



# WHAT'S NEW IN PATHOLOGY?

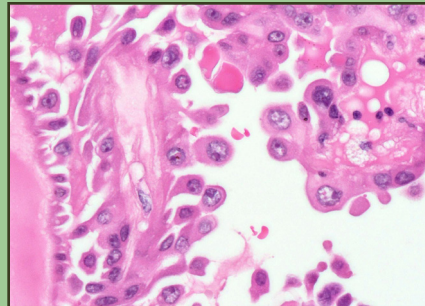
## THE LATEST NEWS IN THYROID

By Andrey Bychkov, M.D., Ph.D

### WHO Classification of Tumours of Endocrine Organs, 4<sup>th</sup> Edition (2017)

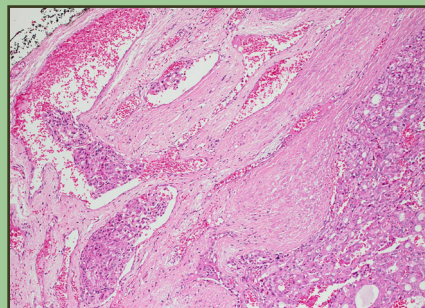
- New entities, expanded genetic profiles (including papillary vs. follicular patterned tumors), updated diagnostic criteria.
- A group of **borderline thyroid tumors** was introduced: **NIFTP** (noninvasive follicular thyroid neoplasm with papillary-like nuclear features), **FT-UMP** (follicular tumor of uncertain malignant potential) and **WDT-UMP** (well differentiated tumor of uncertain malignant potential).
- Borderline tumors are equivalent to carcinoma in situ in other organs; they are placed between follicular adenoma and follicular carcinoma or follicular variant of papillary carcinoma.
- The **hobnail variant** is the latest variant of papillary thyroid carcinoma. It is clinically aggressive, has a

micropapillary growth pattern and its cells have a hobnail appearance (Figure 1).



**Figure 1:** Papillary thyroid carcinoma, hobnail variant has hobnail cells with apically placed nuclei and bulging of the apical surface

- Follicular thyroid carcinomas are divided into **minimally invasive** (capsular invasion only), **angioinvasive** (grossly encapsulated with vascular invasion) and **widely / grossly invasive**.
- Hürthle cell (oncocyctic) tumors are reintroduced as a separate entity. These tumors include **Hürthle cell adenoma and carcinoma**; they previously were classified as oncocyctic variants of follicular adenoma and carcinoma (Figure 2).



**Figure 2:** Angioinvasive Hürthle cell carcinoma

• **Turin criteria** were adopted for the diagnosis of poorly differentiated thyroid carcinoma:

- (a) Presence of a solid/trabecular/insular growth pattern,
- (b) Absence of the conventional nuclear features of papillary carcinoma and
- (c) Presence of at least one of the following: convoluted nuclei,  $\geq 3$  mitoses per 10 high powered fields, tumor necrosis ([Am J Surg Pathol 2007;31:1256](#)).

• Micromedullary carcinoma vs. nodular C cell hyperplasia: suspect invasion if C cell proliferation plus stromal desmoplasia; collagen type IV IHC recommended to identify invasion of basement membrane.

• Reference: [Am J Surg Pathol Rev Rep 2017;22:209](#).

### NIFTP updates

• Additional exclusion criteria are:

1. **No true papillae** (no more 1% cutoff).
2. **No *BRAFV600E* or *TERT* promoter mutations**
3. No distant metastasis.

• References: [Hum Pathol 2018;74:1](#), [Pathol Int 2018;68:327](#).

• **NIFTP is not staged** by AJCC; only size, location and margin status should be reported.

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- NIFTP is accommodated in the new edition of the Bethesda system (see below).
- NIFTP is relatively common in Western practice (15%) but exceedingly rare in Asia (1%).
- Reference: [Endocr Pathol 2018 Feb 23 \[Epub ahead of print\]](#).

## Thyroid cancer staging, AJCC, 8<sup>th</sup> Edition (2017)

### I. Differentiated thyroid cancer:

- Age cutoff for staging was increased from 45 to 55 years at diagnosis.
- **Minimal extrathyroidal extension** detected only on histologic examination was removed from the definition of pT3 disease and therefore has **no impact** on either pT category or overall stage.
- pT3 has two new subcategories: **pT3a for tumors > 4 cm confined to the thyroid gland** and **pT3b for tumors of any size demonstrating gross extrathyroidal extension into strap muscles**.
- N1 disease no longer upstages a patient to stage III; if the patient's age is < 55 years at diagnosis, N1 disease is stage I; if age is ≥ 55 years, N1 disease is stage II.
- **Identification of a psammoma body in a cervical lymph node meets the definition of pN1 disease**, whether or not malignant cells are present.
- **Microscopically positive margins (R1) have no prognostic significance** (equal to negative margins, Ro); only grossly positive margins (R2) carry higher

risks of recurrence and disease specific mortality.

### II. Anaplastic cancers:

- Unlike previous editions in which all anaplastic thyroid cancers were classified as pT4, anaplastic cancers will **now use the same pT definitions as differentiated thyroid cancer**.

### III. Medullary thyroid cancer:

- Has its own chapter, but **most staging parameters are the same as differentiated / anaplastic thyroid carcinoma**.
- Staging is age independent.
- References: [Amin: AJCC Cancer Staging Manual, 8<sup>th</sup> Edition, 2017, CA Cancer J Clin 2018;68:55, Am J Surg Pathol Rev Rep 2018;23:145](#).

## Bethesda System for Reporting Thyroid Cytopathology, 2<sup>nd</sup> Edition (2018)

*\* Only minor updates: 6 diagnostic categories remain the same.*

- Risks of malignancy (ROM) recalculated based on post-2010 data.
- ROM is based on when NIFTP is not considered a malignancy and when NIFTP is still considered a carcinoma.
- The “usual management” of atypia (or follicular lesion) of undetermined significance (AUS/FLUS) and follicular nodules / suspicious for follicular nodules (FN/SFN) now has the option of molecular testing.
- Diagnostic criteria for category IV (FN/SFN) are revised in light of NIFTP by including cases with mild

nuclear papillary thyroid carcinoma (PTC)-like changes.

- Diagnostic criteria for PTC subset of the malignant category is limited to cases with “classical” features of PTC.
- Optional notes may be used to acknowledge NIFTP for subsets of categories IV-VI with cytologic features suggestive of follicular variant of PTC/NIFTP.
- Reference: [Ali: The Bethesda System for Reporting Thyroid Cytopathology, 2<sup>nd</sup> Edition, 2018](#).

### Meet the Author



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Dr. Bychkov is the thyroid pathology editor for PathologyOutlines.com. He has authored more than 40 journal articles and book chapters, serves as a peer reviewer for several journals, and is regularly invited to speak at various Asian pathology meetings. His current research interests are thyroid histopathology and digital pathology.