July 27, 2000

Dear Sirs:

I would like to take this time to express a very serious concern I have ~~As of late~~, instruction from my management for promotion of products in the Wyeth-Ayerst female healthcare division has included information outside of labeling In other words, core presentations for promotion contain information that the FDA has not granted approval for use with these drugs. Specifically, the use of Premarin for the prevention/treatment of cardiovascular disease, Alzheimer's disease, and colorectal cancer. Additionally, the issue of potential side effects has been minimized such as the risk of breast cancer.

My most critical concern is the lives of the women in Alabama and Georgia that may be at risk due to these promotions. Many physicians have been persuaded to use these products for uses that have not been proven on the masses of postmenopausal women in Alabama and Georgia. I fear that their lives may be placed in serious potential danger. It is this matter that forces me to address this issue with the appropriate personnel.

Somewhere along the way, the goal of providing the most credible information for physicians and the best pharmaceuticals to them and the women of Alabama and Georgia became a mute point. It appears the only thing that matters now is increasing market share. I can no longer stand by silently and allow this to continue, for I one day will be faced with the decision of choosing to take hormones and I want to know my physician has been provided with facts not fabrication or maybes.

Very Sincerely,

Cynthia L. Waldrep
Territory Specialist
23AWD

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P.O Box 501
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2. Arti
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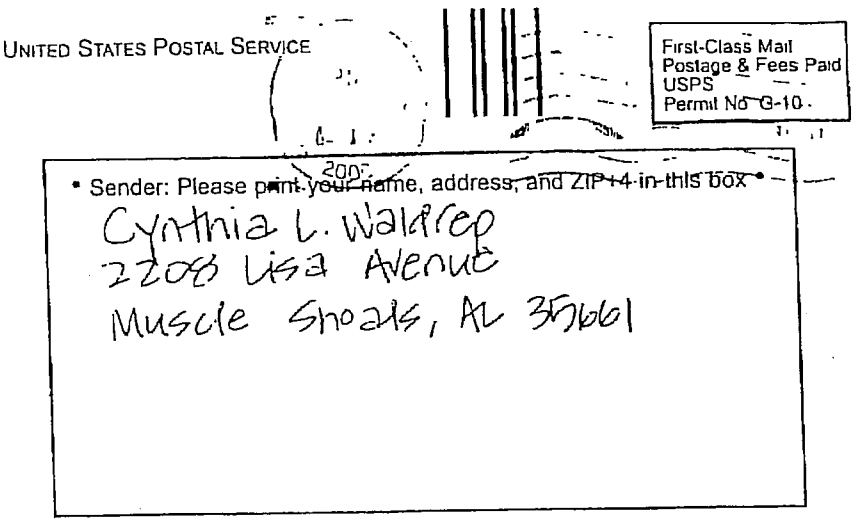
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