Parathyroid Hormone Analogs

Evenity (romosozumab-aqqg), Forteo (teriparatide), Tymlos (abaloparatide)

		Member and	l Medication Information		
		* ir	ndicates required field		
*N	/lem	ber ID:	*Member Name:		
*[OB:		*Weight:		
*Medication Name/Strength:			☐ Do Not Substitute. Authorizations will be processed fo the preferred Generic/Brand equivalent unless specified		
* D	irec	tions for use:			
			ider Information dicates required field		
*R	lequ	esting Provider Name:	*NPI:		
* A	ddre	2SS:	·		
*Contact Person:			*Phone #:		
*F	ax #		Email:		
			y Billed Information		
*C	Diagr	nosis Code:	field for all medically billed products *HCPCS Code:		
*[osir	g Frequency:	*HCPCS Units per dose:		
Se	ervic	ing Provider Name:	NPI:		
Se	ervic	ing Provider Address:			
Fá	acilit	y/Clinic Name:	NPI:		
Fá	acilit	y/Clinic Address:			
	F		luding: laboratory results, chart notes and/or updated t 855-828-4992 , to prevent processing delays.		
Cri	iteri	a for Approval (all of the following must b	pe met):		
	Dia	agnosis of one of the following:			
	☐ Postmenopausal osteoporosis. Chart note page #:				
		Osteoporosis due to sustained systemic glu	ucocorticoid therapy. (Forteo only). Chart note page #:		
		Osteoporosis due to primary hypogonadism in males. (Forteo only). Chart note page #:			
	Very High risk for fracture defined as:				
	☐ Intolerance to antiresorptive therapy (bisphosphonates, denosumab) OR				
	☐ Osteoporotic fracture while on antiresorptive therapy OR				
		□ Previous osteoporotic fracture OR			
		≥ 40 years old with one of the following:			
	_	☐ T-score ≤ -2.5 at the femoral neck or spir	ne year probability of major osteoporotic fracture ≥ 20% or hip		

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UTAH MEDICAID PHARMACY PRIOR AUTHORIZATION REQUEST FORM

Note:

♦ **Boxed warning:** Evenity (romosozumab-aqqg) may increase the risk of myocardial infarction, stroke, and cardiovascular death. It should not be initiated in patients who have had a myocardial infarction or stroke within the preceding year. Consider whether the benefits outweigh the risks in patients with other cardiovascular risk factors. If a patient experiences a myocardial infarction or stroke during therapy, therapy should be discontinued.

Initial Authorization: Up to 12 months for Eve Re-authorization: None, lifetime limits apply.	nity, up to 24 months for Forteo and Tymlos.	
PROVIDER CERTIFICATION I hereby certify this treatment is indicated, nece	essary and meets the guidelines for use.	
Prescriber's Signature	Date	

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