patients to the study cohort.

This study was approved by the Muhimbili University of Health and Allied Sciences (MUHAS) Institutional Review Board.

2.2. Treatment Protocol

Since ORCI does not currently offer surgical services, all surgeries for thyroid cancer are performed at Muhimbili National Hospital and other referral and regional hospitals in Tanzania. According to our institutional protocol, repeat ¹³¹I treatments were administered at 6-8 month intervals until ablation of all functioning tumor tissue had been achieved which were preceded by surgery whenever indicated. After total ablation, all patients were maintained on maximum tolerated thyroxine therapy.

The outcome of RAI therapy was assessed by neck ultrasound, thyroglobulin levels and ¹³¹I whole body scan at six months post ablation and then every 6-8 months. Those who had incomplete ablation after the first ablation with ¹³¹I 100 mCi (3700 mBq) at six months, they underwent surgical remnant removal then received a second dose of 150 mCi (5550 mBq) and were re-evaluated after another six month. If a complete ablation was not attained then a third dose of either 150mCi (5550 mBq) or 200mCi (7400mBq) depending on the site of metastasis was given post remnant surgical removal whenever accessible.

2.3. Data Collection

Clinicopathologic data was abstracted from the medical record, including age, gender, clinical presentation, type of surgery, and the presence or absence of distant metastasis(es) prior to RAI therapy. Treatment data, including number and reasons for repeated surgeries, dose(s) of RAI therapy, dose of thyroxine, and frequency and dose of repeated RAI therapies were abstracted from the medical records. Detailed follow-up information regarding detection of incomplete or complete ablation and the patient’s vital status were collected every six months for a follow-up period of two years following initial treatment. The clinical outcomes were defined as complete remission, incomplete ablation, or death. Complete remission was defined as complete ablation six months post RAI therapy and at any six month interval during follow up. Incomplete ablation was defined as residual or disease progression.

2.4. Statistical Analysis

All data were entered into Excel files. Statistical Package for Social Science (SPSS) version 20 was used for all statistical analyses. Potential risk factors and clinical outcome were considered as dependent variables. Sociodemographic factors, disease profile, and treatment doses were considered as independent variables. Descriptive statistics were used. Associations between risk factors and treatment outcome are calculated by chi square.

3. Results

A total of 52 patients who met the inclusion criteria and consented to participate in the study were recruited.

3.1. Sociodemographic and Clinical Characteristics of Study Participants

As summarized in Table 1 the median age of patients was 46 years (range 17-77), 87% of patients were female, and papillary carcinoma was the most common pathologic subtype (56%). Patients with a follicular subtype were more likely than patients with papillary or Hurthle cell subtypes to have distant metastases at time of diagnosis (60% vs. 30% vs. 0%, p=0.02).
Table 1. Demographic and Clinical Characteristics of Patients at Presentation (N=52).

| characteristics          | Histological subtypes | Total | p-value |
|--------------------------|-----------------------|-------|---------|
|                          | Follicular | Papillary | Hurthle cell |       |         |
| Age                      |            |           |              |       |         |
| >45                      | 13        | 12        | 1            | 26    | 0.2     |
| <45                      | 7         | 17        | 2            | 26    |         |
| Sex                      |            |           |              |       |         |
| Male                     | 4         | 3         | 0            | 7     | 0.4     |
| Female                   | 16        | 26        | 3            | 45    |         |
| Family history of DTC    |            |           |              |       |         |
| Present                  | 7         | 8         | 1            | 16    |         |
| Absent                   | 13        | 21        | 1            | 35    | 0.7     |
| Clinical presentation    |            |           |              |       |         |
| prior to surgery         |           |           |              |       |         |
| Necker mass without     | 17        | 27        | 2            | 46    | 0.4     |
| compression              |           |           |              |       |         |
| Necker mass with        | 3         | 2         | 1            | 6     |         |
| compression              |           |           |              |       |         |
| Local extension          |           |           |              |       |         |
| Present                  | 9         | 12        | 1            | 22    |         |
| Absent                   | 11        | 16        | 2            | 29    | 0.9     |
| Capsular invasion        |           |           |              |       |         |
| Present                  | 11        | 12        | 3            | 26    |         |
| Absent                   | 9         | 16        | 0            | 25    | 0.1     |
| Metastasis at presentation |         |           |              |       |         |
| Present                  | 12        | 8         | 0            | 20    |         |
| Absent                   | 8         | 21        | 3            | 32    | 0.02    |

At the initial six-month follow-up, two patients (4%) were deceased, and seven (13%) were lost to follow-up. Of 43 evaluable patients who underwent an initial assessment at six-month follow-up, 49% (n=21) were determined to have an incomplete ablation, and 51% (n=22) were determined to have a complete ablation (Table 2). Of the 21 patients with incomplete ablation, 71% (n=15) received a second dose of either 150mCi (5550mBq) or 200 mCi (7400 mBq) at 12-month follow-up, and 10% (n=2) received a second dose at 18-month follow-up. Seven patients (32%) received a third dose of 200mci (7400 mBq) at 18 months. At the end of two years, 39 (75%) patients were evaluable for disease status and vital status (Table 3).

