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.

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Licensee 1. Alliance HealthCare Services, Inc.		In accordance with letter dated April 29, 2020.	4. Expiration Date: September 30, 2021	
2. 330 Harper Park Drive, Beckley, WV 25801	Suite C	3. License number: 47-25570-01 is amended in its entirety to read as follows:	5. Docket No.: 030-35774 Reference No.:	
Byproduct, source, and/or special nuclear material	7. Chemical and/or physical fo	Maximum amount that licer may possess at any one tir under this license		
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 Sour	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 Sour	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.	

NRC FORM 374A	C FORM 374A U.S. NUCLEAR REGULATORY COMMISSION		PAGE 2 OF 10 PAGES
MATERIALS LICENSE	License Number 47-25570-01	Docket or Reference Number 030-35774	
SUPPLEMENTARY SHEET	Amendment No. 24		

- 10. A. Licensed material listed in Subitem No. 6.A. incident to mobile nuclear medicine activities may 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 use of licensed material, including areas of exclusive Federal jurisdiction within Agreement States.
 - 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, may be stored at the licensee's facilities located at 525 S. Gould Street, Owasso, 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, may be stored at the licensee's facilities located at 525 S. Gould Street, Owasso, Michigan, for up to one year pending transfer to an authorized recipient in accordance with 10 CFR 30.41.
- 11. Licensed material shall only be used by, or under the supervision of:
 - A. Individuals permitted to work as an authorized user 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
Irfan Ahmad, M.D.	10 CFR 35.200
Afzal Ahmed, M.D.	10 CFR 35.200
Mark J. Akers, M.D.	10 CFR 35.200
Paul D. Akers, M.D.	10 CFR 35.200
Paul J. Alfieri, M.D.	10 CFR 35.200

FORM 374A	U.S. NUCLEAR REGULATO	ORY COMMISSION	PAGE 3 OF 10 PAGES
MATERIALS LICENSE	License Number 47-25570-01	Docket or Reference Number 030-35774	
SUPPLEMENTARY SHEET	Amendment No. 24		
Authorized User(M.D.,D.O.,etc.)	Material and Use		
Syed I. Ali, M.D.	10 CFR 35.200		
Daniel Altman, M.D.	10 CFR 35.200 R REC	311,	
Alok Azad Anand, M.D.	10 CFR 35.200	LA.	
Marsha Anderson, M.D.	10 CFR 35.200	BULAZ	
lbad U. Ansari, M.D.	10 CFR 35.200		
James Baek, M.D.	10 CFR 35.200	1	
Indraneel Banerji, M.D.	4 10 CFR 35.200		
Marc R. Beck, M.D.	10 CFR 35.200		
Martin Black, M.D.	10 CFR 35.200		
Richard R. Black, D.O.	10 CFR 35.200		
Rodger Blake, M.D.	10 CFR 35.200		
Paul H. Blom, M.D.	10 CFR 35.200		
Robert L. Bridges, M.D.	10 CFR 35.200	MMISSIMM	
James M. Browne, M.D.	10 CFR 35.200		
Douglas A. Bruns, D.O.	10 CFR 35.200	N	
James Paul Carl, M.D.	10 CFR 35,200		
Christopher Carrel, M.D.	10 CFR 35.200		
Peter Chirico, M.D.	10 CFR 35.200		
Jesse A. Cole, M.D.	10 CFR 35.200		
Ricky J. Compton, M.D.	10 CFR 35.200		
John Phillip Cox, D.O.	10 CFR 35.200		
Robert J. Cure, M.D.	10 CFR 35.200		
Michael T. Czuba, M.D.	10 CFR 35.200		
Ryan Daily, M.D.	10 CFR 35.200		

