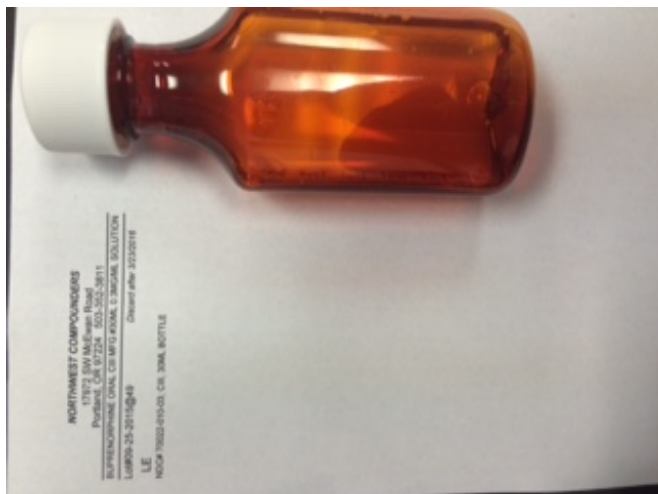


**BUPRENORPHINE- buprenorphine solution**  
**PAYLESS COMPOUNDERS, LLC**

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**BUPRENORPHINE**

buprenorphine solution

**Product Information**

<b>Product Type</b>	ANIMAL COMPOUNDED DRUG	<b>Item Code (Source)</b>	NDC:70022-010
<b>Route of Administration</b>	ORAL, TRANSMUCOSAL	<b>DEA Schedule</b>	CIII
<b>Reporting Period</b>	20151002-20160402		

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
BUPRENORPHINE HYDROCHLORIDE (UNII: 56W8MW3EN1) (BUPRENORPHINE - UNII:40D3SCR4GZ)	BUPRENORPHINE	0.3 mg in 1 mL

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
METHYL PARABEN (UNII: A2I8C7HI9T)	
PROPYL PARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0K00R)	
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		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		10/02/2015	

**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