Pharmacological and Non-Pharmacological Interventions for Smoking Cessation: A Review

Abstract
Dentists are in a unique position to advise the tobacco users to quit the habit through various methods and with effective counseling. The present review provides an outline of the different pharmacological and non-pharmacological interventions for cessation of smoking. Smokers who receive assistance—behavioral, pharmacologic, or both—can experience quit rates of around 20% at least 6 months after quitting. Given the nature of tobacco dependence and the associated difficulty in quitting, pharmacotherapy should be advocated, particularly in patients for whom it is not contraindicated and for whom prior unassisted quit attempts have been unsuccessful. For smokers who are dissonant, physicians should use motivational strategies, such as discussing barriers to cessation and their solutions. For smokers ready to quit, the physician should show strong support and help set a cessation date. Physician counseling for smoking cessation is among the most cost-effective clinical interventions.

Key Words
smoking, cessation, tobacco, interventions, pharmacological, non-pharmacological

INTRODUCTION
Smoking and use of tobacco increases the disease burden and death causing serious health, economic, environmental and social effects. According to the World Health Organization (WHO), use of tobacco in smoking form is the single largest cause of disease and premature death, claiming one life every 8 seconds and killing one of 10 adults globally, which can be preventable. A survey of WHO in 2011 states that there were 100 million premature deaths due to use of tobacco in 20th century, and if this continues by 21st century the number is expected to increase to 1 billion. In India the use of tobacco grows at 2-3% per annum, and by 2020 it will account for 13% of all deaths in the country. Smoking and use of tobacco is a global epidemic causing death of more people than HIV/AIDS, malaria and tuberculosis combined. Necessary steps must be taken to prevent this manmade epidemic globally and in the home country. The authors in this review discussed the important steps taken for prevention of smoking giving more emphasis on pharmacological and non-pharmacological methods of smoking cessation.

PHARMACOLOGICAL METHODS FOR SMOKING CESSATION
Giving up smoking is the easiest thing in the world. I know because I’ve done it thousands of times - Mark Twain. According to the Clinical Practice Guideline for Treating Tobacco Use and Dependence, all smokers trying to quit the tobacco use, must be encouraged to use one or more effective pharmacological agents for cessation except in some special circumstances. The following are the pharmacological agents that are used for smoking cessation (Table 1).

Nicotine Replacement Therapy (NRT)
NRT acts on nicotine receptors in the ventral tegmental area of the brain due to which dopamine is released into the nucleus accumbens. The rationale of using NRT for smoking cessation is twofold. First, it reduces the physical withdrawal symptoms associated with nicotine abstinence among dependent smokers. Second, while alleviating the physiologic symptoms of
Table 1: Pharmacological Agents Used for Smoking Cessation

First line agents
1) Nicotine replacement therapy (NRT): nicotine gum, transdermal patch, nasal spray, oral inhaler and lozenge.

Second line agents
1) Nortriptyline
2) Clonidine

Combination Therapy
1) Nicotine replacement therapy and sustained-release Bupropion
2) Nicotine replacement therapy and nortriptyline

Herbal therapies
1) Lobeline, Dianicline

Emerging therapies
1) Anxiolytic agents - buspirone, diazepam
2) selective serotonin reuptake inhibitors - fluoxetine, paroxetine, sertraline,
3) Mecamylamine
4) Rimonabant
5) Varenicline

