OPTIMA HEALTH PLAN

PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST

<u>Directions</u>: <u>The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request</u>. 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. <u>If information provided is not complete, correct, or legible, authorization may be delayed.</u>

<u>Drug Requested: Topical Immunomodulators (check applicable box below)</u>

	Zyclara [®] (imiquimod) 2.5% Pump: 1 pump per 28 day fill; 2 fills per year		•	(imiquimod) 3.75% Packets/Pump: ox per 28 day fill; 2 fills per year		
	imiquimod 3.75% packets/pump:			(ingenol mebutate) 0.015%/0.05% gel:		
	1 pump/box per 28 day fill; 2 fills per year			30 day fill; 2 fills per year		
	Klisyri® (tirbanibulin) 1% ointment:		1	1 1		
	1 box per year					
D	DRUG INFORMATION: Authorization may be delayed if incomplete.					
Dr	Drug Form/Strength:					
Dosing Schedule:			Length of Therapy:			
Diagnosis:						
CLINICAL CRITERIA: 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.						
<u>Fo</u>	r Actinic Keratosis: Requested product: Klisyri® 1% ointment		W	r External Genital and Perianal arts/Condyloma Acuminata: Requested Product: Zyclara® 3.75% Packets/Pump		
	 □ Picato® gel □ Zyclara® 2.5% or 3.75% pump/packets □ imiquimod 3.75% packets/pump 			Patient has a diagnosis of external genital and/or perianal warts/condylomata acuminata		
	Patient has a diagnosis of Actinic Keratosis			AND		
	Patient has had a 30 day trial and inadequate response or clinically significant adverse reaction to two of the following medications: (Chart notes must be submitted) □ imiquimod (generic Aladara) 5% cream; QL = 48 packets per year			Patient has a documented trial and inadequate response or clinically significant adverse reaction to imiquimod 5% cream (Chart notes must be submitted) OR		
	 □ Topical diclofenac (generic Solaraze) 3% gel; QL= 100 gm per year □ Topical 5-fluoruracil 5 % cream, 2 % solution or 5% solution; QL= 10 mL or 40 gm per year 			Patient has a documented trial and inadequate response or clinically significant adverse reaction to topical podofilox (Chart notes must be submitted)		

(Continued on next page; signature page is required to process request.)

(Please ensure signature page is attached to form.)

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. *

Date of Birth:		
Date:		
Fax Number:		

REVISED/UPDATED: 40/23/2015; 12/22/2015; 12/20/2016; 8/19/2017; (Reformatted) 6/19/2019; 1/22/2020; 6/30/2021

^{*}Approved by Pharmacy and Therapeutics Committee: 8/20/2015