



Batch Master Record

CLEANOUT:

BATCH NUMBER: 880-CM-624

DATE DATE: 7/20/20

08-20-20

FINISHED GOOD: 2000-00000

PRINT DATE: 8/20/20

AMOUNT TO PRODUCE: 5,450,000 lb

WAREHOUSE: 00-1A

LOT NUMBER: T80-LOT

HRIS RATING: A- 1 1-2 B 1 00-1

CONTAINER MEASURE (WEIGHT): 100 LB

Remarks: Print on Batch Ticket

TOTAL RAW MATERIAL REQUIREMENTS:

PRODUCT CODE	PRODUCT NAME / PACKAGE DESCRIPTION	LOT NUMBER/LOC	TOTAL WEIGHT QUANTITY (LBS)	TOTAL VOL. (GALLONS) USED
700-00000	Whylene Glycol	T80-LOT	2,410,000 lb	291.0004
	1.00 lb BULK	T80-LOC		
200-20000	Green Dye	T80-LOT	5,000,000 g	0.7700
	25.00 lb Can	T80-LOC		
200-00000	Ethyl Alcohol	T80-LOT	877,000 lb	105.5004
	1.00 lb BULK	T80-LOC		
000-00000	Methanol	T80-LOT	117,000 lb	14.4007
	1.00 lb BULK	T80-LOC		
000-00000	Ice (A)	T80-LOT	2,040,000 lb	208.0011
	1.00 lb BULK	T80-LOC		

STEP BY STEP INSTRUCTIONS

STEP: 1-0000 WORK CENTER: 1-Blending

EQUIPMENT: 01-Blending

MATERIALS:	TOTAL QUANTITY
2000-00000	5,450,000 lb
Whylene Glycol	
2000-20000	5,000,000 g
Green Dye	
2000-00000	877,000 lb
Ethyl Alcohol	
000-00000	2,040,000 lb
Ice (A)	
0000-00000	117,000 lb
Methanol	

INSTRUCTIONS: Unlimited Instructions - Heat, mix for 2 hours. QC, Filter, Package

START TIME: _____ STOP TIME: _____ INITIALS: _____

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Albert A Gayle



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Handbook of Pharmaceutical Manufacturing Formulations Sarfaraz K. Niazi, 2016-04-19 Providing methodologies that can serve as a reference point for new formulations the second volume covers uncompressed solids which include formulations of powders capsules powders ready for reconstitution and other similar products Highlights from Uncompressed Solid Products Volume Two include the fundamental issues of good manufacturing **Active**

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biological drugs and alternative medicines *Data Integrity and Compliance* José Rodríguez-Pérez,2019-05-08 Data integrity is a global mandatory requirement for the regulated healthcare industry It is more than a mere expectation it s a basic element of good documentation practices one of the most fundamental pillars of a quality management system Robustness and accuracy of the data submitted by manufacturers to regulatory authorities when bringing a medical product to market are crucial The purpose of this book is to consolidate existing data integrity principles and expectations from several regulatory sources including the U S Food and Drug Administration World Health Organization and European Medicines Agency into a single and handy document that provides detailed illustrative implementation guidance It serves as a means of understanding regulatory agencies position on good data management and the minimum expectation for how medical product manufacturers can achieve compliance **Federal Register** ,1978-09 Control Procedures in Drug Production William Leonard Blockstein,1966 **United States of America V. Article of Drug Labeled "White Quadrisect"** ,1972 The United States pharmacopeia ,2012 *United States Pharmacopeia Dispensing Information* ,1998 **Pharmaceutical Dosage Forms, Parenteral Medications** Kenneth E. Avis,Leon Lachman,Herbert A. Lieberman,1984 *Production Part Approval Process (PPAP)* ,2000 Food, Drug, Cosmetic Law Reporter ,1963 Grist from the Mill ,1978 **Vaccine Manual** Noel Mowat,Mark M. Rweyemamu,Food and Agriculture Organization of the United Nations,1997 This manual represents a response to a recommendation arising from an expert consultation on vaccine production and quality control held in 1991 It attempts to birng together information on the state of the art in veterinary vaccine development *McGraw-Hill Encyclopedia of Science & Technology* ,1987 *European Plastics News* ,1986 **Electrochemical and Metallurgical Industry** ,1960 **Hazards of Medication** Eric Wentworth Martin,1971 **Computer Operations** Don B. Medley,Jerry L. Koory,1988

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