U.S. NUCLEAR REGULATORY COMMISSION

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.

	Licens	see			with letter dated	4. Expir	ation Date: November 30, 2036
1.	Alliance HealthCare Servi	ices,		October 05, 20	EGULA	5 Dock	et No.: 030-35774
2.	18201 Von Karman Ave. Ste. 600 Irvine, CA 92612		ES AC		: 47-25570-01 is its entirety to read as		rence No.:
6.	Byproduct, source, and/or special nuclear material	7.	Chemical and/or physical fo	8.	Maximum amount that lice may possess at any one til under this license		Authorized use
A.	Any byproduct material permitted by 10 CFR 35.200	A.	Any	A.	As Needed	A .	For use in imaging and localization studies permitted by 10 CFR 35.200.
B.	Strontium-82	B.	Any Except Sealed Source	ces B.	200 millicuries total	B.	For decay in storage only in accordance with 10 CFR 35.92.
C.	Strontium-85	C.	Any Except Sealed Source	ces C.	1 curie total	C.	For decay in storage only in accordance with 10 CFR 35.92.
D.	Germanium-68	D.	Sealed Sources	→ D.	50 millicuries total	D.	For storage only, limited to one year, incident to transfer in accordance with 10 CFR 30.41.

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	CONDITIONS	3	
of the licensee anywhere in the U	A. Licensed material listed in Subitem No. 6.A. incident to mobile nuclear medicine activities shall be used or stored at temporary job sites of the licensee anywhere in the United States where the U.S. Nuclear Regulatory Commission maintains jurisdiction for regulating the		

If the jurisdiction status of a Federal facility within an Agreement State is unknown, the licensee should contact the Federal agency controlling the job site in question to determine whether the proposed job site is an area of exclusive Federal jurisdiction. Authorization for use of radioactive materials at job sites in Agreement States not under exclusive Federal jurisdiction shall be obtained from the appropriate state regulatory agency.

- B. Licensed material listed in Subitem Nos. 6.B and 6.C., limited to strontium waste generated incident to mobile nuclear medicine activities, shall be stored at the licensee's facilities located at 525 S Gould St., Owosso, Michigan, for decay in storage in accordance with 10 CFR 35.92.
- C. Licensed material listed in Subitem No. 6.D., limited to germanium-68 sealed sources transferred from the licensee's mobile nuclear medicine vans, shall be stored at the licensee's facilities located at 525 S. Gould Street, Owosso, Michigan, for up to one year pending transfer to an authorized recipient in accordance with 10 CFR 30.41.
- 11. The Radiation Safety Officer for this license is Kay Kassel, M.S., C.N.M.T.
- 12. Licensed material shall only be used by, or under the supervision of:
 - A. Individuals permitted to work as authorized users in accordance with 10 CFR 35.13 and 10 CFR 35.14.
 - B. The following individuals are authorized users for the material and medical uses as indicated:

Authorized User (M.D.,D.O.,etc.) Material and Use
David Abramowitz, M.D. 10 CFR 35.200

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Authorized User (M.D.,D.O.,etc.)	Material and Use	,	
Jennifer Jayne Adams, M.D.	10 CFR 35.200		
Irfan Ahmad, M.D.	10 CFR 35.200 R REG	11,	
Afzal Ahmed, M.D.	10 CFR 35.200		
Mark J. Akers, M.D.	10 CFR 35.200	ULAX	
Paul D. Akers, M.D.	10 CFR 35.200	No.	
Syed I. Ali, M.D.	10 CFR 35.200	1	
Rajaa M. Almestady, M.D.	44 10 CFR 35.200		
Daniel Altman, M.D.	10 CFR 35.200		
Marsha Anderson, M.D.	10 CFR 35.200		
Mark C. Arvin, M.D.	10 CFR 35.200	SS/WM/SS/	
Akhtar Ashraf, M.D.	10 CFR 35.200		
James Baek, M.D.	10 CFR 35.200		
Indraneel Banerji, M.D.	10 CFR 35.200		
David Bauer, M.D.	10 CFR 35.200		
James K. Benjamin, M.D.	10 CFR 35.200	40	
James R. Bergh, M.D.	10 CFR 35.200	*	
Richard R. Black, D.O.	10 CFR 35.200	V	
Rodger Blake, M.D.	10 CFR 35.200		
Paul Henry Blom, M.D.	10 CFR 35.200		
Suzanne Bosman, M.D.	10 CFR 35.200		
Joel A. Brake, M.D.	10 CFR 35.200		
James M. Browne, M.D.	10 CFR 35.200		
Douglas A. Bruns, D.O.	10 CFR 35.200		
David J. Burkart, M.D.	10 CFR 35.200		

