

Kaiser Permanente Health Plan of Mid-Atlantic States, Inc.
KALYDECO (Ivacaftor) Prior Authorization (PA)
Pharmacy Benefits Prior Authorization Help Desk
Length of Authorizations: Initial- 12 months; Continuation- 12 months

Instructions:

This form is used by Kaiser Permanente and/or participating providers for coverage of **KALYDECO** (Ivacaftor. Please complete all sections, incomplete forms will delay processing. Fax this form back to Kaiser Permanente within 24 hours fax: 1-866-331-2104. If you have any questions or concerns, please call 1-866-331-2103. Requests will not be considered unless all sections are complete.

KP-MAS Formulary can be found at: http://www.providers.kaiserpermanente.org/mas/formulary.html

1 - Patient Information									
Patient Name:	Kaiser Medical ID#:	Date of Birth:							
	2 – Prescriber Information								
Prescriber Name:	Specialty:	NPI:							
Prescriber Address:									
Prescriber Phone #:	Prescriber Fax #:								
3 – Pharmacy Information									
Pharmacy Name:	Pharmacy NPI:								
Pharmacy Phone #	Pharmacy Fax #:								
	4 – Drug Therapy Requested								
Drug 1: Name/Strength/Formulation:									
Drug 2: Name/Strength/Formulation:									
1									

					5- Diag	nosis/Clinic	ai Criteria			
2. 3. 4.	□ In India Is th □ No Mer	itial therapeate the Me member of Yes mber is NC to Yes the mem	oy lember's di r ≥6 month oT homozyg nber have a	agnosis for s of age? A gous for the t least one	ng therapy? ontinuing the the reques ND e F508del m of the follo	utation in the wing mutati	e date: ion: ne CFTR gene ons in the CF	TR gene?	2789+5G → A	
		R74W	G178R	R352Q	S945L	A1067T	F1074L	S1255P	3272-26A→G	4
		D110E	E193K	A455E	S977F	G1069R	D1152H	D1270N	3849+10kbC→T	
		D110H	L206W	S549N	F1052V	R1070Q	G1244E	G1349D	711+3A → G	
								E56K		
 OR- 6. Members with a R117H mutation in the CFTR gene who have clinically significant disease (patients with RII7H and the 5T form of the poly-T tract, but not 7T or 9T) No □ Yes For Continuation of Therapy, Please Respond to Additional Questions Below: 1. Was there documentation of positive clinical response? AND No □ Yes 2. Did the specialist follow-up occur in the past 12 months? AND AST, ALT, bilirubin and ophthalmic changes (patients up to 17 years) are monitored at least annually No □ Yes 										
						Prescriber Si				
Provide	e any	additiona	l supportir	ng informat	tion that sh	ould be take	en into consi	deration:	re applicable to this	request.
I certify that the information provided is accurate. Supporting documentation is ava Prescriber Signature:								Date:		
private a	nd lega	ally protected	by law, includii	ng HIPAA. If yo	u are not the int	tended recipient,	, you are hereby r	notified that any	c individual and purpose. The disclosure, copying, distributes not intended for receipt	ution or taking of