SENTARA HEALTH PLANS

PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

<u>Directions</u>: <u>The prescribing physician must sign and clearly print name (preprinted stamps not valid)</u> on this request. All other information may be filled in by office staff; <u>fax to 1-800-750-9692</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 the information provided is not complete, correct, or legible, the authorization process can be delayed.</u>

Glucagon-like peptide (GLP-1) receptor agonists

<u>Drug Requested</u>: (select **ONE** of the following)

	Bydureon BCise® (exenatide ER)		Ozempic® (semaglutide)	
	Byetta® (exenatide ER)		Rybelsus® (semaglutide)	
	Liraglutide (generic Victoza®)		Trulicity® (dulaglutide)	
	Mounjaro® (tirzepatide)			
M	IEMBER & PRESCRIBER INFORMATIO	N:	Authorization may be delayed if incomplete.	
Me	mber Name:			
	mber Sentara #:			
Pre	escriber Name:			
Pre	escriber Signature:		Date:	
Off	fice Contact Name:			
	one Number:			
NP	I #:			
	RUG INFORMATION: Authorization may be			
Drı	ug Name/Form/Strength:			
Dosing Schedule:			Length of Therapy:	
Diagnosis:			ICD Code, if applicable:	
We	eight (if applicable):		Date weight obtained:	
D	and an along water D			

Provider please note: Requests received for any target drug above, prescribed solely for chronic weight management will be **DENIED** as these drugs have **NOT** been FDA approved for this indication.

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			nember be discontinuing a previously n if approved for requested medication	prescribed glucagon-like peptide (GLP-1) receptor agonist n?			
				□ Yes OR □ No			
		-	ase list the medication that will be dis along with the corresponding effective	continued and the medication that will be initiated upon e date.			
Mo	Medication to be discontinued:		on to be discontinued:	Effective date:			
				Effective date:			
suppo	ort e	ach		nat apply. All criteria must be met for approval. To uding lab results, diagnostics, and/or chart notes, must be			
			er has a diagnosis of Type 2 Diabetes it documentation):	Mellitus as confirmed by a history of ONE of the following			
		Hemoglobin A1c (A1C) greater than or equal to 6.5%					
☐ Fasting plasma glucose (FPG) greater than or equal to 126 mg/dL (after fast				an or equal to 126 mg/dL (after fasting for at least 8 hours)			
			nour plasma glucose greater than or e 5 g oral glucose after fasting for at lea	qual to 200 mg/dL as part of an oral glucose tolerance test st 8 hours)			
	☐ Member must meet <u>ONE</u> of the following:						
		☐ Hemoglobin A1c (A1C) greater than or equal to 9%					
	 Member has tried and failed, has a clinically significant contraindication or intolerand (verified by chart notes and/or pharmacy paid claims) 						
			ember has atherosclerotic cardiovascu lowing conditions or past medical his	alar disease (ASCVD) as defined by one or more of the story (check all that apply):			
			Acute coronary syndrome				
			Coronary artery disease (CAD)				
			History of myocardial infarction (M	Π			
			Stable or unstable angina				
			History of coronary or other arterial	revascularization			
			History of stroke				
			History of transient ischemic attack	(TIA)			
			Peripheral arterial disease (PAD)				
			ember has been established on reques ectiveness via a lowered hemoglobin	ted drug for at least 90 days <u>AND</u> has demonstrated A1C (A1C) from baseline			

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For Byetta, Bydureon BCise & liraglutide (generic Victoza®) Requests: Member has tried and failed a least 30 days of therapy with TWO (2) of the following:				
☐ Mounjaro [®]	□ Ozempic [®]			
□ Rybelsus®	□ Trulicity®			

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.