



Brafv600e and Ctnb1 Mutational Study in Rathke's Cleft Cysts

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Abstract

Aim: Rathke's cleft cysts and craniopharyngiomas typically involve sellar region and their histogenetic relationship is still matter of debate. Clinical and histopathologic differentiation of cystic lesions from the sellar region, that is, craniopharyngiomas (CPs) and Rathke cleft cysts (RCCs), is challenging and has great importance with respect to variable clinical manifestation and adapted surgical treatment strategies in both entities. The recent acquisition that adamantinomatous and papillary craniopharyngiomas bear distinct molecular alterations i.e., β -catenin (CTNNB1) and BRAFV600 mutations respectively, has suggest to screen for such alteration a series of Rathke cyst to seek a possible relation with one of the two craniopharyngioma type.

Methods: Seven Rathke's cleft cysts were analyzed for BRAF and CTNNB1 mutational status by sequencing and immunohistochemistry. Radiological, clinical and histological features were performed.

Results: None of the 7 Rathke's cleft cysts harbor BRAFV600E mutation. No CTNNB1 mutation was found. Radiological, clinical and histological re-evaluation of the cases confirmed the diagnosis of Rathke's cleft cysts.

Conclusion: BRAFV600E and CTNNB1 mutations appeared, as most reliable factor for the differentiation between purely cystic CPs and RCCs, whereas tumor location, tumor size, and radiological parameter of the tumor were less consistent parameters. This study again confirms that craniopharyngiomas (CPs) and Rathke cleft cysts (RCCs), are associated with distinct pathogenic pathways.

Keywords: BRAFV600E mutation; Rathke's cleft cysts; Craniopharyngioma; Sequencing

Introduction

Rathke's cleft cyst (RCC) and craniopharyngiomas (CP) involve the sellar region and sometimes histological evaluation is challenging especially when small tissue specimens are available for analysis. Both lesions are considered to arise from the remnants of Rathke's diverticulum [1].

Some authors have suggested that these lesions are part of a disease spectrum extending from RCC to craniopharyngioma [2]. Intracystic hemorrhage or inflammation may induce metaplasia in the single cuboidal epithelium of RCC to form stratified squamous epithelia resulting in the induction of craniopharyngioma [3]. Besides the presence of ciliated and goblet cell in some craniopharyngiomas are believed to provide evidence for this [4,5]. Moreover most of the CPs are larger than 20 mm, whereas most of RCCs are smaller than 20 mm in diameter. However, these parameters are insufficient to rule out small cystic CP variants, which do not exhibit any other difference in computed tomography (CT), or magnetic resonance imaging (MRI) [6].

Although CPs and RCCs are histologically benign, they exhibit a different clinical behavior and thus require different surgical treatment strategies. The rates of recurrence and progression, respectively and the interval of progression-free survival in CPs is obviously depending on the extent of surgical removal [7]. The recurrence rate is significantly lower in patients with RCC even after subtotal or partial removal [7]. The outcome is much better and one might conclude from previous data that radical excision is not necessary in these lesions. On the basis of these data distinction between the lesions is truly important. However, the small tissue specimens available for analysis and the similar histological features cause a tricky interpretation.

Previous work has shown that mutations in CTNNB1 with nuclear accumulation of β -catenin are a reliable marker for the identification of craniopharyngiomas of the adamantinomatous subtype [5f]. Whereas papillary craniopharyngiomas are characterized by BRAF V600E mutations as genetic hallmark and consistently don't show nuclear β -catenin expression [8].

On the basis of these data we investigated 7 Rathke's cleft cysts to determine the presence of BRAF V600E and beta-catenin mutations and its suitability as diagnostic marker for the differentiation of cystic lesions of the sellar region. For each case clinical, radiological and

histopathological parameters were also re-evaluated in order to exclude features typical for craniopharyngoma.

Materials and Methods

Patient selection and immunohistochemistry

Surgical specimens from 7 patients with Rathke's cleft cysts were selected retrospectively on the basis of the availability of formalin-fixed paraffin embedded material (FFPE) from two different institutions, the Mediterranean Neurologic Institute (Neuromed), Pozzilli, Italy, and the Pathology Service of University Sapienza, Rome, Italy.

All Rathke's cleft cysts were reviewed by two neuropathologist MA and FG and diagnosed according to the AFIP Atlas of Tumor Pathology (Tumors of the Central Nervous System, Series 4 Fascicle 7).