Table 2: Various pharmacological agents used in NRT

<table>
<thead>
<tr>
<th>Product</th>
<th>Nicotine Gum</th>
<th>Nicotine Lozenge</th>
<th>Nicotrol Transdermal Patch</th>
<th>Nicotrol NS® Matedry spray (0.5 mg nicotine in 50 mL aqueous nicotine solution)</th>
<th>Nicotrol®, 10 mg cartridge delivers 4 mg inhaled nicotine vapor</th>
</tr>
</thead>
<tbody>
<tr>
<td>Precautions</td>
<td>Pregnancy, lactation, respiratory conditions for nasal spray and inhaler</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
| Dosing                           | 1) For patients smoking 25 or more cigarettes/d: 4 mg
2) For patients smoking fewer than 25 cigarettes/d: 2 mg
a) Weeks 1 to 6: one piece every 1 to 2 hours
b) Weeks 7 to 9: one piece every 2 to 4 hours
c) Weeks 10 to 12: one piece every 4 to 8 hours
1) For patients who smoke their first cigarette ≤ 30 minutes after waking: 4 mg
2) For patients who smoke their first cigarette >30 minutes after waking: 2 mg
a) Weeks 1 through 6: one lozenge every 1 to 2 hours
b) Weeks 7 through 9: one lozenge every 2 to 4 hours
c) Weeks 10 through 12: one lozenge every 4 to 8 hours
1) More than 10 cigarettes/d:
   i) 15 mg/d × 6 wks
   ii) 10 mg/d × 2 wks
   iii) 5 mg/d × 2 wks
b) Who smoke 10 cigarettes/d or fewer:
   Not recommended
1) More than 10 cigarettes/d:
   i) 21 mg/d × 6 weeks
   ii) 14 mg/d × 2 weeks
   iii) 7 mg/d × 2 weeks
b) Who smoke 10 cigarettes/d or fewer:
   i) 14 mg/d × 6 weeks
   ii) 7 mg/d × 2 weeks
1) Initial recommended dose: one to two doses/h (one dose=one spray in each nostril), increasing as needed for symptom relief
2) Recommended duration of therapy: 3 to 6 months
1) Six to 16 cartridges daily (approximately one cartridge every 1 - 2 hours)
2) Each cartridge delivers 4 mg of nicotine over 80 to 100 inhalations (approximately 20 minutes of active puffing).
3) Recommended duration of therapy is up to 6 months (taper dosage during final 3 months of treatment)

Adverse effects
Common adverse effects associated with all NRT agents are mouth soreness, hiccups, dyspepsia, hypersalivation, and jaw ache, headache, flatulence, insomnia, nasal/throat irritation (hot, peppery, or burning sensation), rhinitis, tearing, sneezing.

* a - Marketed by GlaxoSmithKline.
* b - Marketed by Pfizer.

withdrawal, the smoker can focus on the behavioral and psychological aspects of quitting before fully abstaining from nicotine. [7] The following are the different agents used for NRT, their dosages, availability (Table 2).

SECOND LINE AGENTS
These agents are not approved by FDA for smoking cessation and are more prone to adverse effects, so should be used in patients who are unable to use first line agents.

Nortriptyline: It is a tricyclic antidepressant. In general the dose is started at 25 mg/d and is gradually increased over 2 weeks to a target dosage of 75 to 100 mg/d. As it causes sedation, daily dosage should be taken at bedtime. The adverse effects with nortriptyline therapy are sedation, dry
Clonidine: Centrally acting α2-adrenergic agonist that reduces sympathetic outflow from the central nervous system. The dosage for smoking cessation is 0.15 to 0.75 mg/d orally and 0.1 to 0.3 mg/d transdermally. Initially therapy is started with 0.1 mg orally twice daily or 0.1 mg/d transdermally and is increased by 0.10 mg/d each week as tolerated. The duration of therapy was differed in various clinical trials, ranging from 3 to 10 weeks. The adverse effects include dry mouth, drowsiness, nervous system. The dosage for smoking cessation is increased by 0.10 mg/d each week as tolerated.

Combination therapy: It uses long-acting formulation (patch) in combination with short-acting formulation (gum, oral inhaler, lozenge, nasal spray). The long-acting formulation prevents onset of withdrawal symptoms. Short acting formulation is helpful to control withdrawal symptoms that occurred during potential relapse conditions (e.g., after meals, when stressed, or when around other smokers).

Rimonabant: It Antagonizes cannabinoid -1 receptors selectively in central nervous system. The various clinical effects of rimonabant are decrease appetite, weight loss, increased HDL cholesterol, decrease triglycerides, smoking cessation, improved glycemic control from favorable insulin action via higher a dinopectin. It also improves abstinence among smokers. Random clinical trials worldwide showed that those who had been on rimonabant 20 mg and were abstinent at 10 weeks were randomized to continue on 20 mg/day, use 5 mg/day, or use placebo. Those who had been abstinent on 5 mg/day were randomized to continue on 5 mg/day or use placebo. The major adverse effects are nasopharyngitis, upper respiratory tract infection, headache, nausea, dizziness, back pain, influenza, and diarrhea.

