

Via Federal Express

August 2, 2004

Geretta Wood
Office of Device Evaluation
510(k) Document Mail Center (HFZ-401)
Center for Devices and Radiological Health
Food and Drug Administration
9200 Corporate Boulevard
Rockville, MD 20850

Reference: **K040904**: Philips HeartStart Home OTC Defibrillator

Dear Ms. Wood:

This letter is to follow-up our conversation on July 29 regarding posting of our panel presentation on FDA's website. You have our permission to post the final version of the presentation that I sent to you by email today.

Please feel free to contact me at 206-664-5290 if there are any questions.

Sincerely,

A handwritten signature in black ink that reads "Teresa Skarr". The signature is written in a cursive, flowing style.

Teresa Skarr
Manager, Regulatory and Medical Affairs