Primary antibody against BRAF p.Val600Glu (clone VE1, 1:100, Spring Bioscience, Pleasanton, CA): positive staining was characterized by diffuse and moderate cytoplasmic staining of the tumor cells.

Primary antibody against β -catenin antigen (BD Pharmingen, 610154, mouse monoclonal, clone 14): were scored as positive, and cases with membranous staining were scored as negative.

Isolation of DNA

DNA was extracted from FFPE tissue specimens with the QIAamp DNA Mini Kit, as described by the manufacturer (Qiagen S.A., Courtaboeuf France), DNA concentration was quantified using the Nanodrop ND-1000 UV-Vis spectrophotometer (Labtech France, Palaiseau, France). Final products were stored at -20°C until use.

Moreover standard diagnostic procedures were performed. Sections were stained with ematoxililn and eosin.

BRAFV600E and CTNNB1 mutation analysis

Mutational analysis was performed amplifying DNA with the primers as follows: BRAF exon 15, 5'-TCATAATGCTTGCTCTGATAGGA-3' (sense) and 5'-GGC CAAAATTTAATCAGTGGA-3' (antisense). The PCR products were purified using the automated system Biomek NXP by Beckman Coulter and Agentcourt AMPure XP reagents. Purified products were submitted to PCR cycle sequencing conditions as follow: denaturation at 95°C for 30 s, annealing at 50°C for 15 s, and extension at 60°C for 240 s. The cycle sequencing products were purified using the same automated system and Agentcourt Clean SEQ reagents. Sequencing analysis was performed using the ABI 3130 XL DNA analyser (Applied Biosystem).

For β -catenin, a 630 base pair genomic PCR fragment spanning codon 1 (exon 2) to codon 92 (exon 4) was amplified using primers primers BCATEX2F (GAAAATCCAGCGTGGACAATG) and BCATEX4R (TCGAGTCATTGCA- TACTGTCC).

Results

Clinical, histological and radiological findings

The series of 7 Rathke's cleft cysts comprised 7 females and 1 male, with a median age of 38. Cases were located within the sella turcica except for one located in the sphenoidal bone (Table 1).

Case	Age (years)/ Sex	RCC histology	BRAFV600 E mutation	β -catenin mutation
1	56 F	Ciliated cuboidal and columnar epithelium with squamous metaplasia	No	No
2*	16 F	Ciliated cuboidal and columnar epithelium	No	No
3	55 F	Ciliated cuboidal and columnar epithelium	No	No
4	33 F	Ciliated cuboidal and columnar epithelium	No	No
5	48 F	Ciliated cuboidal and columnar epithelium with squamous metaplasia	No	No
6	29 F	Ciliated cuboidal and columnar epithelium	No	No
7	32 M	Ciliated cuboidal and columnar epithelium	No	No

Table 1: RCC histology and mutations in different cases.

We further added two cases of craniopharyngiomas, one papillary and one adamantinomatous, in which the histological and radiological features were typical. The histological examination showed cuboidal or columnar ciliated epithelium on a fibrous tissue. In three of 7 Rathke cleft cysts (n.1 n.2 n.5) co-existence of squamous cell metaplasia and cilia with scattered goblet cells could be identified. The intrasphenoidal case (n.2) showed columnar ciliated epithelium with few mucinous cells (Figure 1).

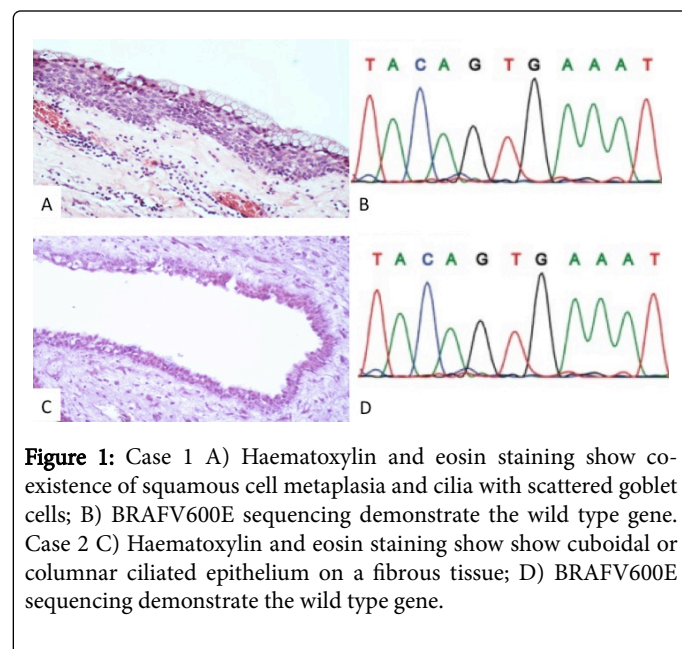


Figure 1: Case 1 A) Haematoxylin and eosin staining show co-existence of squamous cell metaplasia and cilia with scattered goblet cells; B) BRAFV600E sequencing demonstrate the wild type gene. Case 2 C) Haematoxylin and eosin staining show cuboidal or columnar ciliated epithelium on a fibrous tissue; D) BRAFV600E sequencing demonstrate the wild type gene.