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Authorized User (M.D.,D.O.,etc.)	Material and Use		
Jeffery S. Cahoon, M.D.	10 CFR 35.200		
James Paul Carl, M.D.	10 CFR 35.200 R REG	11,	
Raj Rocky Chinnappan, M.D.	10 CFR 35.200	LA.	
Peter Chirico, M.D.	10 CFR 35.200	ULAZ	
Corey W. Chopra, M.D.	10 CFR 35.200		
Jesse A. Cole, M.D.	10 CFR 35.200	1	
Ricky J. Compton, M.D.	4 10 CFR 35.200		
John Phillip Cox, D.O.	10 CFR 35.200		
Robert J. Cure, M.D.	10 CFR 35.200		
Anthony R. D'Amico, M.D.	10 CFR 35.200	Ald S	
Ryan Daily, M.D.	10 CFR 35.200		
Kyle L. Dale, M.D.	10 CFR 35.200	S	
Daniel J. Daunhauer, M.D.	10 CFR 35.200		
Sarsfield Patrick Dougherty, M.D.	10 CFR 35.200		
Hans G. Dransfeld, M.D.	10 CFR 35.200	N	
Joseph W. Dransfeld, M.D.	10 CFR 35.200	**	
Nathaniel D. Dueker, M.D.	10 CFR 35.200		
Douglas M. Dunco, M.D.	10 CFR 35.200		
Rodney A. Dunseath, D.O.	10 CFR 35.200		
George M. Dwyer, M.D.	10 CFR 35.200		
Paul H. Eikens, M.D.	10 CFR 35.200		
Mark W. Elliott, M.D.	10 CFR 35.200		
Susannah G. Ellsworth, M.D	10 CFR 35.200		
Hilary Ann Evans, M.D.	10 CFR 35.200		

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Authorized User (M.D.,D.O.,etc.)	Material and Use		
Thomas H. Farquhar, M.D., Ph.D.	10 CFR 35.200		
Angelo Steven Ferraro, M.D.	10 CFR 35.200 R REG	11,	
Luke Gerges, D.O.	10 CFR 35.200		
Joshua Dale Gibson, M.D.	10 CFR 35.200	ULAZ	
Kendall Goldschmidt, M.D.	10 CFR 35.200		
Allison A. Griffiths, M.D.	10 CFR 35.200	1	
Lee Corey Haikal, M.D.	4 10 CFR 35.200	188	
Nathan R. Hatfield, M.D.	10 CFR 35.200		
David Damion Hazlett, Jr., M.D.	10 CFR 35.200		
Gregory Alan Henkle, M.D.	10 CFR 35.200		
Jeffrey A. Hicklin, M.D.	10 CFR 35.200		
Robert Hills, D.O.	10 CFR 35.200		
Leszek J. Jaszczak, M.D.	10 CFR 35.200	S	
Ronald D. Jenkins, M.D.	10 CFR 35.200		
John Kalabat, M.D.	10 CFR 35.200	N	
Craig S. Kamen, M.D.	10 CFR 35.200	**	
Prasanta K. Karak, M.D.	10 CFR 35.200	V	
Kevin Matthew Kavanaugh, M.D.	10 CFR 35.200		
Jeffrey Kaye, M.D.	10 CFR 35.200		
Imran Kazem, M.D.	10 CFR 35.200		
Stephen Joowhan Kim, M.D.	10 CFR 35.200		
Philip Kohanski, M.D.	10 CFR 35.200		
Kenneth L. Koontz, M.D.	10 CFR 35.200		
Michael V. Korona, Jr., M.D.	10 CFR 35.200		

