Smoking cessation pharmacotherapy: an update for health professionals
Smoking cessation pharmacotherapy: an update for health professionals is intended to serve as a reference and a resource for health professionals providing advice for smoking cessation and should be treated as an addendum to the Smoking cessation guidelines for Australian general practice 2004 edition. It should be noted that while the guidelines include a range of therapies, neither the RACGP nor the endorsing organisations hold a commercial interest in any product being appraised. Treatment should be based on the selection of the best pharmacotherapy option available to a patient, taking into account their individual circumstances. Any part of the publication may be reproduced without seeking copyright permission from the Royal Australian College of General Practitioners provided there is appropriate acknowledgement.

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Associate Professor Matthew Peters is a member of the varenicline advisory board and has received honoraria for speaking at meetings for Pfizer Pty Limited.

Dr John Litt has provided smoking cessation advice and training for Pfizer Pty Limited and is a member of the varenicline advisory board for Pfizer Pty Limited.

Endorsements
The Royal Australian College of General Practitioners
The Pharmacy Guild of Australia
Australian Practice Nurses Association
QUIT Group
Action on Smoking and Health
The Cancer Council Australia
Pharmaceutical Society of Australia
The Thoracic Society of Australia and New Zealand
While the rate of smoking in Australia has declined over recent decades, smoking remains the risk factor with the highest levels of disease and death.¹

Smoking cessation advice and support from health professionals are key aspects of a comprehensive approach to tobacco control. Since the publication of the *Smoking cessation guidelines for Australian general practice* in June 2004, there have been important developments in the understanding of nicotine addiction and the pharmacotherapies available to assist cessation, including a new first-line medicine for the treatment of nicotine dependence and a broader indication for the use of nicotine replacement therapy (NRT).

This update aims to inform a range of health professionals – doctors, nurses, pharmacists, dentists, Quitline counsellors and other health professionals – to enable them to provide up-to-date advice to patients on medicines to assist smoking cessation.
Tobacco smoking is a worldwide threat to human health. The World Health Organization estimates that around five million people die prematurely from tobacco related disease each year, and that this number will increase to 10 million by 2020. Seventy percent of these deaths will occur among people in the developing world.2

Australia is part of the Framework Convention on Tobacco Control, a worldwide effort to control the effects of tobacco smoking on human health.4 With the advent of national tobacco control policies and programs, the prevalence of smoking in Australia is among the lowest of any nation.3 Nevertheless, smoking still causes a higher burden of disease than any other behavioural risk factor, representing 9.5% of the total burden in men and 6% in women.4 More than one in six Australians continue to smoke every day, and tobacco smoking is responsible for the deaths of about 16 000 Australians each year.5

Smoking is a known cause of many serious diseases including cancer, heart disease, stroke and chronic obstructive pulmonary disease. Smoking also has adverse effects in pregnancy both for the mother and the developing foetus, and exposure to environmental tobacco smoke damages the health of children and adults6,7 (Figure 1).

Figure 1. Health effects of smoking (from the Smokescreen Program)7,8
According to the 2004 National Drug Strategy Household Survey conducted between June and November 2004:

- Nineteen percent of Australians aged 14 years and over were daily or weekly smokers (20.6% of men and 17.5% of women).
- The prevalence of daily smoking was 17.4% (18.6% of men and 16.3% of women).
- The prevalence by state is similar, with the exception of the Northern Territory, which has the highest rate (27.3% daily smokers).
- The highest rate of smoking is in the 20–29 years age group (23.5%).
- The prevalence is higher among people from lower socio-economic groups.
- Some Australians have higher prevalence rates, e.g., Aboriginal and Torres Strait Islander people (56% of men and 52% of women) and people of south-east Asian background (Vietnam, Lao, and Cambodia) (around 50% in men, but low in women).
- The rate of smoking in adolescents has fallen to 10.7% daily smoking in people aged 14–19 years.

