AETNA BE	TTER HEALTH®		* ae	etna [®]
Coverage	Policy/Guideline			
Name:	Eligard		Page:	1 of 4
Effective Date: 8/7/2025			Last Review Date:	7/2025
Applica	⊠Illinois	□Florida	⊠New Jersey	
Applies to:	□Maryland	□Michigan	□ Florida Kids	
	⊠Pennsylvania Kids	□Virginia	⊠Kentucky PRMD	

Intent:

The intent of this policy/guideline is to provide information to the prescribing practitioner outlining the coverage criteria for Eligard under the patient's prescription drug benefit.

Description:

The indications below including FDA-approved indications and compendial uses are considered a covered benefit provided that all the approval criteria are met and the member has no exclusions to the prescribed therapy.

FDA-Approved Indication¹

Treatment of advanced prostate cancer

Compendial Uses

- Prostate cancer²
- Androgen receptor positive salivary gland tumors^{2,8}
- Gender dysphoria (also known as transgender and gender diverse [TGD] persons)³⁻⁶

All other indications are considered experimental/investigational and not medically necessary.

Applicable Drug List:

Eligard

Policy/Guideline:

Prescriber Specialties⁷

For gender dysphoria, the medication must be prescribed by or in consultation with a provider specialized in the care of transgender youth (e.g., pediatric endocrinologist, family or internal medicine physician, obstetrician-gynecologist) that has collaborated care with a mental health provider for members less than 18 years of age.

Coverage Criteria

Prostate Cancer¹⁻²

Authorization of 12 months may be granted for treatment of prostate cancer.

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AETNA BETTER HEALTH®				
Coverage	Policy/Guideline			
Name:	Eligard		Page:	2 of 4
Effective Date: 8/7/2025			Last Review Date:	7/2025
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Gender Dysphoria³⁻⁵

Authorization of 12 months may be granted for pubertal hormonal suppression in an adolescent member when all of the following criteria are met:

- The member has a diagnosis of gender dysphoria.
- The member is able to make an informed decision to engage in treatment.
- The member has reached Tanner stage 2 of puberty or greater.
- The member's comorbid conditions are reasonably controlled.
- The member has been educated on any contraindications and side effects to therapy.
- The member has been informed of fertility preservation options.

Authorization of 12 months may be granted for gender transition when all of the following criteria are met:

- The member has a diagnosis of gender dysphoria.
- The member is able to make an informed decision to engage in treatment.
- The member will receive the requested medication concomitantly with genderaffirming hormones.
- The member's comorbid conditions are reasonably controlled.
- The member has been educated on any contraindications and side effects to therapy.
- The member has been informed of fertility preservation options.

Salivary gland tumor^{2,8}

Authorization of 12 months may be granted for treatment of recurrent, unresectable, or metastatic salivary gland tumor as a single agent or in combination with abiraterone and prednisone when the tumor is androgen receptor positive.

Continuation of Therapy

Salivary Gland Tumor

Authorization of 12 months may be granted for continued treatment of salivary gland tumor in members requesting reauthorization when there is no evidence of unacceptable toxicity or disease progression while on the current regimen.

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AETNA BETTER HEALTH®				
Coverage	Policy/Guideline			
Name:	Eligard		Page:	3 of 4
Effective Date: 8/7/2025			Last Review Date:	7/2025
Analiaa	⊠Illinois	□Florida	⊠New Jersey	
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Prostate Cancer

Authorization of 12 months may be granted for continued treatment of prostate cancer in members requesting reauthorization who are experiencing clinical benefit to therapy (e.g., serum testosterone less than 50 ng/dL) and who have not experienced an unacceptable toxicity.

Gender Dysphoria⁶

Authorization of 12 months may be granted for continued treatment for pubertal hormonal suppression in adolescent members requesting reauthorization when all of the following criteria are met:

- The member has a diagnosis of gender dysphoria.
- The member is able to make an informed decision to engage in treatment.
- The member has previously reached Tanner stage 2 of puberty or greater.
- The member's comorbid conditions are reasonably controlled.
- The member has been educated on any contraindications and side effects to therapy.
- Before the start of therapy, the member has been informed of fertility preservation options.

Authorization of 12 months may be granted for continued treatment for gender transition in members requesting reauthorization when all of the following criteria are met:

- The member has a diagnosis of gender dysphoria.
- The member is able to make an informed decision to engage in treatment.
- The member will receive the requested medication concomitantly with genderaffirming hormones.
- The member's comorbid conditions are reasonably controlled.
- The member has been educated on any contraindications and side effects to therapy.
- Before the start of therapy, the member has been informed of fertility preservation options.

Other

Per state regulatory guidelines around gender dysphoria, age restrictions may apply.

Approval Duration and Quantity Restrictions:

Approval: 12 months

AETNA DE	TTER HEALTH®		* ac	etna [®]
	Policy/Guideline			
Name:	Eligard		Page:	4 of 4
Effective Date: 8/7/2025			Last Review Date:	7/2025
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