

Fax form to: 866-930-0019 No prior authorization requests for Buprenorphine-containing products will be taken by phone.

Member Information	Name:			DOB:	
	ID:			Sex:	
	Address: City, State, Zip:			Phone:	
	Name:				
	Office Contact:			XDEA:	
	NPI: Address: City, State, Zip: Phone: Diagnosis:		Fax:		
Draduat name	Diagnosisi		ICD-10:		
Product name:	4-1-1-4	D			
Buprenorphine 2 mg			rphine 8 mg tablet	late 9 mg/2 mg	
	xone tablets 2 mg/0.5 mg	Buprenoi	rphine/naloxone tab	lets 8 mg/2 mg	
***Other ***(Any request for BRAI allergic reaction, or thera	NDED product will require a MedWa	tch form, copy of receipt c	of submission to Med	Watch, and chart note documenti	ng the adverse reaction,
Dose & Frequency:		Quantity:	Daily Dose:		
Buprenorphine (without	naloxone) Tablet Requests Only	(Must meet clinical criter	ia AND ONE of the follo	owing)	
Check One: 🗋 Member Is Pregnant 🗍 Up to 1st 7 Days of induction to therapy 🗍 Hypersensitivity to Naloxone					
Induction Requests Only	 Prescriber certifies they are treating the patient for opioid use disorder through the member's benefit/health plan and billing the plan for the services, and member has signed an informed consentYesNo Is this an induction request?Yes If yes, Date of InductionNo (Please skip to Continuation Request Section) Prescriber certifies that the patient is being referred or already started receiving behavioral and psychosocial therapy servicesYesNo Prescriber certifies that the required state controlled substance report (OARRS, KASPER, etc.) has been reviewed no earlier than 2 days prior to the date of this requestYesNo For patients with concurrent behavior health disorders, the prescriber certifies referral for mental health assessment and/or treatment as indicatedYesNo 				
<u>Continuation Requests</u> <u>Only</u>	 Prescriber certifies they are treating the patient for opioid use disorder through the member's benefit/health plan and billing the plan for the services and member has assigned informed consent Yes No Prescriber certifies that the required state controlled substance report (OARRS, KASPER, etc.) has been reviewed no earlier than 2 days prior to the date of this requestYes No (Attach documentation of reason for any opioid, stimulant or benzodiazepine listed on the report.) The prescriber certifies that the patient has at least one negative urine test for opiates within last 3 monthsYes No The prescriber certifies that the patient has at least one positive urine test for Buprenorphine and/or Norbuprenorphine within last 3 monthsYes No The prescriber certifies that the patient has at least one positive urine test for Buprenorphine and/or Norbuprenorphine within last 3 monthsYes No The prescriber certifies that regular medication compliance checks occur and there have been no abnormal findings in the pill countYes No The prescriber certifies that he/she has completed an evaluation and has documented clinical reasoning for continuation of therapyYes No 				
I attest, by signature, that	the above information is true and acc	urate to the best of my know	wledge and has been d	locumented appropriately in the me	mber's medical records.
Prescriber Signature				Date	