

**U.S. NUCLEAR REGULATORY COMMISSION****Date:** 8/20/09**TELEPHONE CONVERSATION RECORD****Time:** 2:40pm**Mail Control  
or Report No(s).**

143249

**License No(s).**

29-06759-001

**Docket No(s).**

030002490

**Name of Licensee:** St Michael's Medical Center**Name of Participant(s):** Raja Subramanian (physicist)**Telephone No.** 718-419-8046**Subject:** Removal of material authorizations**(NOTE: This will be used as the  
Documents Title in ADAMS)**

The licensee is requesting to remove P-32, Sr-90, and 31.11 (in vitro studies) from their license. I informed Mr. Subramanian that I reviewed the inspection reports from 2003 and 2005 indicated that P-32 IVB was conducted in the past, however, it has not been conducted in over 3 years. Therefore, any remaining P-32 would have decayed. A review of inspection records also indicated that the Sr-90 IVB device was never possessed.

As for 31.11 (in vitro studies), the licensee stated that I-125, I-131, and Co-57 was used in the past. Co-57 has a half-life of greater than 120 days, so an Environmental Assessment/FRN would be necessary and could not be completed prior to the turnover. I informed the licensee that they should prepare their historical assessment and surveys for submission to NJ in the future to support removal.

**Action Required:****Document Availability:****Publicly Available****Non-Publicly Available****Non-Sensitive****Non-Sensitive Copyright****Sensitive- Proprietary** **Sensitive – Privacy Act (includes PII)****Sensitive – Internal****Sensitive – Security-Related** **Immediate Release****Normal Release Date:****Delayed Release Date:****SUNSI Review Completed By:** Michelle Simmons / RA /**Document Accession #:** \_\_\_\_\_