BUPRENORPHINE- buprenorphine solution
PAYLESS COMPOUNDERS, LLC

BUPRENORPHINE

buprenorphine solution

Product Information			
Product Type	ANIMAL COMPOUNDED DRUG	**Item Code (Source)**	NDC:70022-004
Route of Administration	INTRAMUSCULAR, SUBCUTANEOUS	**DEA Schedule**	CIII
Reporting Period	20151006-20160405		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BUPRENORPHINE HYDROCHLORIDE (UNII: 56W8MW3EN1) (BUPRENORPHINE - UNII:40D3SCR4GZ)	BUPRENORPHINE	0.6 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ANHYDROUS DEXTROSE (UNII: 5SL0G7R0OK)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
MANNITOL (UNII: 3OWL53L36A)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**

1	NDC:70022-004-10	10 mL in 1 VIAL, MULTI-DOSE; Number of Units = 10		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		10/06/2015	

Labeler - PAYLESS COMPOUNDERS, LLC (031728341)

Establishment

Name	Address	ID/FEI	Business Operations
PAYLESS COMPOUNDERS, LLC		604160239	outsourcing animal drug compounding

Revised: 10/2015 PAYLESS COMPOUNDERS, LLC