Varenicline: It was approved in 2006 for cessation of smoking by FDA. It is a selective alpha-4-beta-2 nicotinic acetylcholine receptor partial agonist. Preliminary data from a phase II clinical trial indicate that in patients randomly assigned to placebo, varenicline (0.5 mg) twice daily, or varenicline (1.0 mg) twice daily, the pooled abstinence rates at weeks 9 through 12 were 12.4%, 45.1%, and 50.6% respectively. Adverse effects observed in more than 10% of patients taking varenicline included nausea, insomnia, headache, and abnormal dreams.

Emerging Therapies: New compounds that have demonstrated encouraging preliminary results include rimonabant and varenicline.

EMERGING THERAPIES

Table 3: The 5 A’s for Facilitating Smoking Cessation

<table>
<thead>
<tr>
<th>Ask about tobacco use</th>
<th>Identify and document tobacco use status for every patient at every visit</th>
</tr>
</thead>
<tbody>
<tr>
<td>Advise to quit</td>
<td>In a clear, strong and personalized manner urge every tobacco user to quit</td>
</tr>
<tr>
<td>Assess willingness to make a cessation attempt</td>
<td>Is the tobacco user willing to make a cessation attempt at this time?</td>
</tr>
<tr>
<td>Assist in cessation attempt</td>
<td>For the patient willing to make a cessation attempt, use counseling and pharmacotherapy to help him or her quit</td>
</tr>
<tr>
<td>Arrange follow-up</td>
<td>Schedule follow-up contact, preferably within the first week after the cessation date</td>
</tr>
</tbody>
</table>

Table 4: The 5 R’s to Enhance Motivation to Quit Smoking

<table>
<thead>
<tr>
<th>Relevance</th>
<th>Identify motivational factors that are relevant for the patient: risk of heart disease, cancer, social situation, second-hand smoke, personal barriers to cessation and prior quit attempts.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Risks</td>
<td>Ask the patient about negative health effects of smoking</td>
</tr>
<tr>
<td>Rewards</td>
<td>Ask the patient about potential benefits of smoking cessation</td>
</tr>
<tr>
<td>Roadblocks</td>
<td>Ask the patient to identify barriers that will make a quit attempt difficult. Provide patient with information on how these barriers can be addressed.</td>
</tr>
<tr>
<td>Repetition</td>
<td>Repeat motivational intervention with each patient encounter.</td>
</tr>
</tbody>
</table>

Table: The 5 R's to Enhance Motivation to Quit Smoking
the smokers who want motivation to quit smoking. R’s (Table 4) practitioner application is useful.\[^4,17\] A successful motivational intervention requires a practitioner who acknowledges patient-specific concerns and previous successful lifestyle changes.\[^18\]

**Enabling the Smoker to Succeed**

Most efficient method for a smoker to successfully quit the habit is combination of pharmacotherapy with non-pharmacological interventions (i.e., advice and behavioral support). Using the both methods in combination multiplied the success rate when used alone.\[^19\] Psychosocial interventions for quitting the smoking is from advice to intensive group or individual counseling. Self-help manuals should be distributed to individuals in large numbers who had the desire and who are highly motivated, confident to quit the smoking and this intervention has the efficacy rate of 5%. Counseling that is delivered in person and interactive telephone counseling are more effective than simply providing educational or self-help materials.\[^20\]

**Complementary and Alternative Therapies**

Other interventional therapies like hypnosis, acupuncture, diet aids and low-level laser therapy have been suggested for smoking cessation. There are no evidences and clinical studies that improved the quit rates with these therapies. When on individual basis these interventions may boost the confidence of the individual towards smoking cessation.\[^21\]

**CONCLUSION**

Many randomized clinical trials and various studies showed that when using both pharmacological and non-pharmacological interventions had a great success in smoking cessation. Besides these interventions other measures like increase of tax on tobacco products, implementation of strict laws on use of tobacco by the governments, health awareness programs among public, incorporating the different topics of tobacco cessation as a syllabus to both medical and dental graduates, conducting various CDE programs, workshops etc are very important for the millions of individuals to quit the habit.

**REFERENCES**

15. Oncken C, Watsky E, Reeves K. Efficacy and safety of varenicline for smoking cessation.
Presented at the National Conference on Tobacco or Health. Chicago, IL. May 6, 2005.


