

Food and Drug Administration Rockville MD 20857

• The Honorable John D. Dingell Chairman Committee on Energy and Commerce House of Representatives Washington, D.C. 20515-2215

NOV 2 7 2007

Dear Mr. Chairman:

Thank you for your letter of August 13, 2007, co-signed by Chairman Bart Stupak, Subcommittee on Oversight and Investigations, Committee on Energy and Commerce, regarding the circumstances surrounding a Food and Drug Administration (FDA) Warning Letter to Cordis Corporation concerning their CYPHERTM Sirolimus-eluting Coronary Stent in April 2004. We are enclosing a partial response to your request that includes several Establishment Inspection Reports (EIRs) and related documents. Additional material is being compiled and will be sent as soon as possible.

Information in the enclosures includes information that is trade secret, commercial confidential or other information protected from disclosure to the public under the Freedom of Information Act (Title 5, *United States Code [U.S.C.]* 552), the Trade Secrets Act (18 U.S.C. 1905), and FDA regulations. The Committee should not publish or otherwise make public any such information. We would be glad to discuss with the Committee staff the protected status of any specific information.

Thank you again for your interest in this matter. If we can be of further assistance, please let us know. A similar letter has been sent to Chairman Stupak, without enclosures.

Sincerely,

Stephen R. Mason

Acting Assistant Commissioner

for Legislation

Enclosures