DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
Rockville MD 20857

FILE COPY

March 18, 2013

Robert Sayre, Ph.D.
President
Rapid Precision Testing Laboratories
8225 Rockcreek Parkway
Cordova, TN 38016

Dear Dr. Sayre:

Your petition to the Food and Drug Administration regarding sunscreen active ingredients as stated in 21 CFR Parts 201 and 310 labeling and effectiveness testing; Sunscreen Drug Products for Over-the counter Human Use, was received by this office on 3/18/2013. It was assigned docket number FDA-2013-P-0323/CP1, and it was filed on 03/18/2013. 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,

Gloria Ortega
Administrative Proceedings Officer
Division of Dockets Management
FDA/Office of the Executive Secretariat (OES)

FDA-2013-P-0323

ACK