

# FALSE NEGATIVES AND REASONS: ERZURUM RESEARCH AND TRAINING HOSPITAL PATHOLOGY DEPARTMENT EXPERIENCE

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False negatives (cytology as benign, histologically as malignant) reduces the sensitivity of the FNA. False-negative cases indicate malignant lesions undetectable by FNAB.

Fine needle aspiration biopsies (FNAB), adequate for examination, reported as benign re-evaluated with histopathological samples. 61 patients, 52 female and 9 male re-evaluated. 41 of them reported as benign. 20 of the patients, 7 female and 3 male reported as papillar carcinoma. 6 of the papillar carcinomas were single focus, 2 multicentric and 12 papillar microcarcinoma. (Table 1)

Benign	41
Papillar (single focus)	6
Papillar (multi centric)	2
Papillar (microcarcinom)	12

**Table 1:** Re-evaluation of false negative FNAB.

6 of the 8 patients malignant lobe was different from the lobe that FNA was made. Multicentric malignancy was detected in 2 cases.

Malignant FNA diagnosis forms 4-8% of all thyroid FNAs, as in literature. Most of them are papillary thyroid carcinoma. In fact papillary thyroid carcinoma cytopathological features are quite obvious, at the level of diagnostic. Patients diagnosed papillar thyroid carcinoma (PTC) with FNAB showed PTC at the rate of 96-100% with histopathological samples. However, false negative value was 32.78% in our patients under observation. This value is over the false-negative rate ranging between 1-21.3% stated in the literature. However, the results of the histopathological report with the results of cytopathological reports examined in detail the majority of the cases with false-negative study (60%) constitute papillary microcarcinoma. This finding suggests that papillar thyroid carcinoma is overlooked with FNAB.

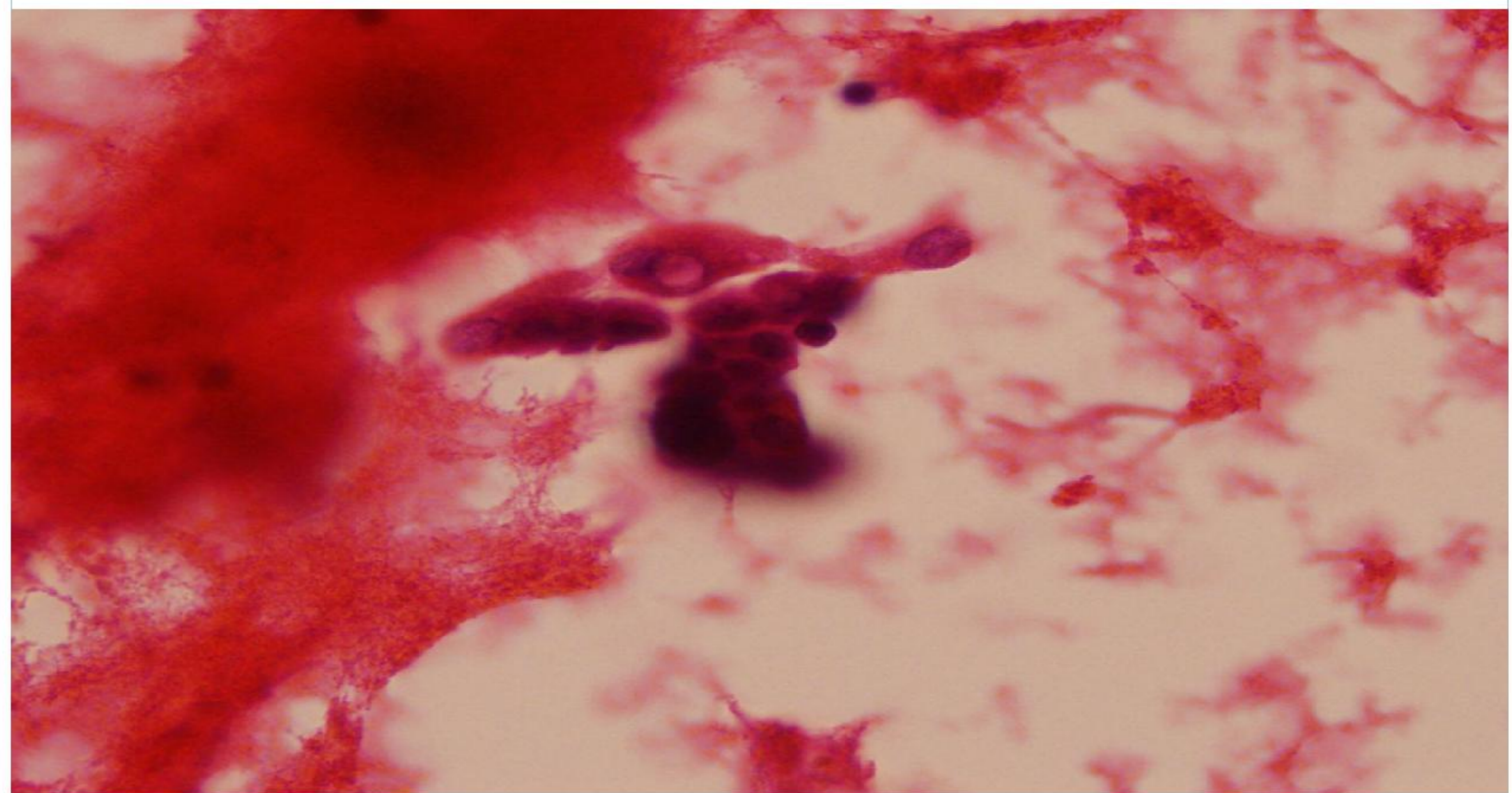
Our false-negative rates are compatible with the literature as PTC excluded. Another reason for our false negative value is the patcy distribution of PTC in the same nodule.

In the study of Amrikach et al. from the 64 benign cytologically benign diagnosed cases, 11 were false negative. 9 of them were histologically papillary carcinoma, 1 with Hurthle cell carcinoma, and 1 follicular carcinoma. 7 of the spreads were re-evaluated and in the 4 of them plenty of benign colloid follicles were monitored. As FNA was performed without ultrasound, false negatives were due to sampling error. When 3 of them re-evaluated its thought to be accurate to diagnosed them as non-diagnostic.

In the study of Chang et al between 129 papillary carcinoma patients 19 of them were diagnosed as benign cytologically. The reason of false negativness is due to smaller tumor size.

In the study of Chow et al there were 3 false negative results. One of them was diagnosed as papillar carcinoma, 1 of them was microcarcinoma in hyperplastic nodule and 1 of them was diagnosed as meduller carcinoma.

As a result, FNAB accompanied with ultrasound, should be applied to all solid, hypoechoic nodules and nodules with microcalcification. Also, because of the presence of PTC showing a patchy distribution in a single nodule, multiple biopsies should be taken from more than one ares, from the large nodules seen sonographically suspicious. False negative results of FNA depends on sample error or pathologist's review error.



**Picture:** First FNA examination malignancy negative patients macroscopic specimen made in spreading imprint, micropapillary intranuclear inclusion with focus on microcarcinoma structure