The survey also identified other groups with smoking rates higher than those of the general population, including:

- People with a mental illness (70–90%).
- People with substance use disorders (74–100%).
- Pregnant women (15–26% of all women and 72% of Indigenous women smoke during pregnancy).
Effectiveness of treating tobacco use and dependence

The benefits of quitting smoking are now established. Successfully quitting smoking can result in an increase in life expectancy of up to 10 years, and there is now substantial evidence that advice from general practitioners and other health professionals helps smokers quit. While spending more time (longer than 10 minutes) advising smokers to quit yields higher abstinence rates than minimal advice, offering brief advice (as little as three minutes) has been shown to have clear benefits (Figure 2).

<table>
<thead>
<tr>
<th>GP time</th>
<th>Estimated abstinence rate</th>
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<tr>
<td>&gt;10 minutes</td>
<td>22.1%</td>
</tr>
<tr>
<td>3–10 minutes</td>
<td>16.0%</td>
</tr>
<tr>
<td>&lt;3 minutes</td>
<td>13.4%</td>
</tr>
<tr>
<td>Nil</td>
<td>6.4%</td>
</tr>
</tbody>
</table>

**Figure 2. Reality pyramid: levels of intervention** (adapted from Litt J et al)

Smoking cessation is both clinically and cost effective compared with other medical and disease preventive measures such as the treatment of hypertension and hypercholesterolaemia, and preventive screening interventions such as mammography and cervical smears. Research has shown that the cost of smoking cessation interventions per life year gained ranged from $500 to $2000, while the median cost per life year gained for other medical interventions was $40 000. Along with childhood immunisation and aspirin use with high risk adults, overall efforts to reduce tobacco smoking are among the most beneficial preventive interventions on human health.

All health professionals should systematically identify smokers, assess their smoking status and offer them advice to stop smoking at every opportunity. Where appropriate, smokers should be referred to more intensive help (eg. Quitline).

Brief interventions for smoking cessation in primary care involve opportunistic advice, encouragement and referral. Interventions should include one or more of the following:

- simple advice to stop smoking
- an assessment of the smoker’s commitment to stop
- an offer of pharmacotherapy and counselling where appropriate
- self help material and referral to more intensive support (eg. Quitline).
Neurobiology of nicotine dependence

‘People smoke for nicotine but die of smoke’
Michael Russell, pioneer in nicotine addiction research

Dependence on smoking is a complex process. It requires a close link in time between the context in which smoking occurs, its rituals, the sensory stimuli of touch, taste and smell, and the extremely rapid delivery of nicotine to the brain that occurs after smoking a modern cigarette. Dependence would not occur without nicotine – but nicotine is not responsible for the harmful effects of smoking.7,18

Greater understanding of the neurobiology of nicotine dependence is improving the use of existing cessation therapies and is helping to develop new compounds to aid smoking cessation. But although most smokers possess some understanding of the health effects of tobacco use, recent research in Australia indicates that the majority of smokers do not take into account a thorough understanding of these effects when making daily decisions about their smoking.19,20

The huge surface area of the lungs means that nicotine is rapidly absorbed into the pulmonary venous circulation. Nicotine is active within the ‘brain reward system’ of the central nervous system within seconds of inhalation.20,21 This is the common pathway for the experience of pleasure from many different social, physical and chemical stimulants, including other drugs of addiction such as cocaine and opiates.

Figure 3. The primary role of the α4β2 nicotinic acetylcholine receptor in the dependence producing effects of nicotine, measured by increased dopamine release in the key brain site for motivation and reward, the nucleus accumbens23.
Neurobiology of nicotine dependence

Within this system, nicotine interacts most critically with specific $\alpha_4\beta_2$ nicotinic acetylcholine (ACh) receptors. This interaction triggers the release of dopamine and the brief but intense activation of the brain reward system. Dopamine levels then decline slowly, with a half life of about 90 minutes. The eventual deactivation of this same reward system leads to sensations of withdrawal or cravings, which may be relieved by smoking another cigarette\(^{22,23}\) (Figure 3).

A long term smoker cycles continuously between pleasure and withdrawal, with the negative effects of nicotine withdrawal being the major factors in a person’s decision to continue smoking.

