PHARMACOLOGIC PRODUCT GUIDE: FDA-APPROVED MEDICATIONS FOR SMOKING CESSATION

		Nicotine	BUPROPION SR	VARENIGLINE			
	GUM	LOZENGE	TRANSDERMAL PATCH	NASAL SPRAY	ORAL INHALER	BUFRUFIUN 3K	VARENICLINE
PRODUCT	Nicorette ¹ , Generic OTC 2 mg, 4 mg original, cinnamon, fruit, mint	Nicorette ¹ , Generic Nicorette1 Mini OTC 2 mg, 4 mg; cherry, mint	NicoDerm CQ¹, Generic OTC (NicoDerm CQ, generic) Rx (generic) 7 mg, 14 mg, 21 mg (24-hr release)	Nicotrol NS ² Rx Metered spray 10 mg/mL nicotine solution	Nicotrol Inhaler ² Rx 10 mg cartridge delivers 4 mg inhaled vapor	Zyban¹, Generic Rx 150 mg sustained-release tablet	Chantix² Rx 0.5 mg, 1 mg tablet
PRECAUTIONS	Recent (≤ 2 weeks) myocardial infarction Serious underlying arrhythmias Serious or worsening angina pectoris Temporomandibular joint disease Pregnancy3 and breastfeeding Adolescents (<18 years)	Recent (≤ 2 weeks) myocardial infarction Serious underlying arrhythmias Serious or worsening angina pectoris Pregnancy3 and breastfeeding Adolescents (<18 years)	Recent (≤ 2 weeks) myocardial infarction Serious underlying arrhythmias Serious or worsening angina pectoris Pregnancy3 and breastfeeding Adolescents (<18 years)	Recent (≤ 2 weeks) myocardial infarction Serious underlying arrhythmias Serious or worsening angina pectoris Underlying chronic nasal disorders (rhinitis, nasal polyps, sinusitis) Severe reactive airway disease Pregnancy3 and breastfeeding Adolescents (<18 years)	Recent (≤ 2 weeks) myocardial infarction Serious underlying arrhythmias Serious or worsening angina pectoris Bronchospastic disease Pregnancy3 and breastfeeding Adolescents (<18 years	Concomitant therapy with medications/conditions known to lower the seizure threshold Hepatic impairment Pregnancy3 and breastfeeding Adolescents (<18 years) Treatment-emergent neuropsychiatric symptoms4 BOXED WARNING REMOVED 12/2016 Contraindications: Seizure disorder Concomitant bupropion (e.g., Wellbutrin) therapy Current or prior diagnosis of bulimia or anorexia nervosa Simultaneous abrupt discontinuation of alcohol or sedatives/benzodiazepines MAO inhibitors in preceding 14 days; concurrent use of reversible MAO inhibitors	Severe renal impairment (dosage adjustment is necessary) Pregnancy3 and breastfeeding Adolescents (<18 years) Treatment-emergent neuropsychiatric symptoms4 BOXED WARNING REMOVED 12/2016
DUSOO	1st cigarette ≤30 minutes after waking: 4 mg 1st cigarette >30 minutes after waking: 2 mg Weeks 1–6: 1 piece q 1–2 hours Weeks 7–9: 1 piece q 2–4 hours Weeks 10–12: 1 piece q 4–8 hours Maximum, 24 pieces/day Chew each piece slowly Park between cheek and gum when peppery or tingling sensation appears (~15–30 chews) Resume chewing when tingle fades Repeat chew/park steps until most of the nicotine is gone (tingle does not return; generally 30 min) Park in different areas of mouth No food or beverages 15 minutes before or during use Duration: up to 12 weeks	1st cigarette ≤30 minutes after waking: 4 mg 1st cigarette >30 minutes after waking: 2 mg Weeks 1–6: 1 lozenge q 1–2 hours Weeks 7–9: 1 lozenge q 2–4 hours Weeks 10–12: 1 lozenge q 4–8 hours • Maximum, 20 lozenges/day • Allow to dissolve slowly (20–30 minutes for standard; 10 minutes for mini) • Nicotine release may cause a warm, tingling sensation • Do not chew or swallow • Occasionally rotate to different areas of the mouth • No food or beverages 15 minutes before or during use • Duration: up to 12 weeks	>10 cigarettes/day: 21 mg/day x 4-6 weeks 14 mg/day x 2 weeks 7 mg/day x 2 weeks 10 cigarettes/day: 14 mg/day x 6 weeks 7 mg/day x 2 weeks • Rotate patch application site daily; do not apply a new patch to the same skin site for at least one week • May wear patch for 16 hours if patient experiences sleep disturbances (remove at bedtime) • Duration: 8–10 weeks	1–2 doses/hour (8–40 doses/day) One dose = 2 sprays (one in each nostril); each spray delivers 0.5 mg of nicotine to the nasal mucosa • Maximum - 5 doses/hour or - 40 doses/day • For best results, initially use at least 8 doses/day • Do not sniff, swallow, or inhale through the nose as the spray is being administered • Duration: 3–6 months	6–16 cartridges/day Individualize dosing; initially use 1 cartridge q 1–2 hours • Best effects with continuous puffing for 20 minutes • Initially use at least 6 cartridges/day • Nicotine in cartridge is depleted after 20 minutes of active puffing • Inhale into back of throat or puff in short breaths • Do NOT inhale into the lungs (like a cigarette) but "puff" as if lighting a pipe • Open cartridge retains potency for 24 hours • No food or beverages 15 minutes before or during use • Duration: 3–6 months	150 mg po q AM x 3 days, then 150 mg po bid • Do not exceed 300 mg/day • Begin therapy 1–2 weeks prior to quit date • Allow at least 8 hours between doses • Avoid bedtime dosing to minimize insomnia • Dose tapering is not necessary • Duration: 7–12 weeks, with maintenance up to 6 months in selected patients	Days 1—3: 0.5 mg po q AM Days 4—7: 0.5 mg po bid Weeks 2—12: 1 mg po bid • Begin therapy 1 week prior to quit date • Take dose after eating and with a full glass of water • Dose tapering is not necessary • Dosing adjustment is necessary for patients with severe renal impairment • Duration: 12 weeks; an additional 12-week course may be used in selected patients • May initiate up to 35 days before target quit date OR may reduce smoking over a 12-week period of treatment prior to quitting and continue treatment for an additional 12 weeks