Table 2. Factors associated with outcomes at six months post RAI (n=43).

| Category                          | Incomplete ablation | Complete ablation | Death | Total | p-value |
|-----------------------------------|---------------------|-------------------|-------|-------|---------|
| Metastasis at diagnosis           |                     |                   |       |       |         |
| Present                           | 10                  | 3                 | 1     | 14    |         |
| Absent                            | 11                  | 18                | 0     | 29    | p=0.02  |
| Age                               |                     |                   |       |       |         |
| >45                               | 9                   | 11                | 0     | 20    |         |
| <45                               | 12                  | 10                | 1     | 23    | p=0.52  |
| Sex                               |                     |                   |       |       |         |
| Male                              | 2                   | 3                 | 0     | 5     |         |
| Female                            | 19                  | 18                | 1     | 38    | p=0.83  |
| Clinical presentation             |                     |                   |       |       |         |
| Necker mass without compression   | 18                  | 20                | 0     | 38    |         |
| Necker mass with compression      | 3                   | 1                 | 1     | 5     | p=0.01  |
| Type of surgery                   |                     |                   |       |       |         |
| Near total thyroidectomy          | 18                  | 13                | 0     | 31    |         |
| Total thyroidectomy               | 3                   | 8                 | 1     | 12    | p=0.06  |

3.2. Factors Affecting Outcomes Post-Therapy

Analysis of factors associated with incomplete ablation at six-month follow-up is summarized in Table 2. The presence of distant metastasis at diagnosis (p=0.02) as well as the presence of compression symptoms associated with a neck mass (p=0.01) were associated with incomplete ablation. An analysis of factors associated with incomplete ablation at two-year follow-up is summarized in Table 3. The presence of metastasis(es) at diagnosis (p=0.02), female gender (p=0.02), and requirement for more than one RAI therapy (p=0.00) were associated with incomplete ablation at 24 months follow-up.
4. Discussion

This two years cohort study done at ORCI the only cancer center offering radioactive iodine treatment to DTC patients in Tanzania presents the outcome of patients treated who underwent post-surgical RAI therapy in Tanzania.

It shows that the occurrence of DTC between the age groups of below 45 and above 45 years of is equal. This is different from what is seen in other studies done elsewhere whereby most of DTC patients where >45 years [13]. This could mean that the DTC in our setting occurs at a relatively younger age and maybe it is not age related. The longstanding endemic goiter which affects many areas of Tanzania [14] maybe the culprit.

Endemic goiter due to longstanding dietary iodine deficiency could also be the cause of the high occurrence of follicular and Hurthle cell histology subtypes [5]. Studies elsewhere have shown the relationship between dietary iodine deficiency and occurrence of follicular and Hurthle cell carcinomas [15]. Further we found out that the family history had low influence on occurrence unlike studies done elsewhere which could also be accounted by endemic goiter for decades [13]. Most of the patients who presented with metastasis were those with follicular subtypes. Follicular subtype is known for being more aggressive [16].

It is known that the more the extensive the surgery is the lower the chances of recurrence and hence the better the prognosis [17]. In this study most patients presented with goiter without compression symptoms. However about 50% of patients had to go for second or third completion thyroidectomy before receiving RAI therapy since successful RAI treatment requires minimal or no thyroid remnant. This further delays RAI therapy and may contribute to poor outcome [18].

Outcome is influenced by pathological subtype whereby follicular subtype is more aggressive and associated with poor prognosis in comparison to papillary [18]. Similarly in this study, we found most patients with metastases at diagnosis were those with follicular subtype. Furthermore about 50% of patients had capsular invasion, a sign of advanced disease at diagnosis, which without aggressive post-surgical RAI therapy may lead to poor outcome.

All patients received the same ablation dose and assessment was done at 6 months to ascertain whether complete ablation has been achieved and factors contributing to it. It was evident that distant metastasis at diagnosis and clinical presentation were associated with poor outcome at six months. This is similar to what is seen elsewhere, whereby those patient who present with metastasis have up to 40% 10 years survival while those who present without metastasis the 10 year survival is >95% [19].