Genetic factors may play a role in determining whether this cycle of dependence develops in a long term smoker. Some characteristics of a nondependent smoker include smoking fewer than seven cigarettes per day, a long interval before the first cigarette of the day, and periods of cessation and recommencement of smoking not associated with withdrawal symptoms.

Assessment of nicotine dependence

Most smokers become nicotine dependent. Assessment of nicotine dependence can help predict whether a smoker is likely to experience nicotine withdrawal upon stopping smoking.

Nicotine withdrawal symptoms commonly include craving, as well as onset of any of the following shortly after stopping:\(^7\)

- depressed mood
- insomnia
- irritability, frustration, anger
- anxiety
- difficulty in concentration
- restlessness
- decreased heart rate
- increased appetite or weight gain.

Characteristics of smokers with nicotine dependence include smoking soon after waking, smoking when ill, difficulty stopping smoking, finding the first cigarette of the day the most difficult to give up, and smoking more in the morning than in the afternoon.\(^{24}\)

A quick assessment of nicotine dependence can be made by asking the smoker:

- how soon after waking do you have your first cigarette?
- how many cigarettes do you smoke each day?
- have you had cravings and withdrawal symptoms when you have tried to quit?

Smoking within 30 minutes of waking, smoking more than 15 cigarettes per day and a history of withdrawal symptoms in previous attempts to quit are all indicators of nicotine dependance (see Quit smoking brief intervention).\(^7\)
Tobacco dependence is a chronic condition that typically requires repeated cessation treatment and ongoing care. A minority of smokers achieve long term abstinence on the first attempt to quit; the majority cycle through multiple attempts with relapse and remission before achieving long term or permanent abstinence. Twenty or more attempts over a period of years is not unusual.

It is important to take every opportunity in a health care setting to identify all smokers, document their smoking status and offer treatment, which may involve counselling by a health professional, referral and pharmacotherapy.

Evidence is now accumulating to show that increasing the number of attempts to quit is the most important strategy for improving cessation rates in the population, and health professionals have a role in prompting and encouraging attempts to quit. The 5As approach provides a structure for identifying all smokers and offering support to help them quit (Quit smoking brief intervention).

The process of quitting smoking (‘stage of readiness to change’) — involving a series of stages from thinking about quitting to making a quit attempt — is a useful framework to identify and provide the appropriate support for a smoker’s level of interest in quitting. This approach has been widely used in guidelines for the management of other risk factors. However, there is some evidence that the likelihood of success in an attempt to quit is unrelated to the smoker’s expressed interest in quitting in the period leading up to the attempt: unplanned attempts to quit are as likely (or even more likely) to be as successful as planned attempts. This means there is benefit in encouraging all smokers to consider quitting whenever the opportunity arises.

The evidence is also compelling that both advice based help and pharmacotherapy can increase the rate of success of quit attempts, and when both are used the benefits are additive.
The 5As structure for smoking cessation

**Quit smoking brief intervention**

1. **ASK**
   - Ask all clients
     - Do you smoke tobacco?
     - Record smoking status (current smoker)

2. **ASSESS**
   - Assess
     - Assess stage of change: “How do you feel about your smoking at the moment?” and “Are you ready to stop smoking now?”
     - Record stage of change
     - Assess nicotine dependence

3. **ADVISE**
   - All smokers should be advised to quit in a way that is clear but nonconfrontational, eg. “The best thing you can do for your health is to quit smoking.”

