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

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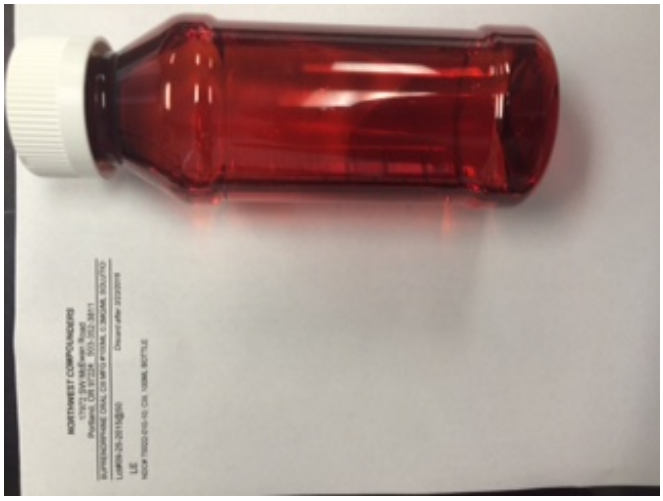

# BUPRENORPHINE

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** | 20151002-20160402 | | |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **BUPRENORPHINE HYDROCHLORIDE** (UNII: 56W8MW3EN1) (BUPRENORPHINE - UNII:40D3SCR4GZ) | 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:70022-010-10 | 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 | | 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