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Authorized User (M.D.,D.O.,etc.)	Material and Use		
Gary W. Kravetz, M.D.	10 CFR 35.200		
Adam Thomas Krompecher, M.D.	10 CFR 35,200 R REG	(1)	
Herbert S. Lambert, M.D.	10 CFR 35.200		
Francisco J. Lammoglia, M.D.	10 CFR 35.200	ULAZ	
Christopher J. Leary, M.D.	10 CFR 35.200		
Terry S. Lee, M.D.	10 CFR 35.200	1	
Eric L. Leonard, M.D.	4 10 CFR 35.200		
Donald Lewis, M.D.	10 CFR 35.200		
Edward J. Maas, M.D.	10 CFR 35.200		
Colleen M. Madden, M.D.	10 CFR 35.200	MM/SS/M/M	
Mrinal Mali, M.D.	10 CFR 35.200		
Michael J. Malnofski, M.D.	10 CFR 35.200		
A. Jane MaLoof, M.D.	10 CFR 35.200	S	
Jack D. Markiewicz, M.D.	10 CFR 35.200		
Phyllis Martin-Simmerman, M.D.	10 CFR 35.200	N	
William Mason, M.D.	10 CFR 35:200	× ·	
Matthew E. Maxwell, M.D.	10 CFR 35.200	Y	
Marco S. Mazzella, M.D.	10 CFR 35.200		
Elvin McCarl, M.D.	10 CFR 35.200		
Richard D. Miller, M.D.	10 CFR 35.200		
Steve Min, D.O.	10 CFR 35.200		
Virginia Molleran, M.D.	10 CFR 35.200		
Craig Moore, M.D.	10 CFR 35.200		
Joshua A. Nepute, M.D.	10 CFR 35.200		

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Authorized User (M.D.,D.O.,etc.)	Material and Use		
Josef R.S. Noga, M.D.	10 CFR 35.200		
Kevin O'Brien, M.D.	10 CFR 35.200 R REG	11,	
Patrick M. O'Toole, M.D.	10 CFR 35.200		
Ademola Michael Obajuluwa, M.D	. 10 CFR 35.200	ULAX	
Dana Olson, M.D.	10 CFR 35.200		
Robert Oostveen, M.D.	10 CFR 35.200	1	
Samir Parikh, M.D.	4 10 CFR 35.200	133	
Bharat Patel, M.D.	10 CFR 35.200		
Grant D. Petty, M.D.	10 CFR 35.200		
Krishna R. Pillai, M.D.	10 CFR 35.200		
Syam P. Reddy, M.D.	10 CFR 35.200		
James Milton Reynolds, M.D.	10 CFR 35.200	Co.	
Reuben Rock, M.D.	10 CFR 35.200		
Daniel Adam Rodgers, M.D.	10 CFR 35.200		
Colin Rose, M.D.	10 CFR 35.200	N	
Heather Rose, M.D.	10 CFR 35,200		
Ronald J. Rosenberg, M.D.	10 CFR 35.200	V -	
Aldo Ruffolo, M.D.	10 CFR 35.200		
Paul Sanchirico, M.D.	10 CFR 35.200		
Mark Shaman, M.D.	10 CFR 35.200		
Paul W. Sheets, M.D.	10 CFR 35.200		
Steven K. Shekut, M.D.	10 CFR 35.200		
Charles W. Siegler, M.D.	10 CFR 35.200		
Justin Sims, M.D.	10 CFR 35.200		

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Authorized User (M.D.,D.O.,etc.)	Material and Use		
Stacy L. Spooner, M.D.	10 CFR 35.200		
Gregory R. Spurling, M.D.	10 CFR 35.200 R REG	11,	
Michael A. Stewart, M.D.	10 CFR 35.200		
LeAnn Stidham, M.D.	10 CFR 35.200	ULAX	
Nathan M. Strabala, M.D.	10 CFR 35.200		
Edward Strauss, M.D.	10 CFR 35.200	1	
Victoria A. Swegles, D.O.	44 10 CFR 35.200		
Sanjay J. Talati, M.D.	10 CFR 35.200		
Shrey K. Thawait, M.D.	10 CFR 35.200		
Smari Thordarson, M.D.	10 CFR 35.200	SS/WW	
Walter Parke Thrush, M.D.	10 CFR 35.200		
Paul E. Timperman, M.D.	10 CFR 35.200		
Gregory T. Turner, M.D.	10 CFR 35.200	S	
Torin P. Walters, M.D.	10 CFR 35.200		
James K. Watson, M.D.	10 CFR 35.200	4	
Ronald R. Weis, M.D.	10 CFR 35.200	× ·	
Jonathan W. Weiss, M.D.	10 CFR 35.200	Y	
Michael Whisenant, M.D.	10 CFR 35.200		
Thomas T. Win, M.D.	10 CFR 35.200		
Milton R. Wolf, M.D.	10 CFR 35.200		
Ehab H. Youssef, M.D.	10 CFR 35.200		
John S. Yungmeyer, M.D.	10 CFR 35.200		
Roy W. "Chip" Zimmer, III, M.D.	10 CFR 35.200		

