Regression of a glioblastoma multiforme: spontaneous versus a potential antineoplastic effect of dexamethasone and levetiracetam

Prakash Peddi,1 Nisha Elizabeth Ajit,2 Gary Von Burton,3 Hazem El-Osta1

SUMMARY
Patients with grade IV astrocytoma or glioblastoma multiforme (GBM) have a median survival of <12 months, increased to 14.6 months by maximal safe resection with radiation and temozolomide. In the absence of chemotherapy, radiotherapy or chemoradiotherapy, spontaneous regression of GBM or regression while only being on dexamethasone (DEX) and levetiracetam (LEV) have seldom been reported. Here, we present a case of a patient who had significant regression of the GBM with DEX and LEV alone. In this study, we hypothesise a plausible antineoplastic role of DEX and or LEV in GBM and highlight molecular, preclinical and clinical studies supporting this role.

BACKGROUND
Patient with GBM has a poor prognosis despite aggressive therapy. This case describes an outlier in the survival curve, doing well currently. A detailed insight into his story offers us plausible explanations for the regression of this tumour. Spontaneous regression of the tumour versus a combined antineoplastic effect of antiepileptic drugs (AEDs) and dexamethasone (DEX) are the leading hypothesis. This case reviews preclinical and retrospective data, which support our hypothesis, and also emphasizes that further clinical studies are a prerequisite for validation and for developing treatment protocols in a tumour with a poor prognosis.

CASE PRESENTATION
A 55-year-old man was admitted on March 2014 for acute-onset lethargy and confusion. His medical history includes biopsy-proven right frontal lobe grade II astrocytoma, diagnosed in April 2000. After undergoing definite radiation, he achieved a long-term remission. In May 2009, MRI of the brain demonstrated a new heterogeneous mass in the right frontal and temporal lobe with minimal mass effect over right anterior horn. As the patient declined surgery, he was empirically treated with oral temozolomide (TMZ) at 150 mg/m² (days 1–5 and repeated every 28 days) from August 2009 to December 2011. As patient stopped taking TMZ, if levetiracetam (LEV) 500 mg orally two times per day) followed by oral dosing of 4 mg two times per day) and levetiracetam (LEV) 500 mg orally two times per day for seizure prophylaxis. Brain biopsy revealed anaplastic fibrillary astrocytes with high mitosis and infiltrating into the adjacent brain parenchyma with associated satellitosis and with large areas of necrosis. Patient was diagnosed with WHO grade IV astrocytoma or GBM (figure 1C). Over the next 2-month period, patient’s cognition gradually improved with resolution of delirium and speech problems, while he remained on oral DEX and LEV. Brain MRI on day 57 revealed mild improvement (figure 1D). On day 99, brain MRI showed further reduction of frontal, parasagittal mass size (figure 1E). On day 120, he was started on concurrent TMZ and radiation followed by the maintenance of TMZ. At the time of writing this manuscript, patient continues to do well with brain MRI showing no signs of recurrence (figure 1F).

DISCUSSION
GBM is the most aggressive and lethal primary brain tumour and carries a dismal prognosis.1,2 Maximal safe resection followed by adjuvant concurrent chemotherapy and radiation with oral TMZ remains the treatment of choice.3 While rare cases of long-term survival with GBM have been described,4 spontaneous regression in GBM setting has never been published. Continuous regression of GBM was noted in our patient while on DEX and LEV, without any cancer-targeted therapy, suggesting that the response may be secondary to DEX and/or LEV.

DEX’s inherent role in altering the natural course of GBM by itself or in congruence with chemotherapy and radiation is unknown.5 In contrary, high-dose DEX (>4.1 mg/day) was proposed to mitigate the effects of chemotherapy resulting in lower overall survival (OS) in patients with GBM.6 Most of the understanding of DEX antineoplastic effects in GBM comes from in vitro studies. Takahashi et al7 demonstrated that DEX inhibits adrenomedullin (AM) induced by inflammatory cytokines interleukin-γ, tumour necrosis factor-α and interleukin-1 in human GBM cell line T98G. Hypoxia and inflammatory cytokines induce AM, which in turn increases vascular endothelial growth fractions through the TGF-β pathway.

To cite: Peddi P, Ajit NE, Burton GV, et al. BMJ Case Rep Published online: [please include Day Month Year] doi:10.1136/bcr-2016-217393

1Department of Hematology
Louisiana State University
Shreveport School of Medicine, Shreveport, Louisiana, USA
2Department of Internal Medicine, Louisiana State University Health Sciences Center Shreveport School of Medicine, Shreveport, Louisiana, USA
3Department of Hematology and Oncology, Louisiana State University Health Sciences Center Shreveport School of Medicine, Shreveport, Louisiana, USA

