

Update No.5 – 2018 Dr Christophe Rosty **Tumour budding in CRC**

Tumour budding is an independent adverse prognostic factor in colorectal carcinoma. It is associated with higher TNM stages, lympho-vascular invasion, lymph node and distant metastasis. International consensus guidelines have been recently published (1).

Definition: A tumour bud is defined as a cluster of 1-4 tumour cells at the invasive front of CRC.

Histological assessment:

- Scan all sections of tumour to identify the area with highest degree of tumour budding at the invasive front ('hot spot').
- Count the number of tumour buds on H&E in the 'hot spot' field at 20x objective.
- Normalise (if necessary) to come up with tumour bud number for a 0.785mm² area (for microscopes used at Envoi, divide number of counted tumour buds by 1.2).

Quantification:

Tumour budding is reporting using a 3-tier system based on the normalised number of tumour buds

- 0 4 buds: Bd 1 low
- 5 9 buds: Bd 2
- 10 or more buds: Bd 3
- high

intermediate



Left: Invasive front of CRC at low magnification.

Right: Tumour buds at 20x magnification.

Clinical significance:		
•	Malignant polyps:	In malignant polyps resected endoscopically, Bd 2 and Bd 3

- Malignant polyps: In malignant polyps resected endoscopically, Bd 2 and Bd 3 are independently associated with *increased risk of lymph node metastasis* (2).
- Stage II CRCs (pT3/T4 N0):

Bd 3 is independently associated with *increased risk of recurrence and mortality* (3).

References

- 1. Lugli A, Kirsch R, Ajioka Y, et al. Recommendations for reporting tumor budding in colorectal cancer based on the International Tumor Budding Consensus Conference (ITBCC) 2016. *Mod Pathol* 2017;30:1299-1311.
- 2. Bosch SL, Teerenstra S, de Wilt JH, Cunningham C, Nagtegaal ID. Predicting lymph node metastasis in pT1 colorectal cancer: a systematic review of risk factors providing rationale for therapy decisions. *Endoscopy* 2013;45:827-834.
- 3. van Wyk HC, Park J, Roxburgh C, et al. The role of tumour budding in predicting survival in patients with primary operable colorectal cancer: a systematic review. *Cancer Treat Rev* 2015;41:151-159.