

Genentech

A Member of the Roche Group

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January 17, 2011

Division of Dockets Management (HFA-305)
United States Food and Drug Administration
5630 Fishers Lane, Room 1061
Rockville, Maryland 20852

Subject: **License No. 1048**
Copy of Genentech Submission to Docket No. FDA-2010-N-0621
Avastin® (bevacizumab) for metastatic breast cancer

RE: Docket No. FDA-2010-N-0621

Dear Sir or Madam:

Please find enclosed a courtesy copy of Genentech's submission to Docket No. FDA-2010-N-0621, which includes data, information and analyses on which Genentech intends to rely at a hearing (requested by Genentech on December 23, 2010) to demonstrate why Avastin should remain approved for metastatic breast cancer.

The complete submission, due to the Agency by January 18, 2011, has been submitted electronically to Docket No. FDA-2010-N-0621 to <http://www.regulations.gov>.

Sincerely,



Michelle H. Rohrer, Ph.D.
Vice President
Regulatory Affairs

FDA-2010-N-0621



From: (650) 225-6463
Michelle Rohrer
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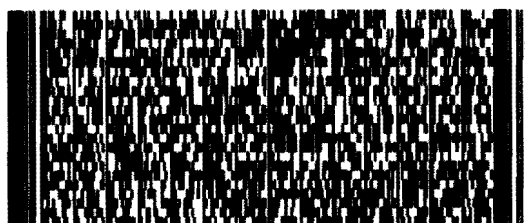
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