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C. The following individuals are authorized users for nonmedical uses as indicated:

Non-Medical Use

Material and Use

Kay Kassel, M.S., C.N.M.T.

Strontium-82/85 for decay-in-storage; Germanium-68 sealed sources for storage only

- 13. The licensee shall conduct a physical inventory every 6 months, or at other intervals approved by the U.S. Nuclear Regulatory Commission, to account for all sealed sources and/or devices received and possessed under the license. Records of inventories shall be maintained for 3 years from the date of each inventory, and shall include the radionuclides, quantities, manufacturer's name and model numbers, and the date of the inventory.
- 14. Sealed sources containing licensed material shall not be opened by the licensee, except as specifically authorized.
- 15. A. Sealed sources shall be tested for leakage and/or contamination at intervals not to exceed the intervals specified in the certificate of registration issued by the U.S. Nuclear Regulatory Commission under 10 CFR 32.210 or by an Agreement State. In the absence of a registration certificate, sealed sources shall be tested for leakage and/or contamination at intervals not to exceed 6 months, or at such other intervals as specified.
 - B. In the absence of a certificate from a transferor indicating that a leak test has been made within the intervals specified in the certificate of registration issued by the U.S. Nuclear Regulatory Commission under 10 CFR 32.210 or by an Agreement State, prior to the transfer, a sealed source received from another person shall not be put into use until tested and the test results received.
 - C. Sealed sources need not be tested if they contain only hydrogen-3; or they contain only a radioactive gas; or the half-life of the isotope is 30 days or less; or they contain not more than 100 microcuries of beta- and/or gamma-emitting material or not more than 10 microcuries of alpha-emitting material.
 - D. Sealed sources need not be tested if they are in storage and are not being used. However, when they are removed from storage for use or transferred to another person, and have not been tested within the required leak test interval, they shall be tested before use or transfer. No sealed source shall be stored for a period of more than 10 years without being tested for leakage and/or contamination.

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- E. The leak test shall be capable of detecting the presence of 185 becquerels (0.005 microcuries) of radioactive material on the test sample. If the test reveals the presence of 185 becquerels (0.005 microcuries) or more of removable contamination, a report shall be filed with the U.S. Nuclear Regulatory Commission in accordance with 10 CFR 30.50(c)(2), and the source shall be removed immediately from service and decontaminated, repaired, or disposed of in accordance with Commission regulations.
- F. Tests for leakage and/or contamination, including leak test sample collection and analysis, shall be performed by the licensee or other persons specifically licensed by the U.S. Nuclear Regulatory Commission or an Agreement State to perform such services.
- G. Records of leak test results shall be kept in units of becquerels (microcuries) and shall be maintained for 3 years.



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representations, and procedures contain those statements, representations, and plicense condition does not limit the license. The U.S. Nuclear Regulatory Commission licensee's application and corresponden regulations.	ed in the documents, including any enclorocedures that are required to be submisee's ability to make changes to the radia on's regulations shall govern unless the size impose on the licensee requirements of Authority dated June 3, 2021, excluding 1299A298)	uct its program in accordance with the statements, closures, listed below. This license condition applies only to nitted in accordance with the regulations. Additionally, this liation protection program as provided for in 10 CFR 35.26. statements, representations, and procedures in the sthat are more restrictive than or in addition to the
	FOR	R THE U.S. NUCLEAR REGULATORY COMMISSION
Date: October 20, 2022		Jason M. Kelly, MPH, Health Physicist Region III