

Certificate of Manufacture / Analysis

Certificate ID: 10924

Drug Product: Cefa	ct: Cefazolin 3 Gram, Bag for Injection		
Batch/Lot ID: 20220	120CEF-1	1	NDC: 71139-7153-1
Strength: 3 grar	n Container:	100 mL bag Volume (/	Amount): <u>115 mL</u>
Production Date: 01/20/	2022	Expirati	ion Date: 03/21/2022
Storage: Refrigerated Temperature, 2°C - 8°C			
Testing Results			
Test/Parameter	Specification	Result	Date of Analysis
Cefazolin Potency / ID	90.0 % - 115.0 % / Positive	95.3 % / Positive	01/27/2022
Endotoxin (USP <85>)	≤ 4.50 EU/mL	< 2.50 EU/mL	01/31/2022
Sterility (USP <71>)	Negative Growth (at 14 Days)	Negative	02/10/2022
Visual Inspection	NMT 0 Units Non-Conforming	Pass	01/20/2022
Filter Integrity	Pressure > 55 psi	NA	NA
*Comments: NA			
Statement of Conformity: IntegraDose Compounding Services, LLC certifies that the above product has successfully passed all testing as per the applicable test specifications. All products manufactured under IntegraDose's Quality System meet the regulatory requirements for 503B Outsourcing Facilities (21 USC 353b: Outsourcing facilities and the FDA Current Good Manufacturing Practice – Guidance for Human Drug Compounding Outsourcing Facilities Under Section 503B of the FD&C Act) and all other applicable internal Quality requirements. All products are manufactured with validated processes to current specifications and are inspected to ensure they meet quality requirements.			
Certificate Preparation Date:02/14/2022 By/Date:			
Quality Review/Date:			

719 Kasota Ave Southeast, Minneapolis, MN 55414

Form QUA-026.2 (06/2020) integradose.org Page 1 of 1