

Food and Drug Administration Rockville MD 20857

November 1, 2006

FILE COPY

Kip Vought Regulus Pharmaceutical Consulting, Inc. 4840 Pearl East Circle Suite 201E Boulder, Colorado 80301

Dear Mr. Vought:

Your petition requesting the Food and Drug Administration to determine whether Mivacron - EQ 2 mg base/mL (NDA 20-098), held by Abbott Laboratories, Inc. has been voluntarily withdrawn from sale for safety or efficacy reasons, was received by this office on 10/31/2006. It was assigned docket number 2006P-0445/CP 1 and it was filed on 10/31/2006. Please refer to this docket number in future correspondence on this subject with the Agency.

Please note that the acceptance of the petition for filing is a procedural matter in that it in no way reflects an agency decision on the substantive merits of the petition.

Sincerely,

/Jennie Butler, Director

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