

## Laboratory Report

**Report prepared for:**

Daniel Pocci  
Elco Corp  
1100 State Road  
Ashtabula, OH 44004  
Phone: 216-767-4001  
Email: [d.pocci@italmatch.com](mailto:d.pocci@italmatch.com), [d.bias@italmatch.com](mailto:d.bias@italmatch.com), [d.morgan@italmatch.com](mailto:d.morgan@italmatch.com)

**Report prepared by:**

Michelle Mccurdy

**Purchase Order:**

70318370

**For further assistance, contact:**

Michelle McCurdy  
Technical Manager  
PO Box 51610  
Knoxville, TN 37950 -1610  
(865) 546-1335 ext. 1826  
[michellemccurdy@galbraith.com](mailto:michellemccurdy@galbraith.com)

Sample: TILX6015820823, 081723117		Received: 2023-08-18			
Analysis	Method	Result	Basis	Sample Amount Used	Date (Time)
<i>454: Nonvolatile residue</i>					
	EP 11.2	< 0.002 %	As Received	84.7 mL	2023-09-12
<i>777: Identification</i>					
	EP 11.2 <sup>1</sup>	Passes Test	As Received	1 mL	2023-08-24
	EP 11.2 <sup>2</sup>	Passes Test	As Received	2 mL	2023-08-24
<i>785: Sulfate</i>					
	EP 11.2	< 20 ppm	As Received	6.4 mL	2023-09-08
	EP 11.2 (matrix spike)	≥ 20 ppm	As Received	6.4 mL	2023-09-08
<i>802: Clarity of Solution</i>					
	EP 11.2	Passes Test	As Received	2 mL	2023-08-28
<i>803: Color of Solution</i>					
	EP 11.2	Passes Test	As Received	2 mL	2023-08-28
<i>f17: Chlorine, Residual Free</i>					
	EP 11.2	< 4 ppm	As Received	15 mL	2023-09-06
<i>j35: Hydrochloric Acid</i>					
	EP 11.2	37.4 %	As Received	1738.23 mg	2023-08-24

1. IDA

2. IDB

**For all samples on this report:**

- The matrix spike analysis was performed to satisfy method requirements. There is no additional charge for the matrix spike result.
- 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 211.

**Signatures:**

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