SENTARA HEALTH PLAN

PHARMACY PRIOR AUTHORIZATION REQUEST*

<u>Directions</u>: <u>The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request.</u> All other information may be filled in by office staff; <u>fax to 1-800-750-9692</u>. No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. <u>If the information provided is not complete, correct, or legible, the authorization process may be delayed.</u>

Drug Requested: sapropterin dihydrochloride (Kuvan®)

ME	MBER & F	PRESCRIBER INFORMATION: Authorization may be delayed if incomplete.
Memb	oer Name: _	
Member Sentara		#: Date of Birth:
Presci	riber Name:	
Prescriber Signatu		re: Date:
Office	Contact Na	me:
Phone Number:		Fax Number:
DEA (OR NPI #: _	
		MATON: Authorization may be delayed if incomplete
,		
		th/Quantity:
		Length of Therapy: ICD Code, if applicable:
RECO	<u>OMMENDEI</u>	DOSING: Initial dose of 10mg/kg/day is recommended, and may be increased to a dose of month of treatment if phenylalanine levels do not decrease below baseline levels.
each l		RITERIA: Check below all that apply. All criteria must be met for approval. To support all documentation, including lab results, diagnostics, and/or chart notes, must be provided denied.
<u>Initi</u>	al Approva	d: 6 months.
	Prescriber is	a metabolic geneticist or a physician knowledgeable in the management of PKU
	Patient has a phenylketon	diagnosis of hyperphenylalaninemia due to tetrahydrobiopterin (BH4)-responsive uria
	Baseline pho	enylalanine labs must be submitted (please attach current labs with level)
	Patient's cur	rrent weight (please note):
		mpliant with a phenylalanine-restricted diet (please submit chart notes documenting enylalanine intake and use of Phe-free medical food supplements)

(Continued on next page)

	Patient does <u>not</u> have hepatic or renal impairment			
	sapropterin dihydrochloride (Kuvan [®]) will <u>NOT</u> be used in combination with Palynziq [™]			
	For brand name Kuvan approval: Member has had trial and intolerable life-endangering adverse event we generic sapropterin dihydrochloride (must submit completed MedWatch form and chart notes to docume adverse event)			
	Is patient a pregnant female? (please note): □ Yes □ No			
To s	uthorization Approval: 1 year. Check below all that apply. All criteria must be met for approval. upport each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be ided or request may be denied.			
	Phenylalanine levels have decreased by at least 30% from baseline levels and have remained below baseline (please attach current labs with level)			
	Patient remains compliant with a phenylalanine-restricted diet (please submit chart notes document current phenylalanine intake and use of Phe-free medical food supplements)			
	Phenylalanine levels will continue to be measured periodically during therapy			
	Patient's current weight:			
	sapropterin dihydrochloride (Kuvan®) will <u>NOT</u> be used in combination with Palynziq [™]			
	For brand name Kuvan approval: Member has had trial and intolerable life-endangering adverse event wit generic sapropterin dihydrochloride (must submit completed MedWatch form and chart notes to documen adverse event)			
	Patient will be maintained on a dose no greater than the FDA-approved maximum of 20mg/kg/day			
	**Length of authorization will be for 1 year if approved for continuation.			
	Yearly reauthorization will be required.**			
Med	lication being provided by a Specialty Pharmacy - PropriumRx			
	Not all drugs may be covered under every Plan			
If	a drug is non-formulary on a Plan, documentation of medical necessity will be required.			

Use of samples to initiate therapy does not meet step edit/preauthorization criteria.

Previous therapies will be verified through pharmacy paid claims or submitted chart notes.