4. **ASSIST**
   - Assist – not ready
     - Discuss the benefits of quitting and risks of continued smoking
     - Provide information about not exposing others to passive smoking
     - Advise that help is available when they’re ready

   - Assist – unsure
     - Do motivational interviewing: “What are the things you like and don’t like about your smoking?”
     - Explore their doubts
     - Explore barriers to quitting
     - Offer written information (eg. Quit Kit) and referral to Quitline 13 7848

   - Assist – ready
     - Affirm and encourage
     - Provide a Quit Kit and discuss a quit plan
     - Recommend pharmacotherapy to nicotine dependent smokers (see Assess)
     - Discuss relapse prevention
     - Offer referral to Quitline 13 7848

5. **ARRANGE FOLLOW-UP**
   - Arrange follow-up
     - For clients attempting to quit, arrange follow up visit, if possible
     - At these visits:
       - congratulate and affirm decision
       - review progress and problems
       - encourage continuance of pharmacotherapy
       - discuss relapse prevention
       - encourage use of support services
     - OR
     - Refer to Quitline 13 7848

   - Successful quitter
     - Congratulate and affirm decision to quit
     - Discuss relapse prevention

   - Relapse
     - Offer support and reframe as a learning experience
     - Explore reasons for relapse and lessons for future quit attempts
     - Offer ongoing support
     - Ask again at future consultations

   - Assess nicotine dependence
     - Nicotine dependence can be assessed by asking:
       1. How many minutes after waking to first cigarette?
       2. Number of cigarettes per day?
       3. What cravings or withdrawal symptoms in previous quit attempts?
     - Smoking within 30 minutes of waking, smoking more than 15 cigarettes per day and history of withdrawal symptoms in previous quit attempts are all markers of nicotine dependence
     - Pharmacotherapy for dependent smokers is proven to double the chances of successfully quitting

   - Assist – action and maintenance
     - Congratulate
     - Discuss relapse prevention
     - Review and reinforce benefits of quitting
     - Offer written information (eg. Quit Kit) and referral to Quitline 13 7848
A number of medicines have been shown to assist smoking cessation in meta-analyses of randomised clinical trials.\textsuperscript{29-32}

No form of pharmacotherapy is a substitute for motivation. A range of nicotine replacement products are available from pharmacies (and supermarkets in some cases) in addition to two prescription-only medicines currently registered in Australia.

Nicotine replacement therapy (NRT) can be used effectively without the intervention of a health care professional; the two prescription-only medicines should be given in combination with counselling support from a health professional or a support service.

Both prescription medicines have shown efficacy in clinical trials in smokers who have regular counselling support. While both advice based help and pharmacotherapy have been shown to increase the rate of success of attempts to quit, when used together the benefits are additive.\textsuperscript{29,30}

**First-line pharmacotherapy options**

First-line options are medicines that have been shown to be effective and are licensed for smoking cessation. In Australia these are NRT, varenicline and bupropion.

From currently available evidence, varenicline is the most effective form of pharmacotherapy.\textsuperscript{33,34} This conclusion is based on a small number of studies, and further research is awaited in different patient populations. Varenicline has been shown to be more effective than bupropion, and although there have been no direct comparative studies with NRT, by inference it is more effective than low dose forms of NRT (which is of similar effectiveness to bupropion). Further research is needed to prove whether varenicline is more effective than high dose forms of NRT or combination therapy (two forms of NRT, or NRT and bupropion).

However, context and patient preference are important in choosing the pharmacotherapy that is most likely to assist the smoker in an attempt to quit: some smokers may prefer a nonprescription medicine such as NRT that they can commence immediately, while others may prefer a non-nicotine option subsidised by the Pharmaceutical Benefits Scheme (PBS) (see Treatment algorithm). The PBS requirement for subsidy is that these medicines are provided only in combination with counselling.

**Nicotine replacement therapy**

Nicotine replacement therapy is available over the counter in pharmacies, and some forms are available in supermarkets. The aim of NRT is to reduce withdrawal symptoms by providing some of the nicotine that would normally be obtained from cigarettes, without providing the harmful components of tobacco smoke. None of the available forms of NRT – transdermal patch, inhaler, lozenge, sublingual tablet and gum – offer the same rapid nicotine delivery of a cigarette.

