Case report: response to the ERK1/2 inhibitor ulixertinib in BRAF D594G cutaneous melanoma

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Melanoma is characterized by oncogenic mutations in pathways regulating cell growth, proliferation, and metabolism. Greater than 80% of primary melanoma cases harbor aberrant activation of the mitogen-activated protein kinase/extracellular-signal-regulated kinase (MEK/ERK) pathway, with oncogenic mutations in BRAF, most notably BRAF V600E, being the most common. Significant progress has been made in BRAF-mutant melanoma using BRAF and MEK inhibitors; however, non-V600 BRAF mutations remain a challenge with limited treatment options. We report the case of an individual diagnosed with stage III BRAF D594G-mutant melanoma who experienced an extraordinary response to the ERK1/2 inhibitor ulixertinib as fourth-line therapy. Ulixertinib was obtained via an intermediate expanded access protocol with unique flexibility to permit both single-agent and combination treatments, dose adjustments, breaks in treatment to undergo surgery, and long-term preventive treatment following surgical resection offering this patient the potential for curative treatment.

Keywords: immunotherapy, melanoma, targeted therapy

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Introduction

Melanoma is associated with aberrant activation of the mitogen-activated protein kinase/extracellular-signal-regulated kinase (MEK/ERK) pathway in more than 80% of cases [1]. Mutations in the BRAF gene are the most common mutations and reported in approximately 50% of melanoma cases [1–3]. Although most BRAF mutations occur at codon 600 (most commonly V600E), non-V600 mutations occur in a low percentage of patients [3].

There has been significant development in targeted therapies for BRAF mutations, including the selective inhibitors of BRAF V600-mutant kinase dabrafenib and vemurafenib, and inhibitors of the downstream kinase MEK such as cobimetinib and trametinib. As a result, numerous landmark studies have reported dramatic improvements in both overall and progression-free survival in patients with melanoma harboring BRAF V600 mutations [4–6]. Given these advances, molecular testing for BRAF mutations in advanced melanoma patients has become standard practice and is recommended by the National Comprehensive Cancer Network and European Society for Medical Oncology. However, despite significant progress in targeting BRAF V600 mutations, options for atypical, non-V600 mutations are severely limited.

BRAF D594G is a class 3 BRAF mutation resulting in a kinase-dead, catalytically inactive BRAF protein that dimerizes with the CRAF protein to result in hyperactivation of ERK [7,8]. BRAF D594G mutations have been shown to confer sensitivity to RAF inhibitors, as well as to MEK inhibitors such as trametinib [9]. Indeed, the phase 2 NCI-MATCH trial reported prolonged stable disease in a BRAF D594G-mutant lung cancer patient treated with trametinib [10]. Here, we present the first case, to our knowledge, of a patient with advanced melanoma harboring the BRAF D594G mutation who has been treated successfully with an ERK inhibitor. Notably, this patient accessed ulixertinib via an expanded access program (NCT04566393) as she was ineligible for active clinical trials at the time. Safety and outcomes data were collected and structured according to an observational research protocol (XCELSIOR, NCT03793088).
Presentation of case
A 43-year-old female patient was diagnosed with pT1a melanoma of the left lateral lower back. Following excision of the melanoma, she was followed for 4 years without incident until she developed left axillary adenopathy measuring 4.7 cm. Biopsy of the adenopathy revealed melanoma. A PET/computed tomography showed FDG-avid lymphadenopathy in the left axillary and subpectoral lymph nodes but no other sites of disease. The disease was deemed unresectable due to axillary vein involvement and muscle wall attachment. MRI of the brain was negative for intracranial metastases.

Molecular profiling of the biopsied tissue through Strata, Ann Arbor, Michigan, USA (https://stratadx.com/) identified the BRAF D594G mutation, a TERT promoter mutation, an inactivating PTEN mutation, and two mutations of unknown significance in MEN1 and CDH20. Microsatellite status was stable and tumor mutational burden was low (4 muts/Mb).

Treatment was initiated with nivolumab 1 mg/kg and ipilimumab 3 mg/kg every 21 days. The patient developed grade 4 immune-related colitis. Due to grade 4 toxicity, immunotherapy was not resumed, and the patient was started trametinib 2 mg daily given the atypical BRAF mutation. She developed diarrhea and a rash and only remained on this treatment for 1 month without response. Her disease remained isolated to the left axillary and subpectoral lymph nodes but remained unresectable. T-VEC, a genetically modified herpes virus injected directly into lesions, was given in combination with pembrolizumab. Following nine T-VEC treatments and four cycles of pembrolizumab 200 mg every 21 days, imaging showed stable disease in the left axilla and subpectoral region, with the largest dominant node still measuring 4.5 cm.