MATERIALS LICENSE SUPPLEMENTARY SHEET License Number 47-25570-01	PAGE 4 OF 10 PAGES
Authorized User(M.D.,D.O.,etc.) Corinne Daurdulian, M.D. Vu Quoc Do, M.D. Hans G. Dransfeld, M.D. Joseph Dransfeld, M.D. Nathaniel D. Dueker, M.D. Adwoa Essel, M.D. Joshua Dale Gibson, M.D. Kendall Goldschmidt, M.D. Amendment No. 24 Material and Use 10 CFR 35.200	
Corinne Daurdulian, M.D. Vu Quoc Do, M.D. Hans G. Dransfeld, M.D. Joseph Dransfeld, M.D. Nathaniel D. Dueker, M.D. Adwoa Essel, M.D. Joshua Dale Gibson, M.D. Kendall Goldschmidt, M.D. 10 CFR 35.200	
Nathaniel D. Dueker, M.D. Adwoa Essel, M.D. Joshua Dale Gibson, M.D. Kendall Goldschmidt, M.D. 10 CFR 35.200 10 CFR 35.200 10 CFR 35.200	
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Nathaniel D. Dueker, M.D. Adwoa Essel, M.D. Joshua Dale Gibson, M.D. Kendall Goldschmidt, M.D. 10 CFR 35.200 10 CFR 35.200 10 CFR 35.200	
Nathaniel D. Dueker, M.D. Adwoa Essel, M.D. Joshua Dale Gibson, M.D. Kendall Goldschmidt, M.D. 10 CFR 35.200 10 CFR 35.200 10 CFR 35.200	
Nathaniel D. Dueker, M.D. Adwoa Essel, M.D. Joshua Dale Gibson, M.D. Kendall Goldschmidt, M.D. 10 CFR 35.200 10 CFR 35.200 10 CFR 35.200	
Joshua Dale Gibson, M.D. Kendall Goldschmidt, M.D. 10 CFR 35.200 10 CFR 35.200	
Kendall Goldschmidt, M.D. 10 CFR 35.200	
Lee Corey Haikal, M.D. 10 CFR 35.200	
Nathan R. Hatfield, M.D. 10 CFR 35.200	
David Damion Hazlett, M.D. 10 CFR 35.200	
Mark R. Heitzman, M.D. 10 CFR 35.200	
Robert Hills, D.O. 10 CFR 35.200	
Craig S. Kamen, M.D. 10 CFR 35.200	
Prasanta K. Karak, M.D. 10 CFR 35.200	
Kastytis C. Karvelis, M.D. 10 CFR 35.200	
Stephen Joowhan Kim, M.D. 10 CFR 35.200	
Philip Kohanski, M.D. 10 CFR 35.200	
Michael V. Korona, Jr., M.D. 10 CFR 35.200	
Gary W. Kravetz, M.D. 10 CFR 35.200	
Adam Thomas Krompecher, M.D. 10 CFR 35.200	
Benjamin Lange, M.D. 10 CFR 35.200	
Christopher J. Leary, M.D. 10 CFR 35.200	
Eric Lawrence Leonard, M.D. 10 CFR 35.200	

C FORM 374A	U.S. NUCLEAR REGULATORY COMMISSION		PAGE 5 OF 10 PAGES
MATERIALS LICENSE	License Number 47-25570-01	Docket or Reference Number 030-35774	
SUPPLEMENTARY SHEET	Amendment No. 24		
Authorized User(M.D.,D.O.,etc.)	Material and Use	,	
Donald Lewis, M.D.	10 CFR 35.200		
Edward J. Maas, M.D.	10 CFR 35.200 R RE	G_{II} ,	
Colleen M. Madden, M.D.	10 CFR 35.200		
Jane MaLoof, M.D.	10 CFR 35.200	GULAX	
Jack D. Markiewicz, M.D.	10 CFR 35.200		
Matthew E. Maxwell, M.D.	10 CFR 35.200	1	
Timothy J. McCue, M.D.	44 10 CFR 35.200		
Chris W. McGary, M.D.	10 CFR 35.200		
Russell Meyer, M.D.	10 CFR 35.200		
Khalid A. Mian, M.D.	10 CFR 35.200	NON	
Thomas E. Miller, M.D.	10 CFR 35.200	3	
Steve Min, D.O.	10 CFR 35.200		
Daniel R. Mitchell, M.D.	10 CFR 35.200	S	
Virginia Molleran, M.D.	10 CFR 35.200		
Craig Moore, M.D.	10 CFR 35.200	40	
Jaochim Mueller, M.D.	10 CFR 35:200	L - X	
AppaRao Mukkamala, M.D.	10 CFR 35.200		
Joshua A. Nepute, M.D.	10 CFR 35.200		
Dana Olson, M.D.	10 CFR 35.200		
Marlo M. Pagano, M.D.	10 CFR 35.200		
Charles Nicholas Pappas, M.D.	10 CFR 35.200		
Samir Parikh, M.D.	10 CFR 35.200		
Bharat Patel, M.D.	10 CFR 35.200		
Joseph Pekala, M.D.	10 CFR 35.200		