Some oral forms of NRT are available in two strengths: 2 mg and 4 mg. The 4 mg version is recommended for more dependent smokers. The 2004 guidelines recommend that pharmacotherapy with NRT should be offered to all smokers who are motivated to quit.\textsuperscript{7} Regular use of NRT beyond 12 months is not generally recommended. Long term use of some forms of NRT has been reported and has not caused ill health effects; it may help some people remain abstinent.\textsuperscript{35}
**Nicotine replacement therapy (NRT)**

**Clinical suitability**
Can be used in all groups of smokers including children and adolescents, pregnant women and patients with cardiovascular disease (check PI).

**Patient choice**
Reasons to prefer:
- Over the counter availability
- Concerns about side effects of varenicline and bupropion
- Can be used in pregnancy.

**Varenicline**

**Clinical suitability**
Not recommended in pregnancy, childhood or with significant intercurrent psychological/psychiatric disease. Nausea in 30% of patients. Reduce dose in severe renal impairment (check PI).

**Patient choice**
Reasons to prefer:
- On current evidence, varenicline is the most effective pharmacotherapy
- PBS subsidy
- Lack of serious adverse effects and interactions.

**Bupropion sustained release**

**Clinical suitability**
Absence of contraindications such as current or past seizures, concurrent monoamine oxidase inhibitors, pregnancy. Caution with other conditions or drugs that lower seizure threshold (check PI).

**Patient choice**
Reasons to prefer:
- PBS subsidy
- Oral non-nicotine preparation
- Relapse in past using NRT
- Evidence of benefit in chronic disease and depression.

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**Smoking cessation pharmacotherapy: an update for health professionals**

**Nicotine replacement therapy (NRT)**

**Clinical suitability**
Can be used in all groups of smokers including children and adolescents, pregnant women and patients with cardiovascular disease (check PI).

**Patient choice**
Reasons to prefer:
- Over the counter availability
- Concerns about side effects of varenicline and bupropion
- Can be used in pregnancy.

- Discuss benefit of follow up GP visits, especially if there are concerns about the medications, eg. common adverse effects such as skin irritation, sleep disturbance
- Encourage use of support services
- Encourage completion of at least 10 weeks of therapy
- Consider combination NRT if withdrawal not controlled
- Consider a further follow up visit if patient needs extra support

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**Varenicline**

- Give initial 4 week script; arrange for return for second script and discussion of progress
- Encourage use of support services
- At follow up, review progress and problems: common adverse effects, nausea and abnormal dreams
- Encourage completion of 12 weeks of therapy
- Consider a further follow up visit if patient needs extra support.

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**Bupropion sustained release**

- Give initial 2 week script; arrange for return for second script and discussion of progress
- Encourage use of support services
- At follow up, review progress and adverse effects: monitor allergy problems (skin rash) and insomnia
- Encourage completion of at least 7 weeks of therapy
- Consider combination treatment if withdrawal not controlled
- Consider a further follow up visit if patient needs extra support
There have been some recent changes to the indications listed for NRT products:\textsuperscript{36,37}

- more than one form of NRT can be used concurrently
  - in Australia, NRT patch and 2 mg gum are licensed for smokers who have relapsed in the past or who experience cravings using only one form of NRT
- NRT can be used by pregnant and lactating smokers
- all forms of NRT can be used by patients with cardiovascular disease
- all forms of NRT can be used by smokers aged 12–17 years
- some forms of NRT are now available in supermarkets; the other NRT products are available over the counter in pharmacies.

Although quitting smoking completely remains the best goal, there is a strong argument for the use of nicotine assisted reduction to help patients quit. New research suggests that reducing smoking with NRT as a first step can increase the number of smokers who quit altogether.\textsuperscript{36}

The available evidence shows that:

- all forms of NRT are equally effective in aiding long term cessation
- NRT helps smokers unwilling or unable to stop smoking to reduce their cigarette consumption (’cut down then stop’)
- there are no real risks of use; using NRT to quit is always safer than continuing to smoke
  - exceptions are pregnant women, for whom there may be a risk to the foetus, and people with acute cardiovascular disease.

There has been little research on the success of combinations of different types of NRT.

