

#### Recorlev

Patient Informati	on:	
Name:	····	
Member ID:		
Address:		
City, State, Zip:		
Date of Birth:		
Prescriber Inforr	nation:	
Name:	iduon.	
NPI:		
Phone Number:		
Fax Number		
Address:		
City, State, Zip:		
Requested Medi	eation	
Rx Name:		
Rx Strength		
Rx Quantity:		
Rx Frequency:		
Rx Route of		
Administration:		
Diagnosis and ICI	Code:	
prescribed a medicate quantities can be proupled upon receipt of the SECTION A: Prequests. Phare medications that	ption benefit requires that we review certain requests for coverage with the prescribe on for your patient that requires Prior Authorization before benefit coverage or coverage vided. Please complete the following questions then fax this form to the toll-free number completed form, prescription benefit coverage will be determined based on the ease note that supporting clinical documentation is required for nacy prior authorization reviews can be subject to trial with additional tare not listed within the criteria. The policies are subject to characteristics, MDH transmittals and updates to treatment guidelines.	e of additional or listed below. plan's rules. <b>ALL</b> PA tional ange base
1 Is this re	uest for initial therapy or for a continuation of therapy?	
[] Initial (	f checked, go to 7)	
[] Contin	ation (If checked, go to 2)	
-	ient currently receiving the requested medication?  Yes to question 7.]	s No

If you have any questions, call: 1-888-258-8250

Does the patient have a previously approved PA on file with the current plan?   Note: If the patient does NOT have a previously approved prior authorization (PA) on file for the requested medication with the current plan, the renewal request will be considered under initial therapy.]   If no, skip to question 7.]  Has the patient been established on therapy for at least 3 months?   If no, skip to question 7.]  Has the patient been established on therapy for at least 3 months?   If no, skip to question 7.]  Has the patient been established on therapy for at least 3 months?   If no, skip to question 7.]  Has documentation been submitted to confirm that the patient has had a significant response to therapy, as determined by the provider? ACTION REQUIRED: Submit supporting documentation. [Note: Improvement of clinical symptoms of Cushing's syndrome and laboratory values showing improvement from baseline.]   No further questions.]    What is the diagnosis or indication?   Endogenous hypercortisolemia due to Cushing's syndrome (If checked, go to 8)   Other (If checked, no further questions)    If the patient 18 years of age or older?   If no, no further questions.]  Does the patient have documentation supporting diagnosis of Cushing's syndrome? ACTION REQUIRED: Submit supporting documentation. [Note: Ahoromal dexamethasone suppression test (DST) or 2 measurements of elevated late night salivary cortisol concentrations.]  Has documentation been submitted to confirm clinical symptoms of Cushing's syndrome?   ACTION REQUIRED: Submit supporting documentation. [If no, no further questions.]    Does the patient have documentation of pretreatment cortisol levels obtained within 90 days of this request? ACTION REQUIRED: Submit supporting documentation. [Note: Urinary free cortisol (UFC) -1.5x ULN.]   If no, no further questions.]    Is the patient a candidate for surgery? ACTION REQUIRED: Submit supporting documentation. [Note: rationale for not being a candidate is required.]   If no, skip to question 15.]	3	Has the patient been receiving medication samples of the requested medication? [If yes, skip to question 7.]	Yes	No
[If no, skip to question 7.]  Has documentation been submitted to confirm that the patient has had a significant response to therapy, as determined by the provider? ACTION REQUIRED: Submit supporting documentation. [Note: Improvement of clinical symptoms of Cushing's syndrome and laboratory values showing improvement from baseline.] [No further questions.]  What is the diagnosis or indication? [I Endogenous hypercortisolemia due to Cushing's syndrome (If checked, go to 8)  [I] Other (If checked, no further questions)  Is the patient 18 years of age or older? [If no, no further questions.]  Does the patient have documentation supporting diagnosis of Cushing's syndrome? ACTION REQUIRED: Submit supporting documentation. [Note: Abnormal dexamethasone suppression test (DST) or 2 measurements of elevated late night salivary cortisol concentrations.] [If no, no further questions.]  Has documentation been submitted to confirm clinical symptoms of Cushing's syndrome? ACTION REQUIRED: Submit supporting documentation. [If no, no further questions.]  Does the patient have documentation of pretreatment cortisol levels obtained within 90 days of this request? ACTION REQUIRED: Submit supporting documentation. [Note: Urinary free cortisol (UFC) -1.5x ULN.] [If no, no further questions.]	4	Does the patient have a previously approved PA on file with the current plan? [Note: If the patient does NOT have a previously approved prior authorization (PA) on file for the requested medication with the current plan, the renewal request will be considered under initial therapy.]	Yes	No
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[] Endogenous hypercortisolemia due to Cushing's syndrome (If checked, go to 8)  [] Other (If checked, no further questions)  8	6	significant response to therapy, as determined by the provider? ACTION REQUIRED: Submit supporting documentation. [Note: Improvement of clinical symptoms of Cushing's syndrome and laboratory values showing improvement from baseline.]	Yes	No
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syndrome? ACTION REQUIRED: Submit supporting documentation. [Note: Abnormal dexamethasone suppression test (DST) or 2 measurements of elevated late night salivary cortisol concentrations.] [If no, no further questions.]  10 Has documentation been submitted to confirm clinical symptoms of Cushing's Yes No syndrome?  ACTION REQUIRED: Submit supporting documentation. [If no, no further questions.]  11 Does the patient have documentation of pretreatment cortisol levels obtained Within 90 days of this request? ACTION REQUIRED: Submit supporting documentation. [Note: Urinary free cortisol (UFC) -1.5x ULN.] [If no, no further questions.]  12 Is the patient a candidate for surgery? ACTION REQUIRED: Submit supporting documentation. [Note: rationale for not being a candidate is required.] [If no, skip to question 15.]	8	·	Yes	No
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documentation. [Note: rationale for not being a candidate is required.] [If no, skip to question 15.]	11	within 90 days of this request? ACTION REQUIRED: Submit supporting documentation. [Note: Urinary free cortisol (UFC) -1.5x ULN.]	Yes	No
13 Did the patient have pituitary surgery? ACTION REQUIRED: Submit supporting Yes No	12	documentation. [Note: rationale for not being a candidate is required.]	Yes	No
	13	Did the patient have pituitary surgery? ACTION REQUIRED: Submit supporting	Yes	No

