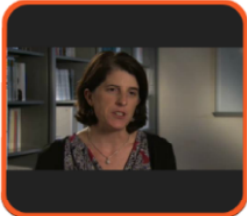



For US journalists only. For non-US journalists, please click [here](#).

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Supporting Documents

Novartis gains FDA approval for Afinitor[®] in advanced breast cancer marking a significant milestone for women battling this disease

- Approval represents the first major advance for US patients with advanced HR+ breast cancer since aromatase inhibitors were introduced more than 15 years ago¹
- In a Phase III trial, Afinitor plus exemestane more than doubled the time women lived before the cancer worsened compared to exemestane alone²
- Afinitor, the first mTOR inhibitor approved for advanced HR+ breast cancer, is given after the disease progresses following prior therapy with letrozole or anastrozole²

East Hanover, N.J., July 20, 2012 – The US Food and Drug Administration (FDA) has approved Afinitor[®] (everolimus) tablets for the treatment of postmenopausal women with advanced hormone receptor-positive, HER2-

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