Correspondence to Dr Prakash Peddi, ppeddi@lsuhsc.edu

Accepted 4 December 2016

Findings that shed new light on the possible pathogenesis of a disease or an adverse effect
Increased VEGF worsens the vicious cycle of more hypoxia and more AM, thereby potentiating angiogenesis, tumour proliferation, invasion and metastasis. Along with above mechanisms, DEX, via the glucocorticoid receptor pathway, is shown to directly inhibit VEGF production in glioblastoma cells. There is also growing body of evidence that DEX inhibits glioma cell proliferation in vitro and tumour growth in vivo. Its capability to inhibit neural stem cells by decreasing cyclin D1 level has been also demonstrated. Inhibition of migration and invasion in various glioma cell lines (C6, U251, U373 and A172) by DEX has been shown by Bauman et al. In U87MG glioma cells, DEX reduces matrix metalloproteinase-2 secretion and thereby tumour invasion. DEX direct cytotoxic effect on glioblastoma cell lines (A172, T98G and 86HG) has been described. Recent preclinical and clinical studies elucidating antitumor efficacy of AEDs in brain tumours demonstrate encouraging results. Valproic acid (VPA) was shown to cause growth arrest and apoptosis in medulloblastoma cell lines by regulating expression of p21Cip1, cyclin-dependent Kinase 4 (CDK4) and c-myc proteins through histone hyperacetylation. Van Nifterik et al. demonstrated increased cytotoxicity in human glioma cells subjected to TMZ and γ-radiation in the presence of VPA. Data from EORTC 26981-22981 and NCIC CE.3 clinical trial database have revealed that the OS of patients who were receiving VPA alone appeared to increase survival benefit from TMZ and radiation compared to those who were on enzyme-inducing AEDs only (HR 0.39, 95% CI 0.24 to 0.63) or those not receiving any AED (HR: 0.67, 95% CI 0.49 to 0.93). Van Nifterik et al demonstrated increased cytotoxicity in human glioma cells subjected to TMZ and γ-radiation in the presence of VPA. Data from EORTC 26981-22981 and NCIC CE.3 clinical trial database have revealed that the OS of patients who were receiving VPA alone appeared to increase survival benefit from TMZ and radiation compared to those who were on enzyme-inducing AEDs only (HR 0.39, 95% CI 0.24 to 0.63) or those not receiving any AED (HR: 0.67, 95% CI 0.49 to 0.93).

Recent preclinical and clinical studies elucidating antitumorous efficacy of AEDs in brain tumours demonstrate encouraging results. Valproic acid (VPA) was shown to cause growth arrest and apoptosis in medulloblastoma cell lines by regulating expression of p21Cip1, cyclin-dependent Kinase 4 (CDK4) and c-myc proteins through histone hyperacetylation. Van Nifterik et al. demonstrated increased cytotoxicity in human glioma cells subjected to TMZ and γ-radiation in the presence of VPA. Data from EORTC 26981-22981 and NCIC CE.3 clinical trial database have revealed that the OS of patients who were receiving VPA alone appeared to increase survival benefit from TMZ and radiation compared to those who were on enzyme-inducing AEDs only (HR 0.39, 95% CI 0.24 to 0.63) or those not receiving any AED (HR: 0.67, 95% CI 0.49 to 0.93). Van Nifterik et al demonstrated increased cytotoxicity in human glioma cells subjected to TMZ and γ-radiation in the presence of VPA. Data from EORTC 26981-22981 and NCIC CE.3 clinical trial database have revealed that the OS of patients who were receiving VPA alone appeared to increase survival benefit from TMZ and radiation compared to those who were on enzyme-inducing AEDs only (HR 0.39, 95% CI 0.24 to 0.63) or those not receiving any AED (HR: 0.67, 95% CI 0.49 to 0.93).
Tumour molecular profiling using Caris Life Sciences (Phoenix, Arizona, USA) revealed a deficiency of the DNA repair system at multiple levels (Table 1): BRCA1/2 mutation, along with MGMT gene silencing by methylation, isocitrate dehydrogenase 1 (IDH1) mutation and 1p/19q codeletion, known for their correlation with improved sensitivity to chemotherapy and other genotoxic stress, were present. It is possible that the impaired DNA repair system has conferred an improved sensitivity to DEX and LEV. This observation needs to be proven through cell lines model harbouring defect at different levels in their DNA repair system and from large clinical database. Although DEX and LEV’s role in GBM regression is supported by few preclinical and retrospective clinical data, randomised clinical trials are still lacking. Further efforts are needed to define the antineoplastic role of DEX and LEV in GBM.

**Learning points**

- Secondary glioblastoma multiforme (GBM) progress from astrocytoma and have a better prognosis than primary GBM.
- Patients with tumours having O6 methylguanine-DNA methyltransferase promoter methylation respond better to alkylating agents such as temozolamide, translating to longer survival.
- Dexamethasone and levetiracetam have been shown in in vitro studies and retrospective studies to play a role in directly inhibiting tumour proliferation and increasing sensitivity to chemotherapy.
- Randomised control trials can validate the role of antiepileptic drugs and dexamethasone in the treatment of GBM.

**Contributors**

PP and NEA conceived and designed the manuscript, and also drafted the manuscript.

**Competing interests**

None declared.

**Patient consent**

Obtained.

**Provenance and peer review**

Not commissioned; externally peer reviewed.

**Open Access**

This is an Open Access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: http://creativecommons.org/licenses/by-nc/4.0/

**REFERENCES**

Findings that shed new light on the possible pathogenesis of a disease or an adverse effect