NRC FORM 374

U.S. NUCLEAR REGULATORY COMMISSION

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MATERIALS LICENSE

Pursuant to the Atomic Energy Act of 1954, as amended, the Energy Reorganization Act of 1974 (Public Law 93-438), and Title 10, Code of Federal Regulations, Chapter I, Parts 30, 31, 32, 33, 34, 35, 36, 37, 39, 40, 70 and 71, and in reliance on statements and representations heretofore made by the licensee, a license is hereby issued authorizing the licensee to receive, acquire, possess, and transfer byproduct, source, and special nuclear material designated below; to use such material for the purpose(s) and at the place(s) designated below; to deliver or transfer such material to persons authorized to receive it in accordance with the regulations of the applicable Part(s). This license shall be deemed to contain the conditions specified in Section 183 of the Atomic Energy Act of 1954, as amended, and is subject to all applicable rules, regulations, and orders of the Nuclear Regulatory Commission now or hereafter in effect and to any conditions specified below.

1	Licensee 1. Straub Clinic & Hospital			In accordance with email dated April 12, 2018.		4. Expiration Date: October 31, 2025	
	Department of Nuclear M	ledicir	ne JCI	EAR REGULA		5. Docket No.: 030-14529	
2.	888 South King Street Honolulu, HI 96813		SNC	 License nur amended in follows: 	mber: 53-18126-01 is a its entirety to read as	Refe	rence No.:
6.	Byproduct, source, and/or special nuclear material	7.	Chemical and/or physical fo	orm	Maximum amount that licer may possess at any one tin under this license		Authorized use
A.	Any byproduct material permitted by 10 CFR 35.100	A.	Any S	A	As Needed	A.	For use in uptake, dilution and excretion studies permitted by 10 CFR 35.100.
B.	Any byproduct material permitted by 10 CFR 35.200	В.	Any File	B.	As Needed	В.	For use in imaging and localization studies permitted by 10 CFR 35.200.
C.	Any byproduct material permitted by 10 CFR 35.300	C.	Any	C. ★★★	1 curie total	C.	For any use permitted by 10 CFR 35.300.
D.	Any byproduct material permitted by 10 CFR 31.11	D.	Prepackaged Kits	D.	5 millicuries total	D.	For use in in-vitro studies.
E.	Yttrium-90		Liquid (IDEC Pharmaceu Zevalin,)	iticals E.	Not to exceed 2 sources 110 millicuries total activ		To be used for dose calibration checks and calibrations and quality assurance testing.

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	MATERIALS LICENSE	License Number 53-18126-01	Docket or Reference Number 030-14529				
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		(Corrected Copy)					
	CONDITIONS						
10.	0. Licensed material may be used or stored only at the licensee's facilities located at 888 South King Street, Honolulu, Hawaii (island of Oahu).						
11.	1. The Radiation Safety Officer for this license is Ronald W. Frick, M.S., CHP, DABR.						
12.	Licensed material shall only be used by, or under the supervision of:						
	A. Individuals permitted to work as authorized users, authorized fluctear pharmacists, and/or authorized medical physicists, in accordance with 10 CFR 35.13 and 10 CFR 35.14.						
	B. The following individuals are authorized	B. The following individuals are authorized users for the material and medical uses as indicated:					
	Authorized User(M.D.,D.O.,etc.)	Authorized User(M.D.,D.O.,etc.) Material and Use					
	Marc Coel, M.D.	135.100; 35.200; 35.300; 31.11	S				
	Brandon Itagaki, M.D.	35.100; 35.200					
	Shay J. Lee, M.D.	35.100; 35.200; 35.300	40				
	Douglas A. Prager, M.D.	35.100; 35.200; 35.300					
	Brian T. Sinclair, M.D.	35.100; 35.200; 31.11					
	Kristi Takaki, M.D.	55.100, 55.200	· · · · · ·				
13.	Sealed sources containing licensed mater specifically authorized.	rial shall not be opened or sources remo	oved from source holders by the licensee, except as				

14. In addition to the possession limits in Item 8, the licensee shall further restrict the possession of licensed material to quantities below the limits specified in 10 CFR 30.72 which require consideration of the need for an emergency plan for responding to a release of licensed material.

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15.	15. Except as specifically provided otherwise in this license, the licensee shall conduct its program in accordance with the statements, representations, and procedures contained in the documents, including any enclosures, listed below. This license condition applies only to those procedures that are required to be submitted to accordance with the regulations. Additionally, this license condition does not limit the licensee's ability to make changes to the radiation protection program as provided for in 10 CFR 35.26. The U.S. Nuclear Regulatory Commission's regulations shall govern unless the statements, representations, and procedures in the licensee's application and correspondence are more restrictive than the regulations. A. Letter with enclosures dated November 27, 2012 (ML12352A137) B. Application dated April 6, 2015 (ML15110A473) C. Letter dated October 8, 2015 (ML15208A424)							
Date	e: July 23, 2018	By:	THE U.S. NUCLEAR REGULATORY COMMISSION					