Radiological re-evaluation in all seven cases demonstrated the typical MR finding of a Rathke's cleft cyst such as dumbbell-shaped cyst, with hypointense signal in T1-weighted images, and hyperintense signal in T2-weighted images, relative to normal brain parenchyma

and they were smaller than 20 mm in diameter. Moreover in three of 7 Rathke cleft cysts (n.1 n.2 n.5) and the intrasphenoidal lesion (n.2) MRI showed findings typical of a cystic lesion.

All reported cases achieve remission, except for a single patient that developed repeated re-accumulation of the mucus and become intractable (Figure 1).

BRAFV600E and β -catenin mutational status by immunohistochemistry

Immunohistochemical for BRAFV600E protein was found negative in all RCCs, as well the cases did not show nuclear expression for β -catenin. To confirm the quality of the immunohistochemistry, we found positive cells for CTNB1 and BRAFV600E in respectively one adamantinomatous craniopharyngioma and one papillary craniopharyngioma (Figure 2).

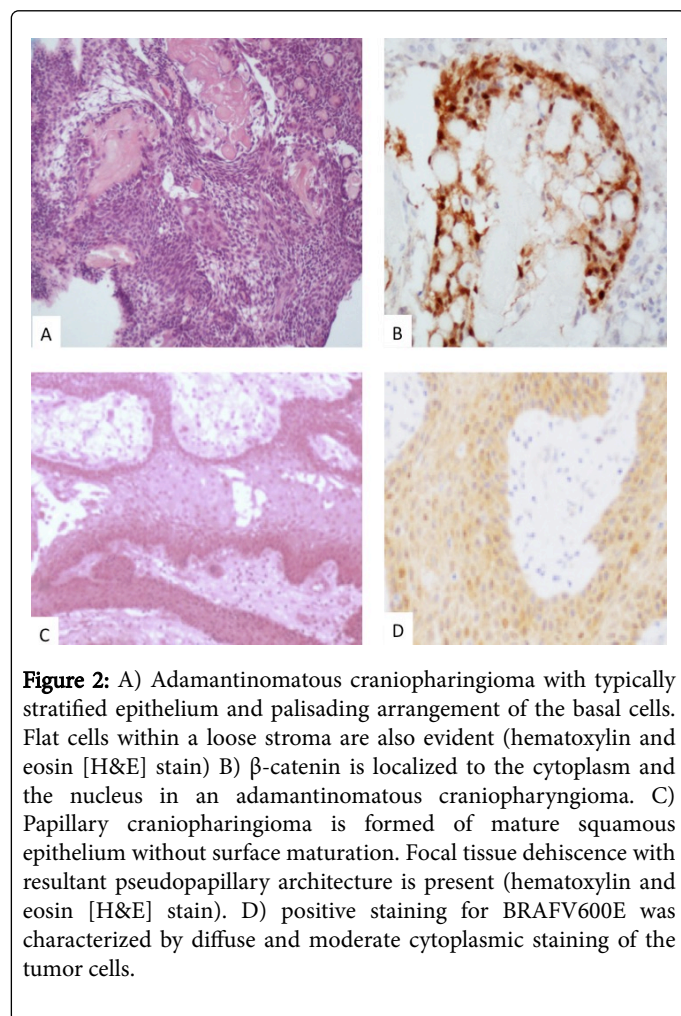


Figure 2: A) Adamantinomatous craniopharyngioma with typically stratified epithelium and palisading arrangement of the basal cells. Flat cells within a loose stroma are also evident (hematoxylin and eosin [H&E] stain) B) β -catenin is localized to the cytoplasm and the nucleus in an adamantinomatous craniopharyngioma. C) Papillary craniopharyngioma is formed of mature squamous epithelium without surface maturation. Focal tissue dehiscence with resultant pseudopapillary architecture is present (hematoxylin and eosin [H&E] stain). D) positive staining for BRAFV600E was characterized by diffuse and moderate cytoplasmic staining of the tumor cells.

