Objectives: To evaluate the clinicopathological features and treatment outcomes of papillary thyroid carcinoma tall cell variant (PTC-TCV) in Saudi population.

RESULTS:
- Mean age of cohort was 52.3 years (range: 46-80), with female predominance (73.8%).
- Mean tumor size was 3.62 cm (range: 0.4-10). Rates of LVSI (66.7%), positive pathological lymph nodes (69.2%), multifocality (42.9%) and extrathyroidal extension, (45.3%).
- Median follow-up was 37.4 months (range: 6-60). Local recurrence rate were seen in 6/42 (14.2%) with female predominance (73.8%).

CONCLUSION: Papillary thyroid carcinoma tall cell variant is associated with aggressive biological behavior.

Methods: This retrospective study were medical records of 776 treated PTC patients between December 2007 and 2015, at King Fahad Medical City and King Khalid University Hospital, Riyadh, Kingdom of Saudi Arabia, was approved for PTC-TCV. Total 42 (5.4%) patients were found to have PTC-TCV, which were investigated for demographic, symptoms, histopathological features, and treatment outcomes locoregional control (LRC), distant metastasis control (DMC), disease free survival (DFS), and overall survival (OS) rates.
Most of papillary thyroid carcinoma (PTC) are associated with an exceptional long-term treatment outcomes including DFS and OS rates; however, some of histopathological variants of PTC have been found with more aggressive biologic behavior and poor perspective. Among these histopathological variants, tall cell variant (TCV) is considered the most aggressive variant of PTC. Mainly, PTC-TCV is characterized by a height greater than twice its width, eosinophilic cytoplasm, basaloid nuclei along with the conventional characteristics of PTC. In 2004, World Health Organization (WHO) defined a PTC as TCV if it contained 50% or more tall cells. Some controversies regarding the proportion of tall cell (30-75%) required for diagnosis of PTC-TCV still persist; however, most of institutes accept the threshold of > 30% to confirm PTC-TCV. TCV has some distinctive differences with classic variant of PTC; A) frequent presence of lymphovascular space invasion (LVSI) and extrathyroidal extension (ETE), B) greater prevalence of pathological positive lymph nodes and distant metastases at the time of diagnosis, and C) its 10-year DFS and OS rates have been reported to be approximately 10-15% lower than that of classic PTC. However, data around PTC-TCV in Saudi population is very scanty. We undertook the present study to evaluate the various clinicopathological characteristics and treatment outcomes of PTC-TCV in Saudi population.

Methods. Patients were reviewed retrospectively. After an authorized approval from the institutional review committee, medical charts of 42 patients with PTC-TCV, who were treated at King Fahad Medical City and King Khalid University Hospital, Riyadh, Kingdom of Saudi Arabia between December 2007-2015, were investigated using database system. Inclusion criteria was patients PTC-TCV histopathology of all ages. Exclusion criteria was defined <10% proportion of tall cells in histopathological specimens. Patients with PTC-TCV were retrieved in the following fashion; clinical variables including age at the diagnosis, gender, and symptoms at the time of diagnosis were reviewed. Histopathological assessment was performed by a skilled histopathologist. Different histopathological variables including the size of primary tumor, presence of LVSI and ETE, multifocality, positive pathological lymph nodes, and tumor lymph node, and metastasis staging were collected. Data regarding different treatment modalities, including the type of thyroidectomy, neck dissection, and adjuvant radioactive iodine (RAI) ablation were also recalled.

Statistical analysis. The primary endpoint was DFS. Secondary endpoints were the locoregional control (LRC), distant metastasis control (DMC) and OS rates. The DFS was defined as, the duration between the date of thyroidectomy and the date of documented all-site disease recurrence, death from cancer and/or last follow-up (censored). All probabilities of LRC, DMC, DFS, and OS rates were shown using the Kaplan-Meier method (comparisons for different survival curves were performed using log rank). All statistical analyses were performed using the the Statistical package for Social Sciences (IBM Corp., Armonk, NY, USA) version 24.

