Exemestane

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Brand name: Aromasin®

IUPAC: (8R,9S,10R,13S,14S)-10,13-dimethyl-6-methylidene-7,8,9,11,12,14,15,16-octahydrocyclopenta[a]phenanthrene-3,17-

dione

FDA approval: Yes Manufacturer Link

Usage:

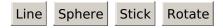
Exemestane (Aromasin®) is used in treatment of advanced breast cancer in postmenopausal women, only after their disease has progressed poorly following tamoxifen therapy. Exemestane is orally administered once daily after a meal.1

• 1Chu, E., & DeVita, V. T. (2015). Physicians' cancerchemotherapy drug manual 2015. Burlington, MA: Jones & Bartlett Learning.

Mechanism:

Exemestane (Aromasin®) binds irreversibly (permanently) to the aromatase enzyme. This stops aromatase from making estrogen by stopping the conversion of adrenal androgens, which prevents the growth of estrogen dependent cancer1

The structure below shows the breast cancer drug Aromasin, also known as Exemestane.



• 1Chu, E., & DeVita, V. T. (2015). Physicians' cancerchemotherapy drug manual 2015. Burlington, MA: Jones & Bartlett Learning.

Side effects:

Common side effects include: hot flashes, nausea, fatigue, eye disturbances, headaches dizziness, insomnia, depression, hypertension, joint pain, limb pains, back pains and osteoarthritis. There are no studies for aromasin in pregnant women yet. It is recommended only for postmenopausal women. If exposed to aromasin while pregnant, patients should contact a physician to assess the potential hazard to the fetus and possible risk of pregnancy loss. 1

• 1Chu, E., & DeVita, V. T. (2015). Physicians' cancerchemotherapy drug manual 2015. Burlington, MA: Jones & Bartlett Learning.