



DEPARTMENT OF HEALTH & HUMAN SERVICES

JUN 3 2014

Food and Drug Administration  
10903 New Hampshire Ave  
Building 51  
Silver Spring, MD 20993

Robert Buchan  
496 Sunset Drive  
Asheville, NC 28804

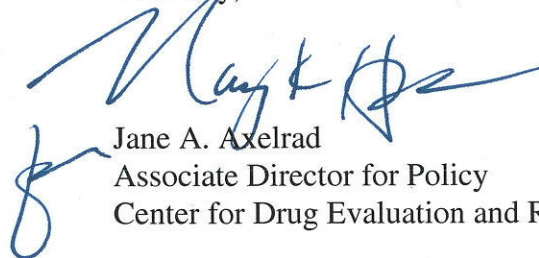
Re: Docket No. FDA-2013-P-1612/CP23

Dear Mr. Buchan:

I am writing to inform you that the Food and Drug Administration (FDA) has not yet resolved the issues raised in your citizen petition received on December 23, 2013. Your petition requests that the Agency (1) overturn the decision to rescind the special protocol assessment (SPA) agreement for the ANCHOR trial conducted with Vascepa (icosapent ethyl) capsules, (2) conduct an independent scientific review of the three outcomes trials cited by the FDA reviewing division that were asserted to reveal substantial scientific issues and were the bases for rescinding the ANCHOR SPA, (3) determine that these three outcomes trials show that patients with high triglyceride levels ( $\geq 200$  milligram/deciliter) and low high-density lipoprotein, the patient population covered by the ANCHOR SPA, had a reduction in cardiovascular disease risks, and (4) delay the PDUFA goal date for Vascepa's supplemental new drug application for the ANCHOR indication (NDA 202057/s005).

FDA has been unable to reach a decision on your petition because it raises complex issues requiring extensive review and analysis by Agency officials. This interim response is provided in accordance with FDA regulations on citizen petitions (21 CFR 10.30(e)(2)). We will respond to your petition as soon as we have reached a decision on your request.

Sincerely,



Jane A. Axelrad  
Associate Director for Policy  
Center for Drug Evaluation and Research