



**Human Research Protection Program  
Committee on Human Research**

**Notification of Expedited Review Approval**

Principal Investigator

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**Type of Submission:** Submission Correction for Modification Form  
**Study Title:** Platelet-Oriented Inhibition in New TIA and minor ischemic stroke (POINT)

**IRB #:** 11-06356  
**Reference #:** 075474

**Committee of Record:** San Francisco General Hospital Panel

**Study Risk Assignment:** Greater than minimal

**Approval Date:** 10/28/2013                      **Expiration Date:** 06/24/2014

**All changes to a study must receive CHR approval before they are implemented.** Follow the [modification request](#) instructions. The only exception to the requirement for prior CHR review and approval is when the changes are necessary to eliminate apparent immediate hazards to the subject (45 CFR 46.103.b.4, 21 CFR 56.108.a). In such cases, report the actions taken by following these [instructions](#).

**Expiration Notice:** The iRIS system will generate an email notification eight weeks prior to the expiration of this study's approval. However, it is your responsibility to ensure that an application for [continuing review](#) approval has been submitted by the required time. In addition, you are required to submit a [study closeout report](#) at the completion of the project.

**Approved Documents:** To obtain a list of documents that were approved with this submission, follow these steps: Go to My Studies and open the study – Click on Submissions History – Go to Completed Submissions – Locate this submission and click on the Details button to view a list of submitted documents and their outcomes.

For a list of all currently approved documents, follow these steps: Go to My Studies and open the study – Click on Informed Consent to obtain a list of approved consent documents and Other Study Documents for a list of other approved documents.

**San Francisco Veterans Affairs Medical Center (SFVAMC):** If the SFVAMC is engaged in this research, you must secure approval of the VA Research & Development Committee in addition to CHR approval and follow all applicable VA and other federal requirements. The CHR [website](#) has more information.