

## Prostate Cancer - Double Vision but Solitary Lesion

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### Abstract

A 53 year old man with a background of castrate-sensitive prostate cancer on intermittent androgen deprivation therapy (ADT) presented with right sixth nerve palsy secondary to a solitary right petroclival lesion involving adjacent dura and bone. The clinical and imaging characteristics of the lesion were consistent with a number of differential diagnoses (including metastatic prostate cancer, meningioma and chondrosarcoma). The patient initially declined biopsy and ADT was recommenced but the lesion continued to enlarge despite an excellent biochemical response. A subsequent biopsy of the petroclival mass demonstrated a WHO grade I meningioma and the patient proceeded to have definitive stereotactic radiotherapy. This case illustrates an unusual solitary skull base lesion in a man with prostate cancer. Whilst bony metastases, usually multiple, in the skeleton are common, solitary skull or brain lesions should be investigated as alternate diagnoses are likely in such circumstance.

**Keywords:** Petroclival lesion; Prostate cancer; Meningioma; Metastasis

**List of Abbreviations:** ADT-Androgen deprivation therapy; CT-Computed tomography; FDG-avid PET-Fludeoxyglucose-avid positron emission tomography; PSA-Prostate specific antigen

### Introduction

Prostate cancer is the most common cancer in Australian men and the third most common cause of cancer deaths, comprising 13% of all cancer deaths in Australian men [1]. One in five Australian men will develop prostate cancer by age 85 with 1 in 25 risk of dying from prostate cancer [1]. However, many cases are indolent and patients often die from competing risks. Whilst the risk of mortality is only 3% in low risk patients, it increases in intermediate risk (7%) and high risk disease (18%) [2].

Approximately 90% of patients with metastatic prostate cancer develop bony metastases, which are characteristically multiple osteoblastic lesions to the axial skeleton [3]. Bone metastases to the skull (8%) and skull base (1.7%) are much less frequent [3,4]. Similarly, patients with parenchymal central nervous system disease are also rare, being reported in less than 1% of patients [5], although the prognosis is usually very poor in such cases [5,6]. Where these cases are reported, most (79%) involve the brain parenchyma itself although some (21%) involve dura only [5]. Importantly, the finding of solitary metastasis to the central nervous system from prostate cancer is exceedingly rare [5].

Here, we report a patient with castrate-sensitive prostate cancer in biochemical remission who presented with progressive right sixth nerve palsy secondary to a solitary right petroclival lesion involving adjacent dura and bone.

### Case Report

A 53-year-old man was referred by his primary physician for the management of an asymptomatic screen-detected prostate cancer, with a prostate specific antigen (PSA) level of 22.5 (institutional reference range 0.0 – 3.5 µg/L). He subsequently underwent a radical prostatectomy and lymph node dissection in March 2010 which revealed a T3bN1 carcinoma (Gleason 4+5) with lymphovascular and perineural invasion, positive margins and extra-capsular extension in one of two involved lymph nodes. Post-

operatively, his PSA nadir level was 4 µg/L but without evidence of macroscopic metastases. He was commenced on ADT with goserelin acetate for 6 months and achieved biochemical complete remission (PSA<0.01) in March 2011. However, treatment was poorly tolerated and an intermittent schedule was adopted.

In July 2011, the patient began experiencing mild diplopia on right gaze which progressed over several months. Examination findings were consistent with mild right-sided sixth nerve palsy. Investigations showed a 10 mm x 3 mm extra-axial, durally-based, homogeneously enhancing mass arising from the petroclival ligament. There was evidence of bony invasion with lysis but no sclerosis (Figure 1a & 1b). Concurrently, his PSA rose from 0.01 µg/L to 0.04 µg/L over 5 months but importantly, re-staging CT and bone scan showed no evidence of other visceral or bony disease. The lesion was only minimally FDG-avid on PET. The patient was offered a biopsy to differentiate between the differential diagnoses of a solitary prostate cancer metastasis or, more likely, an unrelated pathology such as meningioma or chondrosarcoma. The patient declined a biopsy and requested recommencement of ADT for presumed petroclival metastasis from prostate cancer.