Results. Among the 776 PTC patients managed in our centers between December 2007-2015, 42 patients

| Variables                                      | N (%)                  |
|-----------------------------------------------|------------------------|
| Age (years)                                   | Mean (range) SD        |
| Mean (range) SD                               | 52.3 (46-80) SD ± 15.7 |
| Gender                                        | 11 (26.2) Female       |
| 31 (73.8) Male                                |
| Tumor size (cm)                               | Mean (range) SD        |
| Mean (range) SD                               | 3.62 (0.4-10) SD ± 2.05|
| Primary T stage                               |                        |
| T1                                            | 13 (31.0)              |
| T2                                            | 10 (23.7)              |
| T3                                            | 13 (31.0)              |
| T4                                            | 6 (14.3)               |
| Positive pathological lymph nodes             |                        |
| N0                                            | 14 (33.3)              |
| N1                                            | 28 (66.7)              |
| Metastasis at time of diagnosis               |                        |
| M0                                            | 39 (92.8)              |
| M1                                            | 3 (7.2)                |
| Multifocality                                 |                        |
| Yes                                           | 18 (42.9)              |
| Unifocal                                      | 24 (57.1)              |
| Lymphovascular space invasion                 |                        |
| Yes                                           | 17 (40.5)              |
| No                                            | 25 (59.5)              |
| Extra-thyroidal extension                     |                        |
| Yes                                           | 19 (45.3)              |
| No                                            | 23 (54.7)              |
| Total thyroidectomy                           |                        |
| 42 (100)                                      |
| Neck dissection                               |                        |
| Lateral                                      | 9 (21.4)               |
| Central                                      | 33 (78.6)              |
| Adjuvant radioactive iodine therapy           |                        |
| No                                           | 3 (7.2)                |
| Yes                                          | 39 (92.8)              |

N - number, SD - standard deviation

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(5.4%) were found to have PTC-TCV. Mean age of cohort was 52.3 years (range: 46-80), with female predominance (73.8%). Different attributes of cohort are shown in Table 1.

All patients underwent total thyroidectomy and near-total thyroidectomy. Most of cohort underwent central and lateral neck dissection (78.6%). Mean tumor size was 3.62 cm (range: 0.4-10). Lymphovascular space invasion was observed in 25/42 patients (59.5%), while positive pathological lymph node were seen in 28/42 patients (66.7%). Positive pathological lymph nodes were seen in 62.5% of patients with tumor size ≤ 2 cm (16 patients) as compared with 28 patients with tumor size > 2 cm (64.3%) with \( p = 0.20 \). Multifocality was seen in 18/42 patients (42.9%). No correlation was found between size of primary with multifocality, ETE \( (p=0.12, 0.17) \) and LVSI \( (p=0.21) \). Triiodothyronine \( (T_3) \) and thyroxine \( (T_4) \) stage was seen in 19 patients \( (45.3\%) \). Two patients (4.8%) had distant metastasis at the time of diagnosis.

Adjuvant RAI was given to most of cases (39 patients, 92.8%). Median delay between surgery and adjuvant RAI was 8.5 weeks (range: 6-13). Doses were, 30 millicurie (mCi) in 5 patients (11.8%) for patients with primary size ≤ 2 cm without other risk factors; 100 mCi in 9 patients (21.4%) with positive margins, ETE and LVSI; 150 mCi in 23 patients (54.7%) with positive pathological lymph nodes and 200 mCi in 2 patients (4.8%) with distant metastasis. Postoperative radiation therapy (PORT) to thyroid bed and regional lymph nodes was given to 4 patients (3 patients with skeletal muscle invasion, one patient with tracheal invasion). Mean dose for PORT was 60 Gy (range: 60-66 Gy), given as daily 2 Gy/fraction, 5 days/week over 6-6.5 weeks (30-33 fractions). Median follow-up was 37.4 months (range: 6-60). Local recurrence rate (LRRs) were seen in 6/42 (14.2%) patients (Table 2). The LRRs were salvaged with neck resection (3 patients) followed by RAI (2 patients). Distant metastasis was seen in 8/42 (19%) patients. Among whom, 5 patients were treated with RAI, one patients with brain metastasis was managed with resection and radiation therapy and one patient with bone lesion was treated with palliative radiation therapy (Figure 1).

At the time of analysis, we found 5 year 82.3% LRC, 77.8% DMC, 69.2% DFS, and 86.7% OS (Figure 2 & 3). Further, univariate and multivariate analyses of different prognostic components for DFS and OS showed PTC-TVC as an important independent prognosticator (Odds ratio: 4.2; 95% confidence interval: 1.79-7.3; \( p = 0.03 \)).

