The Medical Letter®

on Drugs and Therapeutics

Last updated: September 13, 2018



Comparison Table: Some Drugs for Maintenance Treatment of Opioid Use Disorder

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Drug	Some Available Formulations	Target Mainten- ance Dosage	Adverse Effects	Drug Interactions	Pregnancy/Lactation	Comments	Cost ²						
Buprenorphine – generic Probuphine (Braeburn) Sublocade (Indivior)	2, 8 mg sublingual tabs 74.2 mg subdermal implant 100 mg/0.5 mL, 300 mg/1.5 mL prefilled syringes for subcutane- ous injection	16 mg once/d 4 implants for 6 mos 100 mg once/mo (300 mg first 2 mos; increase to 300 mg once/mo if clinical response			➤ Considered safe and effective for use in pregnant or breast- feeding women	Buprenorphine is a schedule III controlled substance; can be prescribed in outpatient setting Generally considered the maintenance treatment of choice; similar in efficacy to methadone at higher doses Probuphine: approved for 6 or 12 months' treatment in patients stabilized on ≤8 mg/d of buprenorphine;	\$1083.60 ⁵ 4950.00 ⁵ 9480.00 ⁵						
Buprenorphine/Naloxone – generic Bunavail (BioDelivery Sciences) Cassipa (Teva) Suboxone (Indivior) Zubsolv (Orexo)	2/0.5, 8/2 mg sublingual tabs 2.1/0.3, 4.2/0.7, 6.3/1 mg buccal films 16/4 mg sublingual films 2/0.5, 4/1, 8/2, 12/3 mg sublingual films 0.7/0.18, 1.4/0.36, 2.9/0.71, 5.7/1.4, 8.6/2.1, 11.4/2.9 mg sublingual tabs	is inadequate) 16/4 mg once/d 8.4/1.4 mg once/d 16/4 mg once/d 16/4 mg once/d 11.4/2.8 mg once/d	 Sedation/respiratory depression (less than methadone), headache, abdominal pain, constipation, nausea, vomiting, insomnia, sweating Probuphine: implant-site pain, pruritus, and erythema; insertion and removal of drug-eluting implants has been associated with nerve injury and implant migration and extrusion Sublocade: injection-site reactions Bunavail: oral mucosal erythema 	 Coadministration of benzodiazepines or other sedating drugs can be dangerous Inducers of CYP3A4 can reduce buprenorphine levels, and inhibitors of CYP3A4 can increase them Concurrent use of other serotonergic drugs can result in serotonin syndrome May interfere with analgesic activity of full opioid agonists 	 ▶ Generally considered safe in pregnant women, but efficacy data are limited; buprenorphine without naloxone is preferred ▶ Formulations without naloxone are preferred for use in breastfeed- ing women 	drug levels are similar to those with 8 mg/d of sublingual buprenorphine; REMS certification is required for prescribers ► Sublocade: approved for treatment of patients who have received transmucosal buprenorphine for ≥7 days with dose adjustment to 8-24 mg/d of buprenorphine sublingual tabs or equivalent; refrigerate during storage (discard if left at room tem for >7 days); steady state buprenorphine levels with target dosage ~10% higher than those with 24 mg/d of sublingual tabs ► Cassipa: approved for use after induction and stabilization of the patient, and titration to a dose of 16 mg of buprenorphine using another product ► Efficacy of naloxone-containing formulations may be reduced in patients with severe hepatic impairment ► Bunavail 4.2/0.7 mg and Zubsolv 5.7/1.4 mg produce systemic buprenorphine levels equivalent to those seen with 8 mg of other buprenorphine formulations	2339.10 ⁵ 2793.60 ⁶ N.A. 2933.30 ⁵ 2989.40 ⁶						

Drug	Some Available Formulations	Target Mainten- ance Dosage	Adverse Effects	Drug Interactions	Pregnancy/Lactation	Comments	Cost ²
Methadone – generic Dolophine (Roxane)	5, 10 mg tabs; 5, 10 mg/ 5 mL oral solution; 10 mg/mL oral concen- trate; 40 mg tabs for oral suspension ² 5, 10 mg tabs ³	80-120 mg once/d (some rapid metabolizers may require more frequent dosing)	➤ Sedation/respiratory depression, constipation, lightheadedness, dizziness, sedation, nausea, vomiting, sweating, QT-interval prolon- gation and (rarely) arrhythmias such as torsades de pointes	 Coadministration of benzodia-zepines or other sedating drugs can be dangerous Inducers of CYP3A4 or 2B6 can reduce methadone levels, and inhibitors of CYP3A4 or 2B6 can increase them Concurrent use of other serotonergic drugs can result in serotonin syndrome Concurrent use of other QT interval-prolonging drugs can cause arrhythmias such as torsades de pointes 	► Generally considered safe and effective for use in pregnant or breastfeeding women	Schedule II controlled substance; only available for treatment of opioid use disorder through opioid treatment programs with supervised dosing Shown to reduce mortality in clinical trials; should be used when buprenorphine is unavailable or ineffective and in patients who would benefit from supervised dosing Drug accumulates for 4-7 days during induction; respiratory depressant effect peaks later and lasts longer than analgesic effect Federal law prohibits initial doses >30 mg or first daily doses >40 mg	\$73.50 ⁴ 850.80
Naltrexone – generic Revia (Duramed) extended-release microspheres – Vivitrol (Alkermes)	50 mg tabs 380 mg ER suspension for injection	50 mg once/d 380 mg IM in alter- nating buttocks q4 wks or once/mo	 ▶ Generally well tolerated; nasopharyngitis, insom- nia, headache, nausea, and toothache are most commonly reported ▶ Rarely: depressed mood/ suicidality (cause and effect not established), hepatotox- icity (occurs frequently in opioid- and alcohol-dependent patients) ▶ Vivitrol: injection-site reactions 	 ▶ Blocks effects of usual doses of opioids, including antidiarrheals and antitussives ▶ Can precipitate severe withdrawal in patients with physiological opioid dependence; patients should be free of dependence for ≥7 days before initiation ▶ Discontinue 72 hours (oral formulation) or 30 days (IM formulation) before elective surgery 	 Data on safety and efficacy in pregnancy are limited; women at high risk for relapse who become pregnant can generally continue treatment Women taking naltrexone should not breastfeed; the drug passes into breast milk to a clinically significant extent 	 Not a controlled substance ER naltrexone is an alternative for highly motivated patients who do not have access to buprenorphine or methadone, or do not want to take an opioid, and for those who also have alcohol use disorder Adherence is poor with oral naltrexone; it has generally only been effective in patients who were legally required to take the drug Has not been shown to reduce mortality Vivitrol: must be refrigerated; use within 7 days of removing from refrigerator (do not expose to temperatures 77°F) 	371.00 1648.90 7854.00

N.A. = cost not yet available
1. Drugs for opioid use disorder. Med Lett Drugs Ther 2017; 59:89.
2. Approximate WAC for 6 months' treatment at the lowest target maintenance dosage. WAC = wholesaler acquisition cost, or manufacturer's published price to wholesalers; WAC represents published catalogue or list prices and may not represent an actual transactional price. Source: AnalySource® Monthly. May 5, 2017. Reprinted with permission by First Databank, Inc. All rights reserved. ©2017. www.fdbhealth.com/policies/drug-pricing-policy. www.fdbhealth.com/policies/drug-pricing-policy. www.fdbhealth.com/policies/drug-pricing-policy.
4. Cost of oral concentrate.
5. Approximate WAC for 6 months' treatment at the target maintenance dosage. Source: AnalySource® Monthly. February 5, 2018.