



MEDICARE FORM

Zoladex® (goserelin acetate) Medication Precertification Request

Page 1 of 2

(All fields must be completed and legible for precertification review.)

For Illinois MMP: FAX: 1-855-320-8445 PHONE: 1-866-600-2139

For other lines of business: Please use other form.

Note: Zoladex is non-preferred. The preferred product is Eligard. Firmagon is also a preferred product.

Please indicate: Start of treatment: Start date / / Continuation of therapy, Date of last treatment / /

Precertification Requested By: Phone: Fax:

A. PATIENT INFORMATION

Form section A: Patient Information. Fields include First Name, Last Name, DOB, Address, City, State, ZIP, Home Phone, Work Phone, Cell Phone, Email, Patient Current Weight, Patient Height, Allergies.

B. INSURANCE INFORMATION

Form section B: Insurance Information. Fields include Aetna Member ID #, Group #, Insured, Medicare status, Medicaid status, and other coverage details.

C. PRESCRIBER INFORMATION

Form section C: Prescriber Information. Fields include First Name, Last Name, Address, City, State, ZIP, Phone, Fax, St Lic #, NPI #, DEA #, UPIN, Provider Email, Office Contact Name, Phone, and Specialty.

D. DISPENSING PROVIDER/ADMINISTRATION INFORMATION

Form section D: Dispensing Provider/Administration Information. Divided into Place of Administration and Dispensing Provider/Pharmacy details.

E. PRODUCT INFORMATION

Form section E: Product Information. Fields include Request is for: Zoladex (goserelin acetate) Dose: and Frequency:

F. DIAGNOSIS INFORMATION - Please indicate primary ICD code and specify any other where applicable.

Form section F: Diagnosis Information. Fields include Primary ICD Code, Secondary ICD Code, and Other ICD Code.

G. CLINICAL INFORMATION - Required clinical information must be completed in its entirety for all precertification requests.

Form section G: Clinical Information. Includes initiation requests and specific clinical criteria for Zoladex 3.6 mg requests only, such as breast cancer, chronic anovulatory uterine bleeding, and endometriosis.

Continued on next page



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Patient First Name	Patient Last Name	Patient Phone	Patient DOB
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G. CLINICAL INFORMATION (continued) – Required clinical information must be completed in its entirety for all precertification requests.

Gender dysphoria

- Yes No Is the requested medication being prescribed for pubertal suppression in an adolescent patient?
 → Yes No Is the patient undergoing gender transition?
 → Yes No Will the patient receive the requested medication concomitantly with gender affirming hormones?
 → Please indicate the Tanner Stage of puberty the patient has reached: Stage I Stage II Stage III Stage IV Stage V Unknown

Preservation of ovarian function

- Yes No Is the patient premenopausal and undergoing chemotherapy?

Prevention of recurrent menstrual related attacks in acute porphyria

- Yes No Is the requested medication being requested to prevent recurrent menstrual related attacks in acute porphyria?
 Yes No Is the requested medication being prescribed by, or in consultation with, a physician experienced in the management of porphyrias?

Prostate cancer

Note: Zoladex is non-preferred. The preferred product is Eligard. Firmagon is also a preferred product.

- Yes No Has the patient had a trial and failure, intolerance, or contraindication to Eligard?

Please explain if there are any other medical reason(s) that the patient cannot use Eligard when indicated for the patient's diagnosis?

Uterine leiomyomata (fibroids)

- Yes No Will the requested medication be given prior to surgery?

For Zoladex 10.8 mg requests only:

Breast cancer

Please indicate the patient's hormone receptor (HR) status: HR-positive HR-negative Unknown

Gender dysphoria

- Yes No Is the requested medication being prescribed for pubertal suppression in an adolescent patient?
 → Yes No Is the patient undergoing gender transition?
 → Yes No Will the patient receive the requested medication concomitantly with gender affirming hormones?
 → Please indicate the Tanner Stage of puberty the patient has reached: Stage I Stage II Stage III Stage IV Stage V Unknown

Prostate cancer

- Yes No Has the patient had an ineffective response, contraindication, or intolerance to Eligard?
 Yes No Has the patient had an ineffective response, contraindication, or intolerance to Firmagon?

For Continuation Requests (clinical documentation required for all requests):

Breast cancer

- Yes No Has the patient experienced clinical benefit while receiving the requested drug?
 Yes No Has the patient experienced an unacceptable toxicity while receiving the requested drug?

Gender dysphoria

- Yes No Is the requested medication being prescribed for pubertal suppression in an adolescent patient?
 → Yes No Is the patient undergoing gender transition?
 → Yes No Will the patient receive the requested medication concomitantly with gender affirming hormones?
 → Please indicate the Tanner Stage of puberty the patient has reached: Stage I Stage II Stage III Stage IV Stage V Unknown

Preservation of ovarian function

- Yes No Is the patient premenopausal and still undergoing chemotherapy?

Prevention of recurrent menstrual related attacks in acute porphyria

- Yes No Has the patient experienced clinical benefit while receiving the requested drug?
 Yes No Has the patient experienced an unacceptable toxicity while receiving the requested drug?

Prostate cancer

- Yes No Has the patient had prior therapy with Zoladex within the last 365 days?
 Yes No Has the patient experienced clinical benefit to therapy while receiving the requested drug (e.g., serum testosterone less than 50 ng/dl)?
 Yes No Has the patient experienced an unacceptable toxicity while receiving the requested drug?

H. ACKNOWLEDGEMENT

Request Completed By (Signature Required): _____ Date: ____/____/____

Any person who knowingly files a request for authorization of coverage of a medical procedure or service with the intent to injure, defraud or deceive any insurance company by providing materially false information or conceals material information for the purpose of misleading, commits a fraudulent insurance act, which is a crime and subjects such person to criminal and civil penalties.

The plan may request additional information or clarification, if needed, to evaluate requests.