**Discussion.** In the present study, PTC-TCV constituted 5.4% of all PTC cases, which is in agreement

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**Table 2 - Treatment outcomes in our tall-cell variant papillary thyroid carcinoma.**

| Outcomes                                  | N (%) |
|-------------------------------------------|-------|
| Locoregional recurrences alone            | 6 (14.2) |
| With RAI                                  | 4/39 (10.3) |
| Without RAI                               | 1/3 (33.3) |
| Thyroid bed                               | 2 (4.8) |
| Regional lymph nodes                      | 4 (9.5) |
| Distant metastasis alone                  | 8 (19.0) |
| With RAI                                  | 7/39 (17.9) |
| Without RAI                               | 1/3 (66.7) |
| Lungs                                     | 5 (11.8) |
| Bones                                     | 2 (4.8) |
| Brain                                     | 1 (2.4) |
| Locoregional recurrence with distant metastasis | 3 (7.1) |
| 5 year locoregional control               | (82.3) |
| 5 year Distant metastasis control         | (77.8) |

RAI - radioactive iodine ablation

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**Figure 1 -** Whole body scintigraphy showing iodine avid bilateral lungs, neck, and right femur uptake.
Tall cell variant PTC ... Al-Qahtani et al

with the reported prevalence of 1.3-13% among all PTC.\(^1\)\(^,\)\(^8\) The reason for this variation in reported prevalence can be explained by difficulty in distinguishing PTC-TCV from other histopathological variants, thus PTC-TCV is believed to be underdiagnosed in most of cases.\(^9\) In our cohort, women accounted for (73.8%) with the female to male ratio 3:1 is also in agreement with reported literature.\(^10\) Similarly, mean age of 52.3 years in our cohort study, is also in agreement with reported data.\(^11\) Further more, the multifocality (42.9%), pathological positive lymph nodes (66.7%) and distant metastasis (7.2%) at the time of diagnosis in our cohort were found to be similar to those reported in various studies.\(^12\) However, in disagreement with other studies, the percentage of ETE was comparatively low in our cohort (45.3% versus reported 63.9%). Mean primary tumor size was comparatively larger in our cohort (3.6 cm versus reported approximately 2.5-3 cm).\(^1\)\(^,\)\(^3\) Michels J et al\(^13\) also reported larger primary tumor size in their study of 56 patients with TCV. Due to these aggressive biological behavior, all patients in our cohort underwent total thyroidectomy. Although neck dissection (central and lateral) was performed in most of patients (78.6%), we recommend routine prophylactic central lymph node dissection at the time of thyroidectomy in PTC-TVC patients.\(^1\)\(^,\)\(^3\) Low occurrence of LRRs and distant metastasis in group, which received adjuvant RAI as part of management in our cohort, supports the notion that adjuvant RAI should be considered for all PTC-TVC patients, regardless of tumor size and lymph node status.\(^6\)\(^,\)\(^14\) In disagreement with similar reported data, PTC-TCV in our cohort had lower 5-year DFS rate (69.2%) after extensive treatment, which can be explained by our high risk cohort itself.\(^15\) In agreement with other related data, we found PTC-TVC as an important negative prognosticator.\(^16\)

There were limitations in our study; A) small sample size, B) BRAF mutations were not analysed in our cohort, as current evidence suggests that PTC-TVC is associated with 80-100% BRAF mutations as compared with 63-83% in classic PTC,\(^17\) and C) rearranged during transcription/PTC (RET/PTC) gene rearrangements were also not examined. Among 13 different RET/PTC oncogenes, RET/PTC1 and RET/PTC3 represent the most common types.\(^18\) Recent data\(^19\) has shown a striking correlation between the RET/PTC3 rearrangement and the PTC-TCV phenotype (35.8% positive), which suggests that molecular pathogenesis in PTC-TCV may be different, which needs further investigations.

In conclusion, PTC-TVC is associated with aggressive biological behavior. Our results were found in agreement with previous studies. However, more molecular studies are warranted to address the role of BRAF mutations, and Rearranged during Transcription/PTC (RET/PTC) gene rearrangements in PTC-TVC.

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