

Association of PD-1 and PD-L1 expression with clinicopathological variables in Papillary Thyroid Cancer

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Introduction

- 10-30% of patients with Papillary thyroid cancer (PTC) → recurrence post total thyroidectomy (TT) → eventually become de-differentiated¹
- Programmed cell death protein (PD-1) and its ligand (PD-L1) interaction → important oncological role
- Aim of the study: Identify the association of PD-1 & PD-L1 with clinicopathological variables in PTC

Methodology

- 29 patients who underwent TT for PTC between April 2016 to March 2017
- PD-1 and PD-L1 immunohistochemistry on surgical specimens
- Expression on Tumor cells and Tumor Infiltrating Lymphocytes (TILs): analysed and correlated with clinico-pathological behaviour

Results

- In tumor cells, PD-1 expression: negative in all specimens

Pathological variables	PD-L1 in tumor cells	PD-1 in TILs	PD-L1 in TILs
All samples	62% positive → 21% strong positive	48%: >1%; 17.2%: 26-40%	24% positive
Classical variant	61.6% moderate/strong positive	69%: >1%; 31%: 26-40%	30.8% positive
High risk variants	Negative	<1%	Negative
ETE	60% positive	60%: >1%	80% negative
PNE	100% positive	50%: >1%	50% negative
LVI	100% positive	25%: >1%	75% negative

Table 1. Clinopathological variables and PD-1 and PD-L1 IHCs

High risk variants: Diffuse sclerosis, tall cell, hobnail, cribriform

ETE- Extrathyroidal extension LVI: Lymphovascular invasion
PNE: Perinodal extension **Bold:** p<0.05

- No association between the strength of PD-L1 expression and ETE

Discussion/ Conclusion

- Various studies ^{2,3} have described the association of these immune markers in aggressive tumors like ATC, PDTC etc.
- Role of PD-1/ PD-L1 still unclear in Papillary thyroid cancer prognosis.³
- PD-L1 expression in tumor cells and TILs is associated with classical variant of Papillary thyroid cancer.
- PD-L1 expression in tumor cells is also associated with aggressive features like Extra thyroidal extension, Perinodal extension and Lymphovascular invasion.
- Studies with larger sample size and correlation with clinical follow up of patients is warranted.

References

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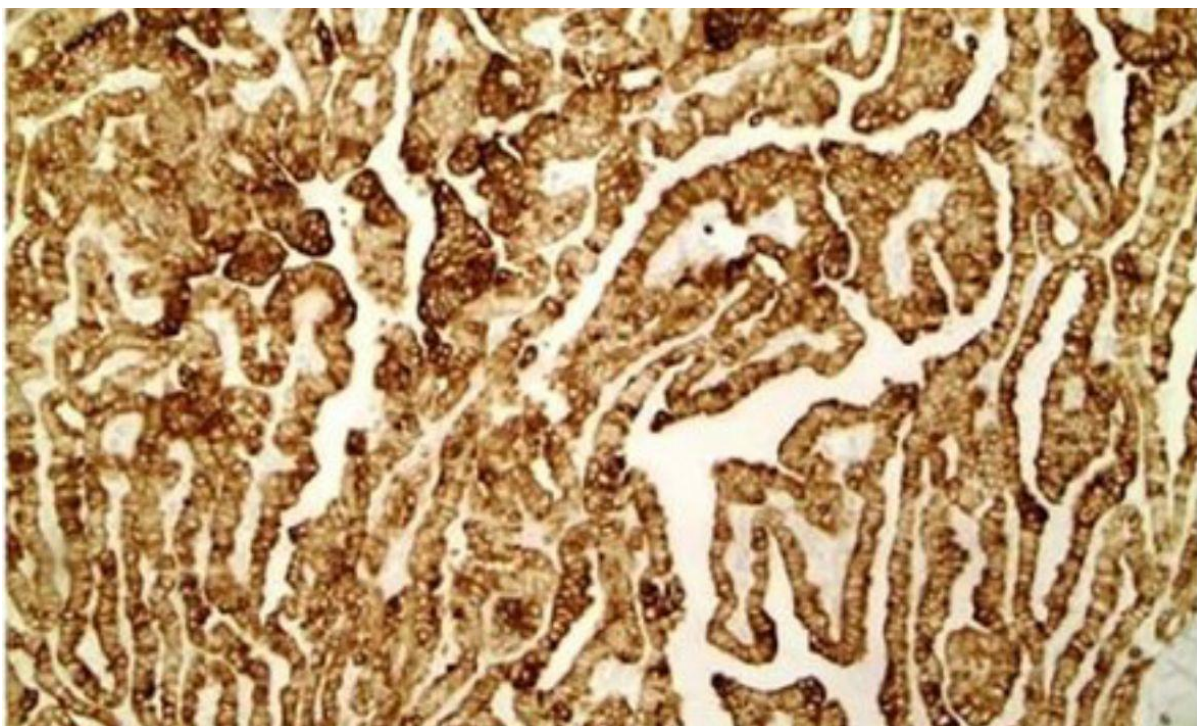