RC FORM 374A	U.S. NUCLEAR REGULA	TORY COMMISSION	PAGE 6 OF 10 PAGES
MATERIALS LICENSE	License Number 47-25570-01	Docket or Reference Number 030-35774	
SUPPLEMENTARY SHEET	Amendment No. 24		
Authorized User(M.D.,D.O.,etc.)	Material and Use	,	
Mark Peterson, M.D.	10 CFR 35.200		
Grant D. Petty, M.D.	10 CFR 35.200 R RE	GULAZ	
Krishna R. Pillai, M.D.	10 CFR 35.200	LA.	
James Milton Reynolds, M.D.	10 CFR 35.200		
Sean Reynolds, M.D.	10 CFR 35.200		
Michael E. Robertello, M.D.	10 CFR 35.200	1	
Ruben Rock, M.D.	4 10 CFR 35.200		
Daniel Adam Rodgers, M.D.	10 CFR 35.200		
Colin Rose, M.D.	10 CFR 35.200	MMISS	
Heather Rose, M.D.	10 CFR 35.200	ALL S	
Ronald J. Rosenberg, M.D.	10 CFR 35.200	3	
Robert M. Salman, M.D.	10 CFR 35.200		
Paul Sanchirico, M.D.	10 CFR 35.200	S	
Gerling Sauter, M.D.	10 CFR 35.200		
Charles Seigler, M.D.	10 CFR 35.200	40	
Marc A. Seltzer, M.D.	10 CFR 35,200	~ **	
Mark Shaman, M.D.	10 CFR 35.200	X V	
Stanley M. Shapiro, M.D.	10 CFR 35.200		
Paul W. Sheets, M.D.	10 CFR 35.200		
Paul D. Shreve, M.D.	10 CFR 35.200		
Ralph E. Shrider, M.D.	10 CFR 35.200		
Alan Siegel, M.D.	10 CFR 35.200		
Justin Sims, M.D.	10 CFR 35.200		
Roshan Sivagnanam, M.D.	10 CFR 35.200		

NRC FORM 374A	U.S. NUCLEAR REGULATORY COMMISSION		
MATERIALS LICENSE	License Number 47-25570-01	Docket or Reference Number 030-35774	
SUPPLEMENTARY SHEET	Amendment No. 24		

Authorized User(M.D.,D.O.,etc.) Material and Use LeAnn Stidham, M.D. 10 CFR 35.200 10 CFR 35,200 Victoria A. Swegles, D.O. 10 CFR 35.200 Sanjay J. Talati, M.D. Smari Thordarson, M.D. 10 CFR 35.200 10 CFR 35.200 Walter Parke Thrush, M.D. 10 CFR 35.200 Boguslaw Uchman, M.D. 10 CFR 35,200 Matthew Waack, M.D. 10 CFR 35,200 Torin P. Walters, M.D. James K. Watson, M.D. 10 CFR 35.200 10 CFR 35,200 Jonathan W. Weiss, M.D. 10 CFR 35,200 Ehab Hassan A. Youssef, MBBCH, M.D.

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

- 12. The Radiation Safety Officer for this license is Kay Kassel, M.S., C.N.M.T.
- 13. In addition to the possession limits in Item 8, the licensee shall further restrict the possession of licensed material to quantities below the minimum limit specified in 10 CFR 30.35(d) for establishing decommissioning financial assurance.

NRC FORM 374A	U.S. NUCLEAR REGULATORY COMMISSION		PAGE 8 OF 10 PAGES
MATERIALS LICENSE	License Number 47-25570-01	Docket or Reference Number 030-35774	
SUPPLEMENTARY SHEET	Amendment No. 24		

- 14. 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.
- 15. Sealed sources containing licensed material shall not be opened by the licensee, except as specifically authorized.
- 16. 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.
 - 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.

NRC FORM 374A	U.S. NUCLEAR REGULATORY COMMISSION		PAGE 9 OF 10 PAGES
MATERIALS LICENSE	License Number 47-25570-01	Docket or Reference Number 030-35774	
SUPPLEMENTARY SHEET	Amendment No. 24		

- 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.
- 17. 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 in 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. Application dated April 5, 2011 [ML110970462]
 - B. Letter dated August 31, 2011 [ML112440261]
 - C. Letter dated June 19, 2015 [ML15195A190]
 - D. Letter dated October 13, 2015 [ML15292A554]
 - E. Letter dated January 4, 2016 [ML16022A219]
 - F. Letter dated January 12, 2016 [ML16022A229]
 - G. Letter dated August 23, 2018 [ML18242A382]
 - H. Letter dated August 31, 2018 (delegation of authority) [ML18262A147]
 - I. Letter dated September 16, 2019 [ML19270E773]
 - J. Letter received November 5, 2019 [ML19322A669]

NRC FORM 374A	U.S. NUCLEAR REGULATORY COMMISSION		PAGE 10 OF 10 PAGES
MATERIALS LICENSE	License Number 47-25570-01	Docket or Reference Number 030-35774	
SUPPLEMENTARY SHEET	Amendment No. 24		

Continued

K. Letter dated December 23, 2019 with attachment [ML20009D990]



FOR THE U.S. NUCLEAR REGULATORY COMMISSION

Date: May 14, 2020 By: ____

Penny Lanzisera Region 1