



BUPRENORPHINE Sublingual Tablets C-III

PRODUCT DESCRIPTION

Round/White Sublingual Tablets



NDC Number	Strength	Unit Size
0054-0176-13	2 mg	Bottle - 30 tablets
0054-0177-13	8 mg	Bottle - 30 tablets

BUPRENORPHINE AND NALOXONE Sublingual Tablets C-III

PRODUCT DESCRIPTION

Round/Speckled Peach to Peach Sublingual Tablets



NDC Number	Strength	Unit Size
0054-0188-13	2 mg / 0.5 mg	Bottle - 30 tablets
0054-0189-13	8 mg / 2 mg	Bottle - 30 tablets

DISKETS® Dispersible Tablets C-II (Methadone Hydrochloride Tablets for Oral Suspension USP)

PRODUCT DESCRIPTION

Light Pinkish-Orange Cross-Scored Tablets. Orange-Pineapple Flavored



NDC Number	Strength	Unit Size
0054-4538-25	40 mg	Bottle - 100 tablets

METHADONE HCl Tablets USP C-II

PRODUCT DESCRIPTION

Round/White Tablets

BRAND EQUIVALENT

DOLOPHINE®



NDC Number	Strength	Unit Size
0054-4570-25	5 mg	Bottle - 100 tablets
0054-4571-25	10 mg	Bottle - 100 tablets

METHADONE HCl Oral Concentration USP C-II

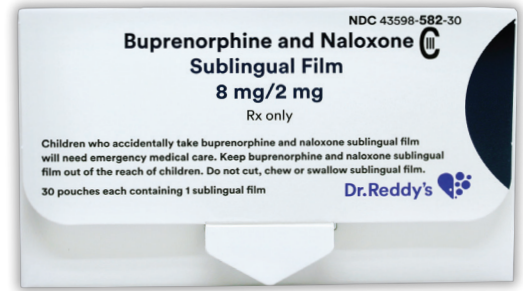
Product Description	NDC Number	Strength	Unit Size
Clear/Colorless Solution - Flavorless	0054-0391-68	10 mg / mL	1000 mL Bottle
Clear/Red Colored Solution - Cherry flavored	0054-0392-68	10 mg / mL	1000 mL Bottle

NOW AVAILABLE

Buprenorphine and Naloxone Sublingual Film

2 mg/0.5 mg, 4 mg/1 mg, 8 mg/2 mg, and 12 mg/3 mg

A therapeutic equivalent generic version of *Suboxone*[®] (buprenorphine and naloxone) sublingual film.



BUPRENORPHINE AND NALOXONE SUBLINGUAL FILM

Strength	Packaging	NDC
2 mg/0.5 mg	30 pouches each containing 1 sublingual film	43598-579-30
4 mg/1 mg	30 pouches each containing 1 sublingual film	43598-580-30
8 mg/2 mg	30 pouches each containing 1 sublingual film	43598-582-30
12 mg/3 mg	30 pouches each containing 1 sublingual film	43598-581-30

INDICATION

Buprenorphine and naloxone sublingual film is indicated for treatment of opioid dependence. Buprenorphine and naloxone sublingual film should be used as part of a complete treatment plan that includes counseling and psychosocial support.

IMPORTANT SAFETY INFORMATION

Prescription use of this product is limited under the Drug Addiction Treatment Act.

CONTRAINDICATIONS

Buprenorphine and naloxone sublingual film is contraindicated in patients with a history of hypersensitivity to buprenorphine or naloxone as serious adverse reactions, including anaphylactic shock, have been reported.

For additional information, contact your Account Manager or Customer Service at 800-631-2174.

Customer Service
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