Looking at outcome at two years whereby those who had had incomplete ablation at six months received additional RAI therapy until complete ablation was achieved. The factors affecting outcome had increased whereby now we have metastasis at diagnosis, gender, number of RAI therapies and cumulative RAI therapies received. Male gender has been associated with more aggressive disease. We see gender being an important contributor of outcome at two years, and not immediately. But also repeated RAI therapies for those with incomplete ablation improves outcome, by attaining ablation after initial treatment [20].

This study gives an overall picture of thyroid DTC management in Tanzania. We saw 14 out of 52 patients had an advanced disease evidenced by distant metastasis on

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**Table 3. Factors associated with outcomes at twenty four months post RAI (N=39).**

| Category                        | Incomplete ablation | Complete ablation | Death | Total | p-value |
|---------------------------------|---------------------|-------------------|-------|-------|---------|
| Metastasis at diagnosis         |                     |                   |       |       |         |
| Present                         | 6                   | 5                 | 1     | 12    | 0.02    |
| Absent                          | 5                   | 22                | 0     | 27    | 0.57    |
| Age                             |                     |                   |       |       |         |
| >45                             | 5                   | 14                | 1     | 20    |         |
| <45                             | 6                   | 13                | 0     | 19    |         |
| Sex                             |                     |                   |       |       |         |
| Male                            | 2                   | 2                 | 1     | 5     | 0.02    |
| Female                          | 9                   | 25                | 0     | 34    |         |
| Nearest presentation            |                     |                   |       |       |         |
| Neck mass without compression   | 8                   | 26                | 1     | 25    |         |
| Neck mass with compression      | 3                   | 1                 | 0     | 3     | 0.08    |
| Type of surgery                 |                     |                   |       |       |         |
| Near total thyroidectomy        | 8                   | 20                | 1     | 29    | 0.83    |
| Total thyroidectomy             | 3                   | 7                 | 0     | 10    |         |
| Total RAI doses received        |                     |                   |       |       |         |
| One                             | 3                   | 20                | 1     | 24    |         |
| Two                             | 2                   | 6                 | 0     | 8     |         |
| Three                           | 6                   | 1                 | 0     | 7     | 0.00    |
| Cumulative amount of RAI Received in mCi |       |                   |       |       |         |
| 100                             | 3                   | 20                | 1     | 24    |         |
| 200                             | 2                   | 6                 | 0     | 8     |         |
| 250                             | 1                   | 0                 | 0     | 1     |         |
| >300                            | 5                   | 1                 | 0     | 6     | 0.01    |

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presentation. This contributes to poor clinical outcome by delaying to achieve complete ablation. We have seen about 15 out of 52 patients had to receive multiple therapies of RAI post repeated and completion surgery to achieve complete ablation. These multiple repeated doses translate to an extra financial burden to the patients, relatives and the government at large which could be significant in a limited resource setting.

This calls for increased awareness to the public and knowledge among clinicians of the importance of early detection, because in such patients complete ablation can be achieved by a single RAI ablation dose.

RAI is purchased and imported from South Africa (SA), where an ablation dose of 100 mCi (3700 mBq) costs about 900USD which most patients cannot afford. Thanks to this study, the government is now aware of this cost implication to the vast majority of Tanzanian, consequently all public patients at ORCI now receive RAI paid for by the government. This further improves the treatment outcome of DTC patients since the patients are guaranteed of timely availability of RAI.

One of the limitations of this study was that patients are referred to ORCI after surgery has been done at either referral or regional hospitals because there are no surgical services at ORCI. Thus it is not possible to stage and ascertain the TNM status of the DTC prior to surgery and treatment; however assessment of metastasis was done at ORCI before RAI therapy. Another limitation was challenges of sending back patients with partial thyroidectomy for completion thyroidectomy because some had surgery in the regional or zonal hospitals far away from ORCI and refereeing them back was costly and the majority could not afford.

5. Conclusion

This study has shown that, the clinical outcome of DTC patients following RAI therapy at ORCI in Tanzania may be determined by presence of distant metastases at presentation, male sex, and extent of initial surgery.

6. Recommendations

The outcome of DTC patients can be improved by early diagnosis, and early referral for RAI post-surgery. This will allow complete ablation to be achieved by low RAI doses and short hospital stay resulting in reduced burden to the government and families for treating patients with DTC.

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Authors Contributions

LS: designed the study, collected the data, made the analysis, wrote the manuscript
TM: supervised study designing, data collection, data analysis and manuscript writing
JM: assisted in study designing, data analysis and critical review
KM: supervised study designing, data collection, data analysis and manuscript writing
BS: supervised study designing, data collection, data analysis and manuscript writing
KVL: data analysis, manuscript writing, critical review

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