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ADVERSE EFFECTS	Mouth/jaw soreness Hiccups Dyspepsia Hypersalivation Effects associated with incorrect chewing technique (due to rapid nicotine release): Lightheadedness Nausea/vomiting Throat and mouth irritation	Mouth irritation Nausea Hiccups Heartburn Headache Sore throat Dizziness	Local skin reactions (erythema, pruritus, burning) Headache Sleep disturbances (abnormal or vivid dreams, insomnia); associated with nocturnal nicotine absorption	Nasal and/or throat irritation (hot, peppery, or burning sensation) Rhinitis Ocular irritation/tearing Sneezing Cough Headache	Mouth and/or throat irritation Cough Headache Rhinitis Dyspepsia Hiccups	Insomnia Dry mouth Nervousness/difficulty concentrating Nausea Dizziness Constipation Rash Seizures (risk is 0.1%) Neuropsychiatric symptoms (rare; see PRECAUTIONS)	Nausea Sleep disturbances (insomnia, abnormal/vivid dreams) Constipation Flatulence Vomiting Neuropsychiatric symptoms (rare; see PRECAUTIONS)
ADVANTAGES	Might serve as an oral substitute for tobacco Might delay weight gain Can be titrated to manage withdrawal symptoms Can be used in combination with other agents to manage situational urges Relatively inexpensive	Might serve as an oral substitute for tobacco Might delay weight gain Can be titrated to manage withdrawal symptoms Can be used in combination with other agents to manage situational urges Relatively inexpensive	Once-daily dosing associated with fewer adherence problems Of all NRT products, its use is least obvious to others Can be used in combination with other agents; delivers consistent nicotine levels over 24 hours Relatively inexpensive	Can be titrated to rapidly manage withdrawal symptoms Can be used in combination with other agents to manage situational urges	Might serve as an oral substitute for tobacco Can be titrated to manage withdrawal symptoms Mimics hand-to-mouth ritual of smoking Can be used in combination with other agents to manage situational urges	Twice-daily oral dosing is simple and associated with fewer adherence problems Might delay weight gain Might be beneficial in patients with depression Can be used in combination with NRT agents Relatively inexpensive (generic formulations)	Twice-daily oral dosing is simple and associated with fewer adherence problems Offers a different mechanism of action for patients who have failed other agents
DISADVANTAGES	Need for frequent dosing can compromise adherence Might be problematic for patients with significant dental work Proper chewing technique is necessary for effectiveness and to minimize adverse effects Gum chewing might not be acceptable or desirable for some patients	Need for frequent dosing can compromise adherence Gastrointestinal side effects (nausea, hiccups, heartburn) might be bothersome	When used as monotherapy, cannot be titrated to acutely manage withdrawal symptoms Not recommended for use by patients with dermatologic conditions (e.g., psoriasis, eczema, atopic dermatitis)	Need for frequent dosing can compromise adherence Nasal administration might not be acceptable or desirable for some patients; nasal irritation often problematic Not recommended for use by patients with chronic nasal disorders or severe reactive airway disease	Need for frequent dosing can compromise adherence Cost of treatment Cartridges might be less effective in cold environments (≤60°F)	Seizure risk is increased Several contraindications and precautions preclude use in some patients (see PRECAUTIONS) Patients should be monitored for potential neuropsychiatric symptoms4 (see PRECAUTIONS)	Cost of treatment Patients should be monitored for potential neuropsychiatric symptoms4 (see PRECAUTIONS)
COST/DAY5	2 mg or 4 mg; \$1.90–\$3.60 (9 pieces)	2 mg or 4 mg; \$3.33—\$3.60 (9 pieces)	\$1.52 - \$2.90 (1 patch)	\$8.72 (8 doses)	\$14.88 (6 cartridges)	\$2.58–\$8.25 (2 tablets)	\$15.14 (2 tablets)

- 1. Marketed by GlaxoSmithKline.
- 2. Marketed by Pfizer.
- 3. The U.S. Clinical Practice Guideline states that pregnant smokers should be encouraged to quit without medication based on insufficient evidence of effectiveness and theoretical concerns with safety. Pregnant smokers should be offered behavioral counseling interventions that exceed minimal advice to quit.
- 4. In July 2009, the FDA mandated that the prescribing information for all bupropion- and varenicline-containing products include a black-boxed warning highlighting the risk of serious neuropsychiatric symptoms, including changes in behavior, hostility, agitation, depressed mood, suicidal thoughts and behavior, and attempted suicide. Clinicians should advise patients to stop taking varenicline or bupropion SR and contact a health care provider immediately if they experience agitation, depressed mood, or any changes in behavior that are not typical of nicotine withdrawal, or if they experience suicidal thoughts or behavior. If treatment is stopped due to neuropsychiatric symptoms, patients should be monitored until the symptoms resolve. Based on results of a mandated clinical trial, the FDA removed this boxed warning in December 2016
- 5. Approximate cost based on the recommended initial dosing for each agent and the wholesale acquisition cost from Red Book Online. Thomson Reuters, July 2018.