tumourigenesis of such lesions. In the present case, no manifestations of immunodeficiency were observed. Furthermore, HHV-8 was only detectable in KS cells. Therefore, the concurrent occurrence of the two neoplastic components seemed more likely coincidental rather than aetiologically related. Clinically, chemotherapy with radiotherapy seems a promising way for the treatment of such cases, and the ABVD regimen looks like a proper choice.

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**CASE REPORT**

A 37-year-old man presented with a painless circumscribed nodule that had been gradually enlarging in the left parotid gland area for six years. No cervical lymphadenopathy was detected. A subtotal parotidectomy was performed and a tumour was excised. No tumour recurrence was observed 18 months after the surgery.

Grossly, the tumour was completely circumscribed within the parotid gland, measuring 20×18×15 mm in size. The cut surface revealed multilocular cystic spaces. No bleeding or necrosis was detected in the tumour.

Histologically, the tumour consisted mainly of multiple cystic formations lined with stratified squamous epithelium lacking a granular layer, but containing keratin material and calcification (figure 1). There was no evidence of cytological atypia or abnormal mitotic activity in the epithelium. No tubules, duct-like glandular structures, or sebaceous differentiation were observed. No evidence of skin appendages was found beneath the epithelium. The stroma of the tumour was predominantly a dense

**Figure 1** H&E stained sections showing the keratocystoma morphology. (A) Characteristic tumour view consisting of variably sized and shaped cystic lesions filled with lamellar keratin material and calcification (×20). (B) The stroma of the tumour showing predominantly a dense lymphoid element (×100). (C) The stratified squamous epithelium lacking a granular layer filled with lamellar keratin material (×200). (D) Solid squamous cell islands exhibiting uniform, bland nuclei and abundant eosinophilic cytoplasm (×400).
lymphoid element. The tumour cells stained negatively with periodic acid-Schiff (PAS) and Alcian blue.

By immunohistochemistry, the tumour cells were diffusely positive for cytokeratins AE1/AE3 and 5/6, focally positive for cytokeratin 7, and stained strongly for p63, but weakly for p53 (figure 2). The basal cells of the stratified epithelium were partially positive for cytokeratin 19. The tumour cells were completely negative for cytokeratins 8/18 and 20. Less than 1% of all tumour cells were Ki-67 positive.

DISCUSSION
Keratocystoma is a rare benign tumour of salivary gland origin whose nomenclature remains to be confirmed. To the best of our knowledge, only four cases, including the present case, have been reported in the English literature.1-4 This type of tumour needs to be differentially diagnosed with several tumours of the parotid gland containing squamous cells, including primary and metastatic squamous cell carcinomas, mucoepidermoid carcinoma, squamous metaplasia in certain benign neoplasias and conditions, and cysts.5 In this case there is no evidence of cytological atypia or abnormal mitotic activity in the squamous cell, which is different from squamous cell carcinoma. The absence of incorporated mucous cells by the mucicarmine stain (Alcian blue) and the presence of marked keratinisation are important for separating this tumour from mucoepidermoid carcinoma. The histology of the tumour is different from Warthin’s tumour, which presents with cystic spaces surrounded by two uniform rows of cells with centrally placed pyknotic nuclei. The first banchial clef cyst can involve the parotid gland. The cystic spaces are usually lined with squamous epithelium with keratin material filling the lumen, but pseudo-stratifed columnar epithelium can also be seen. Sebaceous lymphadenoma and non-sebaceous lymphadenoma of the parotid gland need to be excluded. Overexpression of p63 and p53 in the tumour support neoplastic proliferation. We believe that this type of tumour is a true cystic neoplasm, not a christoma or trichoadenoma, based on its morphological and immunohistochemical findings.

**Take-home messages**

- A rare case of a benign tumour of the parotid gland, presenting as a keratocystoma, is described.
- It is believed that only four cases, including the present case, have been reported in the English literature.
- Histologically, the tumour consisted mainly of variably sized and shaped cystic lesions filled with lamellar keratin material and calcification.
- By immunohistochemistry, the tumour cells were positive for cytokeratins, p63 and p53.

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**Figure 2** Immunohistochemical findings in the keratocystoma specimen. The tumour cells strongly stained for AE1/AE3 (A), CK5/6 (B), and p63 (C), or weakly stained for p53 (D) in the keratocystoma case. Inset images show positive cells in the region. All sections are counterstained with haematoxylin (×200).
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BOOK REVIEW

Biopsy interpretation of the thyroid

Edited by Scott L Boerner, Sylvia L Asa. Published by Lippincott Williams and Wilkins, Philadelphia, Pennsylvania, USA, 2009, pp 288, hardback, £100, ISBN: 978-0781772044.

This represents another offering from the excellent Biopsy Interpretation series that covers most organs.

This is a superb book organised into 13 chapters spanning the normal thyroid gland and covering various morphological/pattern-based entities that the surgical pathologist is likely to encounter. The index is functional and user friendly. The integration of fine needle aspiration cytology with histological interpretation is a particular strength; one that has been interwove cleanly and unequivocally. The comments and diagnostic criteria are well set out and leave the reader in no doubt how to interpret what they may encounter in histological sections and as well as with cytological preparations.

I used it as a bench book beside my microscope and was thrilled to see as much as three illustrations of the rather uncommon lymphoepithelial cyst of the thyroid!

The illustrations are magnificent throughout and for this the authors should be complimented. The book illustrations are supplemented by electronic images.

In summary, this book blends fine needle aspiration cytology findings with the histological appearances in a cogent, clear and authoritative fashion. I would thoroughly recommend every pathologist and trainee who encounters thyroid pathology to read this book, and have it within easy access of your microscope.

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