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	documentation. [If no, no further questions.]		
14	Has documentation been submitted to show pituitary surgery was not curative? ACTION REQUIRED: Submit supporting documentation. [If no, no further questions.]	Yes	No
15	Has documentation been submitted to show baseline laboratory testing obtained within 90 days of this request for the following: A) Liver tests (ALT/AST/total bilirubin), B) Electrocardiogram (ECG), C) Potassium, and D) Magnesium levels? ACTION REQUIRED: Submit supporting documentation. [If no, no further questions.]	Yes	No
16	Has the patient been treated with mitotane within 6 months of this request? [If yes, no further questions.]	Yes	No
17	Does the patient have any clinical or radiological signs of compression of the optic chiasm? [If yes, no further questions.]	Yes	No
18	Does the patient have a contraindication to oral Ketoconazole tablets? [If yes, no further questions.]	Yes	No
19	Has the patient failed at least 90 days of therapy with maximally tolerated dosages of Ketoconzazole in the last 12 months? ACTION REQUIRED: Submit supporting documentation. [If yes, skip to question 21.]	Yes	No
20	Has documentation been provided to confirm that the patient has experienced intolerance, adverse side effect, or treatment failure to Ketoconazole? ACTION REQUIRED: Submit supporting documentation. [If no, no further questions.]	Yes	No
21	Has the patient failed at least 90 days of therapy with Signifor/Signifor LAR in the last 12 months? ACTION REQUIRED: Submit supporting documentation. [If no, no further questions.]	Yes	No
22	Has documentation been provided to confirm that the patient has experienced intolerance, adverse side effect, or treatment failure to Signifor/Signifor LAR? ACTION REQUIRED: Submit supporting documentation. [If no, no further questions.]	Yes	No
23	Is the requested medication being prescribed by or in consultation with an endocrinologist? [If no, no further questions.]	Yes	No
24	Does the provider attest that Recorlev daily dosage of the requested medication will not exceed 1,200 mg?	Yes	No

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Please document the diagnoses, symptoms, and/or any other information important to this review:				
SECTION B: Physician Signature				
SECTION D. I Hysician Signature				
DUMOIOLANI CIONATUDE	DATE			
PHYSICIAN SIGNATURE	DATE			

#### **FAX COMPLETED FORM TO: 1-833-896-0656**

**Disclaimer:** An authorization is not a guarantee of payment. Member must be eligible at the time services are rendered. Services must be a covered Health Plan Benefit and medically necessary with prior authorization as per Plan policy and procedures.

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