

**Brand Name:** Casodex

**Generic:** bicalutamide

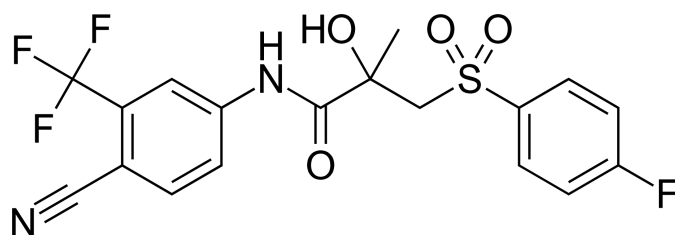
**Type:** small molecule

**Year Accepted/Phase:** 1995

### **Mechanism:**

Casodex is classified as an antiandrogen. It works by competitively inhibiting the action of androgens (male hormones), particularly testosterone, at the androgen receptor in prostate cells.

### **Chemical Structure:**



### **Indication:**

Casodex is indicated for the treatment of advanced prostate cancer, either alone (monotherapy) or in combination with other treatments like surgical castration or luteinizing hormone-releasing hormone (LHRH) analog therapy.

## **Clinical trials:**

### **Casodex 150 Study Group (Phase III)**

**Pubmed:** <https://pubmed.ncbi.nlm.nih.gov/12234503/>

**Purpose:** Evaluate the efficacy and safety of bicalutamide 150 mg as monotherapy compared to castration (surgical or medical) in patients with non-metastatic prostate cancer.

**Dates:** Conducted from 1995 to 1999.

**Results:** The trial showed that bicalutamide 150 mg provided similar overall survival rates compared to castration. However, it had a better side effect profile, with fewer instances of hot flashes and impotence.

**Impact:** This study supported the use of bicalutamide as a monotherapy option for patients with non-metastatic prostate cancer.

### **EPC (Early Prostate Cancer) Program (Phase III)**

**Pubmed:** <https://pubmed.ncbi.nlm.nih.gov/15931272/>

**Purpose:** Assess the efficacy of bicalutamide 150 mg in addition to standard care (radiotherapy or watchful waiting) in patients with localized or locally advanced prostate cancer.

**Dates:** Conducted from 1995 to 2001.

**Results:** The trials demonstrated that adding bicalutamide to standard care improved progression-free survival and reduced the risk of disease progression compared to standard care alone.

**Impact:** These findings led to the approval of bicalutamide as an adjuvant treatment in localized or locally advanced prostate cancer.

### **Casodex Combination Trial (Phase III)**

**Pubmed:** <https://pubmed.ncbi.nlm.nih.gov/9428389/>

**Purpose:** Evaluate the combination of bicalutamide 50 mg with an LHRH analog (such as leuprolide) versus castration in patients with advanced prostate cancer.

**Dates:** Conducted from 1992 to 1996.

**Results:** The combination of bicalutamide with an LHRH analog showed comparable efficacy to castration in terms of overall survival and progression-free survival. The combination also offered a better side effect profile.

**Impact:** This trial established the combination therapy of bicalutamide with an LHRH analog as an effective treatment option for advanced prostate cancer.