



2/20/2026

Prior Authorization

Internal Use Only

JOHNS HOPKINS HEALTH PLANS

Crenessity - 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 **1-410-424-4607**.
Please contact Johns Hopkins Health Plans at **1-888-819-1043** with questions regarding the Prior Authorization process.

When conditions are met, we will authorize the coverage of Crenessity - Priority Partners MCO.

Drug Name (select from list of drugs shown)

Crenessity (crinecerfont)

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: _____

Diagnosis: _____

ICD Code: _____

Comments: _____

Please circle the appropriate answer for each question.

1. Will the requested medication be used for any indications that are not Food and Drug Administration (FDA)-approved or guideline-supported?

Y N

[If yes, no further questions.]

2. Does the patient have severe renal impairment or end-stage renal disease?	<input type="checkbox"/> Y <input type="checkbox"/> N
[If yes, no further questions.]	
3. Is the patient less than 4 years of age?	<input type="checkbox"/> Y <input type="checkbox"/> N
[If yes, no further questions.]	
4. Has the plan authorized the requested 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 6.]	
5. Does the patient have documentation showing a positive response to treatment, evidenced by at least ONE of the following: A) Reduction in glucocorticoid usage, B) Decrease in body mass index standard deviation, C) Improved insulin resistance, or D) Reduction in hirsutism?	<input type="checkbox"/> Y <input type="checkbox"/> N
NOTE: Submission of medical records is required.	
[No further questions.]	
6. Does the patient have a diagnosis of classis congenital adrenal hyperplasia (CAH)?	<input type="checkbox"/> Y <input type="checkbox"/> N
NOTE: Submission of medical records is required.	
[If no, no further questions.]	
7. Is there documentation of a genetic test confirmation of pathogenic variants in CYP21A2 gene?	<input type="checkbox"/> Y <input type="checkbox"/> N
NOTE: Submission of medical records is required.	
[If yes, skip to question 9.]	
8. Is there documentation of laboratory tests confirming 21-hydroxylase deficiency (such as baseline morning serum 17-hydroxyprogesterone (17-OHP), measured by liquid chromatography-tandem mass Spectrometry (LC-MS/MS), cosyntropin (ACTH) stimulation test, or adrenal steroid profile)?	<input type="checkbox"/> Y <input type="checkbox"/> N
NOTE: Submission of medical records is required.	
[If no, no further questions.]	
9. Will the patient continue to receive concurrent treatment with glucocorticoids, and has been stable for at least one month?	<input type="checkbox"/> Y <input type="checkbox"/> N
NOTE: Submission of medical records is required.	

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