**Varenicline (Champix\textsuperscript{®})**

Varenicline is a new class of drug for first-line pharmacotherapy. It was developed specifically for smoking cessation by targeting the nicotinic acetylcholine (ACh) receptor in the reward centres in the brain. Varenicline binds with high affinity at the $\alpha_4\beta_2$ nicotinic ACh receptor, where it acts as a partial agonist to alleviate symptoms of craving and withdrawal. At the same time, if a cigarette is smoked, the drug prevents inhaled nicotine from activating the $\alpha_4\beta_2$ receptor sufficiently to cause the pleasure and reward response (Figure 4).

![Figure 4. Mode of action: varenicline's partial agonism at the $\alpha_4\beta_2$ nicotinic acetylcholine receptor can simultaneously decrease craving and withdrawal symptoms](image-url)
In two clinical studies with identical study designs, varenicline in combination with counselling produced a continuous abstinence rate from week 9 through to 1 year of 21.9% (8.4% in placebo group \(p < 0.001\) and 16.1% in bupropion group \(p = 0.057\)) and 23% (10.3% in placebo group \(p < 0.001\) and 14.6% in bupropion group \(p = 0.004\)) respectively. The abstinence rate for varenicline was significantly better than both bupropion and placebo long term. As yet there have been no direct comparison studies with varenicline and NRT alone or in combination (with other NRT or bupropion).

Prolonged use of varenicline has also been shown to reduce relapse. In subjects who stopped smoking at the end of 12 weeks of treatment, an additional 12 weeks of treatment was more beneficial than placebo in maintaining abstinence to the end of treatment and to one year from the start of treatment. The difference in continuous abstinence for weeks 13–52 between intervention and control groups was small, however. The benefit appears to be maintained only for the period of use of varenicline.

Nausea is the most common adverse effect of varenicline and was reported in the studies in almost 30% of smokers, although less than 3% discontinued treatment due to nausea. Abnormal dreams were also more common in the varenicline group (13.1%) than either the bupropion (5.9%) or placebo groups (3.5%). No clinically meaningful drug interactions have been identified.

Varenicline is excreted almost entirely by the kidneys. Dose adjustment is required for severe renal impairment; dosing should begin at 0.5 mg once daily for the first 3 days, and then be increased to 1 mg once daily. No dose adjustment is needed for patients with hepatic impairment.

There is a lack of evidence of the efficacy and safety of varenicline in patients with significant psychiatric problems, and therefore use in such patients is not recommended. There have been reports from the United States of depressed mood, agitation, changes in behaviour, suicidal ideation and suicides in patients attempting to quit smoking using varenicline. Smoking cessation itself can be associated with exacerbation of underlying psychiatric disease. The relationship of these symptoms, if any, to varenicline is being investigated by regulatory authorities. Prescribers should ask patients to report any mood or behaviour changes.

Varenicline is available in Australia on the PBS as short term adjunctive therapy for nicotine dependence. The first script is a starter pack lasting 4 weeks, followed by a maintenance batch for the remainder of the 12 week course. The patient should set a quit date 1–2 weeks after commencing varenicline. The medicine can be taken whole with water and food to help reduce nausea.

The recommended dose of varenicline is 1 mg twice daily following a 1 week titration as follows:

<table>
<thead>
<tr>
<th>Days</th>
<th>Dose</th>
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<tr>
<td>1–3</td>
<td>0.5 mg once daily</td>
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<tr>
<td>4–7</td>
<td>0.5 mg twice daily</td>
</tr>
<tr>
<td>8</td>
<td>1 mg twice daily</td>
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</table>

**Bupropion sustained release (Zyban®)**

Originally developed as an antidepressant, bupropion is a non-nicotine oral therapy which reduces the urge to smoke and reduces symptoms from nicotine withdrawal.
Since 2001, bupropion sustained release has been available in Australia as a PBS authority item once per year. It is a short term adjunctive therapy for nicotine dependence in conjunction with counselling with the goal of maintaining abstinence.

