



4/2/2026
Prior Authorization
Internal Use Only
JOHNS HOPKINS HEALTH PLANS Voranigo - Priority Partners MCO
This fax machine is located in a secure location as required by HIPAA regulations. Complete/review information, sign and date. Fax signed forms to Johns Hopkins Health Plans at <b>1-410-424-4607</b> . Please contact Johns Hopkins Health Plans at <b>1-888-819-1043</b> with questions regarding the Prior Authorization process. When conditions are met, we will authorize the coverage of Voranigo - Priority Partners MCO.

Drug Name (select from list of drugs shown) Voranigo Tablets (vorasidenib)
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Quantity	Frequency	Strength
Route of Administration	Expected Length of Therapy	

Patient Information	
Patient Name:	_____
Patient ID:	_____
Patient Group No.:	_____
Patient DOB:	_____
Patient Phone:	_____

Prescribing Physician	
Physician Name:	_____
Physician Phone:	_____
Physician Fax:	_____
Physician Address:	_____
City, State, Zip:	_____

<b>Diagnosis:</b> _____	<b>ICD Code:</b> _____
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Comments: _____
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<b>Please circle the appropriate answer for each question.</b>	
1. Will the requested medication be used for any indications that are not Food and Drug Administration (FDA)-approved or guideline-supported?	<input type="checkbox"/> Y <input type="checkbox"/> N
[If yes, no further questions.]	

2. Has the plan authorized this medication in the past for this patient (i.e., previous authorization is on file under this plan)?	<input type="checkbox"/> Y <input type="checkbox"/> N
NOTE: The use of physician samples, or manufacturer product discounts, does not guarantee coverage under the provisions of the medical and/or pharmacy benefit. All pertinent criteria must be met in order to be eligible for benefit coverage.	
[If no, skip to question 4.]	
3. Is there documentation showing that the patient has not experienced unacceptable toxicity or disease progression while on treatment?	<input type="checkbox"/> Y <input type="checkbox"/> N
NOTE: Submission of medical records is required.	
[No further questions.]	
4. Does the patient have central nervous system (CNS) cancer?	<input type="checkbox"/> Y <input type="checkbox"/> N
NOTE: Submission of medical records is required.	
[If no, no further questions.]	
5. Does the patient have a susceptible isocitrate dehydrogenase-1 (IDH1) or isocitrate dehydrogenase-2 (IDH2) mutation?	<input type="checkbox"/> Y <input type="checkbox"/> N
NOTE: Submission of medical records is required.	
[If no, no further questions.]	
6. Will the requested medication be used as monotherapy adjuvant treatment of WHO grade 2 astrocytoma or WHO grade 2 oligodendroglioma?	<input type="checkbox"/> Y <input type="checkbox"/> N
NOTE: Submission of medical records is required.	
[If yes, skip to question 8.]	
7. Will the requested medication be used as monotherapy treatment of progressive or recurrent disease for ANY of the following types of CNS cancers: A) H3-mutated high-grade glioma, B) High-grade astrocytoma with piloid features (HGAP), C) WHO grade 3 Pleomorphic xanthoastrocytoma (PXA), D) WHO grade 2 or 3 oligodendroglioma and Karnofsky Performance Status (KPS) greater than or equal to 60, OR E) WHO grade 2 astrocytoma and KPS greater than or equal to 60?	<input type="checkbox"/> Y <input type="checkbox"/> N
NOTE: Submission of medical records is required.	
[If no, no further questions.]	
8. Is the patient 12 years of age or older?	<input type="checkbox"/> Y <input type="checkbox"/> N

I attest that the medication requested is medically necessary for this patient. I further attest that the information provided is accurate and true, and that the documentation supporting this information is available for review if requested by the claims processor, the health plan sponsor, or, if applicable a state or federal regulatory agency.

**Prescriber (Or Authorized) Signature and Date**