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Bevacizumab: a review of its use in advanced cancer. Keating GM(1). Author information: (1)Springer, Private Bag 65901, Mairangi Bay, 0754, Auckland, New Zealand. demail@springer.com. The humanized monoclonal antibody bevacizumab (Avastin(®)) has been available in the EU since 2005.

Bevacizumab: a review of its use in advanced cancer.

Bevacizumab (Avastin) interferes with growth of the cancerous cells. It is a humanized monoclonal antibody drug which blocks angiogenesis by inhibiting Vascular Endothelial Growth Factor A (VEGF-A). Researchers have shown the advantages of this drug in metastatic breast cancer, in blindness, ovarian cancer and many more.

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This review provides an overview of the clinical experience and lessons learned since bevacizumab's initial approval, and highlights how this knowledge has led to the investigation of novel combination therapies. In the past 15 years, our understanding of VEGF's role in the tumor microenvironment has evolved.

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A review of bevacizumab in the treatment of malignant pleural mesothelioma. Brosseau S(1)(2), Assoun S(1), Naltet C(1), Steinmetz C(3), Gounant V(1), Zalcman G(1)(2). Author information: (1)Department of Thoracic Oncology & CIC 1425/CLIP2 Paris-Nord, Bichat-Claude Bernard Hospital, APHP, Paris, France.

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Drug Approval Package: Avastin (Bevacizum) NDA #125085

A recombinant humanized monoclonal antibody directed against the vascular endothelial growth factor (VEGF), a pro-angiogenic cytokine. Bevacizumab binds to VEGF and inhibits VEGF receptor binding, thereby preventing the growth and maintenance of tumor blood vessels.

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Bevacizumab is a monoclonal antibody that functions as an angiogenesis inhibitor. It works by slowing the growth of new blood vessels by inhibiting vascular endothelial growth factor A (VEGF-A),

in other words anti-VEGF therapy. Bevacizumab was approved for medical use in the United States in 2004.

Bevacizumab - Wikipedia

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Bevacizumab Biosimilar Under Review by FDA. The FDA has accepted a Biologics License Application for MYL-14020, a proposed biosimilar to bevacizumab, according to a press release from co-developers Biocon and Mylan. The BLA is seeking approval for the biosimilar as a treatment for multiple types of cancer and the FDA has set an action date goal of December 27, 2020, for a decision on the BLA.

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Samsung Bioepis has announced that the European Medicines Agency (EMA) has accepted for review its market authorization application for SB8, a proposed bevacizumab biosimilar referencing Avastin. The biosimilar was studied in a phase 3 clinical trial in 763 patients with metastatic or recurrent nonsquamous, non-small cell lung cancer (NSCLC).

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