Bupropion doubles the cessation rate compared with placebo: over 12 months, data from two randomised controlled trials showed 9% and 19% of smokers had not smoked for the 12 months following placebo and bupropion therapy respectively. It has been shown to be effective in a range of patient populations including smokers with depression, cardiac disease and respiratory diseases including chronic obstructive pulmonary disease. It has also been shown to improve short term abstinence rates for people with schizophrenia.

Bupropion is contraindicated in patients with a history of seizures, eating disorders and patients taking monoamine oxidase inhibitors. It should be used with caution in people taking medications that can lower seizure threshold, such as antidepressants and oral hypoglycaemic agents.

Seizures are the most clinically important adverse effect (0.1% risk). Common adverse effects are insomnia, headache, dry mouth, nausea, dizziness and anxiety.

Bupropion is available as a starter pack of 30 tablets and a continuation pack of 90 tablets. The dose of bupropion sustained release is 150 mg once a day for the first 3 days and then increased to 150 mg twice a day. The patient should stop smoking in the second week of treatment. Bupropion can be used in combination with NRT, but blood pressure should be monitored.

Clinical trials have shown that bupropion is not as effective as varenicline, and it is likely to be the treatment of choice only in cases where varenicline is not appropriate (patient choice or as a result of severe side effects) or for smokers with depression or schizophrenia.

Second-line pharmacotherapy options

Nortriptyline

The tricyclic antidepressant nortriptyline has been shown to approximately double cessation rates compared to placebo (OR=2.1). A recent systematic review shows that the use of nortriptyline for smoking cessation resulted in higher prolonged abstinence rates after at least 6 months compared to placebo treatment. The efficacy of nortriptyline does not appear to be affected by a past history of depression; it is, however, limited in its application by its potential for serious side effects. These include dry mouth, constipation, nausea, sedation and headaches, and a risk of arrhythmia in patients with cardiovascular disease. It can be dangerous in overdose.

Nortriptyline is not registered for smoking cessation in Australia. The dose of nortriptyline used for smoking cessation is approximately 75 mg per day for 12 weeks. Further information about dose titration can be obtained from the New Zealand smoking cessation guidelines.

Possible future options

There are a number of tobacco cessation therapies and nicotine vaccines in development. The selective type 1 cannabinoid receptor antagonist rimonabant and the nicotine receptor partial agonist cysteine have demonstrated some efficacy in studies, but as yet there is insufficient evidence for their use in tobacco cessation. Further data and research are needed before any of these methods of treatment for tobacco dependency can be recommended.
There are several population groups for whom there are particular implications regarding nicotine dependence and the effects of smoking, and the use of medicines for smoking cessation.

Many of these groups – children and adolescents, pregnant and lactating women, people with mental illnesses, people with substance use disorders and people with smoking related diseases – have not been studied in clinical trials of pharmacotherapy for smoking cessation. However, the same guidelines for quitting smoking apply to all groups: every opportunity should be taken to offer all smokers advice and support to stop smoking.

Counselling and behavioural interventions may be modified to be appropriate for the individual smoker. In addition, all smokers should be offered pharmacotherapy with the following guidance (Table 1).

### Table 1. First-line pharmacotherapy in special populations

<table>
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<tr>
<th>Special group</th>
<th>Varenicline</th>
<th>Bupropion</th>
<th>Nicotine replacement therapy</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pregnant and lactating women</td>
<td>ND</td>
<td>ND</td>
<td>✔(^a)</td>
</tr>
<tr>
<td>Children and adolescents (12-18 years)</td>
<td>ND</td>
<td>ND</td>
<td>✔</td>
</tr>
<tr>
<td>People with smoking related diseases</td>
<td>ND</td>
<td>✔</td>
<td>✔</td>
</tr>
<tr>
<td>• Cardiovascular disease</td>
<td>ND</td>
<td>✔</td>
<td>✔</td>
</tr>
<tr>
<td>• Chronic obstructive pulmonary disease</td>
<td>ND</td>
<td>✔</td>
<td>✔</td>
</tr>
<tr>
<td>• Diabetes</td>
<td>ND</td>
<td>ND</td>
<td>✔(^b)</td>
</tr>
<tr>
<td>• Severe renal impairment</td>
<td>✔(^c)</td>
<td></td>
<td>✔(^d)</td>
</tr>
<tr>
<td>• Moderate to severe hepatic impairment</td>
<td>✔</td>
<td></td>
<td>✔(^d)</td>
</tr>
<tr>
<td>People with mental illness</td>
<td>ND</td>
<td>✔</td>
<td>✔</td>
</tr>
<tr>
<td>• Depression</td>
<td>ND</td>
<td>✔</td>
<td>✔</td>
</tr>
<tr>
<td>• Schizophrenia</td>
<td>ND</td>
<td>✔(^e)</td>
<td>✔</td>
</tr>
<tr>
<td>People with substance use disorders</td>
<td>ND</td>
<td>✔(^f)</td>
<td>✔</td>
</tr>
<tr>
<td>Contraindications (apart from hypersensitivity to active ingredient)</td>
<td>Yes(^g)</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