BRAFV600E and CTNNB1 (β -catenin) mutation analysis

We additionally performed sequencing in all cases. None of the seven RCC, showed mutations in BRAF (Figure 1) and CTNNB1 (β -catenin).

Discussion

Rathke's cleft cysts and craniopharyngiomas typically arise in the sellar region and on histological examination by some authors they are considered part of a disease spectrum [2].

Histological evidence are shown by Park, et al. [4] which described the occurrence of a primary tumor with microscopic features of RCC, followed by a progression diagnosed as a CP with nuclear accumulation of beta-catenin, arguing for the possibility that CPs may develop from RCCs directly due to beta-catenin mutations.

Moreover other cases of RCC with smooth transitions from single cuboidal epithelium to squamous and/or stratified epithelium in Rathke's epithelia and ciliated and goblet cell craniopharyngioma are believed to provide evidence for this spectrum [5,8].

Moreover, in practice, purely cystic CPs is extremely difficult to distinguish from RCCs during preoperative analysis and by microscopical evaluation. Nevertheless, differentiating both entities is important because the operative management and follow-up is different. One major factor is the risk of persisting endocrine deficits after extensive resection of CP masses that can be avoided in less aggressive partial resection of the RCC wall.

The recent acquisition that adamantinomatous and papillary craniopharyngiomas harbour distinct molecular alterations i.e., β -catenin (CTNB1) [5] and BRAFV600 mutations [8] respectively stimulate us to screen a series of seven cases of RCC for both mutations using immunohistochemistry and DNA sequencing. Moreover we re-evaluated clinical, radiological and histopathological parameters.

Radiological re-evaluation in all seven cases demonstrated a dumbbell-shaped cyst, with hypointense signal in T1-weighted images, and hyperintense signal in T2-weighted images, relative to normal brain parenchyma. The diameter of all cases was smaller than 20 mm. Moreover in three of 7 Rathke cleft cysts and the intrasphenoidal lesion MRI showed findings typical of a cystic lesion, causing a challenge in differential diagnosis with CP.

The histological parameters of the cases showed columnar mucin-producing, ciliated cells with a focal area of squamous epithelium. The intrasphenoidal case showed columnar epithelial layer with cilia and rare goblet cells: the peculiarity of the latter case was the unusual location in the sphenoidal bone, which although rare, one case has been reported [9].

All cases were negative for BRAFV600E and CTNB1 mutations. On the other hand the two craniopharyngioma adamantinomatous and papillary tested showed respectively BRAFV600E and CTNB1 immunopositivity. Published series of RCC do not harbour the BRAF V600E mutation [10] except for a recent paper by Schweizer, et al. [11] in which they analysed for BRAF status 33 RCC, and found 3 cases harbouring BRAFV600E mutation. The authors performed immunohistochemistry in all cases and pyrosequencing in 8 out 33. Pyrosequencing results of the VE1 positive cases clearly indicated the presence of a BRAF V600E mutation whereas in one case a low level mutation was observed.

The histological re-evaluation of the three positive cases showed unusual morphological features for RCC and also clinical parameters were unusual especially the adult ages of patients that were typically of RCC. This re-evaluation prompt the authors to re-diagnosed the BRAF positive case as papillary craniopharyngiomas. The paper by Schweizer,

et al. focuses on the importance of a careful histological evaluation of cystic sellar lesion for an accurate distinction between RCC with squamous metaplasia and papillary craniopharyngioma.

Our study was carried out in a series similar to those of Schweizer, et al. with respect to number of sequenced cases. However, the histological and radiological re-evaluation of the two BRAFv600E mutated RCC cases confirmed that they were misdiagnosed papillary craniopharyngioma, being composed of lesions showed a collagenous cyst wall layered by flat squamous epithelium mixed with columnar cells with cilia and goblet cells. Moreover the clinical parameters were consistent with this diagnosis such as the young age of the case to allow us to exclude a craniopharyngioma rare in young. Also MRI showed features consistent with the diagnosis of RCC such as a dumbbell-shaped cyst, with hypointense signal in T1 and hyperintense signal in T2-weighted images.

In conclusion the issue of a possible spectrum between papillary craniopharyngiomas and Rathke's cleft cysts, as supposed by others, was not supported in our study, both on histological and genetic level. Finally, in extensively cystic CP, difficult to differentiate from RCC, the pathological evaluation associated with molecular findings could definitively define the diagnosis.

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