

Laboratory Report

Report prepared for:

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Purchase Order:

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Sample: Batch: HOKX1329040924 Lot: 093024117	Received: 2024-10-02
Lab ID: 2024-U-9099	

Analysis	Method	Result	Basis	Sample Amount Used	Date (Time)
<i>118: Residue on Ignition (ROI)</i>					
	JP XVIII Supplement I	< 1.0 mg	As Received	10.0 mL	2024-10-18
<i>774: Br/I, Free Br/Cl, SO4/SO3</i>					
	JP XVIII Supplement I ¹	Passes Test	As Received	Direct	2024-10-23
	JP XVIII Supplement I ²	Passes Test	As Received	Direct	2024-10-23
	JP XVIII Supplement I ³	Passes Test	As Received	Direct	2024-10-23
<i>777: Identification</i>					
	JP XVIII Supplement I ⁴	Passes Test	As Received	Direct	2024-10-22
	JP XVIII Supplement I ⁵	Passes Test	As Received	1 mL	2024-10-23
<i>785: Sulfate</i>					
	JP XVIII Supplement I	Passes Test	As Received	15 mL	2024-10-23
<i>j35: Hydrochloric Acid</i>					
	JP XVIII Supplement I	37.5 %	As Received	3357.5 mg	2024-10-22

1. Bromine or chlorine
2. Bromide or iodide
3. Sulfite
4. ID 1
5. ID 2

For all samples on this report:

6. These analyses were performed in general compliance with the Laboratory sections of Current Good Manufacturing Practices for bulk pharmaceuticals as defined in 21 CFR 210 and .

Signatures:

Modified By:	michelle.mccurdy	2024-10-29T15:25:20.92-04:00
Inspected By:	tammy.saylor	2024-10-29T16:14:16.017-04:00
Created By:	michelle.mccurdy	2024-10-25T20:50:54.35-04:00
Published By:	tammy.saylor	2024-10-29T16:14:19.48-04:00

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- ␣ "Inspected By" signature indicates QA review and approval.
- ␣ "Published By" signature indicates authorized release of data.