

May 8, 2008

Dear Member:

The manufacturer Actavis Totowa, has announced its recall of the prescription heart drug Digitek® from the market. Actavis Totowa announced all Digitek® currently in pharmacies will be recalled. The FDA supports this recall. This recall is due to the chance that the tablets have twice the amount of the drug.

Our records indicate that you had prescriptions filled for Digitek® over the past 6 months. If you are still taking Digitek®, you should contact your physician. Your doctor was notified and is expecting to hear from you. There are other manufacturers of digoxin, the active ingredient in Digitek®. Your doctor may want you to change to another brand of digoxin. He or she may want you to come into the office so your digoxin blood level can be checked. You may obtain additional information on this recall from www.actavis.us.

If you have any questions, please feel free to contact Member Services at the number listed on your card.

Thank you,



Deno E. Sebastian, R.Ph., M.S.A.
Director of Pharmacy