



Icahn School of Medicine at Mount Sinai  
Mount Sinai Beth Israel  
Mount Sinai Brooklyn  
The Mount Sinai Hospital  
Mount Sinai Queens  
New York Eye and Ear Infirmary  
of Mount Sinai  
Mount Sinai St. Luke's  
Mount Sinai West

**Program for the Protection  
of Human Subjects**  
*Institutional Review Boards*  
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## **APPROVAL OF RESEARCH**

Date: 8/9/2018

To: **Richard Whelan, MD (Richard.Whelehan@mountsinai.org)**

On **8/7/2018**, an Institutional Review Board of the Mount Sinai School of Medicine, in accordance with Mount Sinai's Federal Wide Assurances (FWA#00005656, FWA#00005651) to the Department of Health and Human Services approved the following human subject research from **8/7/2018** until **8/6/2019** inclusive:

Type of Review:	<b>Continuing Request for Approval</b>
Project Title:	<b>Tumor, perioperative blood sample, and clinical data banking for cancer, physiology, immunology, and other research studies</b>
Investigator:	<b>Richard Whelan, MD (Dept: SU - Surgery)</b>
Project Information:	<b>HS#: 15-00817 GCO#1: 16-2619(0001) Icahn School of Medicine at Mount Sinai</b>
Sites:	<b>Mount Sinai West</b>
IND or IDE (if any):	<b>No INDs;No IDEs;</b>
Submission Details (if any):	<b>Personnel changes: Karan Jatwani was added to study team on 6/18/18. Carl Wincler was removed from study personnel.</b>

Between **6/20/2019** and **6/25/2019**, or within 30 days prior to study close, whichever is earlier, you are to submit a completed FORM HRP-212: Continuing/Final Review Progress Report and required attachments, in order to request continuing IRB approval or study closure. If IRB continuing review approval is not granted before the expiration date of **8/6/2019**, IRB approval of this research expires on that date.

□□□□□□ The IRB has determined that this research involves GREATER THAN minimal risk. Minimal risk means that the probability and magnitude of harm or discomfort anticipated in the research are not greater in and of themselves than those ordinarily encountered in daily life or during the performance of routine physical or psychological examinations or tests (45CFR.46.102; 21CFR50.3k).

In conducting this research you are required to follow the requirements listed in the **Investigator Manual**. If stamped approved consent forms are attached, use copies of these forms to document consent. IRB approval does not constitute or imply institutional support for the conduct of this research. Additionally, all required local committee approvals at each **research affiliate** site must be obtained prior to initiation.

cc: Study Contact(s): Xiaohong (Peter) Yan (peter.yan@mountsinai.org)

**Yonnette Joseph**

IRB Analyst I

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