

# **CERTIFICATE OF ANALYSIS**

### DRAXIMAGE MAA MACROSALB (10 vials/kit)

2.5 mg of Stannous Albumin Macro Aggregate / vial

# Certificate:	2022FE25-7			
Batch Number:	1K141	Manufacturing Date:	2022JA24	
Lot Number (FP):	1K141	Expiry Date:	01/2024	
Product Number:	3000001020	Specification Number and Version:	3000001020_v01	
Standard of Testing:	House, USP <sup>1</sup>	Date of Release: (Date with QA Initials)	2022 FERS MBI	

		Tests on the Lyophilized product	
Test Description	Analytical Method	Specifications	Results
Description	10019	A white freeze-dried plug or powder, clean and free of foreign matter. The flip-off cap is blue.	Conforms
Resuspendability	10079	A white suspension which may separate on standing.	Conforms
рН	10005	5.2 – 6.0	5.4
Loss on Drying	10021	≤ 5%	1%
Particle Density	10088	3 - 8 x 10 <sup>6</sup> aggregated albumin particles in each vial	6x10E6 part./vial
Particle Size	10080	Particle Size < 10 μm: ≤ 10%	4%
		Particle Size ≥ 10 μm - ≤ 70 μm: ≥ 90%	96%
		Particle Size > 100 µm: ≤ 0.2%	< 0.1%
		Particle Size > 150 μm: none	None
MAA Identification	10000	A blue color develops.	Conforms
SnCl <sub>2</sub> .2H <sub>2</sub> O Assay	10039	≥ 0.06 mg/vial	0.11 mg/vial
Total Tin Assay	10040	≤ 0.12 mg	0.09 mg/vial
Stannous Albumin Macro Aggregated Complex	10082	2.2 - 3.0 mg/vial	2.4 mg/vial
Human Serum Albumin	10068	3.5 – 6.5 mg/vial (or alternative method 10089)	5.5 mg/vial
Sodium Chloride	10084	0.96 – 1.44 mg/vial	1.38 mg/vial
Residual Solvents¹	USP<467>	Meets USP requirements (No test required)	Conforms
Sterility <sup>1,2</sup>	10007	Sterile	Conforms
Bacterial Endotoxins	10008	≤ 16.5 EU/vial	< 5.2 EU/vial (

<sup>&</sup>lt;sup>2</sup> Outside testing

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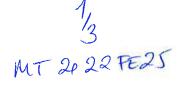
#### A Jubilant Pharma Company



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	Te	ests on the Reconstituted product	
Test Description	Analytical Method	<u>Specifications</u>	Results
Radiochemical Purity	10043	Not less than 90% of the total radioactivity is found as aggregated albumin at the point of application, at 15 – 30 minutes post labeling.	
		Not less than 90% of the total radioactivity is found as aggregated albumin at the point of application, at least 12 hours post labeling.	100%
Centrifugation Procedure	10087	Not more than 10% of the total radioactivity is found in the supernatant liquid, at 15 – 30 minutes post labeling.	4%
		Not more than 10% of the total radioactivity is found in the supernatant liquid, at least 12 hours post labeling.	3%
Biological Distribution	10044	In not less than 2 of 3 animals, at 15 – 30 minutes post labeling, not less than 80% of the radioactivity is found in the lungs of the animal at 15 minutes post injection.	
		In not less than 2 of 3 animals, at 15 – 30 minutes post labeling, not more than a total of 5% of the radioactivity is found in the liver of the animal at 15 minutes post injection.	1: 2% 2: 2% 3: 2%
		In not less than 2 of 3 animals, at 15 – 30 minutes post labeling, not more than 5% of the radioactivity is found in the kidneys of the animal at 15 minutes post injection.	1: 1% 2: 2% 3: 1%
		In not less than 2 of 3 animals, at 12 -24 hours post labeling, not less than 80% of the radioactivity is found in the lungs of the animal at 15 minutes post injection.	1: 95% 2: 96% 3: 98%
		In not less than 2 of 3 animals, at 12 - 24 hours post labeling, not more than a total of 5% of the radioactivity is found in the liver of the animal at 15 minutes post injection.	1: 2% 2: 2% 3: 2%
		In not less than 2 of 3 animals, at 12 – 24 hours post labeling, not more than 5% of the radioactivity is found in the kidneys of the animal at 15 minutes post injection.	1: 1% 2: 1% 3: 1%

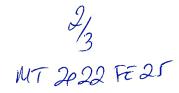
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This batch of product has been tested by Jubilant DraxImage Quality Control Laboratory under Canadian Establishment License Number **101869-A** and complies with the specification requirements.

MUSTAPHA TOLLABI

OC Sr. Supervisor

NATHALIE BRUNET

M. Tollah

2022 FE25

Verified by:

Name and Title

Signature

Date

OA, ASSOCIATE
Approved by:

Name and Title

Signature

Mathelie Brunet

Date

2022FE25

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OURVALUES



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