



חטיבת טכנולוגיות רפואיות, מידע ומחקר
המכון לביקורת ותקנים של חומרי רפואה
The Institute for Standardization and Control of Pharmaceuticals

אישור יצרן / יבואן תכשירים רפואיים MANUFACTURER'S / IMPORTER'S AUTHORIZATION

Authorization number	מספר האישור
MIA 158/2025/B	
Name of authorization holder	שם בעל האישור
Isotopia Molecular Imaging Ltd.	איזוטופיה מולקולר אימג'ינג בע"מ
Address of site	כתובת האתר
39 Alexander Yanay St., Segula Ind. Zone, Petach Tikva, 4927735, Israel	רח' אלכסנדר ינאי 39, א.ת. סגולה, פתח תקווה 4927735
Legally registered address of authorization holder	כתובת בעל האישור
As written above	כרשום לעיל
Scope of authorization and dosage forms	תחומי האישור וצורות המינון
See Annexes 1, 2	ראה נספחים 1, 2
Legal basis of authorization	הבסיס החוקי לאישור
Pharmacist Regulations [Good Manufacturing Practice] 2008	תקנות הרוקחים [תנאי ייצור נאותים] תשס"ח
Responsible officer of Israeli Ministry Of Health granting the authorization	בעל התפקיד ברשות האחראי למתן האישור
Michael Carmi, Pharmacist GMP inspector	מיכאל כרמי, רוקח רכז ארצי בקרת מפעלים
Signature, stamp and date	חתימה, חותמת ותאריך
e-mail: michael.carmi@moh.gov.il phone: office 972-2-6551795 cell 972-50-6242452 3.7.25 מיכאל כרמי ע. מפקח ארצי תנאי ייצור נאותים	



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SCOPE OF AUTHORIZATION : ANNEX 1 – MEDICINAL PRODUCTS

Name and address of the site **Isotopia Molecular Imaging Ltd.**
39 Alexander Yanay St., Segula Ind. Zone, Petach Tikva, 4927735, Israel

☒ Human Medicinal Products

AUTHORIZED OPERATIONS

- ☒ Manufacturing Operations (according to part 1)
☒ Importation of Medicinal Products (according to part 2)

Part 1 - MANUFACTURING OPERATIONS

1.1	Sterile products
	1.1.2 Terminally sterilized
	1.1.2.5 Other terminally sterilized prepared products (<i>Lutetium-177 sterile radionuclides</i>)
	1.1.3 Batch certification
1.5	Packaging
	1.5.2 Secondary packing
1.6	Quality control testing
	1.6.1 Microbiological: sterility (<i>performed at another site of the firm</i>)
	1.6.2 Microbiological: non-sterility (<i>performed at another site of the firm</i>)
	1.6.3 Chemical/Physical (<i>the Gamma-ray spectrometry test is outsourced</i>)



מיכאל כרמי
ע. מפקח ארצי
תנאי יצור נאותים

Any restrictions or clarifying remarks related to the scope of these manufacturing operations

The commercial products are Lutetium-177 carrier added & Lutetium-177 non carrier added, sterile radionuclides.



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Part 2 - IMPORTATION OF MEDICINAL PRODUCTS	
2.2	Batch certification of imported medicinal products
	2.2.1 Sterile Products
	2.2.1.1 Aseptically prepared (lyophilized powder in glass vial, for diagnostic use)
	2.2.4 Other importation activities
	2.2.4.1 Radiopharmaceuticals/Radionuclide generators (radiopharmaceutical precursor, sterile solution)
2.3	Other importation activities
	2.3.1 Site of physical importation : <i>site of contract agent</i>
	2.3.2 Importation of intermediate which undergoes further processing: <i>the imported products are reconstituted before use</i>

Any restrictions or clarifying remarks related to the scope of these importing operations:

The imported products are stored at a contract manufacturer's warehouse

3.7.25

מיכאל כרמי
ע. מפקח ארצי
תנאי יצור נאותים

MINISTRY OF HEALTH JERUSALEM
GOV. INSTITUTE
FOR
CONTROL AND STANDARD
OF
PHARMACEUTICALS
משרד הבריאות



SCOPE OF AUTHORIZATION: ANNEX 2 - INVESTIGATIONAL MEDICINAL PRODUCTS

Name and address of the site **Isotopia Molecular Imaging Ltd.**

39 Alexander Yanay St., Segula Ind. Zone, Petach Tikva, 4927735, Israel

☒ Human Investigational Medicinal Products for phases I, II clinical trials

AUTHORIZED OPERATIONS

☒ Manufacturing Operations (according to part 1)

Part 1 - MANUFACTURING OPERATIONS

1.1	Sterile products
	1.1.2 Terminally sterilized
	1.1.2.5 Other terminally sterilized prepared products (Terbium-161 sterile radionuclides)
	1.1.3 Batch certification
1.5	Packaging
	1.5.2 Secondary packing
1.6	Quality control testing
	1.6.1 Microbiological: sterility (performed at another site of the firm)
	1.6.2 Microbiological: non-sterility (performed at another site of the firm)
	1.6.3 Chemical/Physical (the Gamma-ray spectrometry test is outsourced)



Any restrictions or clarifying remarks related to the scope of these manufacturing operations

The IMP is Terbium-161 sterile radionuclide. It is manufactured in a dedicated room (# 23). This activity was authorized after an "inspection for cause" that was conducted on 3 June 2025. The authorization is valid for phases I,II clinical trials

Investigational products are not included in the CAA Agreement between the EU & Israel