## DEPARTMENT OF HEALTH & HUMAN SERVICES



Food and Drug Administration Rockville MD 20857

September 05, 2001

Mr. Charles H. Kyper Kyper & Associates LLC 103 Nolen Lane Chapel Hill, NC 27516

Dear Mr. Kyper:

Your petition requesting the Food and Drug Administration to revoke Compliance Program 7383.003 for Class III 510(k) Pre-amendment Devices was received by this office on 09/05/01. It was assigned docket number 01P-0389/CP 1 and it was filed on 09/05/01. 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,

len K. Harri

Helen K. Harris Dockets Management Branch

01P-0389

ACKI