OPTIMA HEALTH PLAN PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

Directions: <u>The prescribing physician must sign and clearly print name</u> (preprinted stamps not valid) on this request. All other information may be filled in by office staff; fax to <u>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.

Drug Name: Afrezza® (insulin human)

- Afrezza® is rapid acting insulin indicated to improve glycemic control in adult patients with diabetes mellitus.
- Afrezza® should not be used in patients who smoke or to treat diabetic ketoacidosis.
- Afrezza® should be administered at the beginning of meals, via oral inhalation. Dosing must individualized.
- Before initiating, perform a detailed medical history, physical examination, and spirometry (FEV₁) in all patients to identify potentiation lung disease.

Boxed Warning: Afrezza® is contraindicated			
	d in patients with chronic lung disease such as as	thma or COP	D
Type 1 diabetes: must be used with long-acting	ng insulin		
dication: (please check the indication that	applies)		
☐ Type 1 diabetes	☐ Type 2 diabetes		
ease complete all boxes below: for	original authorization (continued authorizat	ion see below	v)
Patient has tried and failed 30 days of therapy	with subcutaneous rapid acting insulin	☐ Yes	□ No
☐ Humalog® ☐ Apidra® ☐ Novolog® (docu	mentation (chart notes) of failure must be attached)		
Patient is at least 18 years of age		☐ Yes	□ No
Patient currently smokes or has quit smoking	within the past 6 months*	☐ Yes	□ No
Patient is diagnosed with chronic obstructive	pulmonary disease (COPD)*	☐ Yes	□ No
Patient is diagnosed with asthma*		☐ Yes	□ No
Pulmonary function tests were completed * □ FEV _{1:} Date:		☐ Yes	□ No
If treating type 1 diabetes : patient is on conc	omitant long acting insulin*	☐ Yes	□ No
1	nd failed 30 days of therapy with at least 2 oral	☐ Yes	□ No
baseline, 6 months of therapy, and annually	over time as measured by FEV ₁ . Assess pulmor thereafter, even in the absence of pulmonary stients with a decline of $\geq 20\%$ of FEV ₁ from be	symptoms.	ning at
LENGTH OF ORIG	GINAL AUTHORIZATION IS 6 MONTHS	_	
*CONTINUED APPROVAL (1 YEAR II	N LENGTH) IS BASED ON RE-SUBMISSIO	N OF AROV	
CRITERIA AND CURRENT SPIROMET	RY RESULTS.	N OF ADOV	Æ
	RY RESULTS. <mark>I be verified through Pharmacy cl</mark>		Æ
			Æ
	l be verified through Pharmacy cla		/E
PAID transaction(s) will item Name:	l be verified through Pharmacy cla	aims.	
PAID transaction(s) will ient Name:	ll be verified through Pharmacy classification of Birth:	aims.	

Fax Number: _____

*Approved by Pharmacy and Therapeutics Committee: 3/19/2015

Phone Number:

REVISED/UPDATED: 4/29/2015; 12/24/2015;

DEA/NPI #: _