

Unusual case of occipital lobe dysembryoplastic neuroepithelial tumour with *GNAI1-BRAF* fusion

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DESCRIPTION

A previously healthy 9-year-old girl presented to otolaryngology clinic due to new onset bilateral sensorineural hearing loss. An MRI brain was ordered as part of her routine workup, and she was found to have an incidental 1.3 cm non-enhancing T1 hypointense, T2 hyperintense mass in the right occipital lobe just superior to the tentorium without associated diffusion restriction. The patient otherwise denied any problems with vision, nausea, vomiting or seizures. Her neurological examination was normal other than the sensorineural hearing loss confirmed on audiometry, which was likely unrelated to the imaging findings. Given the imaging appearance, a low-grade glioma was suspected, and she was monitored closely with sequential MRI. However, follow-up imaging 2 years later showed a slight increase in tumour size with new heterogeneous enhancement (figure 1). At that time, the patient underwent gross total resection. Permanent sections showed a proliferation of bland, occasionally process-forming cells in a largely myxoid background with microcyst formation, scattered 'floating' neurons within pools of mucin and increased mitotic activity. Findings were most consistent with a dysembryoplastic neuroepithelial tumour (DNET).¹ Immunohistochemistry demonstrated NeuN-positive scattered neurons corresponding to the floating neurons noted on H&E staining, diffusely positive glial fibrillar acidic

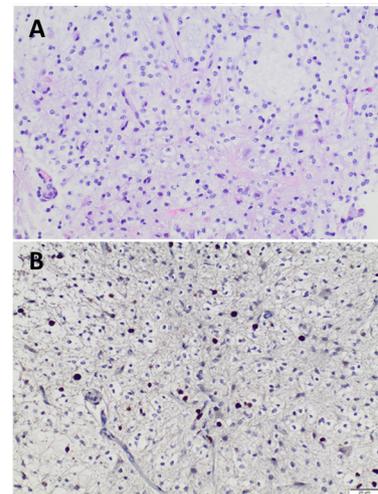


Figure 2 H&E stained sections showing a proliferation predominantly of bland, occasionally process-forming cells in a largely myxoid background with microcyst formation and scattered 'floating' neurons with mitotic activity noted (A), and Ki-67 staining showing ~3%–5% nuclear labelling (B).

protein (GFAP) staining and 3%–5% Ki67 positivity (figure 2). Next-generation sequencing with solid tumour mutation panel analysis revealed a *GNAI1-BRAF* fusion, and a microarray demonstrated chromothripsis of chromosome 7 with nine segmental gains and breakages at 7q34 in *KIAA1549*.

DNETs are WHO grade I low-grade gliomas more commonly seen in children that often present with focal epilepsy.² *BRAF* p.V600E mutations, *FGFR1* alterations and *BRAF-KIAA1549* fusion are more commonly reported, although the incidences vary among different studies.^{3–5} In addition, copy number alterations, most frequently copy number gains in chromosomes 5 and 7, are reported in DNETs, although they do not correlate with specific histological subtypes.^{1 6} We report an unusual occipital low-grade glioma with histological characteristics of a DNET and genetics demonstrating a *GNAI1-BRAF* fusion which has been reported at very low frequencies compared with the more common *KIAA1549-BRAF* fusions in cerebellar juvenile pilocytic astrocytoma. However, our reported case of *GNAI1-BRAF* has not been previously described in DNETs. The rare *GNAI1-BRAF* fusion represents an activating mutation that may potentially be amenable to targeted therapies.⁷ Thus, molecular stratification with next-generation sequencing can aid in tumour classification and further characterise phenotype–genotype correlation to better guide clinical management.

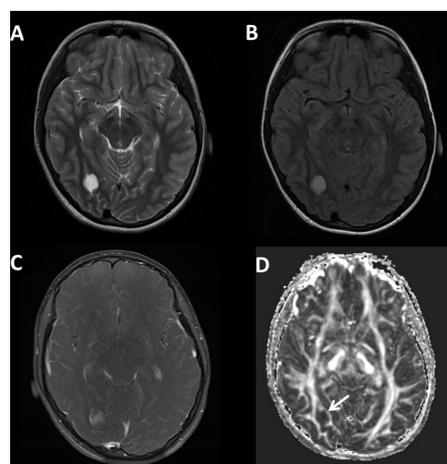


Figure 1 MRI brain demonstrates a well-circumscribed, T2 hyperintense (A) and T2 FLAIR hyperintense (B) mass in the right occipital lobe, with post-contrast showing heterogeneous enhancement (C), and fractional anisotropy on diffusion tensor imaging showing fibre displacement of the right optic radiation on both sides of the lesion (D).



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Learning points

- ▶ Dysembryoplastic neuroepithelial tumours (DNETs) are predominantly paediatric low-grade gliomas that may or may not present with focal neurological symptoms.
- ▶ *GNAI1-BRAF* fusion has not been previously reported in patients with DNETs, thus adding to the genotypic variation of this heterogeneous group of tumours.
- ▶ Next-generation sequencing can provide additional tumour characterisation and guide future studies in phenotype–genotype correlations.

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REFERENCES

- 1 Suh Y-L. Dysembryoplastic neuroepithelial tumors. *J Pathol Transl Med* 2015;49:438–49.
- 2 Chassoux F, Landré E, Mellerio C, *et al.* Dysembryoplastic neuroepithelial tumors: epileptogenicity related to histologic subtypes. *Clin Neurophysiol* 2013;124:1068–78.
- 3 Ryall S, Zapotocky M, Fukuoka K, *et al.* Integrated molecular and clinical analysis of 1,000 pediatric low-grade gliomas. *Cancer Cell* 2020;37:569–83.
- 4 Chappé C, Padovani L, Scavarda D, *et al.* Dysembryoplastic neuroepithelial tumors share with pleomorphic xanthoastrocytomas and gangliogliomas BRAF(V600E) mutation and expression. *Brain Pathol* 2013;23:574–83.
- 5 Gierke M, Sperveslage J, Schwab D, *et al.* Analysis of IDH1-R132 mutation, BRAF V600 mutation and KIAA1549-BRAF fusion transcript status in central nervous system tumors supports pediatric tumor classification. *J Cancer Res Clin Oncol* 2016;142:89–100.
- 6 Chiang JC, Ellison DW. Molecular pathology of paediatric central nervous system tumours. *J Pathol* 2017;241:159–72.
- 7 Jain P, Silva A, Han HJ, *et al.* Overcoming resistance to single-agent therapy for oncogenic *BRAF* gene fusions via combinatorial targeting of MAPK and PI3K/mTOR signaling pathways. *Oncotarget* 2017;8:84697–713.

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