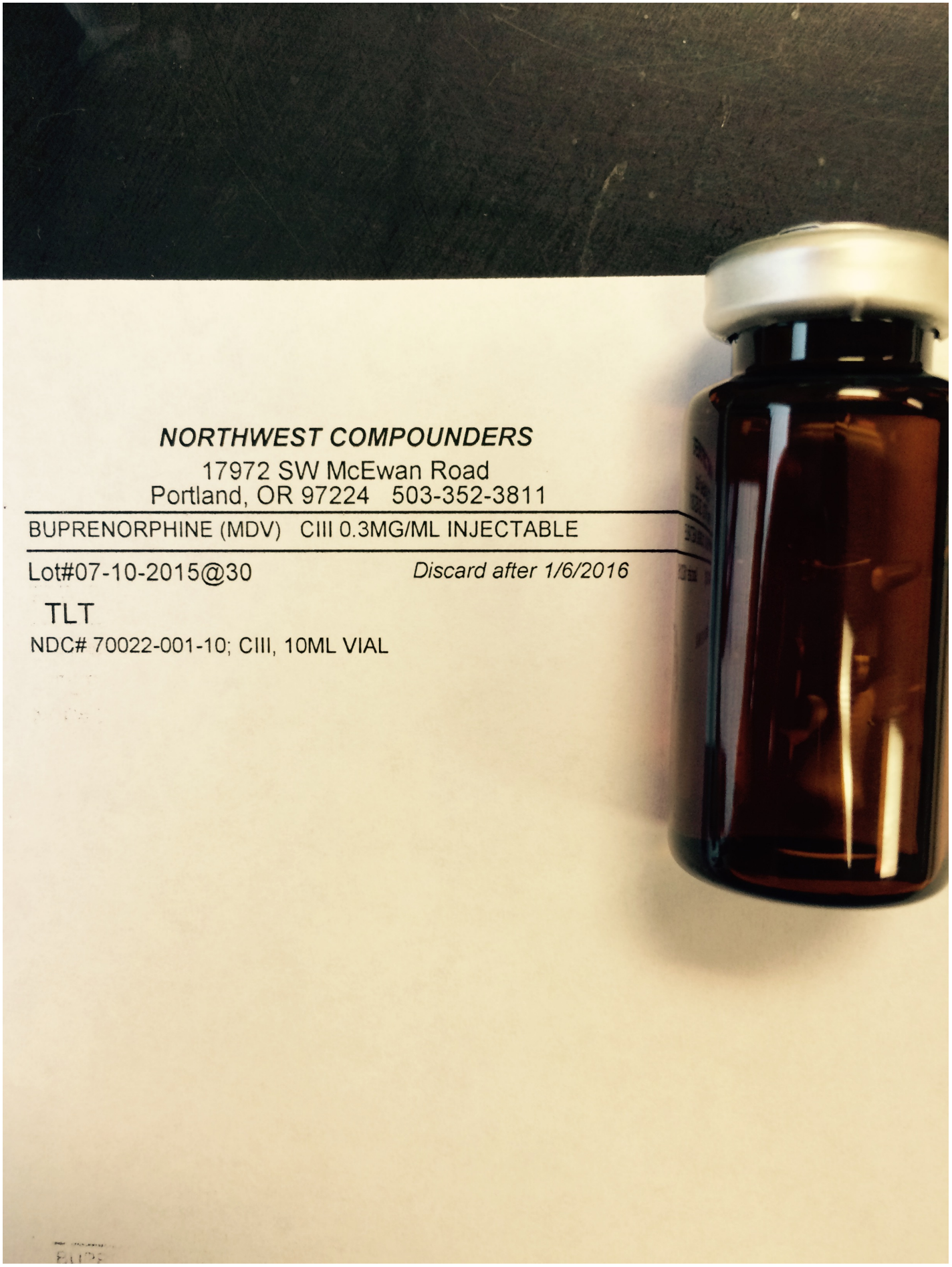
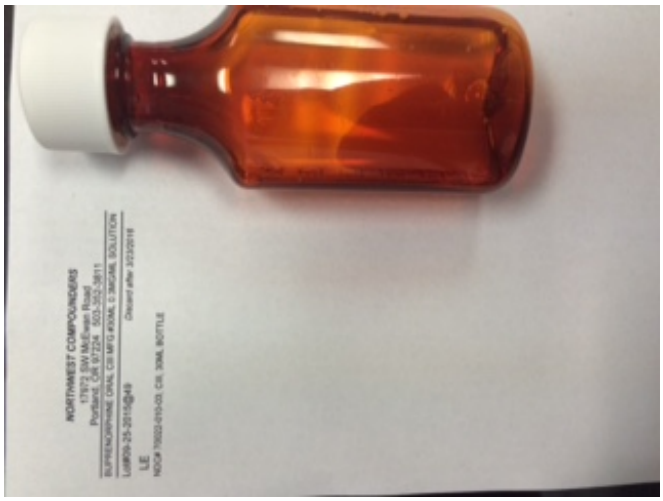
BUPRENORPHINE- buprenorphine injection, solution
BUPRENORPHINE- buprenorphine solution
PAYLESS COMPOUNDERS, LLC
Reference Label Set Id: 8081ccb8-1b38-4673-a9dd-0fb18fa6038a


NORTHWEST COMPOUNDERS
17972 SW McEwan Road
Portland, OR 97224 503-352-3811
BUPRENORPHINE (MDV) CIII 0.3MG/ML INJECTABLE
Lot#07-10-2015@30
Discard after 1/6/2016
TLT
NDC# 70022-001-10; CIII, 10ML VIAL



BUPRENORPHINE

buprenorphine injection, solution

Product Information			
Product Type	ANIMAL COMPOUNDED DRUG	**Item Code (Source)**	NDC:70022-001

Route of Administration	INTRAMUSCULAR, SUBCUTANEOUS	**DEA Schedule**	CIII
Reporting Period	20240630-20241231		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BUPRENORPHINE HYDROCHLORIDE (UNII: 56W8MW3EN1) (BUPRENORPHINE - UNII:40D3SCR4GZ) (Source NDC: 38779-0888)	BUPRENORPHINE	0.3 mg in 1 mL

Inactive Ingredients

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70022-001-10	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			

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	20240630-20241231		

Active Ingredient/Active Moiety

Ingredient Name	Basis of	Strength

Ingredient Name	Basis of Strength	Strength
BUPRENORPHINE HYDROCHLORIDE (UNII: 56W8MW3EN1) (BUPRENORPHINE - UNII:40D3SCR4GZ) (Source NDC: 38779-0888)	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			

BUPRENORPHINE

buprenorphine solution

Product Information

Product Type	ANIMAL COMPOUNDED DRUG	**Item Code (Source)**	NDC:70022-010
Route of Administration	ORAL, TRANSMUCOSAL	**DEA Schedule**	CIII
Reporting Period	20240630-20241231		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BUPRENORPHINE HYDROCHLORIDE (UNII: 56W8MW3EN1) (BUPRENORPHINE - UNII:40D3SCR4GZ) (Source NDC: 38779-0888)	BUPRENORPHINE	0.3 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	

WATER (UNII: 059QF0KO0R)	
SUCROSE (UNII: C151H8M554)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70022-010-03	30 mL in 1 BOTTLE, PLASTIC; Number of Units = 30		
2	NDC:70022-010-10	100 mL in 1 BOTTLE, PLASTIC; Number of Units = 100		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other			

Labeler - PAYLESS COMPOUNDERS, LLC (031728341)

Establishment

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

Revised: 6/2024 PAYLESS COMPOUNDERS, LLC