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Pati Pati	ient Name: ient ID: ient Group No:  vsician Office Address:	NPI#:	_ Date: _ Patient Date Of Birth: Patient Phone:	8/12/2024  Physician Name: Specialty: Physician Office Telephone				
Dru	g Name (specify drug)							
Qua	antity:		_					
Route of Administration: Diagnosis:			_ Expected Length of Therapy: . ICD Code:					
Cor								
	• • •	e answer for each applica	ble question.					
1.	What is the diagnosis? Phenylketonuria (If ch	ecked, go to 2)						
	Biopterin metabolic defects - Autosomal dominant guanine triphosphate cyclohydrolase deficiency (Segawa disease) (If checked, go to 2)							
	Biopterin metabolic defects - Autosomal recessive guanine (GTP) cyclohydrolase deficiency (If checked, go to 2)							
	Biopterin metabolic defects - 6-pyruvoyl-tetrahydropterin synthase (6-PTS) deficiency (If checked, go to 2)							
	Biopterin metabolic defects - Sepiapterin reductase deficiency (If checked, go to 2)							
	Biopterin metabolic de checked, go to 2)	efects - Dihydropteridine red	uctase (DHPR) deficiency (If					
	Biopterin metabolic de called primapterinuria	efects - Pterin-4a-carbinolan ) (If checked, go to 2)	nine dehydratase deficiency (also					
	Other, please specify	. (If checked, no further ques	stions)					
2.	ACTION REQUIRED: If	rmed by an enzyme assay, of Yes, attach supporting char Submit supporting docume	genetic testing, or phenylalanine level? t note(s) or test results.	Y		N		
3.	Is this request for contin	uation of therapy with the re	quested medication?	Y		N		
4.	Is the requested medica	tion being requested for a b	iopterin metabolic defect?	Y		N		
5.	What is the patient's bas level?	seline (with dietary interventi	ons alone) blood phenylalanine (Phe)					
	Greater than or equal go to 6)	to 6 mg/dL (greater than or	equal to 360 micromol/L) (If checked,					
	Less than 6 mg/dL (le	ess than 360 micromol/L) (If	checked, no further questions)					
	No baseline blood Ph	e level (If checked, no furthe	er questions)					
6.	Will the requested medion phenylketonuria?	cation be initiated in a patier	nt currently receiving Palynziq for	Υ		N		

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7.	Will Palynziq be discontinued after an appropriate period of overlap?	Y		N	
8.	What is the diagnosis?				
	Phenylketonuria (If checked, go to 9)				
	Biopterin metabolic defects - Autosomal dominant guanine triphosphate cyclohydrolase deficiency (Segawa disease) (If checked, go to 11)				
	Biopterin metabolic defects - Autosomal recessive guanine (GTP) cyclohydrolase deficiency (If checked, go to 11)				
	Biopterin metabolic defects - 6-pyruvoyl-tetrahydropterin synthase (6-PTS) deficiency (If checked, go to 11)				
	Biopterin metabolic defects - Sepiapterin reductase deficiency (If checked, go to 11)				
	Biopterin metabolic defects - Dihydropteridine reductase (DHPR) deficiency (If checked, go to 11)				
	Biopterin metabolic defects - Pterin-4a-carbinolamine dehydratase deficiency (also called primapterinuria) (If checked, go to 11)				
	Other, please specify. (If checked, no further questions)				
9.	Which of the following has the patient demonstrated following the therapeutic trial with the requested medication?				
	Reduction in blood phenylalanine (Phe) level of greater than or equal to 30% from baseline (If checked, go to 10)				
	Phenylalanine (Phe) levels in an acceptable range (less than 360 micromol/L or 6 mg/dL) (If checked, go to 10)				
	Improvement in neuropsychiatric symptoms (If checked, go to 10)				
	None of the above (If checked, no further questions)				
10.	Will the requested medication be used concomitantly with Palynziq for phenylketonuria?	Y		N	
11.	Is the patient experiencing benefit from therapy as evidenced by disease stability or disease improvement?				
	Yes, disease stability (If checked, no further questions)				
	Yes, disease improvement (If checked, no further questions)				
	No, neither disease stability nor disease improvement (If checked, no further questions)				

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

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