Fig 1. PD-L1 diffuse positive in tumor cells

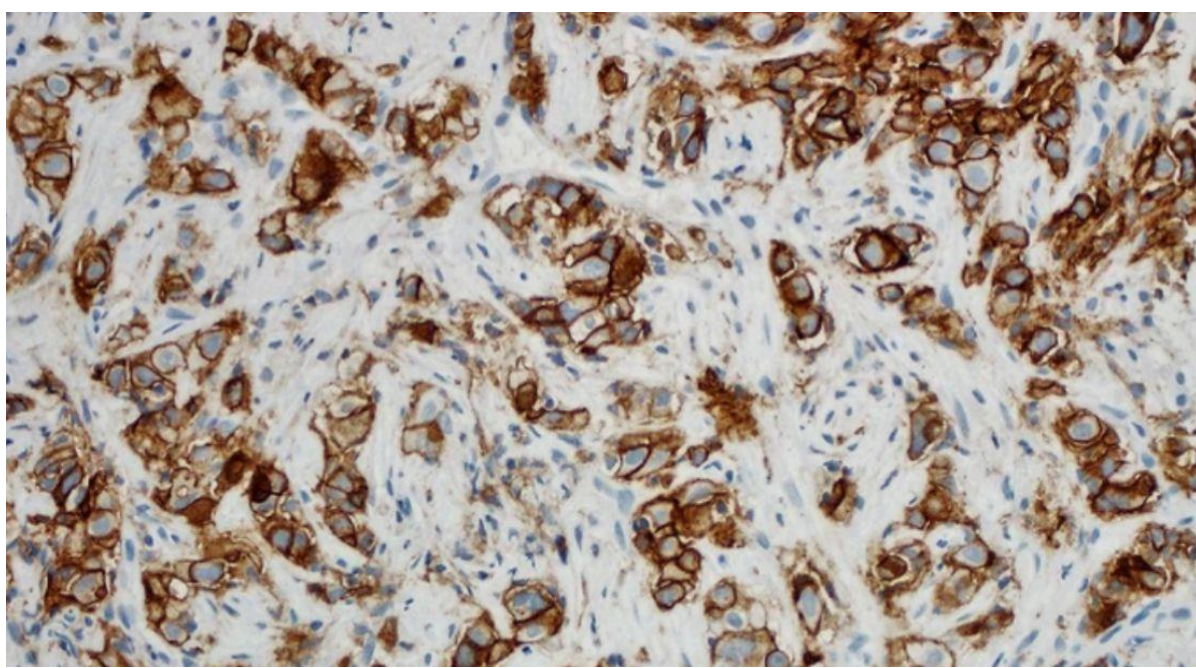


Fig 2. Positive PD-L1 in diffuse sclerosis variant

Results

- Mean age: 36.45±11.46 years
- Age and sex distribution of the study population is depicted in Figure 1

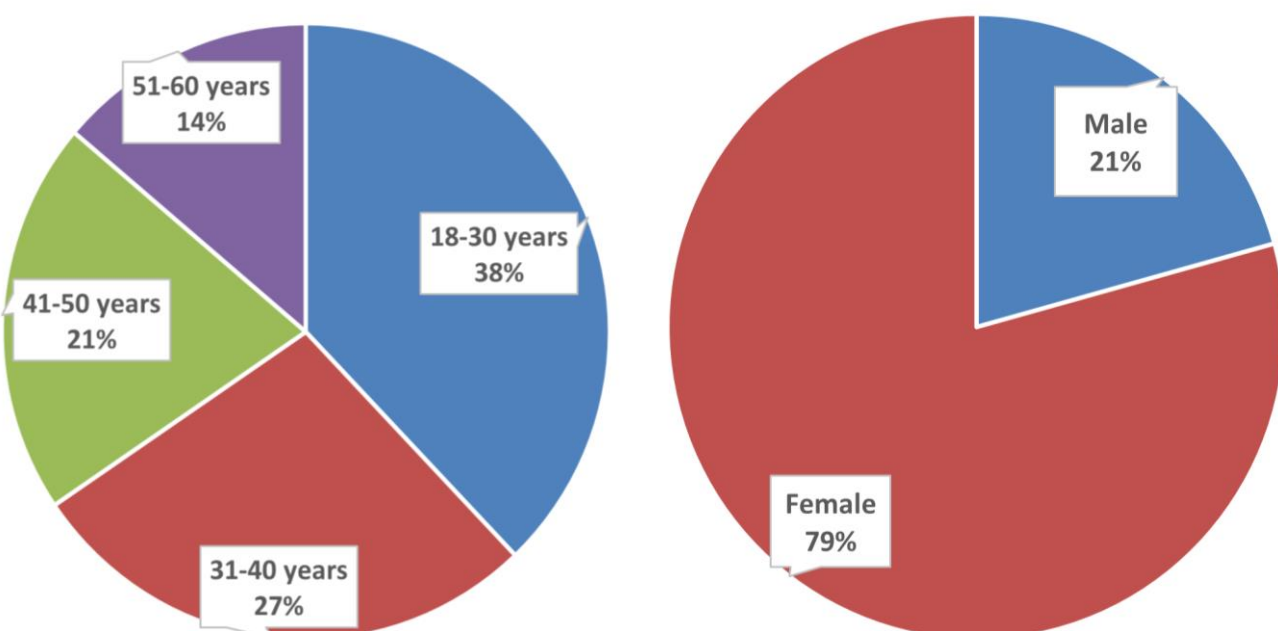


Fig 3. Age and sex distribution of the study population