SENTARA COMMUNITY PLAN (MEDICAID)

MEDICAL PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

<u>Directions:</u> The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request. All other information may be filled in by office staff; <u>fax to 1-844-305-2331</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 information provided is not complete</u>, correct, or legible, authorization can be delayed.

<u>Drug Requested</u>: iDose[®] TR (travoprost intracameral implant) 75 mcg (J2508) (Medical)

MEMBER & PRESCRIBER INFORMATION: Authorization may be delayed if incomplete.			
Member Name:			
Member Sentara #:		Date of Birth:	
Prescriber Name:			
Prescriber Signature:			
Office Contact Name:			
Phone Number:			
DEA OR NPI #:			
DRUG INFORMATION: Authoriz	ation may be delayed if incor	nplete.	
Drug Form/Strength:			
osing Schedule: Length of Therapy:			
Diagnosis:	ICD Code, if applicable:		
Weight:	Date:		
□ Left Eye	□ Right Eye	□ Both Eyes	
☐ Standard Review. In checking this box or the member's ability to regain maxim			

Dosing Limits:

- iDose® is an intracameral implant containing 75 mcg travoprost, pre-loaded in a single-dose inserter
- Maximum of 1 single implant per eye per lifetime. Do not readminister to an eye that has received a prior implant
- Re-Treatment of Previously Treated Eye(s). NOT COVERED. iDose® is approved for one-time use in each treated eye. Repeat administration in previously treated eye(s) will NOT be approved

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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.

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	Member is 18 years of age or older		
	Prescribed by, or in consultation with, an Ophthalmologist		
	Member has ONE of the following diagnoses:		
	☐ Open-Angle Glaucoma (OAG)		
	☐ Ocular Hypertension (OHT)		
	Member has NO history of any of the following:		
	 Prior corneal or endothelial cell transplants 		
	 Active or suspected ocular/periocular infection or corneal endothelial cell dystrophy 		
	Absent or ruptured posterior lens capsule		
	• Any eye/laser surgeries within the past 6 months in the affected eye(s)		
	Member has documented treatment failure, intolerance, or contraindication of <u>TWO</u> ophthalmic prostaglandin analogs (e.g., bimatoprost, latanoprost, or travoprost) (verified by chart notes and/or pharmacy paid claims)		
	Member has documented treatment failure, intolerance, or contraindication of at least <u>TWO</u> ophthalmic products (either as monotherapy or as concomitant therapy) from two different pharmacological classes for the treatment of OAG or OHT (verified by chart notes and/or pharmacy paid claims):		
	☐ Alpha-agonist (e.g., brimonidine)		
	☐ Beta-blockers (e.g., betaxolol, timolol)		
	☐ Carbonic anhydrase inhibitors (e.g., brinzolamide, dorzolamide)		
	☐ Rho kinase inhibitor (e.g., netarsudil)		
	Member has documented treatment failure, intolerance, or contraindication to Durysta [™] (verified by chart notes and/or medical claims)		
Medication being provided by (check applicable box(es) below):			
□ P	Physician's office OR		
For urge	ent reviews: Practitioner should call Sentara Health Pre-Authorization Department if they believe a		

standard review would subject the member to adverse health consequences. Sentara's definition of urgent is a lack of treatment that could seriously jeopardize the life or health of the member or the member's ability to regain maximum function.

**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. *

^{*}Approved by Pharmacy and Therapeutics Committee: 5/23/2024 REVISED/UPDATED/REFORMATTED: 5/8/2024: 6/5/2024