

Histological tumour grade (Core)

Reason/Evidentiary Support

The histologic (microscopic) grading of salivary gland carcinomas has been shown to be an independent predictor of behaviour and plays a role in optimizing therapy. Further, there is often a positive correlation between histologic grade and clinical stage.¹⁻⁴ However, as alluded to above, most salivary gland carcinoma types have an intrinsic biologic behaviour and attempted application of a universal grading scheme is not recommended.³ Thus by assigning a histologic type the tumour grade itself is often implied. Thus a generic grading scheme is no longer recommended for salivary gland carcinomas.⁵

Carcinoma types for which grading systems exist and are relevant are incorporated into histologic type. The major categories that are amenable to grading include adenoid cystic carcinoma, mucoepidermoid carcinoma, and adenocarcinoma, not otherwise specified.^{2,3,6} Additionally, with the new World Health Organization (WHO) classification, polymorphous adenocarcinoma is another tumour type that is to be graded,⁷ with the understanding that a validated grading scheme has not yet been established.

In adenoid cystic carcinoma histologic grading is based on growth pattern.⁶ Those adenoid cystic carcinomas showing 30% or greater of solid growth pattern are considered to be histologically high grade carcinomas. However, recent studies suggest that any solid component may still be of prognostic relevance.⁸ The histologic grading of mucoepidermoid carcinoma includes a combination of growth pattern characteristics (e.g. cystic, solid, neurotropism) and cytomorphologic findings (e.g. anaplasia, mitoses, necrosis).⁹⁻¹¹ Adenocarcinomas, not otherwise specified, do not have a formalized grading scheme and are graded intuitively based on cytomorphologic features.³ Similarly, as the concept of grading polymorphous adenocarcinomas will be a new one,⁷ as these also lack a formalized grading scheme. Currently, the recommendation is to grade these intuitively based on cytomorphologic features, acknowledging that the majority will be low grade.

High grade transformation has evolved into an important concept of tumour progression in salivary gland carcinomas. Historically designated as 'dedifferentiation', it describes progression of a typically monomorphic carcinoma into a pleomorphic high grade carcinoma.¹² The importance of this phenomenon is that tumours demonstrating high grade transformation show an aggressive clinical course that deviates drastically from the usual behaviour for a given tumour type, thus alerting to the potential need for more aggressive clinical management. Tumours for which this phenomenon is well characterized include acinic cell carcinoma, adenoid cystic carcinoma, and epithelial-myoepithelial carcinoma. Mammary analogue secretory carcinoma and polymorphous adenocarcinoma also rarely undergo high grade transformation.^{13,14}

References

- 1 Spiro RH, Thaler HT, Hicks WF, Kher UA, Huvos AH and Strong EW (1991). The importance of clinical staging of minor salivary gland carcinoma. *Am J Surg* 162(4):330-336.
- 2 Spiro RH, Huvos AG and Strong EW (1982). Adenocarcinoma of salivary origin. Clinicopathologic study of 204 patients. *Am J Surg* 144(4):423-431.
- 3 Seethala RR (2011). Histologic grading and prognostic biomarkers in salivary gland carcinomas. *Adv Anat Pathol* 18(1):29-45.
- 4 Kane WJ, McCaffrey TV, Olsen KD and Lewis JE (1991). Primary parotid malignancies. A clinical and pathologic review. *Arch Otolaryngol Head Neck Surg* 117(3):307-315.
- 5 Lydiatt WM, Mukherji SK, O'Sullivan B, Patel SG and Shah JP (2017). Major Salivary Glands. In: *AJCC Cancer Staging Manual 8th ed*, Amin MB et al (eds), Springer, New York.
- 6 Szanto PA, Luna MA, Tortoledo ME and White RA (1984). Histologic grading of adenoid cystic carcinoma of the salivary glands. *Cancer* 54(6):1062-1069.
- 7 Fonseca I, Assaad A, Katabi N, Seethala RR, Simpson RHW, Skalova A, Weinreb I and Wenig BM (2017). Polymorphous Adenocarcinoma. In: *WHO Classification of Tumours of the Head and Neck*, El-Naggar AK, Chan JK, Grandis JR, Ohgaki H and Slootweg P (eds), IARC, Lyon, France, 167-168.
- 8 Xu B, Drill E, Ho A, Ho A, Dunn L, Prieto-Granada CN, Chan T, Ganly I, Ghossein R and Katabi N (2017). Predictors of Outcome in Adenoid Cystic Carcinoma of Salivary Glands: A Clinicopathologic Study With Correlation Between MYB Fusion and Protein Expression. *Am J Surg Pathol* 41(10):1422-1432.
- 9 Seethala RR, Dacic S, Cieply K, Kelly LM and Nikiforova MN (2010). A reappraisal of the MECT1/MAML2 translocation in salivary mucoepidermoid carcinomas. *Am J Surg Pathol* 34(8):1106-1121.
- 10 Brandwein MS, Ivanov K, Wallace DI, Hille JJ, Wang B, Fahmy A, Bodian C, Urken ML, Gnepp DR, Huvos A, Lumerman H and Mills SE (2001). Mucoepidermoid carcinoma: a clinicopathologic study of 80 patients with special reference to histological grading. *Am J Surg Pathol* 25(7):835-845.
- 11 Auclair PL, Goode RK and Ellis GL (1992). Mucoepidermoid carcinoma of intraoral salivary glands. Evaluation and application of grading criteria in 143 cases. *Cancer* 69(8):2021-2030.

- 12 Costa AF, Altemani A and Hermsen M (2011). Current concepts on dedifferentiation/high-grade transformation in salivary gland tumors. *Patholog Res Int* 2011:325965.
- 13 Skalova A, Vanecek T, Majewska H, Laco J, Grossmann P, Simpson RH, Hauer L, Andrie P, Hosticka L, Branzovsky J and Michal M (2014). Mammary analogue secretory carcinoma of salivary glands with high-grade transformation: report of 3 cases with the ETV6-NTRK3 gene fusion and analysis of TP53, beta-catenin, EGFR, and CCND1 genes. *Am J Surg Pathol* 38(1):23-33.
- 14 Simpson RH, Pereira EM, Ribeiro AC, Abdulkadir A and Reis-Filho JS (2002). Polymorphous low-grade adenocarcinoma of the salivary glands with transformation to high-grade carcinoma. *Histopathology* 41(3):250-259.