The patient was then enrolled in an Expanded Access Program for the ERK1/2 inhibitor ulixertinib (NCT04566393). She started single-agent ulixertinib 600 mg twice daily. After 6 days of treatment, she noticed a decrease in the bulkiness of the left axillary lymph nodes. Due to grade 1 mucositis, grade 1 rash, grade 1 diarrhea, and intermittent grade 2 visual floaters, the ulixertinib dose was adjusted numerous times over the course of several weeks and ultimately held for a period of 2 weeks, after which it was resumed at 150 mg twice daily with no further adverse effects.

Imaging after 2 months of ulixertinib showed persistent disease in the left axilla and subpectoral region, again measuring 4.5 cm. However, the nodal mass demonstrated central necrosis and had detached from the chest wall, musculature, and vessels. The disease was deemed resectable. Ulixertinib was held for 1 week before left axillary lymph node dissection with en bloc pectoralis minor muscle resection of levels I, II, and III. Pathology revealed a histologic response to treatment, with extensive tumoral melanosis and very few preserved tumor cells remaining in two lymph nodes (about 20 cells in each node) (Fig. 1). The patient’s disease was deemed high risk for local recurrence based on the bulky disease, and so, she received postoperative radiation to the left axillary lymph node basin (3000 cGy).

Notably, despite the histologic response to treatment, the lymph node size had not changed significantly after treatment with ulixertinib compared with after treatment with T-VEC and pembrolizumab. A biopsy was not performed between these lines of therapy. Therefore, we could not determine how much of the treatment effect was due to immunotherapy versus the ulixertinib. For this reason, she completed 1 year of adjuvant therapy with both pembrolizumab 200 mg every 21 days and ulixertinib 150 mg twice daily. She is currently disease-free 1 year after surgery.

Discussion
This is the first case report demonstrating the use of an ERK1/2 inhibitor in a patient with BRAF-mutant melanoma.

The efficacy of targeted treatments in BRAF-mutant cancers varies, with some patients showing no response and others demonstrating a dramatic response. Moreover, while there has been significant development in options for BRAF V600-mutant tumors, options for atypical mutations outside of BRAF V600 remain limited. This patient’s tumor had an atypical BRAF mutation that results in a kinase-dead, catalytically inactive BRAF protein that dimerizes with the CRAF protein to lead to hyperactivation of ERK [7,8]. Her tumor also had a TERT promoter mutation. The combination of BRAF V600E and TERT promoter mutations has been reported in several tumor types and has been correlated with more aggressive disease and poor outcomes [11–13]. BRAF V600E has been shown to signal through the MAPK pathway to promote the formation of the GABPA-GABPB transcriptional complex to upregulate TERT expression in TERT promoter-mutant cells [14,15]. While the combination of the BRAF and TERT mutations leads to a more aggressive phenotype, the effects of this combination on response to therapy remain elusive. BRAF V600E and TERT promoter mutations were associated with better response to BRAF V600E-targeted therapies such as dabrafenib and trametinib in one study but showed a predictive value for treatment failure in a separate study [15,16]. Furthermore, it is unclear how an atypical BRAF mutation fits into this picture, as activation of the MEK/ERK pathway occurs through a mechanism distinct from that of BRAF V600E. Understanding the BRAF-TERT paradigm may prove relevant in understanding a patient’s disease and in predicting their therapeutic outcomes.

Ulixertinib (BVD-523) is a highly potent, selective, reversible ERK1/2 inhibitor that has shown antitumor activity in both BRAF- and RAS-mutant preclinical
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models [17]. A phase 1 study in patients with advanced solid tumors reported clinical activity of single-agent ulixertinib in both BRAF V600- and non-V600-mutant malignancies, including objective responses in a glioblastoma patient and a lung cancer patient with brain metastases, demonstrating also that ulixertinib penetrates the CNS [18]. Ulixertinib is currently under investigation in multiple phase 1 and phase 2 clinical trials in several different tumor types, including a phase 2 clinical trial currently enrolling patients with advanced solid tumors harboring atypical (non-V600) BRAF mutations or alterations in MEK1/2 (NCT04488003).

A unique aspect of this patient’s care was the flexibility in which the physician was able to treat her with an investigational therapeutic. Under the intermediate expanded access protocol NCT04566393, we were able to make dose adjustments when necessary, pause treatment for...
surgery and radiation, and combine ulixertinib with an immunotherapy agent as part of personalized care. Ulixertinib provided a clinically meaningful benefit by rendering her disease resectable, and she currently remains disease-free at 1 year.

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Conflicts of interest
J.C.F., S.G., B.F., T.J.S., and M.S. are employees of and own stock options of xCures, Inc. J.B. and D.K. are employees of BioMed Valley Discoveries. For the remaining authors, there are no conflicts of interest.

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