

**Letter of Request for Post-Approval Change/s  
through Collaborative Review Procedure**

<NAME OF HEAD OF AGENCY>  
<Designation of Head of Agency>  
Food and Drug Administration  
Civic Drive, Filinvest Corporate City  
Alabang, Muntinlupa City

<b>DTN</b>	
Date	

Attention:      Licensing and Registration Division  
                         Center for Drug Regulation and Research

Sir/Madam,

We would like to submit our application for Post-approval Change/s through the Collaborative Review Procedure for the following product:

Product Name/Strength and Form	CPR Validity/Drug Registration Number	Current	Proposed Change/s	Classification/ Specific Type of PAC/s
				e.g. AVG MaV-1

For your approval.

Very truly yours,

*Company representative name and signature*  
*Position*