✔ = suitable  ND = lack of safety data

\(^a\) = intermittent dosing products preferable  \(^b\) = but closely monitor blood sugar levels
\(^c\) = dosing adjustment required \(^d\) = with caution
\(^e\) = with close follow up \(^f\) = caution with alcohol abuse
\(^g\) = seizures, anorexia, bulimia, central nervous system tumours, monoamine oxidase inhibitor treatment within 14 days
Supporting quality use of smoking cessation pharmacotherapies

Even motivated smokers relapse. Most relapses occur in the first few weeks of an attempt to quit. It often takes a number of attempts to quit smoking successfully, and smokers should be encouraged to keep trying to quit even if they have returned to smoking.

All available first-line medicines for stopping smoking (NRT, varenicline and bupropion) have been shown to double the chance of quitting, but only if they are used correctly and for the right length of time. Optimum use of these medicines involves prescribing them in conjunction with patient counselling and support. Integrating behavioural therapy (face-to-face, internet and telephone counselling and coaching) increases the quit rate significantly. Many different health professionals (physicians, nurses, dentists, psychologists, pharmacists) are effective in increasing abstinence rates.

Besides the support offered to smokers by health professionals, other support options include:

- individual face-to-face support (doctor or other health professional with special training to help people to quit smoking)
- telephone advice (Quitline)
  - the national Quitline number is 13 QUIT (13 7848) and calls are charged at the cost of a local call (about 25c, mobile phones extra). All Quitline services in Australia have agreed to national minimum standards of service delivery
  - research based proactive counselling service
  - fax referral to Quitline (smokers can be referred to the Quitline for extended support using the fax referral sheet)
  - Quitline adviser, course leader or coach
  - self help books (eg. www.quitbecauseyoucan.org.au)
- support programs offered by pharmaceutical companies, such as:
  - www.click2quit.com.au (NRT, GlaxoSmithKline Consumer Healthcare)
  - Champix ‘My Time to Quit’ program (varenicline, Pfizer).

Internet resources

RACGP publications (including Smoking cessation guidelines for Australian general practice)
www.racgp.org.au/guidelines
www.racgp.org.au/guidelines/smokingcessation

Quit Victoria
www.quit.org.au

Federal, state and territory initiatives (provides links to other tobacco control sites)
www.quitnow.info.au

Action on Smoking and Health (ASH) Australia
www.ashaust.org.au
Conclusion

Tobacco dependence is a chronic condition which leads to disease and death for smokers, and substantial costs for the public health system.

There are now three first-line medical options available in Australia to assist smokers to quit:
- nicotine replacement therapy
- varenicline
- bupropion.

These pharmacotherapies have been shown to be effective in increasing long term abstinence. Other agents for tobacco dependence treatment are in development, further increasing the potential options for smokers who want to quit smoking.

It is important for health professionals to maximise the opportunities for smoking cessation by encouraging all smokers to quit and by supporting all patients attempting to quit. Supporting a quit attempt involves recommending pharmacotherapy that suits the individual smoker, and ensuring all smokers have access to appropriate counselling and support.

References


References