With ADT, the patient's PSA fell steadily over the next 6 months but unfortunately, his diplopia worsened and the lesion continued to enlarge (Figure 1c & 1d). He also developed subtle signs of right trigeminal nerve involvement, manifested by reduced corneal sensation. The patient agreed to a biopsy and limited decompression of the lesion. Histopathology demonstrated a grade 1 meningioma with an elevated ki67 index, without evidence of carcinoma (Figure 1e & 1f). The patient underwent definitive stereotactic radiotherapy 50 Gy in 20 fractions prescribed to 90% isodose (5 fractions per week over 4 weeks), with stabilisation of his neurological deficits

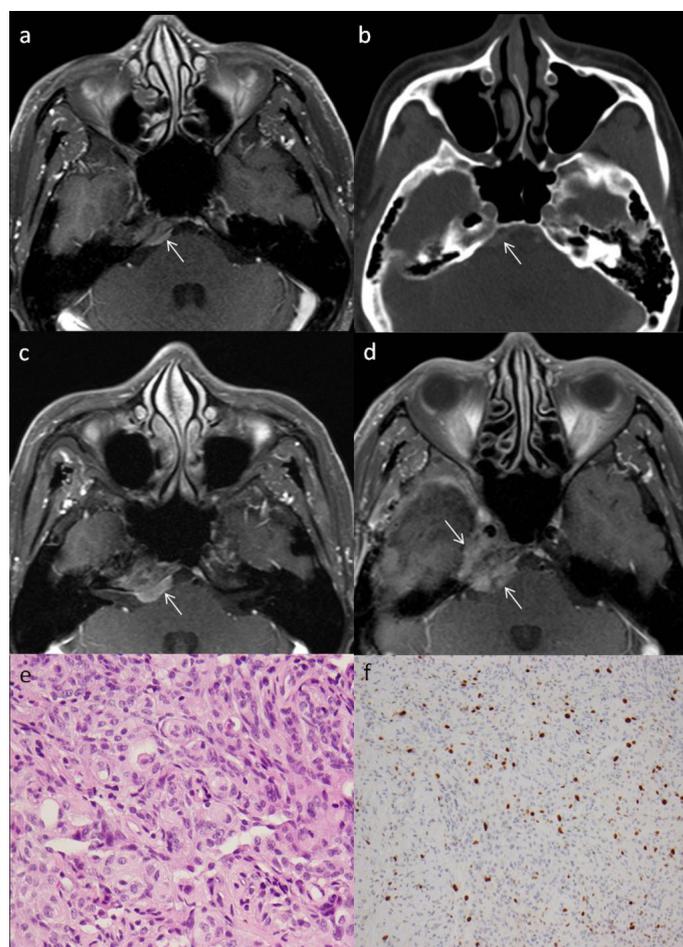


Figure 1: (a) Axial post contrast T1 MR. (b) Bone window CT images demonstrate a durally-based, homogeneously enhancing petroclival mass (arrow) with associated bone lysis when compared to the bone density on the left. (c) Corresponding MRI 10 months and (d) 18 months later with increase in lesion size and progressive bone invasion. (e) High power images (400x) showing characteristic morphological features of meningioma with syncytia, nests and whorls of meningothelial cells. (f) A ki67 stain of the lesion showing moderately elevated ki67 proliferative index.

## Discussion

This case highlights the diagnostic challenge of a solitary base of skull lesion in a patient with prostate cancer presenting with a sixth nerve palsy. Although cases of diplopia from cranial nerve palsies [6,7] or ocular rectus muscle invasion [8] from metastatic prostate cancer have been reported, these are relatively rare. Whilst it would have been tempting to make a presumptive diagnosis

of metastatic prostate cancer, a number of features in this case raised the possibility for an alternate diagnosis: the general rarity of solitary metastatic lesions in the brain or skull; the relatively low absolute levels of PSA; and the radiological appearance of the osteolytic focus in the petroclival region. It should be noted that a negative PSA does not exclude the presence of brain metastasis [9,10]. However, a diagnostic biopsy was initially declined by the patient on several occasions. The most common causes of clival lesions are primary tumours such as chordomas or chondrosarcomas (83%) with the rest being meningiomas and metastatic disease from sites such as the lung, prostate, liver or skin (melanoma) [11].

Our patient was ultimately diagnosed and treated for a meningioma. The diagnosis of meningioma was considered in his initial differential diagnosis but the rarity of skull base meningiomas (2.8%) [12] and the rapid rate of change argued for a diagnosis of metastatic disease, given that most meningiomas tend to remain asymptomatic and are slow growing (one series showed 35 of 57 patients displayed no growth over a 29-month follow-up period) [13,14]. There is also sparse literature showing that metastatic prostate cancer can simulate meningiomas [15-17]. One intriguing possibility is the hypothesis that the unusual behaviour of the meningioma in this case may be the result of using luteinising hormone releasing hormone (LHRH) agonists as there is emerging data that some meningiomas may be regulated by the LHRH receptor expression [18-20].

## Conclusion

This is a rare case of a concomitant atypical meningioma, possibly further stimulated by the use of ADT for prostate cancer treatment. Solitary lesions in the brain or skull of patients with a history of prostate cancer should be aggressively investigated given their rarity and the range of alternate diagnoses which can present similarly. The prognosis and management of many of these alternate diagnoses also differ substantially from those of metastatic prostate cancer and a high index of suspicion would aid an earlier diagnosis.

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