

## BlueShield. SODIUM PHENYLBUTYRATE Federal Employee Program. PRIOR APPROVAL REQUEST

Send completed form to: Service Benefit Plan Prior Approval P.O. Box 52080 MC 139 Phoenix, AZ 85072-2080 Attn. Clinical Services Fax: 1-877-378-4727

Additional information is required to process your claim for prescription drugs. Please complete the patient portion, and have the prescribing physician complete the physician portion and submit this completed form.

P	atient Infor	mation (required		Provider Information (required)			
Date:				Provider Name:			
Patient Name:				Specialty:	NPI:		
Date of Birth: Sex: □Male □		□Female	Office Phone:	Office Fax:	Office Fax:		
Street Address:				Office Street Address:			
City:		State:	Zip:	City:	State:	Zip:	
Patient ID: <b>R</b>	1 1	1 1 1	, , ]	Physician Signature:		•	
PHYSICIAN COMPLETES							
NOTE: Form must be completed in its <b>entirety</b> for processing							
Please select medication:							
□Buphenyl tablet, powder for solution (sodium phenylbutyrate) □Pheburane oral pellets (sodium phenylbutyrate) □Olpruva packets for oral suspension (sodium phenylbutyrate)							
**Check www.fepblue.org/formulary to confirm which medication is part of the patient's benefit							
Is this request for brand or generic? ☐ Brand ☐ Generic							
1. What is the patient's diagnosis?							
☐ Urea cycle disorders due to argininosuccinic acid synthetase (AS) deficiency							
☐ Urea cycle disorders due to carbamylphosphate synthetase (CPS) deficiency							
☐ Urea cycle disorders due to ornithine transcarbamylase (OTC) deficiency							
$\Box$ Other (pl	ease specify):						
2. Has the patient been on this medication continuously for the last <b>6 months</b> excluding samples? □Yes □No*							
*If NO, please answer the following questions:							
a. Has there been a failure to control ammonia levels with dietary restrictions and/or amino acid supplementation? $\square Yes$							
	b. Is the prescribing physician experienced in the management of urea cycle disorders (UCDs)?  \(\sigma\)Yes \(\sigma\)No						
c. Does	the patient have	e acute hyperammo	onemic encephalo	opathy? □Yes □No			
3. Does the prescriber agree to monitor electrolytes at baseline and as clinically indicated? □Yes □No							
4. Will this medication be used in combination with dietary protein restrictions? □Yes □No							



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Message:

Attached is a Prior Authorization request form.

For your convenience, there are 3 ways to complete a Prior Authorization request:

Electronically Online (ePA) Results in 2-3 minutes FASTEST AND EASIEST	Now you can get responses to drug Prior Authorization requests <b>securely</b> online. <b>Online</b> submissions may receive <b>instant</b> responses and do not require faxing or phone calls.  Requests can be made 24 hours a day, 7 days a week. For more information on electronic prior authorization (ePA) and to register, go to <b>Caremark.com/ePA.</b>
Phone (4-5 minutes for response)	The FEP Clinical Call Center can be reached at (877)-727-3784 between the hours of 7AM-9PM Eastern Time. A live representative will assist with the Prior Authorization, asking for the same information contained on the attached form. Please review the form and have your answers ready for faster service.  The process over the phone takes on average between 4 and 5 minutes.
Fax (3-5 days for response)	Fax the attached form to (877)-378-4727. Requests sent via fax will be processed and responded to within 5 business days. The form must be filled out completely, if there is any missing information the Prior Authorization request cannot be processed.  Please only fax the completed form once as duplicate submissions may delay processing times.

