AvMed

PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

<u>Directions</u>: <u>The prescribing physician must sign and clearly print name (preprinted stamps not valid)</u> on this request. All other information may be filled in by office staff; <u>fax to 1-305-671-0200</u>. No additional phone calls will be necessary if all information (<u>including phone and fax #s</u>) on this form is correct. <u>If the information provided is not complete, correct, or legible, the authorization process can be delayed.</u>

<u>Drug Requested</u>: Furoscix[®] (furosemide)

ME	MBER & PRESCRIBER I	NFORMATION: Authorization may be delayed if incomplete.
Meml	ber Name:	
Member AvMed #:		
Presc	riber Name:	
		Date:
Office	e Contact Name:	
Phone Number:		Fax Number:
NPI #	:	
		orization may be delayed if incomplete.
Drug	Form/Strength:	
Dosing Schedule:		
Diagnosis:		ICD Code:
Weight (if applicable):		Date weight obtained:
<u>Quai</u>	ntity Limit: 6 on-body infusor	s per 90 days
supp		below all that apply. All criteria must be met for approval. To ntation, including lab results, diagnostics, and/or chart notes, must be
Init	ial Authorization: 3 months	
	Medication is being prescribed by	by or in consultation with a cardiologist
	Member is 18 years of age or old	der
٥	Member has a diagnosis of New York Heart Association (NYHA) Class II, III or IV chronic heart failure	
	Member is experiencing congestion due to fluid overload that is <u>NOT</u> considered to be an emergency situation	
	Member has a clinical reason fo as bumetanide, furosemide, or to	r requiring Furoscix (e.g., reduced responsiveness to oral diuretics such orsemide)

(Continued on next page)

- Prescriber agrees that member will use Furoscix for short-term use only **AND** will replace with oral diuretics as soon as practical
- ☐ Member does <u>NOT</u> have any of the following: hypersensitivity to furosemide or medical adhesives; anuria; hepatic cirrhosis or ascites; or acute pulmonary edema
- ☐ Prescriber attests the member will be monitored outpatient for fluid, electrolyte, and metabolic abnormalities

Second Authorization for continued therapy after initial authorization approval: 3 months

☐ Member meets <u>ALL</u> initial authorization criteria and requires re-treatment due to persistent reduced response to oral diuretics (**submit documentation**)

<u>Reauthorization</u>: For use beyond 6 months. Length of authorization is 12 months and will be reassessed annually. Check below all that apply. All criteria must be met for approval. To support each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be provided or request may be denied.

- ☐ Member must have disease improvement and/or stabilization <u>OR</u> improvement in the slope of decline (e.g., improvement in signs/symptoms of fluid overload edema, dyspnea, rapid weight gain)
- ☐ Member has <u>NOT</u> experienced any treatment-restricting adverse effects (e.g., fluid, electrolyte, or metabolic abnormalities, worsening renal function, ototoxicity, acute urinary retention)
- ☐ Member is considered to be refractory to oral diuretics, and continued use of Furoscix is medically necessary for stabilization of their condition

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. *